Identification and characterization of RomX and RomY, two novel motility regulators in *Myxococcus xanthus*

**Dissertation**
zur Erlangung des akademischen Grades
des Doktors der Naturwissenschaften
(Dr. rer. nat.)

dem Fachbereich Biologie
der Philipps-Universität Marburg
vorgelegt

von

**Dobromir Szadkowski**
aus Nowy Dwor Mazowiecki, Polen

Marburg an der Lahn, 2018
Die Untersuchungen zur vorliegenden Arbeit wurden von August 2013 bis April 2018 am Max-Planck-Institut für terrestrische Mikrobiologie in Marburg unter der Leitung von Prof. Dr. MD Lotte Søgaard-Andersen durchgeführt.

Vom Fachbereich Biologie der Philipps Universität Marburg als Dissertation angenommen am ___.___.___

Erstgutachter: Prof. Dr. MD Lotte Søgaard-Andersen
Zweitgutachter: Prof. Dr. Martin Thanbichler
Weitere Mitglieder der Prüfungskommission:

Prof. Dr. Victor Sourjik
Prof. Dr. Gert Bange

Tag der mündlichen Prüfung: ___.___.___
Die während der Promotion erzielten Ergebnisse sind zum Teil in folgenden Originalpublikationen veröffentlicht worden:

**MglC, a Paralog of Myxococcus xanthus GTPase-Activating Protein MglB, Plays a Divergent Role in Motility Regulation,**

**The RomX/RomR MglA GEF complex establishes front-rear polarity for efficient motility in M. xanthus**
Dla Rodzicow
# Table of contents

## 1. INTRODUCTION

1.1 Spatial organization of bacterial cells .................................................... 13
1.2 Characteristics of small eukaryotic GTPases .......................................... 16
1.3 Myxococcus xanthus as a model organism ........................................... 18
1.4 Type IV pilus motility .............................................................................. 19
1.5 Gliding motility ...................................................................................... 21
1.5.1 Motor cargo complex .......................................................................... 25
1.5.2 The focal adhesion complexes model ............................................... 26
1.6 Slime ........................................................................................................ 28
1.7 Cell polarity and motility regulation – spatial control of motility .......... 29
1.7.1 Spatial organization of cell polarity by MglA, MglB and RomR ............ 29
1.7.2 Frz chemosensory system .................................................................... 32
1.7.3 Regulation of cell polarity by MglC, an MglB homolog ....................... 35
1.7.4 Regulation of motility by the small GTPase SofG ............................. 36
1.8 Scope of this study .................................................................................. 37

## 2. RESULTS

2.1 Identification of RomX and RomY ............................................................ 39
2.2 Characterization of RomX in Myxococcus xanthus ............................... 40
2.2.1 RomX is important for both motility systems .................................... 42
2.2.2 RomX acts in the same pathway as RomR, MglA and MglB ............... 46
2.2.3 RomX localizes dynamically to the cell poles .................................... 49
2.2.4 RomX is polar targeting determinant of MglA .................................. 54
2.2.5 Formation of focal adhesion depends on RomX and RomR in the presence of MglB .......................................................... 57
2.2.6 Focal adhesion immobility depends on RomX and RomR ............... 62
2.2.7 RomX and RomR localize to focal adhesion complexes .................... 64
2.2.8 RomX incorporation into the focal adhesion complexes depends on RomR .......................................................... 66
2.2.9 RomR incorporation into focal adhesion complexes is independent of RomX .......................................................... 67
2.2.10 RomX interacts with MglA and RomR in the bacterial two hybrid system .......................................................... 70
2.2.11 RomX/RomR complex interacts with MglA-GTP ................................ 71
2.2.12 RomX and/or RomR do not inhibit MglB GAP and MglA GTPase activity .......................................................... 74
2.2.13 RomX/RomR complex stimulates MglA nucleotide exchange .......................................................... 75
2.3. CHARACTERIZATION OF ROMY IN MYXOCOCCUS XANTHUS ................................................................................ 77
  2.3.1 RomY is important for both motility systems .................................................................................. 78
  2.3.2 RomY acts in the same pathway as MglA, MglB, RomR and RomX ................................................... 79
  2.3.3 RomY localization studies .................................................................................................................. 82
  2.3.4 RomY interacts with RomX and MglA in bacterial two hybrid system ............................................... 85
2.4 CHARACTERIZATION OF THE IMPORTANCE OF ROMR FOR REVERSALS .................................................................... 86
  2.4.1 ΔromR and ΔromX mutants respond to IAA ................................................................................... 86
  2.4.2 Mimicking and blocking of possible RomR phosphorylation site has no effect on the motility ........... 89
3. DISCUSSION ..................................................................................................................................... 93
  3.1 ROMX REGULATES CELL POLARITY TOGETHER WITH MGLA, MGLB AND ROMR ...................................................... 93
  3.2 ROMR/ROMX COMPLEX ACTS AS AN MGLA GEF ............................................................................................. 95
  3.3 FORMATION OF FOCAL ADHESION COMPLEXES DEPENDS ON ROMX AND ROMR ...................................................... 96
  3.4 ROMX AND ROMR LOCALIZE TO THE FOCAL ADHESION COMPLEXES ................................................................. 97
  3.5 ROMX AND ROMR ARE IMPORTANT FOR ATTACHMENT, STABILITY AND DIRECTIONALITY OF THE FOCAL ADHESION
  COMPLEXES .................................................................................................................................................... 97
  3.5 ROMX/RomR ESTABLISHES FRONT-REAR POLARITY, CELL ASYMMETRY AND GLIDING MOTILITY BY ITS TRIPLE FUNCTION.. 99
  3.6 ROMY REGULATES CELL POLARITY TOGETHER WITH MGLA, MGLB, ROMR AND ROMX .......................................... 100
  3.7 ROMX AND ROMR ARE NOT ESSENTIAL FOR GENERATING AN OUTPUT FROM THE FRZ SYSTEM ................................. 102
4. MATERIALS AND METHODS ........................................................................................................... 105
  4.1 CHEMICALS, EQUIPMENT AND SOFTWARE ..................................................................................................... 105
  4.2 MEDIA ................................................................................................................................................... 107
  4.3 MICROBIAL METHODS ....................................................................................................................................... 109
  4.3.1 E. coli strains used in this study ........................................................................................................ 109
  4.3.2 M. xanthus strains used in this study ............................................................................................... 110
  4.3.3 Cultivation and storage of E. coli and M. xanthus ........................................................................... 112
  4.3.4 Bacterial Two Hybrid Assay (BACHT) ............................................................................................... 113
  4.3.5 Motility assays for M. xanthus .......................................................................................................... 114
  4.3.6 Reversal frequency assay for M. xanthus on 1.5% agar, 0.5% CTT .................................................. 114
  4.3.7 Reversal frequency assay for M. xanthus moving by the T4P-dependent motility ........................................... 114
  4.3.8 Trypan blue and congo red dyes binding assay ............................................................................... 115
  4.3.9 Epifluorescence microscopy ............................................................................................................. 115
  4.3.10 Tirf microscopy ....................................................................................................................................... 116
  4.4 MOLECULAR BIOLOGY METHODS ................................................................................................................. 116
  4.4.1 Plasmids and oligonucleotides ......................................................................................................... 116
  4.4.2 Plasmids construction ...................................................................................................................... 119
  4.4.3 Generation of in-frame deletion mutants ........................................................................................ 120
  4.4.4 DNA isolation from E. coli and M. xanthus ...................................................................................... 121
4.4.5 Polymerase Chain Reaction (PCR) ................................................................. 121
4.4.6 Agarose gel electrophoresis ................................................................. 122
4.4.7 DNA restriction and ligation ................................................................. 123
4.4.8 Preparation and transformation of chemically competent E. coli cells ................................................................. 123
4.4.8 Preparation and transformation of electrocompetent M. xanthus cells ................................................................. 123
4.5 BIOCHEMICAL METHODS .............................................................................. 124
4.5.1 SDS-polyacrylamide gel electrophoresis (SDS-PAGE) ......................................................... 124
4.5.2 Determination of total protein concentration in cell extracts ......................... 124
4.5.3 Immunoblot analysis .................................................................................. 125
4.5.4 Proteins purification .................................................................................. 125
4.5.5 GTPase assay .......................................................................................... 129
4.5.6 Pull down experiment ................................................................................ 129
4.5.6 Nucleotide exchange experiments ................................................................ 130
5. REFERENCES .................................................................................................. 132
ACKNOWLEDGEMENTS ..................................................................................... 140
CURRICULUM VITAE .......................................................................................... 141
ERKLÄRUNG ....................................................................................................... 143
EINVERSTÄNDNISERKLÄRUNG ........................................................................ 144
Abstract

Well-defined front-rear cell polarity is essential for directional cell movement. The rod-shaped *Myxococcus xanthus* cells move using two motility systems and with defined front-rear polarity. Both systems are polarized, i.e. type IV pili assemble at the leading pole while the Agl/Glt gliding motility complexes assemble at the leading, translocate rearward to propel the cell, and disassemble at the lagging pole. During cellular reversals, which are induced by the Frz chemosensory system, polarity of the motility systems is inverted. The Ras-like GTPase MglA together with MglB, the cognate MglA GTPase activating protein (GAP) and the RomR response regulator constitute a module that determine front-rear polarity. MglA-GTP and MglB localize to and define the leading and lagging pole, respectively. MglA-GTP and MglB depend on polarly localized RomR for correct polar targeting. During the Frz system-induced reversals, MglA, MglB and RomR switch poles.

Here, using a comparative genomics approach together with experimental work, we identify RomX and RomY as integral components of the polarity module. RomX localizes asymmetrically to the poles with a large cluster at the lagging pole. *In vitro* data analyses demonstrated that RomX not only interacts directly with RomR alone and MglA-GTP alone but that RomR, RomX and MglA together form a heteromeric RomR/RomX/MglA-GTP complex. In this complex, RomX is sandwiched between RomR and MglA-GTP. Moreover, our data provide evidence that the RomR/RomX complex has MglA GEF activity. *In vivo*, polarly localized RomR recruits RomX and, in turn, the RomR/RomX complex recruits MglA-GTP to the leading cell pole. Thus, the RomR/RomX complex has dual functions in establishing front-rear polarity for motility in *M. xanthus*, it is a GEF that stimulates the accumulation of MglA-GTP, the active form of MglA, and it is a polar recruitment factor that recruits MglA-GTP to the leading cell pole. Both activities contribute to a high local concentration of MglA-GTP at this pole. At the leading pole, the RomR/RomX/MglA-GTP complex stimulates the assembly of Agl/Glt gliding motility complexes and is also incorporated into these complexes. However, in the absence of MglB, RomR/RomX is not essential for assembly of these complexes. Importantly, the Agl/Glt complexes assembled in the absence of RomR/RomX are less stable and transfer less directionally towards the lagging pole. These results suggest that RomR/RomX at the leading and MglB at the lagging cell pole establish leading lagging polarity axis for efficient gliding motility.
RomY localizes unipolarly with a cluster at the lagging cell pole. *In vivo* experiments demonstrated, that RomY regulates reversals and cell polarity similarly to MglB. Moreover, RomY localization depends on MglB suggesting a functional connection between these two proteins. Protein-protein interaction analyses suggested that RomY directly interacts with MglA and RomX. Moreover, RomX and RomR are not essential for gliding motility in the absence of RomY. Based on these data we suggest that RomY either stimulates MglB GAP activity or MglA GTPase activity.
Zusammenfassung


Abbreviations

ATP/ADP  adenosin tri-/diphosphate
BACTH  bacterial Adenylate Cyclase-based Two Hybrid
bp  base pair
BSA  bovine serum albumin
cAMP  3',5'-cyclic monophosphate
cDNA  single-stranded complementaty DNA
CTT  casitone Tris medium
CR  congo red
DNA  deoxyribonucleic acid
DMSO  dimethyl sulfoxide
DTT  dithiothreitol
EDTA  ethylenediaminetetraacetic acid
EPS  exopolysaccharides
GAP  GTPase activating proteins
GEF  guanine nucleotide exchange factor
GFP  green fluorescent protein
GTP/GDP  guanosine tri-/diphosphate
h  hours
HPK  histidine protein kinase
IM  innermembrane
IPTG  isopropyl β-D-1-thiogalaktopyranoside
kDa  kilodalton
LPS  lipopolysaccharides
MBP  maltose binding protein
mGTP/mGDP  (2’-(or-3’)-O-(N-Methylanthraniloyl) Guanosine Tri-/diphosphate
min  minutes
MOPS  3-(N-morpholino) propanesulfonic acid
OD  optical density
OM  outermembrane
PMF  proton motive force
pN  piconewton
SD  standard deviation
SDS-page  sodium dodecyl sulfate polyacrylamide gel electrophoresis
T4P  type IV pili
TB  trypan blue
TEMED  N,N,N',N'-Tetramethylethane-1,2-diamine
TIRF  total internal reflection fluorescence microscope
YFP  yellow fluorescent protein
WT  wild type
X-gal  5-Brom-4-chlor-3-indoxyl-β-D-galactopyranosid
1. Introduction

1.1 Spatial organization of bacterial cells

Until recently, bacterial cells were thought of as compartmented sacks of proteins localizing diffusely in the cytoplasm, inner membrane, periplasm or outer membrane. Development of microscopy techniques such as electron and fluorescence microscopy allowed studying in more details not only the morphology of bacterial cells but also the spatial organization of their content. The observation that bacterial chemoreceptors localize to the cell poles and that the division protein FtsZ localizes to the division site changed our thinking about bacteria, from unorganized sacks of proteins to highly spatially organized cells with proteins localizing to specific subcellular regions (Bi & Lutkenhaus, 1991, Alley et al., 1992, Maddock & Shapiro, 1993). Since then, many studies of the spatial organization of bacterial cells have shown that protein localization can be highly dynamic and change in response to external or internal signals or, alternatively, this localization is cell cycle dependent. Cell polarity with proteins localizing asymmetrically within a cell is important for many processes in bacteria including growth, division, cell cycle regulation, motility and signal transduction (Shapiro et al., 2009). Major questions in bacterial cell biology are how proteins find their correct localization and in some cases change this localization over time. Eukaryotic cells have developed sorting machineries based on vesicle transport that deliver protein cargo to the proper destinations. Bacterial cells lack vesicle sorting- and transport systems. Instead, protein localization in bacteria is typically mediated by a diffusion and capture mechanism. In this mechanism, proteins diffuse rapidly throughout the three dimensions of the cytoplasm or in the two dimensions of the membranes. These diffusing proteins can then recognize and bind to a cellular cue or specific landmark proteins.
Well studied examples of cellular cues for protein localization include geometrical cues and membrane lipids. An example of a protein that binds to a geometrical cue is DivIVA from *Bacillus subtilis*. The DivIVA protein localizes at bacterial cell poles and division sites by directly recognizing high negative curvature of a membrane (Lenarcic *et al*., 2009, Ramamurthi & Losick, 2009) (Figure 1, A). Proteins can also recognize positive curvature of the membrane. For instance, SpoVM recognizes and binds to membranes with positive curvature created during endospore formation in *B. subtilis* (Ramamurthi *et al*., 2009) (Figure 1, B). It has also been proposed that not only membrane curvature but also certain membrane lipids can serve as a spatial cue for protein localization in bacteria. Cardiolipin-rich domains have been identified at the poles and division sites of *B. subtilis* (Kawai *et al*., 2004) and *Escherichia coli* (Mileykovskaya & Dowhan, 2000). Localization of the mechanosensitive channel MscC and transporter ProP in *E. coli* correlates with subcellular cardiolipin composition (Romantsov *et al*., 2010). These observations suggest that cardiolipins may act as a recognition cue for certain proteins. However, it remains unknown whether this is due to direct interactions between a protein and cardiolipin.

Protein localization involving a landmark protein recruiting another protein is the best-studied mechanism leading to specific cellular protein localization. These landmark proteins can be divided into polymer-forming proteins and non-polymer-forming proteins. DivIVA is a well-studied example of a polymer-forming landmark protein. As mentioned, DivIVA binds to membrane regions with negative curvature. At the division site, DivIVA binds and directly interacts and recruits MinJ, that recruits the cell division inhibitory MinCD complex (Bramkamp *et al*., 2008, Patrick & Kearns, 2008, Gregory *et al*., 2008) (Figure 2, A).
Another example of polymer-forming proteins that function as landmarks are the bactofilins (Kuhn et al., 2010). In *Caulobacter crescentus* the bactofilins BacA and BacB in a cofactor independent manner polymerize in vitro and recruit the peptidoglycan synthase PbpC to the stalked pole in vivo (Kuhn et al., 2010). *Myxococcus xanthus* possesses four bactofilins, which also spontaneously polymerize in vitro to form long filaments (Kuhn et al., 2010, Koch et al., 2011, Bulyha et al., 2013). BacP localizes to large patches in the two subpolar regions and functions as a landmark for the small GTPase SofG (Bulyha et al., 2013). SofG, in turn, is important for the polar localization of the ATPases PilB and PilT that are important for type IV pili (T4P) function and motility (Bulyha et al., 2013)(Figure 2, B).
The integral membrane protein HubP from *Vibrio cholerae* is a well-studied example of a non-polymer forming landmark protein that localizes to the cell poles. The N-terminal peptidoglycan binding LysM domain is necessary and sufficient for HubP localization to the cell poles. HubP in turn, recruits and directly interacts with the ParA1 ATPase, which plays a role in chromosome 1 segregation, and with the ParA ATPase FlhG, which is involved in flagella assembly regulation (Yamaichi *et al.*, 2012, Fogel & Waldor, 2006, Correa *et al.*, 2005) (Figure 2, C). While it is relatively well-understood how polymer forming proteins such as DivIVA (Lenarcic *et al.*, 2009, Ramamurthi & Losick, 2009) and PopZ from *C. crescentus* (Bowman *et al.*, 2008, Ebersbach *et al.*, 2008) become localized, much less is known about how non-polymer forming landmarks eventually become localized.

1.2 Characteristics of small eukaryotic GTPases

In eukaryotes, small Ras-like GTPases, also known as small GTP-binding proteins, have important functions in protein localization and regulation of cell polarity (Jaffe & Hall, 2005, Charest & Firtel, 2007, Kortholt & van Haastert, 2008, Chiou *et al.*, 2017). Recently, small Ras-like GTPases have also emerged as being important for protein localization and regulation of cell polarity in bacteria (Schumacher & Søgaard-Andersen, 2017). Small Ras-like GTPases function as molecular switches. The GDP-bound form represents the inactive protein, while the GTP-bound form represents the active form and interacts with downstream effectors to activate downstream pathways. GDP as well as GTP are tightly bound to these GTPases and, moreover, their intrinsic GTPase activity is low (Bourne *et al.*, 1991). Generally, the activity of small GTPases is regulated by guanine nucleotide exchange factors (GEFs) and GTPase activating proteins (GAPs). GEFs promote GDP to GTP exchange in that way stimulating the accumulation of the active form. GAPs promote GTP hydrolysis in that way stimulating accumulation of the inactive form (Bos *et al.*, 2007) (Figure 3). Certain small GTPases are prenylated on their C-terminus, which provides the attachment of their active form to endomembranes. These GTPases can be displaced from the membranes by guanidine dissociation inhibitors (GDIs) that bind to the C-terminal lipid to maintain the small GTPase in an inactive complex (Cherfils & Zeghouf, 2013).
Figure 3. Regulation of small GTPases by GEF, GAP and GDI.

GEF catalyses the exchange of GDP to GTP. GAP activates GTPase activity and hydrolysis of GTP to GDP. GDI proteins affect nucleotide dissociation and inhibit GAP activity.

Structural characterization of GEFs revealed that GEFs share mechanistic hallmarks but they showed a stunning diversity in amino acid sequence and structure (Cherfils & Zeghouf, 2013). Nevertheless, GEFs of different families are thought to follow common reaction schemes (Bos et al., 2007). GEFs catalyze the dissociation of the nucleotide from the small GTPase by modifying the nucleotide-binding site leading to a decreased affinity for the nucleotide. In the current model for how GEFs function, the GEF first associates with the GDP-bound small GTPase. GDP dissociates from this complex leaving the GEF bound to the nucleotide-free GTPase. Next, the GTPase binds GTP, promoting GEF dissociation from the complex and leaving the GTPase in the active form (Vetter & Wittinghofer, 2001, Hodgkin & Kaiser, 1979). According to this model, a stable complex only exists between the GEF and the nucleotide-free GTPase. In vitro studies have also shown that the affinity of the small GTPase for GTP and GDP is similar and the GEF does not favor rebinding of GDP or GTP (Bos et al., 2007). Because the GTP concentration in vivo is higher than the GDP concentration, GTP binding to the nucleotide-free GTPase is favored. However, for some GEFs interactions with cognates GTPases in the GTP bound were shown. Detailed biochemical studies revealed that the interaction is part of a positive feedback loop and activates GEF, what in turn leads to accumulation of GTPase-GTP (Margarit et al., 2003, Menetrey et al., 2007, Richardson et al., 2012, Cohen et al., 2007, Chen et al., 2010, Lin et al., 2006).

Similar to the GEFs of small GTPases, the GAPs of these GTPases are structurally diverse and belong to different subfamilies (Bos et al., 2007). Structural studies of pairs of cognate GTPases and GAPs have revealed that the general mechanism of GTPase
stimulation of a GAP is to complete the GTPase active site by providing the so-called arginine finger into the GTPase active site (Rittinger et al., 1997, Scheffzek et al., 1997).

1.3 *Myxococcus xanthus* as a model organism

*M. xanthus* is a rod-shaped, Gram-negative soil bacterium. *M. xanthus* has a complex life cycle in which cells grow and divide in the presence of nutrients. If placed on a solid surface, *M. xanthus* cells move to form swarming colonies in the presence of nutrients. In the absence of nutrients, *M. xanthus* cells initiate a developmental program that leads to the formation of multicellular fruiting bodies inside which the rod-shaped motile cells differentiate to spherical spores. Cell motility and its regulation are required for formation of swarming colonies as well as for fruiting body formation (Kroos et al., 1988).

*M. xanthus* cells do not possess flagella, and thus they are not able to swim. However, they are able to move on a solid surface along their long axis. To facilitate motility, *M. xanthus* cells use two genetically independent motility systems, gliding, also referred to as adventurous (A) motility and type IV pili-dependent motility, also referred to as social (S) motility. The gliding motility system was initially referred to as adventurous because it allows single cell movement, while the type IV pili-dependent system was initially referred to as social because it is generally used by cells to move in groups (Hodgkin & Kaiser, 1979). Inactivation of one motility system, leaves cells motile by means of the second system, while inactivation of both systems leads to non-motile cells (Hodgkin & Kaiser, 1979). Additionally, *M. xanthus* cells change direction of movement on average every 10–15 minutes (Blackhart & Zusman, 1985). During these events, which are referred to as reversals, cells change polarity with the old lagging cell pole becoming the new leading cell pole and *vice versa*. 
1.4 Type IV pili motility

Type IV pili- (T4P) dependent motility in *M. xanthus* is similar to twitching motility in *Pseudomonas aeruginosa*. T4P are widespread among bacteria and play a role in natural transformation (Dubnau, 1999), pathogenesis (Craig & Li, 2008), biofilm formation (O'Toole & Kolter, 1998) and predation (Evans et al., 2007). In *M. xanthus*, T4P-dependent motility is favoured on wet and soft surfaces (Shi & Zusman, 1993) and depends on T4P (Kaiser, 1979) and exopolysaccharides (Yang et al., 2000). T4P extend from the leading cell pole, attach to a surface, and then retract, pulling a cell forward (Kaiser, 1979, Skerker & Berg, 2001, Zhou & Nan, 2017). During retractions, T4P generate a force up to 150 pN (Merz et al., 2000). In *M. xanthus*, retraction of T4P is thought to be stimulated by exopolysaccharides (Li et al., 2003).

T4P function depends on a conserved set of 11 proteins (Wall & Kaiser, 1999) (Figure 5, A). The T4P machinery (T4PM) that supports extension and retraction of T4P is a multi-layered structure that spans from the outer membrane to the cytoplasm (Chang et al., 2016). It consists of the PilQ secretin that forms an outer membrane pore for the pilus; TsaP that forms a periplasmic ring around PilQ; a mid-periplasmic ring formed by the periplasmic domains of PilQ and PilP; a lower periplasmic ring formed by PilO and
PilN that connect across the inner membrane to the PilM protein that forms a cytoplasmic ring. The inner membrane protein PilC form the cytoplasmic dome inside the PilM ring. The pilus is formed by PilA, the major subunit of the pilus (Chang et al., 2016) (Figure 5, B and C). The cytoplasmic ATPases PilB and PilT provide the energy for T4P extension and retraction, respectively (Jakovljevic et al., 2008) and associate and interact directly with PilC and PilM (Bischof et al., 2016, Chang et al., 2016). During T4P extension, PilA subunits are extracted from the outer leaflet of the inner membrane and inserted at the base of the growing pilus in a process that is powered by PilB (Jakovljevic et al., 2008, Chang et al., 2016). During retractions, the PilA subunits are removed from the base and reinserted into the inner membrane in a process powered by PilT (Jakovljevic et al., 2008, Chang et al., 2016).

Assembly of the T4PM starts from PilQ in the outer membrane (OM) and proceeds inwards (Friedrich et al., 2014). PilQ localization to the outer membrane depends on the Tgl pilotin (Friedrich et al., 2014) Tgl is an OM lipoprotein and in its absence, PilQ does not form oligomeric form. Next, PilP, PilN, PilO and PilM are incorporated into the T4PM and in parallel TsaP is incorporated. Following PilPNOM incorporation, PilC in the inner membrane is incorporated (Figure 6). Finally, PilB and PilT can associate with

**Figure 5. Genetic map of pili locus and architectural model of type IV pili complex.**

(A) Genetic map of pil locus. All the pil genes are clustered at the same locus, but not all pil genes are shown, tsaP is not a part of pil cluster. Figure modified from Friedrich et al., 2014.

(B) Schematic model of piliated (left) and non piliated (right) type IV pili machinery.

(C) Architectural models of piliated (left) and non piliated (right) type IV pili basal bodies. The colour code for the proteins is similar to the one used for the genes. B and C are reproduced from Chang et al., 2016.
the base of this machinery to stimulate extension or retraction of T4P (Friedrich et al., 2014).

Figure 6. Model of assembly of the *M. xanthus* type IV pili machinery.
Description in text. Colour code for the proteins as in Figure 5. Figure modified from Friedrich et al., 2014.

T4P assemble only at the leading cell pole in *M. xanthus* and during reversals, the pole at which T4P are assembled switches (Kaiser, 1979, Mignot et al., 2005). The mechanism underlying this switch in polarity of T4P depends on dynamic protein localization to the cell poles. T4PM proteins can be divided into two groups. One group includes the proteins that are localized to both cell poles and remain at the cell poles during a reversal (TsaP, PilQPNOCM). The second group is composed of the two ATPases PilB and PilT. PilB primarily localizes to the leading cell pole while PilT primarily localizes to the lagging cell pole (Friedrich et al., 2014, Nudleman et al., 2006, Bulyha et al., 2009). Notably, during a reversal, PilB and PilT are released from the cell poles and then associate with new leading and lagging cell pole, respectively. Therefore, the mechanism underlying T4P polar switching during a reversal involves dynamic localization of PilB and PilT. As described in more details below, the polarity regulation involved in sorting PilB and PilT to the correct cell poles involves the two small GTPases, MglA and SofG (Bulyha et al., 2013).

1.5 Gliding motility

Gliding motility, which is favoured on hard and dry surfaces, is generally used by *M. xanthus* to move as single cells (Shi & Zusman, 1993). The exact mechanism of
gliding motility in *M. xanthus* remains unclear. Bioinformatics together with *in vivo* protein localization and protein–protein interaction studies have revealed that the machinery involved in gliding motility also spans from the outer membrane to the cytoplasm.

### Cluster G1

![Diagram of G1 cluster genes](image)

Luciano *et al.* identified three genomic regions referred as G1, G2 and M1 that encode components of the gliding motility machinery (Figure 7) (Luciano *et al.*, 2011). In addition to the proteins encoded in these three clusters, the cytoplasmic protein AglZ (Yang *et al.*, 2004, Mignot *et al.*, 2007) together with the actin-like protein MreB (Mauriello *et al.*, 2010b, Treuner-Lange *et al.*, 2015) and the small GTPase MglA (Leonardy *et al.*, 2010, Zhang *et al.*, 2010) are important for gliding motility. The proteins encoded by the G1 and G2 clusters are thought to make up structural components of the gliding machinery (Luciano *et al.*, 2011, Jakobczak *et al.*, 2015, Faure *et al.*, 2016) and localize to the cell envelope and cytoplasm as shown in Figure 8. By contrast, the three proteins encoded by the M1 cluster make up the motor of the machinery and localize to the inner membrane (Sun *et al.*, 2011, Nan *et al.*, 2013) (Figure 8).
Experiments with carbonyl cyanide-m-chlorophenylhydrazone (CCCP) has shown that the proton motive force (PMF) is the source of energy for gliding motility in *M. xanthus*. The PMF arises from gradients in both the chemical potential energy, in the form of a pH difference across the inner membrane, and electrical potential energy, caused by a voltage difference across the inner membrane. Nigericin that reduces the pH gradient without changing membrane potential abolishes gliding motility whereas, valinomycin that destroys the membrane potential without changing the magnitude of the pH gradient has no effect on the gliding motility. Based on these data, it was concluded that the pH gradient across the membrane is a source of energy for gliding motility.

The M1 cluster encodes the component of a proton conducting channel: AglQ, AglR, AglS. Bioinformatics analysis shows that AglR is a TolQ/ExbB/MotA homolog, whereas AglQ and AglS are TolR/ExbD/MotB homologs. The MotA/MotB proteins build a stator part of the flagellar rotary motor, the TolQ/TolR complex is important for the outer membrane stability (Bernadac *et al.*, 1998) and the ExbB/ExbD proteins are involved in the activation of TonB-dependent transporters in the outer membrane (Postle, 2007). All three protein complexes form a proton channel in the inner membrane. Moreover, in all three systems energy from the proton flux can be converted to a mechanical output with a change in protein conformation.

Assembly and disassembly of the gliding motility machinery depend on the small Ras like GTPase MglA. MglA is absolutely required for both gliding and T4P-dependent
motility (Hodgkin & Kaiser, 1979) and function as a nucleotide-dependent molecular switch to stimulate motility (Mauriello et al., 2010b, Leonardy et al., 2010, Patryn et al., 2010, Zhang et al., 2010, Miertzschke et al., 2011). MglA in its active GTP-bound form is incorporated into the gliding motility complexes (Treuner-Lange et al., 2015). MglA-GTP interacts directly interact with MreB (Mauriello et al., 2010b, Treuner-Lange et al., 2015). MglA also interacts directly with AglZ; however, it is not known whether this interact is nucleotide-dependent (Mauriello et al., 2010b). MglA, MreB and AglZ are thought to form a complex that interacts with GltI in the cytoplasm in that way connecting to the gliding motility machinery (Treuner-Lange et al., 2015, Faure et al., 2016). Also, the MglA, MreB and AglZ complex stimulates formation of the gliding motility complexes at the leading cell pole (Treuner-Lange et al., 2015) and MglA-GTP has also been suggested to regulate directionality of the gliding machinery (Nan et al., 2011).

1.5.5 Two models of gliding motility

Two models have been proposed to explain how the gliding motility complex assembles to generate movement, the motor cargo model and the focal adhesion model (Figure 9).

**Figure 9. Schematic of the two models of the gliding motility machinery.**

(A) Motor cargo model in which motor proteins (green dots) tracking on a helical cytoskeleton deform peptidoglycan and OM in order to create traction.  
(B) Model of focal adhesion mechanism in which multi-protein complexes (green dots) span the inner and outer membrane and attach to the substrate.
1.5.5.1 Motor cargo complex

The motor cargo complex model was proposed based on the localization of GltD protein (Nan et al., 2011). GltD was proposed to localize to both the cytoplasm and periplasm (Nan et al., 2010). 3D reconstructions of GltD-mCherry fluorescence showed that GltD-mCherry forms a twisted endless looped helix that spans the length of the cells (Figure 10). Additionally, the distance between adjacent nodes is nearly identical to that of MreB helices, 0.45±0.09 and 0.47±0.1 µm, respectively (Nan et al., 2011, Mauriello et al., 2010b).

Figure 10. GltD forms a twisted endless looped helix.
3D reconstruction of the GltD helix from three individual cells. Scale bar: 1 µm. Figure reproduced from Nan et al., 2011.

Moreover, the GltD helix rotates as cells move on a 1.5% agar surface and changes direction of rotation when cells reverse their direction. Furthermore, this helix rotates clockwise when viewed from the lagging cell pole. In addition, rotation depends on the PMF and polymerization of MreB. The calculated linear velocity of the GltD helix (4.4-9.6 µm/min) is in agreement with the maximum velocity of a gliding cell (≈2-4 µm/min) (Sun et al., 1999). GltD localizes with a higher concentration at the leading cell pole and when the cell reverses, GltD relocalizes to the new leading cell pole. AglR also forms a helix that spans the length of the cells (Figure 11) (Nan et al., 2013). The pitch of the AglR helix (1.34±0.51 µm) is similar to that of GltD. Moreover, the velocity of the rotating helices of AglR and GltD is similar. By tracking AglR, Nan et al. (2013) discovered that AglR moves along the cell width and cell lengths in zigzag trajectories in two dimensions suggesting that AglR molecules move in helical trajectories in three dimensions (Nan et al., 2013). Additionally, AglR molecules tended to slow down near the ventral side of cells, i.e. the part of the cell surface where a cell is in contact with the substratum during gliding. Slowing down of the motor units depends on the hardness of the surface suggesting that the motor units slow down due to resistance of the underlying surface. Like for GltD, AglR movement depends on PMF and MreB polymerization.
Additionally, helical movement of AglR depends on GltD and in the absence of GltD, helical movement of AglR is replaced by linear motion along the long axis of the cell. By contrast, in the absence of AglZ, AglR moves faster but loses its directionality (Nan et al., 2013). Finally, TIRF images of cells expressing cytoplasmic GFP placed on glass microscope slide revealed intensity variations similar to the periodicity of MreB and GltD with a period of 0.83±0.23 µm (Nan et al., 2011).

Figure 11. AglR forms a twisted endless helix.
Structured illumination microscopy of AglR-pamCherry in two fixed cells. For each cells, the area in which void fields are covered with helical fluorescence signal. Helical tracks are shown on a magnified section of each cell. Scale bar: 1 µm. Figure reproduced from Nan et al., 2011.

Based on these data the motor cargo model was proposed. In this model, the motor proteins (AglQRS) together with other proteins important for gliding motility are hypothesized to form a complex and move on an endless looped helix that spans the length of cells. When these complexes are at the ventral side of a cell, it is speculated that they slow down because of increased resistance, in that way creating nearly stationary clusters along the cell length. At the positions where these clusters form, they are thought to push on and distort the cell envelope resulting in directed cell movement (Figure 9, A).

1.5.5.2 The focal adhesion complexes model

The focal adhesion model was proposed by Mignot et al. and is based on AglZ localization. In fully motile cells, AglZ localizes at the leading cell pole and in ordered
clusters spanning the cell length. As cells move forward, AglZ clusters stay in fixed positions with respect to the substratum. The only clusters that move with a cell are those at the leading cell pole. Moreover, clusters disassembled close to the lagging cell pole (Mignot et al., 2007) (Figure 12).

**Figure 12.** AglZ localizes in clusters along the cell length and stay in fixed position relative to the substratum in moving cells.
AglZ-YFP localization in moving cell. White arrowheads highlight the position of stationary AglZ-YFP clusters. Scale bar: 1 µm. Figure reproduced from Mignot et al., 2007.

The number of AglZ clusters correlates with cell length (Mignot et al., 2007) and the localization of AglZ clusters correlates with the site of force generation (Sun et al., 2011). Importantly, cephalaxin treated cells, which elongate up to 10 times of normal length, has almost normal gliding motility suggesting that the force for gliding is distributed along the cell body. Based on these observations, it was suggested that the force for motility is generated at the sites of the AglZ clusters (Mignot et al., 2007). Because the clusters do not move relative to the substratum they were referred to as focal adhesion complexes (Mignot et al., 2007) by analogy to eukaryotic focal adhesion complexes (Hoiczyk & Baumeister, 1998).

Since the initial model was proposed for focal adhesion complexes as the site of force generation for gliding motility, most Glt and Agl proteins have been shown to (co-)localize in focal adhesion complexes (Mignot et al., 2007, Nan et al., 2010, Sun et al., 2011, Jakobczak et al., 2015, Treuner-Lange et al., 2015, Faure et al., 2016). In the current model for assembly of the Agl/Glt complexes for gliding motility, assembly occurs at the leading cell pole leading to formation of the complex that spans from the cytoplasm to the outer membrane (Figure 9, B). The assembly is stimulated by the AglZ/MglA/MreB complex. Upon assembly, the complex adheres to the substratum and generates a force that propels a cell forward. As an Agl/Glt gliding motility complex approaches the lagging cell pole, it disassembles (see details below). Interestingly, recent
data suggest that the Agl/Glt complexes do not only move between poles but also move across the cell width following a helical path. In agreement with counterclockwise rotation of AglZ clusters, Faure et al. have found that a moving cell is rotating clockwise during cell propulsion and angular velocity is proportional to the speed of the cell (Faure et al., 2016). These findings suggest that surface adhesion of a focal adhesion complex leads to clockwise rotation and forward drive of the cell.

Finally, Balagam et al. simulated a head-to-side collision between two cells moving on crossing paths with parameters fitted to the motor cargo and the focal adhesion complexes model. In the simulation for the motor cargo model both cells changed direction. By contrast, in the focal adhesion model, the new common direction was the same as the direction of the primary cell before collision. To identify which of the two scenarios simulate the behaviour of colliding M. xanthus cells, experiments with colliding cells were performed. In the experiments, the direction of the primary cells did not change after collision. This behaviour is similar to the simulations of the focal adhesion model of gliding motility. Thus, comparison of simulations with experimental observation indicates that the focal adhesion complexes model of the gliding motility is correct (Balagam et al., 2014). From hereon, I discuss gliding motility within the framework of the focal adhesion complex model.

1.6 Slime

Gliding M. xanthus cells deposit slime. The composition of slime is not known in details; however, it can be stained with fluorescently labelled concavalin A demonstrating that it contains polysaccharide (Ducret et al., 2012). Slime was proposed to be secreted at the focal adhesion sites to promote adhesion of focal adhesion complexes to the substrate. However, cells that lack gliding motility (aglQ, gltD and gltE mutants) still deposit slime underneath the cell body, showing that slime export to the cell surface and deposition do not require an active Agl/Glt machinery. Furthermore, a wza0 mutant, in which all of putative exopolisacharides (EPS) export machineries are lacking, a difA mutant, which is reduced in EPS production, and a sasA mutant, which is reduced in LPS O-antigen synthesis, are still moving by gliding motility and deposit slime. Therefore, these results suggest that the polysaccharide component of slime is neither EPS nor LPS (Ducret et al., 2012). Similarly, the mechanism involved in slime synthesis remains unknown.
1.7 Cell polarity and motility regulation – spatial control of motility

Both motility systems in *M. xanthus* are polarized with T4P assembling only at the leading cell pole and focal adhesion complexes assembling at the leading and disassembling at the lagging cell pole. Moreover, this polarity is dynamic and changes during reversals. A protein module consisting of the Ras-like GTPase MglA, its cognate GTPase activating protein (GAP) MglB, and the response regulator RomR establishes polarity of the two motility systems, while signaling by the Frz chemosensory system inverses the polarity (Leonardy *et al.*, 2010, Zhang *et al.*, 2010, Miertzschke *et al.*, 2011, Patryn *et al.*, 2010, Keilberg *et al.*, 2012, Zhang *et al.*, 2012).

1.7.1 Spatial organization of cell polarity by MglA, MglB and RomR

MglA is absolutely essential for movement by both motility system in *M. xanthus* (Hartzell & Kaiser, 1991) and functions as a nucleotide-dependent molecular switch to stimulate motility (Leonardy *et al.*, 2010, Zhang *et al.*, 2010, Miertzschke *et al.*, 2011, Patryn *et al.*, 2010). As other members of the Ras superfamily of small GTPases, MglA is active and stimulates motility in its GTP-bound form and is inactive in the GDP-bound state. Patryn *et al.* reported a $K_D$ of 11.8 µM for GTP and 9.6 µM for GDP (Patryn *et al.*, 2010). Using MglATt from *Thermus thermophilus*, which is 62/81% identical/similar to MglAMx from *M. xanthus*, Leonardy *et al.* reported $K_D$'s of 24.6 nM and 7.3 nM for GTP and GDP, respectively (Leonardy *et al.*, 2010). Moreover, Miertzschke *et al.* suggested that the affinity of MglATt for GTP could be even 100-fold lower than for GDP (Miertzschke *et al.*, 2011). Nevertheless, in the current model for regulation of MglA activity, it is hypothesized that GDP to GTP exchange would be stimulated by a so far unknown guanine nucleotide exchange factor (GEF) (Figure 13).

![Figure 13. GTPase cycle of MglA.](image)

Figure reproduced from Schumacher & Søgaard-Andersen, 2017.
MglB is not essential for motility per se but important for regulation of motility (Leonardy et al., 2010, Zhang et al., 2010). Structural analyses of MglB from *T. thermophilus* demonstrated that MglB is a Roadblock/LC7 protein (Miertzschke et al., 2011). Moreover, the crystal structure of an MglA–MglB complex revealed that they interact with a 1:2 stoichiometry, and in contrast to other Ras-like GTPases and their cognate GAPs, no residues from MglB reach into the active site of MglA, confirming the absence of any conserved potential catalytic residue (Miertzschke et al., 2011) (Figure 14).

![Figure 14. The MglA-MglB complex.](image)

Structure of MglA-GppNHp (blue) bound to the MglB (green). Dotted lines show flexible loops that were not visible in electron density. Figure reproduced from Miertzschke et al., 2011.

RomR is essential for gliding motility and important for T4P-dependent motility (Zhang et al., 2012, Keilberg et al., 2012). Sequence analysis showed that RomR possesses an N-terminal receiver domain and C-terminal output domain. The output domain of RomR contains a conserved C-terminal region (RomR-C) and an unstructured proline-rich region that links the two conserved regions (Keilberg et al., 2012) (Figure 15).

![Figure 15. Schematic representation of RomR.](image)

RomR contains three distinct regions: N-terminal response regulator domain (REC), an unstructured proline rich linker domain (Linker), and a conserved C-terminal part (RomR-C). Numbers correspond to amino acid sequence of RomR. Figure reproduced from Keilberg et al., 2012.
MglA, MglB as well as RomR localize dynamically to the cell poles (Figure 16). In the active GTP-bound form, MglA localizes to the leading cell pole while the inactive form MglA-GDP is diffusely localized throughout the cytoplasm. MglB localizes primarily at the lagging cell pole. Finally, RomR localizes in a bipolar asymmetric pattern with the bigger cluster at the lagging cell pole (Zhang et al., 2012, Keilberg et al., 2012). RomR has been suggested to have two pole targeting determinants, one is the linker region in the output domain and the second is RomR-C (Figure 15) (Keilberg et al., 2012). Furthermore, both regions are necessary for motility (Keilberg et al., 2012). The RomR receiver domain failed to localize to the poles but has been suggested to be important for cellular reversals (Leonardy et al., 2007).

![Figure 16. Model for spatial dynamic regulation of motility in M. xanthus.](image)

Localization of MglA, MglB and RomR at the leading and lagging cell poles before and after a cellular reversal.

Localization of MglA, MglB are RomR is mutually dependent. In the absence of MglB, MglA as well as RomR localize in bipolar symmetric patterns. Similar bipolar localization was observed for MglA locked in the GTP-bound form. In the absence of MglA, MglB as well as RomR localize unipolarly (Zhang et al., 2010, Keilberg et al., 2012, Zhang et al., 2012, Leonardy et al., 2010). Importantly, localization studies of an MglA variant locked in the GTP-bound form have shown that RomR is polar targeting determinant for MglA-GTP (Keilberg et al., 2012, Zhang et al., 2012). Finally, in the absence of RomR, MglA localizes diffusely to the cytoplasm and MglB becomes more unipolar. MglA, MglB and RomR are not only mutually dependent on each other for localization but also directly interact (Miertzschke et al., 2011, Keilberg et al., 2012). How the three proteins become asymmetrically localized to the cell poles is not well-
understood except that (1) MglB excludes MglA-GTP from the lagging cell pole by stimulating the conversion of MglA-GTP to MglA-GDP at this pole; and, (2) RomR is a polar targeting determinant of MglA-GTP at the leading cell pole (Patryn et al., 2010, Leonardy et al., 2010, Zhang et al., 2010, Miertzschke et al., 2011, Keilberg et al., 2012, Zhang et al., 2012).

During the Frz-induced reversals, MglA-GTP, MglB and RomR are released from their respective poles and rebind to the relevant opposite cell poles (Figure 16) (Leonardy et al., 2007, Keilberg et al., 2012, Zhang et al., 2012, Zhang et al., 2010, Leonardy et al., 2010). Time-lapse microscopy analyses have suggested that MglA is the first protein to be released and rebind to the new leading cell pole (Leonardy et al., 2010, Zhang et al., 2010). These observations were confirmed by colocalization studies in which it was observed that MglA and MglB colocalize at the old lagging cell pole immediately before cells reverse (Zhang et al., 2010).

The output of the MglA/MglB/RomR module is generated by MglA-GTP (Zhang et al., 2012, Keilberg et al., 2012). Moreover, deletion of mglB restored gliding in a ΔromR strain (Keilberg et al., 2012). Additionally, MglA locked in the GTP form restored gliding in a romR mutant (Keilberg et al., 2012). How MglA-GTP stimulates motility at the leading pole is not understood for the T4P-dependent system except that correct polar localization of PilB, PilT and FrzS depends on MglA-GTP (Bulyha et al., 2013, Zhang et al., 2012). In the case of the gliding motility system, MglA-GTP at the leading cell pole interacts with AglZ and MreB to stimulate assembly of the Agl/Glt motility machinery in that way stimulating formation of the focal adhesion complexes (Treuner-Lange et al., 2015). MglA-GTP not only stimulates formation of these complexes but is also incorporated into them. By incorporating MglA-GTP into the gliding motility complexes, these complexes become sensitive to the MglB GAP activity at the lagging pole and disassemble at this cell pole. Consistently, a ΔmglB mutant and a strain containing MglA locked in the GTP-bound form hyperreverse because the gliding motility complexes are not disassembled at the lagging cell pole (Leonardy et al., 2010, Zhang et al., 2010, Miertzschke et al., 2011, Treuner-Lange et al., 2015).

**1.7.2 Frz chemosensory system**

The reversal frequency is regulated by the Frz chemosensory system (Blackhart & Zusman, 1985). The components of this signal transduction system show high similarity to the components of chemosensory systems involved in chemotaxis in other bacteria.
(McBride et al., 1989, Trudeau et al., 1996) (Figure 17). In details, the Frz system consists of the following components: FrzCD, a cytoplasmic MCP (methyl-accepting chemotaxis protein) homolog, two CheW homologs, FrzA and FrzB, a methyltransferase FrzF, which methylates FrzCD, a methylesterase FrzG, which demethylases FrzCD, a CheA histidine kinase with a CheY-like receiver domain, FrzE, and a response regulator with two CheY-like receiver domains, FrzZ (Figure 17). The specific signals that induce cellular reversals are not known. It has been proposed that the intercellular C-signal, which is important for development, (Jelsbak & Søgaard-Andersen, 2002, Søgaard-Andersen & Kaiser, 1996) and exopolysaccharide (Zhou & Nan, 2017) may regulate Frz activity. However, it is not know how these extracellular compounds would regulate Frz activity in the cytoplasm. Current data suggests that these signal(s) could be sensed by FrzCD or by FrzF (Scott et al., 2008, Bustamante et al., 2004).

Figure 17. Schematic representation of the Frz pathway from M. xanthus.
Frz proteins are blue with the core Frz proteins (FrzCD, FrzA and FrzE kinase domain) shown in dark blue. Experimentally confirmed phosphotransfers from FrzE kinase are indicated with black arrows. FrzE is suggested to phosphorylated FrzE$^{CheY}$ at low signalling level (thin line) and FrzZ at high signalling level (thick line) Phosphorylated FrzE$^{CheY}$ inhibits FrzE kinase activity. Phosphorylated FrzZ inhibits phosphotransfer to FrzE$^{CheY}$, stimulates kinase activity or both (dashed lines). FrzE may phosphorylate RomR (green pentagon) or not known response regulator (purple pentagon) at the higher signalling level (thick grey arrows). Coloured arrows at the bottom show the Frz output. Proteins labelled with single letters have the Frz prefix. Figure reproduced from Schumacher & Søgaard-Andersen, 2017.
Upon receiving a signal, FrzE autophosphorylates at a conserved histidine residue in the histidine phosphotransfer (Hpt) domain (Inclan et al., 2007, Inclan et al., 2008). In vitro experiments showed direct phosphotransfer from FrzE Hpt domain to both CheY-like domains of the FrzZ (Inclan et al., 2007). Inclan et al. suggested that FrzE CheY-like domain inhibits autophosphorylation of the Hpt domain. Alternatively, the FrzE CheY-like domain may act as a phosphosink for the kinase (Inclan et al., 2008). Further studies on FrzZ confirmed in vitro phosphorylation of both CheY-like domain, with Asp52 as the preferential site of phosphorylation. Moreover, the reversal frequency is directly correlated with the amount of phosphorylated FrzZ present in a cell with a higher pool of phospho-FrzZ correlating with a higher reversal frequency (Kaimer & Zusman, 2013). Kaimer & Zusman also showed that FrzZ phosphorylation depends on cell-surface contact (Kaimer & Zusman, 2013). Furthermore, FrzZ phosphorylation is independent of cell movement, i.e. cells with defective gliding and T4P-dependent motility still accumulate phospho-FrzZ. Phospho-FrzZ localizes to the leading cell pole and its localization depends on MglA and FrzE CheY-like domain (Kaimer & Zusman, 2013, Kaimer & Zusman, 2016).

Interestingly, FrzE overexpression resulted in inhibition of FrzZ phosphorylation and reduced reversals (Kaimer & Zusman, 2016). This inhibitory effect of FrzE is mediated by the C-terminal CheY-like domain (Kaimer & Zusman, 2016). Furthermore, substitution of the conserved Asp residue (Asp709) in the FrzE CheY-like domain to a non-phosphorylatable amino acid residue (FrzE<sup>D709A</sup>) resulted in hyper-reversing cells and a high level of FrzZ phosphorylation independent of the FrzE<sup>D709A</sup> level (Kaimer & Zusman, 2016). This indicates that phosphorylation of the CheY-like domain in FrzE is required for this inhibitory effect (Kaimer & Zusman, 2016). It was suggested that FrzZ transfers the signal from the Frz system to the downstream effectors (Kaimer & Zusman, 2013). Surprisingly, a double mutant <i>frzZ frzE<sup>D709A</sup></i> showed wild type reversal behaviour (Kaimer & Zusman, 2016). Furthermore, a <i>frzZ</i> mutant was still able to respond to isoamyl alcohol, which is an artificial activator of the Frz system, unlike a kinase (<i>frzE</i>) or MCP (<i>frzCD</i>) mutant, suggesting that FrzZ is not required for signal transduction under these conditions. Observation of cells moving by T4P-dependent motility revealed rare reversals in a <i>frzZ</i> mutant, suggesting that FrzZ is important for reversals in the T4P-dependent motility system but not strictly required (Guzzo et al., 2015). In the current model for the Frz system induced reversals, FrzZ acts as a stimulator of reversals. Moreover, the response regulator RomR (see details below) and/or a yet unknown
response regulator would contribute to transducing the reversal signal to the MglA/MglB/RomR module (Figure 17). The output response of the Frz system is a reversal. At the cellular, this involves an inversion of the leading-lagging cell pole with the relocation of MglA, MglB and RomR. How the Frz system connects to the MglA/MglB/RomR polarity module remains unknown. Of note, RomR was proposed as a connector between Frz and the polarity module (Guzzo et al., 2015, Leonardy et al., 2007, Keilberg et al., 2012) because a RomR variant that mimics the phosphorylated form of RomR (RomR<sup>DS3E</sup>) caused a hyper-reversing and RomR with blocked phosphorylation (RomR<sup>DS3N</sup>) a hypo-reversing phenotype. Moreover, the effect of the RomR<sup>DS3E</sup> variant was observed in cells lacking FrzZ or several components of the Frz system (Leonardy et al., 2007, Keilberg et al., 2012). However, until now a kinase involved in RomR phosphorylation has not been identified.

1.7.3 Regulation of cell polarity by MglC, an MglB homolog

Most mglA genes are located next to an mglB gene (Wuichet & Søgaard-Andersen, 2014). However, M. xanthus encodes an orphan paralog of MglB, named MglC (McLoon et al., 2015). MglB and MglC from M. xanthus only share 8%/17% identity/similarity. However, structural prediction of MglC showed that MglC is predicted to have Roadblock/LC7 fold similar to the MglB homolog from T. thermophilus (McLoon et al., 2015, Miertzschke et al., 2011). In-frame deletion of mglC causes a defect in both motility systems. In the case of the gliding motility system, this defect was traced down to a defect in reversals while it still remains unknown whether the defect in T4P-dependent motility is caused by a reversal defect. Epistasis analysis have shown that MglC acts in the same pathway as MglA, MglB and RomR. MglC localizes predominantly at the lagging cell pole with occasional localization of a smaller cluster at the leading cell pole. Localization of MglC is dynamic and switches pole during a cellular reversal. In the absence of MglA, localization of MglC shows highly asymmetric pattern. In the ΔmglB mutant, MglC shows predominantly bipolar symmetric localization. Thus, MglB is not necessary for polar MglC localization but is important for correct polar asymmetry. In contrast, in the absence of RomR, MglC shows diffuse localization. Thus, MglC polar localization completely depends on RomR. Analysis of direct protein-protein interaction have shown that MglC directly interacts with MglB and RomR. Because of the opposite reversal frequencies of ΔmglB and ΔmglC strains and the direct MglB-MglC interaction, it was hypothesized that MglC could act as an inhibitor of the GAP activity of MglB on MglA.
However, in a GTPase assay no inhibitory effect of MglC on the GAP activity of MglB was observed (McLoon et al., 2015). It remains unknown how MglC regulates reversals. However, it was suggested that by interacting with both, RomR and MglB, MglC could function between Frz system and RomR or between RomR and MglB.

1.7.4 Regulation of motility by the small GTPase SofG

As described, PilB and PilT are polarly localized (Bulyha et al., 2009) and sorted to the opposite cell poles by MglA (Bulyha et al., 2013). However, MglA is not necessary for the PilB and PilT polar localization. PilB and PilT polar localization is also independent of the stationary T4PM components (Friedrich et al., 2014).

Two or more small GTPases often function in parallel or in the same pathway to regulate polarity or motility in eukaryotes (Charest & Firtel, 2007, Heasman & Ridley, 2008). Based on these observations, Bulyha et al. identified an orphan small GTPase, named SofG, that regulates T4P-dependent motility. SofG shows 34% identity and 50% similarity to MglA and, similarly to MglA, contains an intrinsic Arg finger, which is important for its function. So far, a GAP and a GEF for SofG have not been identified (Bulyha et al., 2013).

SofG itself localizes in a subpolar cluster and is important for polar PilB and PilT localization (Bulyha et al., 2013). Moreover, SofG localization depends on BacP (Bulyha et al., 2013), a bactofilin cytoskeletal protein that polymerizes in vitro independently of any cofactors and in vivo localizes to two subpolar patches (Kuhn et al., 2010). The SofG cluster is highly dynamic and shuttles back and forth between the subpolar localization and the cell pole over one of the BacP patches. Importantly, dynamics of the SofG cluster depends on GTP hydrolysis, SofG variant that is impaired in GTP hydrolysis does not show dynamics of the WT protein. Additionally, SofG GTP hydrolysis is important for the polar localization of PilT and PilB (Bulyha et al., 2013).
Based on these results Bulyha et al. (2013) proposed a model in which the dynamic localization of PilB and PilT is regulated by a cascade of two small GTPases. First, SofG localizes to one of the BacP patches. This subpolar SofG cluster shuttles and brings PilT and PilB to the same pole. Next, MglA sorts PilB and PilT to the opposite cell poles and sets up correct T4P asymmetry (Bulyha et al., 2013) (Figure 18).

1.8 Scope of this study

*M. xanthus* is a model organism for studying bacterial multicellularity, cell-cell communication, development and motility. Regulation of motility combines a typical bacterial chemosensory system (Frz system) with a polarity module (MglA/MglB/RomR) that incorporates typical eukaryotic components for polarity. However, it is not known how the Frz system connects to the MglA/MglB/RomR polarity module. Similarly, we
lack a clear understanding of the importance of the MglA GTPase cycle because an MglA GEF has not been identified so far. In this study, by using a comparative genomic approach, we identified RomX and RomY as candidate components of the polarity module. By using fluorescence live-cell imaging, protein-protein interaction analyses, \textit{in vitro} analyses of proteins, we demonstrate that RomX and RomY are integral components of the polarity module.
2. Results

2.1. Identification of RomX and RomY

*romR* has a limited genomic distribution and, generally, co-occurs with MglA and MglB (Keilberg *et al.*, 2012). Therefore, in order to identify novel proteins that play a role in regulation of polarity and motility, we searched for proteins with the same genomic distribution as *romR* in 1609 completely sequenced prokaryotic genomes (personal communication, Kristin Wuichet).

![Figure 19. RomX and RomY have the same genomic distribution as RomR.](image)

Each column represents the presence (color) or absence (white) of MglA, MglB, RomR, RomX, RomY and Frz system (Kristin Wuichet, personal communication).

The analysis revealed two proteins, MXAN_3350 (from hereon RomX) and MXAN_5749 (from hereon RomY) that co-occur with RomR. *romX* and *romY* are conserved in 28 out of 31 genomes containing *romR* (Figure 19). RomX is encoded in 28 out of 31 genomes encoding RomR and in one genome that does not encode RomR (Figure 19). Seven genomes encode two RomX homologs. Among the genomes that
encode RomX as well as RomR, one does not encode MglA and MglB and one does not encode MglB. By contrast, \textit{romY} only co-occurs with \textit{romR}. Moreover, all genomes that encode RomR and RomY also encode MglA, indicating a possible connection between RomR, RomY and MglA.

\textit{romX} and \textit{romY} were not identified in genetic screens to identify regulators of motility or reversals. However, based on the bioinformatic analyses, we hypothesized that RomX and RomY are new motility regulatory proteins.

2.2. Characterization of RomX in \textit{Myxococcus xanthus}

All 36 RomX homologs share a domain that does not match any currently characterized domain models. Seven genomes encode two RomX homologs (Figure 20, A). In the case of these proteins, one of the paralogs contain an N-terminal response regulator receiver domain in addition to the conserved RomX domain.
Figure 20. Bioinformatics analysis of RomX

(A) Alignment of identified RomX homologs. Gene accession number showed in the same colour are coming from the same species.

(B) Conservation of \textit{romX} neighbourhood in selected myxobacterial genomes. Genes are depicted as arrows. Arrow orientation indicates coding direction. Homologous proteins are indicated in same colours. Numbers in arrows in \textit{M. xanthus} indicate gene number. Numbers in arrows (outside of \textit{M. xanthus}) indicate identity and similarity of their encoded protein to their \textit{M. xanthus} homolog. Numbers were obtained from Pairwise Sequence Alignment of protein using EMBOSS Needle. Right part shows predicted proteins functions based on the NCBI BLAST and conserved domain analysis (Marchler-Bauer \textit{et al.}, 2017).
The *romX* locus is conserved in myxobacterial genomes (Figure 20, B). *rluA* in *M. xanthus*, which is predicted to encode a pseudouridine synthase, was identified in a transposon mutagenesis screen as potentially important for gliding motility (Youderian *et al*., 2003); however, none of the remaining genes flanking *romX* have been implicated in motility.

### 2.2.1 RomX is important for both motility systems

To test the role of *romX* in motility, we generated an in-frame deletion of *romX*. To evaluate T4P-dependent and gliding motility in the *romX* mutant, motility was tested on soft (0.5%) agar, which is favorable to T4P-dependent motility, and hard (1.5%) agar, which is favorable to gliding motility. T4P-dependent motility is evident by the formation of flares at the colony edge. Gliding motility is evident by the presence of single cells at the edge of a colony.
Figure 21. ΔromX mutant shows defect in T4P-dependent motility, no gliding motility and normal EPS accumulation.

(A) Motility assays showing colonies of indicated mutants after 24 hours incubation on agar plates favoring T4P-dependent motility (0.5% agar) and gliding motility (1.5% agar), respectively. Bars, 1000 μm (T4P-dependent motility), 500μm (gliding motility – left panel) and 50 μm (gliding motility – right panel).

(B) Genomic neighborhood of romX. Genes are depicted as arrows. Arrow orientation indicates coding direction. Numbers below indicate distance between genes. Arrow above shows genomic fragment used as a promoter for the ectopic expression of romX.
(C) Immunoblot of RomX accumulation. Cells were grown in liquid culture, harvested, and total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using α-RomX. Calculated molecular masses of RomX and RomX-YFP are indicated. Immunoblot with α-PilC serves as loading control.

(D) EPS accumulation in WT and selected mutants. Aliquots of 20 µl cell suspensions at 7 × 10^9 cells/ml were spotted on 0.5% agar supplemented with 0.5% CTT and 40 µg/ml congo red (Cr) or 20 µg/ml trypan blue (Tb) and incubated at 32 °C for 24 hours.

WT formed the flares characteristic of T4P-dependent motility on 0.5% agar. The ΔpilA mutant that lacks the major pilin and therefore does not display T4P-dependent motility served as a negative control. The ΔromX mutant formed less and shorter flares in comparison to WT. In the case of gliding motility, WT showed single cells at the colony edge on hard agar while the ΔaglQ mutant, which lacks an essential component of the gliding machinery and served as a negative control, did not. The ΔromX mutant only displayed small groups of cells at the colony edge but no single cells. To more carefully test for defects in the two motility systems, we inactivated pilA or aglQ in the ΔromX mutant. The double deletion strain ΔromXΔaglQ on 0.5% agar formed short flares that were undistinguishable from the flares of the ΔromX mutant. Surprisingly, the ΔromXΔaglQ mutant still moved on 1.5% agar in a similar fashion to the ΔromX mutant. We hypothesize that this movement is performed by T4P-dependent motility. The ΔpilAΔromX double mutant showed a flat colony edge under both conditions (Figure 21, A). Because RomX was identified as co-occurring with RomR, we compared the motility characteristics of the three ΔromX mutants to that of the corresponding ΔromR mutants. The three ΔromX mutants and the three ΔromR mutants have similar motility phenotypes (Figure 21, A).

To confirm that the motility defects in the ΔromX mutant was caused by lack of RomX, complementation strains in which romX and romX-YFP were ectopically expressed under the control of the native promoter (Figure 21, B) were generated. α-RomX antibodies failed to detect a protein of the calculated size of RomX (10.3 kDa) in immunoblot analysis of total cell extracts from WT and the complementation strain expressing romX ectopically from the Mx8 attB site; by contrast, these antibodies detected the ectopically expressed RomX-YFP protein (Figure 21, C) suggesting that this protein is either overexpressed or that native RomX cannot be detected by immunoblotting due to its small size. Both complementation strains displayed WT motility (Figure 21, A). Based on these experiments, we conclude that RomX is important for T4P-dependent motility and necessary for gliding motility.

Defects in T4P-dependent motility can be caused by lack of EPS accumulation (Li et al., 2003, Lu et al., 2005). To test the possibility that the defect in T4P-dependent
motility in the ΔromX mutant was caused by lack of EPS, EPS accumulation was checked using colorimetric assays. Cells were inoculated on solid medium containing nutrients and supplemented with congo red or trypan blue. We observed no difference in congo red and trypan blue binding for WT and the ΔromX mutant. In this experiment, the ΔpilA mutant served as a negative control and no binding of congo red and trypan blue was observed for this strain (Figure 21, D) (Black et al., 2006). Thus, the defect in T4P-dependent motility in the ΔromX strain is not caused by lack of EPS accumulation. As expected, the ΔromR mutant also bound congo red and trypan blue similarly as WT (Figure 21, D). Thus, the defect in T4P-dependent motility in the ΔromR strain is also not caused by lack of EPS accumulation.

To further examine the effect of RomX on T4P-dependent motility, we analysed the motility characteristics of single cells moving by means of T4P using an assay in which single cells are submerged in a 1% methylcellulose solution and moving on a polystyrene surface (Hu et al., 2011). As shown in Figure 22, A, cells of the WT and the ΔaglQ strain with inactivated gliding motility moved with similar velocity. Therefore, under these experimental conditions, cells were moving by T4P-dependent motility. Importantly, the velocity of the ΔromXΔaglQ and ΔromRΔaglQ cells were not statistically significantly different from that of the WT and the ΔaglQ cells (Figure 22, A). Moreover, because the ΔromX and ΔromR mutants accumulate EPS as WT (Figure 21, B), the observed single cell movement on 1% methylcellulose (in this assay) was not due to bypass of a lack of EPS by the methylcellulose. However, under the same conditions, the ΔromXΔaglQ and ΔromRΔaglQ mutants reversed much less frequently than WT and the ΔaglQ strain (Figure 22, B). In total, these data demonstrate that RomX and RomR are not important for T4P-dependent motility per se but they are important for reversals in the T4P-dependent motility systems.

Altogether, these observations demonstrate that RomX has two functions in motility. First, RomX is important for reversals in T4P-dependent motility but is not important for T4P-dependent motility per se as previously described for RomR (Keilberg et al., 2012, Zhang et al., 2012); and, second, RomX is essential for gliding motility as previously described for RomR (Keilberg et al., 2012, Zhang et al., 2012). Consistently, a ΔfrzEΔaglQ double mutant had the same motility characteristics as the ΔromRΔaglQ and ΔromXΔaglQ double mutants on hard and soft agar (Figure 21, A).
2.2.2 RomX acts in the same pathway as RomR, MglA and MglB

We tested whether RomX acts together with MglA, MglB and RomR in the same pathway. To test the relationship between RomX and MglA, MglB as well as RomR, genetic epistasis experiment in which motility phenotypes were analyzed was performed.
Figure 23. RomX acts in the same pathway as MglA&B and RomR to regulate motility.

(A) Motility assays as described in Figure 21 comparing single and double mutants of romX, mglA, mglB and romR. The numbers indicate the increase in colony diameter in mm ± standard deviation after 24h. Scale bars, 1000 μm (T4P-dependent motility), 500μm (gliding motility – left panel) and 50 μm (gliding motility – right panel).

(B) Velocity of WT and selected mutants on 1.5% agar, 0.5% CTT. Single moving cells were tracked for 15 min. Graph presents velocity of the cells in μm/min ± standard deviation. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>20 for each of two independent experiment. Number in each column represents percent of moving single cells. n>450 for each of two independent experiment. NA – not applicable; ΔaglQ, ΔromX, ΔromR were not moving as single cells in the tested conditions.

(C) Reversal frequency of WT and selected mutants on 1.5% agar, 0.5% CTT. Shown are boxplots of the measured reversals of isolated cells for 15 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>50 for each of two independent experiment. NA – not applicable; ΔaglQ, ΔromX, ΔromR were not moving as single cells in the tested conditions.
RESULTS

ΔmglA and ΔmglAΔromX mutants showed flat colony edges under both tested conditions (Figure 23, A). This suggests that MglA acts downstream of RomX. The double ΔmglBΔromX mutant showed short, misformed flares similarly to the single ΔmglB and ΔromX mutants on 0.5% agar. Importantly, no additive effect on T4P-dependent motility of the two single deletions were observed in the double ΔmglBΔromX deletion. For gliding motility, we observed that the colony edge of ΔmglBΔromX mutant was similar to that of the ΔmglB mutant. However, the increase in colony diameter of the ΔmglBΔromX mutant was significantly smaller than the increase of the ΔmglB mutant (t-test p<0.05). Additionally, ΔmglBΔromX cells spread significantly more than cells of ΔromX mutant (t-test p<0.05) (Figure 23, A). Thus, from these data it is not clear whether MglB and RomX act in the same pathway. Single deletions of ΔromR and ΔromX caused similar motility phenotype on both tested conditions. Moreover, no additive effect of the two single deletions was observed in the double deletion ΔromRΔromX (Figure 23, A). These observations suggest that RomR and RomX act in the same pathway.

To better understand the gliding motility defect in the ΔmglBΔromX mutant, analysis of single cells movement on 1.5% agar was performed. WT single cells were moving with an average velocity 1.53 µm/min and 79% of observed cells were moving (Figure 23, B). The ΔaglQ mutant lacks gliding motility, served as a negative control and did not display single cell movement. 81% of the ΔmglB cells moved and they moved faster than WT (Figure 23, B). As mentioned the ΔromX and ΔromR mutants lack gliding motility (Figure 21, A and Figure 23, B). Importantly, in both the ΔmglBΔromX and ΔmglBΔromR mutant gliding motility was partially restored. However, these two strains moved significantly slower than single ΔmglB mutant. Moreover, only approx. 30% of the ΔmglBΔromX and ΔmglBΔromR mutants cells displayed single cell gliding (Figure 23, B). Thus, gliding motility of the ΔromX and ΔromR mutants is partially restored in the absence of mglB.

Additionally, reversal frequencies were quantified. Under the tested conditions, WT reversed on average 1.28 times in 15 min. In agreement with previous observations (Leonardy et al., 2010, Zhang et al., 2010), the ΔmglB mutant hyperreversed with on average 3.14 reversals in 15 min (Figure 23, C). The ΔmglBΔromX mutant as well as the ΔmglBΔromR mutant was also hyper-reversing in comparison to WT. However, ΔmglBΔromX and ΔmglBΔromR reversed significantly less than the ΔmglB mutant (t-test p<0.001).
In total, the epistasis experiments suggest that \textit{romX} acts in the same pathway as \textit{mglA}, \textit{mglB} and \textit{romR}. Moreover, analysis of single cells moving by gliding motility and reversals revealed that RomX and RomR have antagonistic function to MglB.

\textbf{2.2.3 RomX localizes dynamically to the cell poles}

MglA, MglB, RomR and RomX were localized using active fluorescent fusion proteins expressed as the only copy. First, the stability of the used fusions was investigated by immunoblot analysis on whole cell lysates.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure24}
\caption{MglA, MglB, RomR and RomX fluorescent fusions are stable. (A) Immunoblot of RomX-YFP. Cells were grown in liquid culture, harvested, and total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using \(\alpha\)-RomX. Calculated molecular masses of RomX and RomX-YFP are indicated. Immunoblot with \(\alpha\)-PilC serves as loading control. (B) Immunoblot of MglA-mVenus. Cells were grown and protein separation as in A. SDS-PAGE analyzed by immunoblot using \(\alpha\)-MglA. MglA and MglA-mVenus with calculated molecular masses are indicated. Immunoblot with \(\alpha\)-PilC serves as loading control. (C) Immunoblot of MglB-mCherry. Cells were grown and protein separation as in A. SDS-PAGE analyzed by immunoblot using \(\alpha\)-MglB. MglB and MglB-mCherry with calculated molecular masses are indicated. Immunoblot with \(\alpha\)-PilC serves as loading control. (D) Immunoblot of RomR-mCherry. Cells were grown and protein separation as in A. SDS-PAGE analyzed by immunoblot using \(\alpha\)-RomR. RomR and RomR-mCherry with calculated molecular masses are indicated. Immunoblot with \(\alpha\)-PilC serves as loading control.}
\end{figure}

We tested whether RomX-YFP is stable in \(\Delta\textit{romX}\) mutant and in double deletion mutants with \(\Delta\textit{mglA}\), \(\Delta\textit{mglB}\), \(\Delta\textit{romR}\) or \(\Delta\textit{romY}\). Native RomX protein was not detectable.
RESULTS

in the WT strain. RomX-YFP was detected in all investigated mutants. No degradation products were observed in the immunoblot analysis (Figure 24, A). Next, accumulation of the endogenous MglA-mVenus fusion was studied. This fusion accumulated at native level and was stable in all investigated strains (Figure 24, B). Endogenous MglB-mCherry fusion also accumulated at native level and it was stable in all tested strains (Figure 24, C). Finally, accumulation of the endogenous RomR-mCherry fusion was tested. This fusion also accumulated in all investigated strains. However, its level of accumulation was lower than native RomR in the wild type as well as in the ΔromX and ΔromY mutants (Figure 24, D).

As mentioned, the RomX-YFP fusion complemented the motility defects of the ΔromX mutant (Figure 21, A). MglB-mCherry was previously reported to be fully functional (Keilberg et al., 2012). Strain that expressed mglA-mVenus as the only copy of mglA showed a minor defect in both T4P-dependent and gliding motility (Figure 25, A). RomR-mCherry was fully functional (Figure 25, B).

Using fluorescence microscopy, we found that RomX-YFP predominantly localized in an asymmetric bipolar pattern with a large cluster at the lagging pole and a small cluster at the leading cell pole. Approx. 50% of the fluorescence signal was collected from the cell body. During reversals, the polar clusters switched polarity (Figure 26, A).
Next, we analyzed the localization of RomX in the absence of MglA, MglB and RomR. In the absence of MglA, RomX localized mainly unipolarly. By contrast, in the absence of MglB, RomX localized more bipolar symmetric than in WT. When examining RomX localization in the absence of RomR, we found that RomX localized diffusely throughout the cytoplasm (Figure 26, B). Therefore, polar asymmetry of RomX depends on MglA and MglB, while polar localization depends on the RomR.

Next, we analyzed the localization of MglA, MglB or RomR in the absence of each other or RomX. As previously observed, MglA localized mainly in a unipolar pattern (Leonardy et al., 2010, Zhang et al., 2010, Patryn et al., 2010) (Figure 26, C). As previously observed (Zhang et al., 2010, Leonardy et al., 2010), MglA localization pattern shifted towards bipolar symmetric in the absence of MglB. Finally, we confirmed that MglA in the absence of RomR localized diffusely throughout the cytoplasm (Keilberg et al., 2012, Zhang et al., 2012). Importantly, MglA also localized mainly diffusely throughout the cytoplasm in the absence of RomX (Figure 26, C). Thus, polar localization of MglA depends on both RomX and RomR. As previously observed (Keilberg et al., 2012, Zhang et al., 2012), MglB localized in bipolar asymmetric pattern. As previously reported (Keilberg et al., 2012, Zhang et al., 2012), MglB localization changed to more bipolar asymmetric with a large fraction of cells with unipolar localization in the absence of MglA. Moreover, MglB was more unipolar in the absence of RomR. In the absence of RomX, MglB also localized mainly unipolar. Thus, polar asymmetry of MglB depends on MglA, RomR and RomX. As previously observed (Leonardy et al., 2007, Keilberg et al., 2012, Zhang et al., 2012), RomR localized in a bipolar asymmetric pattern similarly to MglB. As reported earlier (Keilberg et al., 2012, Zhang et al., 2012), RomR in the absence of MglA localized predominantly in a unipolar pattern. Moreover, we confirmed that RomR localized more symmetric in the absence of MglB (Keilberg et al., 2012, Zhang et al., 2012). In the absence of RomX, localization of RomR changed towards unipolar (Figure 26, C). Therefore, RomR polar asymmetry depends on MglA, MglB and RomX.

Altogether, these data suggest that RomX, similarly to RomR, is important for polar localization of MglA-GTP. Because RomR is important for polar localization of RomX but not vice versa, these observations strongly suggest that RomR is a polar targeting determinant of RomX, which in turn helps to bring about polar localization of MglA-GTP. Moreover, polar asymmetry of MglB and RomR depends on RomX. We speculate that these latter effects are indirect and caused by the lack of polar localization of MglA-
GTP in the absence of RomX because MglB and RomR are also shifted towards unipolar localization in the absence of MglA (Keilberg et al., 2012, Zhang et al., 2012).
Figure 26. RomX localizes in bipolar asymmetric pattern.

(A) Localization of RomX-YFP. Upper left panel shows snapshot, lower left panel shows time-lapse fluorescence microscopy at 30 s intervals. Cells were transferred from liquid culture to a thin agar pad on a microscope slide and imaged by fluorescence microscopy. The fluorescence signal for each pole and cytoplasm was calculated and plotted in the upper right graph. Cyan dot represents the mean. To distinguish between different localization patterns, the $\omega$ value that represents asymmetry between the polar clusters was calculated from the equation:

$$\omega = \frac{\text{total fluorescence at pole 1} - \text{total fluorescence at pole 2}}{\text{total fluorescence at pole 1} + \text{total fluorescence at pole 2}}$$

By definition, pole 1 is the pole with the highest fluorescence. The $\omega$ value is between 0 (bipolar symmetric localization) and 1 (unipolar localization). The localization patterns are binned from the $\omega$ values as follows: unipolar ($\omega > 0.9$), bipolar asymmetric ($0.9 > \omega > 0.2$) and bipolar symmetric ($\omega < 0.2$). Diffuse localization was determined when no polar signal was detected. The localization patterns observed are indicated in the schematics. n>200. Scale bar: 5 µm.

(B) Localization of RomX-YFP in indicated strains. Left panels show fluorescent microscopy image. Cells were treated as in A. Percent of fluorescence signal for each pole and cytoplasm were calculated and plotted in the left graph. Cyan dot represents the mean. The localization patterns observed are indicated in the schematics. n>200. Scale bar: 5 µm.

(C) Localization of MglA-mVenus, MglB-mCherry and RomR-mCherry in indicated strains. Cells were treated as in A. Left panel shows fluorescent microscopy image. Percent of fluorescence signal for each pole and cytoplasm were calculated and plotted on the upper graph. Cyan dot represents the mean. The localization patterns observed are indicated in the schematics. n>200. Scale bar: 5 µm.

2.2.4 RomX is polar targeting determinant of MglA

Lack of RomX or RomR causes diffuse localization of MglA. Similarly, an inactive, GDP-locked form of MglA localizes diffusely (Leonardy et al., 2010). Additionally, RomX displays diffuse localization in the absence of RomR. So, RomX and RomR act together to stimulate polar localization of MglA-GTP. We thought of four models to explain how RomX and RomR could bring about polar localization of MglA-GTP: (1) RomX together with RomR or alone acts as an MglA GEF; (2) RomX together with RomR or alone inhibits MglB GAP activity; (3) RomX is an MglA-GTP polar targeting determinant and is sandwiched between RomR and MglA-GTP; (4) combinations of these activities.

To explore these possibilities, we determined localization of MglA$^{Q82A}$-mVenus, which is locked in the GTP-bound, in the presence and absence of RomX or RomR. First, motility assays with mglA$^{Q82A}$ and mglA$^{Q82A}$-mVenus mutants showed no differences between the two strains (Figure 27, A). Moreover, MglA$^{Q82A}$-mVenus accumulated in all investigated mutants at similar level as the native fusion protein in WT (Figure 27, B).
Figure 27. MglA_Q82A-mVenus is a functional fluorescent fusion protein.

(A) Motility assay of mglA_Q82A-mVenus. Motility assay performed as in Figure 21, A. Scale bars, 1000 μm (T4P-dependent motility), 500 μm (gliding motility – left panel) and 50 μm (gliding motility – right panel).

(B) Immunoblot of MglA_Q82A-mVenus. Cells were grown in liquid culture, harvested, and total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using α-MglA. MglA and MglA-mVenus with calculated molecular masses are indicated. Immunoblot with α-PilC serves as loading control.

As previously reported (Miertzschke et al., 2011, Keilberg et al., 2012, Treuner-Lange et al., 2015), MglA_Q82A localized in a bipolar symmetric pattern with a cluster that “oscillates” between the cell poles (Figure 28, A). The “oscillating” cluster is a large complex of the Agl/Glt motility machinery. Because this machinery contains the GTP-locked MglA_Q82A variant and, therefore, is insensitive to MglB at the lagging cell pole, cells move one cell length before reversing leaving the impression of an oscillating cluster (Miertzschke et al., 2011, Keilberg et al., 2012, Treuner-Lange et al., 2015). By contrast, MglA_Q82A-mVenus in the absence of RomR or RomX mostly localized to the “oscillating” cluster and the cytoplasm (Figure 28, A). Quantitative localization analysis revealed that difference in the cytoplasmic signal was statistically significant (t-test p<0.001) (Figure 28, B). These localization patterns suggest that one of the functions of RomX is to act as a polar targeting determinant for MglA-GTP, i.e. RomR recruits RomX to a pole and RomX recruits MglA-GTP.
Figure 28. Polar recruitment of MglA-GTP depends on RomX and RomR.

(A) Snapshots and time-lapse microscopy of MglA<sup>Q82A</sup>-mVenus in indicated strains. Cells were treated as in Figure 26 and imaged by time-lapse fluorescence microscopy at 30s intervals. Scale bar: 5 µm for snapshots and 1 µm for time-lapses.

(B) Analysis of the cytoplasmic fluorescence of MglA<sup>Q82A</sup>-mVenus in indicated strains. Shown are boxplots of the measured integrated cytoplasmic fluorescence, boxes enclose the 25<sup>th</sup> and 75<sup>th</sup> percentile, whiskers represent the 10<sup>th</sup> and 90<sup>th</sup> percentile, and dots outliers. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>250.

(C) mglA<sup>A82A</sup> is epistatic to romX and romR. Motility assays as described in Figure 21 comparing single mutant mglA<sup>Q82A</sup>, ΔromX and ΔromR to double mglA<sup>Q82A</sup>ΔromX and mglA<sup>A82A</sup>ΔromX mutants. Bars, 1000 µm (T4P-dependent motility), 500 µm (gliding motility – left panel) and 50 µm (gliding motility – right panel).
RESULTS

(D) Velocity of WT and selected mutants on 1.5% agar, 0.5% CTT. Single moving cells were tracked for 15 min. Graph presents velocity of the cells in μm/min ± standard deviation. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>20 for each of two independent experiment. Number in each column represents percent of moving single cells. n>450 for each of two independent experiment. NA – not applicable; ΔaglQ, ΔromX, ΔromR were not moving as single cells in the tested conditions.

(E) Reversal frequency of WT and selected mutants on 1.5% agar, 0.5% CTT. Shown are boxplots of the measured reversals of isolated cells for 15 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>50. NA – not applicable; ΔaglQ, ΔromX, ΔromR mutants were not moving as single cells in the tested conditions.

Additionally, we tested whether MglA locked in the GTP bound form can bypass the motility defects caused by lack of RomR or RomX. In agreement with previous observations (Miertzschke et al., 2011), the mglA<sup>Q82A</sup> mutant hyper-reversed and formed short flares on 0.5% agar and displayed no single cell at the colony edge after 24 hours on 1.5% agar (Figure 28, C). As previously reported (Keilberg et al., 2012), the double mglA<sup>Q82A</sup>ΔromR mutant showed motility phenotype similar to mglA<sup>Q82A</sup> mutant. Furthermore, the double mglA<sup>Q82A</sup>ΔromX mutant also had the phenotype of the mglA<sup>Q82A</sup> mutant (Figure 28, C). Next, analysis of single cells moving by gliding motility was performed. WT moved with a velocity of 1.46 μm/min and 70% of single cells were moving. As previously observed, ΔaglQ, ΔromX and ΔromR mutants did not move as a single cell on 1.5% agar. The mglA<sup>Q82A</sup> mutant moved faster than WT with an average velocity of 2.09 μm/min and 84% of single cells were moving. Furthermore, the mglA<sup>Q82A</sup>ΔromX and mglA<sup>Q82A</sup>ΔromR mutants moved on 1.5% agar with higher velocity than WT (Figure 28, D). Moreover, the percent of moving cells in the case of mglA<sup>Q82A</sup>ΔromX and mglA<sup>Q82A</sup>ΔromR mutants was comparable to WT and the mglA<sup>Q82A</sup> mutant (Figure 28, D). The mglA<sup>Q82A</sup> mutant hyper-reversed with on average 4.8 reversals per 15 minutes (Figure 28, E). Furthermore, the mglA<sup>Q82A</sup>ΔromX and mglA<sup>Q82A</sup>ΔromR mutants showed a similar hyper-reversing phenotype (Figure 28, E). Thus, RomX and RomR are not required for gliding motility when MglA is locked in the GTP-bound form. Additionally, the mglA<sup>Q82A</sup> mutation is epistatic to the ΔromX and ΔromR mutations. Altogether these observations suggest that MglA-GTP acts downstream of RomX and RomR.

2.2.5 Formation of focal adhesion depends on RomX and RomR in the presence of MglB

MglA-GTP is an integral part of the gliding motility machinery and localizes to the focal adhesion complexes (Treuner-Lange et al., 2015). To further investigate hypothesis
that RomX and RomR influence nucleotide state of MglA, we focused on the localization of MglA in focal adhesion complexes. To visualize MglA in the focal adhesion complexes we used TIRF microscopy to only visualize fluorescent fusions proteins that are localize close to the cell-substratum contact surface.

Figure 29. MglA localization to focal adhesion complexes depends on RomX and RomR.
TIRF microscopy of MglA-mVenus in indicated strains. First column presents pictures done in in TIRF microscopy, second shows TIRF time-lapse microscopy at 20 s intervals. Scale bar for snapshots: 5µm; for time lapses montages: 1 µm.
RESULTS

MglA-mVenus showed localization in the focal adhesion complexes. When cells were moving, these clusters stayed stationary with respect to the substratum. Additionally, MglA-mVenus clusters assembled at the leading cell pole and disassembled near the lagging cell pole (Figure 29). In the ΔmglB mutant, MglA-mVenus in the focal adhesion complexes was also visible. By contrast, in the ΔromX, ΔromR and double ΔromXΔromR mutants, MglA-mVenus did not localize to the focal adhesion complexes (Figure 29). Importantly, MglA-mVenus localized to focal adhesion complexes in ΔmglBΔromX, ΔmglBΔromX and triple deletion ΔmglBΔromRΔromX mutants (Figure 29). Previously, we showed that MglA locked in the GTP-bound form localizes in a single focal adhesion complex independently of RomX and RomR (Figure 28, A). These observations support the idea that RomX and RomR are necessary for assembly of the focal adhesion complexes only when nucleotide state of MglA is regulated by MglB. Moreover, the data support the idea that RomX together with RomR have the opposite function on MglB, and thus, may act as an MglA GEF or inhibit MglB GAP activity.

To further analyse the formation of focal adhesion complexes in the absence of RomX and/or RomR, we localized AglZ, which is an integral part of the gliding machinery (Mignot et al., 2007). First, AglZ-YFP accumulation in the investigated strains was tested. AglZ-YFP accumulated at similar levels in the absence and presence of MglB, RomX and RomR in single, double and triple deletion mutants (Figure 30).

Figure 30. AglZ-YFP is stable in the absence or presence of MglB, RomX and RomR in single, double and triple deletion mutants.

Immunoblot of AglZ-YFP. Cells were grown and protein separation as in Figure 24. SDS-PAGE analyzed by immunoblot using α-GFP. AglZ-YFP with calculated molecular mass is indicated. Immunoblot with α-PilC serves as a loading control.

AglZ localized unipolarly with clusters distributed along the cell length. Additionally, when cells were moving these clusters stayed stationary with respect to the substratum. Moreover, clusters were assembled at the leading cell pole. In the absence of
MglB, we observed more AglZ-YFP clusters. Additionally, these clusters stayed stationary with respect to the substratum and were assembled at the leading cell pole. By contrast, and as observed for MglA, we did not observe AglZ-YFP clusters along the cell body in the absence of RomX or RomR. Importantly, in the ΔmglBΔromX, ΔmglBΔromR and ΔromXΔromRΔmglB mutants, AglZ-YFP was observed in the focal adhesion complexes when visualized by TIRF microscopy. Detailed comparisons indicated that although superficially similar, the focal adhesion complexes in these three mutants displayed three notable differences to those formed in WT. First, they “slipped” the contact to the substratum and moved within a cell, i.e. they were not stationary with respect to the substratum. Second, they moved in opposite directions within a cell. Third, they disassembled away from the lagging cell pole (Figure 31). To tally these differences, we quantified the movement of Agl/Glt complexes within cells and the lifetime of individual Agl/Glt complexes. As shown in Figure 32, in the ΔmglB ΔromX, ΔmglB ΔromR and ΔromXΔromRΔmglB mutants, the Agl/Glt complexes moved a significantly longer distance per min within cells and had significantly shorter lifetimes than in WT and ΔmglB cells. Altogether, these findings are consistent with the observation that the ΔmglB mutation only partially suppresses the gliding defects in the ΔromX and ΔromR mutants, and the ΔmglBΔromX and ΔmglBΔromR cells move with a significantly lower speed than WT and ΔmglB cells (Figure 23, B). We conclude that RomX and RomR are not only important for formation of focal adhesion complexes but also for their correct adhesion and stability. Furthermore, the data suggest that RomX and RomR also play a role in setting the directionality or polarity of the focal adhesion complexes. We speculate, that we did not observe lack of polarization and stability of the focal adhesion complexes in the ΔmglBΔromX, ΔmglBΔromR and ΔmglBΔromXΔromR mutants for MglA-mVenus, because this fusion protein is only partially functional.
Figure 31. AglZ localization to focal adhesion complexes depends on RomX and RomR.
TIRF microscopy of AglZ-YFP in indicated strains. First column presents pictures done in in TIRF microscopy, second (and third for selected strains) shows TIRF time-lapse microscopy at 20 s intervals. Scale bar for snapshots: 5 µm; for time lapses montages: 1 µm.
RESULTS

2.2.6 Focal adhesion immobility depends on RomX and RomR

The focal adhesion complexes formed in the absence of MglB as well as RomR and/or RomX are more mobile within cells. To further investigate this phenomenon, we localized MglA<sup>Q82A</sup>-mVenus, which is locked in the GTP-bound form, using TIRF microscopy in the presence and absence of RomX or RomR (Figure 33). We observed multiple clusters of MglA<sup>Q82A</sup>-mVenus along the cell body. MglA<sup>Q82A</sup>-mVenus in the absence of RomX or RomR also localized to focal adhesion complexes along the cell body. Some of these clusters remained stationary with respect to the substratum when cells were moving, while other clusters “slipped” the contact to the substratum and moved within a cell, i.e. they were not stationary with respect to the substratum or clusters moved in opposite directions within a cell (Figure 33). We quantified these differences as described above and found that the focal adhesion complexes formed in the absence of RomX or RomR moved a significantly longer distance per min within cells (Figure 34, A) but had the same lifetime as in the presence of RomX and RomR (Figure 34, B). In addition, the focal adhesion complexes formed in the presence of MglA<sup>Q82A</sup> and RomR as well as RomX had the same mobility within cells as focal adhesion complexes formed in the presence of MglA<sup>WT</sup> protein (compare Figure 32, A and Figure 34, A). We conclude
that RomX and RomR are important for immobility of the focal adhesion complexes while MglA locked in the GTP-bound form has no effect on the mobility of the focal adhesion complexes. Moreover, RomX and RomR are not important for stability of the focal adhesion complex in strains with MglA locked in the GTP bound form. Finally, our data suggest that RomX and RomR are giving correct directionality to the focal adhesion complexes.

Figure 33. Polarization of the focal adhesion complexes depends on RomX and RomR.
TIRF microscopy of MglA<sup>Q82A</sup>-mVenus in indicated strains. First column presents pictures done in in TIRF microscopy, second shows TIRF time-lapse microscopy at 20 s intervals. Scale bar for snapshots: 5 µm; for time lapses montages: 1 µm.
2.2.7 RomX and RomR localize to focal adhesion complexes

The data presented so far are consistent with the idea that RomR and RomX in addition to recruiting MglA-GTP to the leading cell pole, may also function to inhibit MglB GAP activity or function as an MglA GEF. Based on quantification of the fluorescence signal of the MglB-mCherry fusion, which is only detected as a full-length protein in immunoblots (Figure 24, C), more than 90% of MglB is localized diffusely in the cytoplasm. To further explore the idea that RomR and/or RomX could inhibit MglB GAP activity or function as an MglA GEF, we hypothesized that RomR and/or RomX would localize to the focal adhesion complexes to replenish MglA-GTP in case of MglB stimulated GTP hydrolysis by MglA. To explore this idea, we colocalized RomX with AglQ, which is a subunit of the Agl motor and localizes to the focal adhesion complexes (Sun et al., 2011).
Figure 35. RomX and RomR localize to the focal adhesion complexes.  
(A) AglQ colocalizes with RomX. Co-localization of AglQ and RomX observed on chitosan coated glass. First column presents pictures done TIRF microscopy, second shows TIRF time-lapse microscopy at 20 s intervals. Scale bar: 5 µm. for time lapses montages: 1 µm  
(B) AglZ colocalizes with RomR. Co-localization of AglZ and RomR observed on chitosan coated glass. First column presents pictures done in TIRF microscopy, second shows TIRF time-lapse microscopy at 20 s intervals. Scale bar for snapshots: 5µm; for time lapses montages: 1 µm.

AglQ-mCherry and RomX-YFP localized to clusters along the cell body when visualized using TIRF microscopy. Moreover, AglQ-mCherry and RomX-YFP colocalized in these clusters. Furthermore, these clusters were assembled at the leading
cell pole and remained stationary with respect to the substratum when in moving cells (Figure 35, A). Similarly, RomR-mCherry colocalized with AglZ-YFP in focal adhesion complexes. We conclude that RomX as well as RomR are incorporated into the focal adhesion complexes.

2.2.8 RomX incorporation into the focal adhesion complexes depends on RomR

As previously observed, RomX-YFP localized in the bipolar asymmetric pattern and in focal adhesion complexes when visualized by TIRF microscopy. As expected, in cells of the ΔmglA mutant, RomX-YFP was only unipolarly localized because the focal adhesion complexes do not assemble in the absence of MglA. In the absence of MglB, RomX-YFP was incorporated into the focal adhesion complexes. Next, we asked if the incorporation of RomX into the focal adhesion complexes depends on RomR. As expected, RomX-YFP was also diffusely localized in the absence of RomR (Figure 36). Importantly, we did not observe RomX-YFP in the focal adhesion complexes in the double ΔmglBΔromR mutant (Figure 36). The data suggests that RomX requires RomR for incorporation into the focal adhesion complexes.
2.2.9 RomR incorporation into focal adhesion complexes is independent of RomX

Subsequently, we asked if the incorporation of RomR into the focal adhesion complexes depends on RomX. As previously observed, RomR-mCherry localized in the bipolar asymmetric pattern as well as in focal adhesion complexes when visualized by TIRF microscopy (Figure 37). As expected, in the ΔmglA mutant, RomR-mCherry did
not localize to the focal adhesion complexes while it still localized to focal adhesion complexes in the $\Delta mglB$ mutant. Moreover, and as expected, RomR was not incorporated into focal adhesion complex in the absence of RomX (Figure 37). Importantly, RomR localized to focal adhesion complexes in the double deletion $\Delta mglB\Delta romX$ mutant (Figure 37). The data suggests that RomR does not depend on RomX for incorporation into the focal adhesion complexes.
Figure 37. RomR localization to focal adhesion complexes depends on MglA and RomX.
TIRF microscopy of RomR-mCherry in indicated strains. First column presents pictures done in in TIRF microscopy, second shows TIRF time-lapse microscopy at 20 s intervals. Scale bar for snapshots: 5 µm; for time lapses montages: 1 µm.
2.2.10 RomX interacts with MglA and RomR in the bacterial two hybrid system

To finally test whether RomX interacts directly with MglA, MglB, and/or RomR, we first used the bacterial two hybrid system (Figure 38). To this end, full length MglA, MglB and RomX as well as the three domains of RomR (Receiver, linker and C-terminal part (see Figure 15) were fused to the T18 and T25 fragments of the *Bordetella pertussis* adenylate cyclase. As a positive control, we used the plasmids pUT18C-zip and pKT25-zip, which express T18 and T25 fragments fused to leucine zippers that provide strong interactions. As a negative control, we used cells co-transformed with one plasmid expressing T18-zip or T25-zip and the other plasmid expressing the bait fused to T25 or T18, respectively.

We confirmed known interactions: MglA–MglB, MglB–RomR as well as self-interaction of MglB (Miertzschke *et al.*, 2011, Keilberg *et al.*, 2012). Moreover, RomR-REC, RomR-Linker and RomR-C were observed to interact. Importantly, RomX was found to self-interact and to interact with MglA and the RomR receiver domain (Figure 38, A). These results are in agreement with the localization data and the hypothesis that RomX acts between RomR and MglA in the regulation of polarity and motility.
Figure 38. RomX interacts with RomR receiver domain and MglA in BACTH.
(A) Interaction between indicated proteins in fusion with T18 or T25 adenyl cyclase fragment. Novel interactions between RomX – RomR-Receiver domain (REC) and RomX – MglA and RomX self-interaction are indicated in red. Blue colonies indicate interaction, while white colonies indicate no interactions.
(B) Model for proteins interactions found in this study.

2.2.11 RomX/RomR complex interacts with MglA-GTP

To further test for direct interacts between MglA, RomR and RomX we turned to in vitro experiments. To this end, we purified C-terminal His6-tagged MglA (MglA-His6),
N-terminal His6-tagged RomX (His\textsubscript{6}-RomX), N-terminal Strep-tagged RomX (Strep-RomX), N-terminal MalE-tagged RomR (MalE-RomR) and MalE (Figure 39).

![Figure 39. SDS-PAGE gels with the purified indicated proteins.](image)

SDS-PAGE gels show selected fractions after the size exclusion chromatography. Molecular size marker and calculated molecular weights of proteins are indicated.

We first tested whether RomX interacts to form a stable complex with RomR or MglA in pull-down experiments. When MalE-RomR was bound to an amylose resin, it bound His\textsubscript{6}-RomX whereas His\textsubscript{6}-RomX neither bound to the resin on its own nor in the presence of MalE (Figure 40, A). We conclude that RomX and RomR interact directly. Moreover, when Strep-RomX was bound to Strep-Tactin beads, it bound MglA-GTP but MglA-GDP whereas MglA-GTP did not bind to the beads (Figure 40, B). We conclude that RomX forms a stable complex with MglA-GTP but not with MglA-GDP.

To test whether RomR, RomX and MglA can form a heteromeric complex, MalE-Rom was bound to the amylose resin with or without His\textsubscript{6}-RomX. In the absence of His\textsubscript{6}-RomX, binding of MglA-GTP and MglA-GDP to MalE-RomR was not observed (Figure 40, C). However, when the amylose resin contained His\textsubscript{6}-RomX bound to MalE-RomR, MglA-GTP but not MglA-GDP was retained (Figure 40, D). Because neither MglA-GTP nor MglA-GDP bind stably to RomR in this assay, these data strongly suggest that MglA-GTP interacts with RomX in the RomX/RomR complex.

Previously, a weak interaction between RomR and MglA in pull-down experiments using a His6-RomR protein and a GST-MglA protein was reported (Keilberg et al., 2012). Here, we used an MglA-His\textsubscript{6} protein because this protein has GTPase activity \textit{in vitro} and this activity is stimulated by MglB (see below). We speculate that the difference between the data reported here and previously are due to the use of different purified proteins. We conclude that MglA-GTP but not MglA-GDP interacts with the RomR/RomX to form a stable complex leading to the formation of a RomR/RomX/MglA-GTP complex (Figure 40, E). These results are in agreement with the observations \textit{in vivo} that RomR recruits RomX to a pole and RomX is a polar targeting determinant of MglA-GTP.
**Figure 40. MglA-GTP interacts with RomR/RomX complex in vitro.**

(A) 1st panel: MalE-RomR can be bound to amylose resin. Purified MalE-RomR was applied to an amylose column. 2nd panel: His6-RomX does not bind to amylose resin. Purified His6-RomX was applied to an amylose column. 3rd panel: MalE-RomR and His6-RomX interact. Purified MalE-RomR and His6-RomX were mixed and applied on an amylose column. 4th panel: His5-RomX does not interact with MalE. Purified MalE and His5-RomX were mixed and applied on an amylose column.

(B) Strep-RomX interacts with MglA-His6-GTP but not MglA-His6-GDP in vitro. Strep-RomX and MglA-His6-GTP or MglA-His6-GDP were mixed and immobilized on the strep-tactin magnetic beads. Proteins were eluted with buffer supplemented with biotin.

(C) MalE-RomR does not interact with MglA-His6 in vitro. MalE-RomR was immobilized on the column, subsequently MglA-His6-GTP or MglA-His6-GDP were applied.
(D) RomR/RomX complex interacts with MglA-GTP. MalE-RomR and His6-RomX were mixed and immobilized on the amylose column, subsequently MglA-His6-GTP or MglA-His6-GDP were applied. Proteins were eluted with buffer supplemented with maltose.

(E) Schematic representation of experiment from D.
For all pull-downs shown are proteins from load, last wash before elution and/or before MglA-His6 loading, MglA-His6 loading (if was loaded) and elution.

2.2.12 RomX and/or RomR do not inhibit MglB GAP and MglA GTPase activity

Our cell biology data suggest that one of the function of RomX and RomR is to stimulate MglA-GTP accumulation in the cell. One possibility to achieve this would be to inhibit MglB GAP or MglA GTPase activity. In order to test the hypothesis, GTPase assay was performed. A low level of intrinsic activity of MglA alone was observed. No GTPase activity for MglB, RomX and RomR were detected. MglB, as expected, stimulated MglA GTPase activity whereas RomX, RomR and RomX/RomR did not. Moreover, RomX and/or RomR pre-incubated with MglB did not interfere with MglB GAP activity (Figure 41, left part). Similarly, when RomR and/or RomX was pre-incubated with MglA-GTP before addition of MglB, we did not observe inhibition of MglB GAP activity (Figure 41, right part). We conclude that RomX and/or RomR do not inhibit MglB GAP activity.

![Figure 41. RomX and/or RomR do not inhibit MglA GTPase and MglB GAP activity.](image)

Specific GTPase activity of 3 µM MglA, 6 µM MglB, 6 µM RomX, 6 µM RomR and combinations of the proteins in the presence of 1 mM GTP. Experiment was performed in triplicates and graph shows mean ± SD.
2.2.13 RomX/RomR complex stimulates MglA nucleotide exchange

To test whether RomX and/or RomR have MglA GEF activity, we monitored GDP to GTP exchange using two different experimental set-ups.

In the first set-up, MglA-His$_6$ was preloaded with the fluorescent GDP analog N-methylanthraniloyl (mant)-GDP before MalE-RomR and/or His$_6$-RomX were added. Subsequently, excess of GTP was added and the decrease in fluorescence intensity associated with release of mantGDP and binding of GTP by MglA-His$_6$ was monitored. MglA spontaneously exchanged mantGDP for GTP slowly and the exchange rate was not affected by BSA, which served as a negative control (Figure 42, A). RomX and RomR independently did not stimulate the nucleotide exchange rate. Importantly, in the presence of both, RomX and RomR, the two proteins synergistically stimulated the nucleotide exchange rate supporting that the RomX/RomR complex has MglA GEF activity (Figure 42, A).

Figure 42. RomX/RomR complex stimulates MglA nucleotide exchange.
(A) MglA was loaded with mGDP and mGDP release was measured in real time (as shown in the scheme above) in the absence and in the presence of RomX, RomR, RomX/RomR or BSA. Arrow indicates a start of the reaction by adding GTP. Fluorescence was measured every second.
(B) MglA was loaded with GDP and mGTP binding was measured in real time (as shown in the scheme above) in the absence and in the presence of RomX, RomR, RomX/RomR or BSA. Arrow indicates start of the reaction by adding MglA-GDP or MglA-GDP and RomX, RomR, RomX/RomR or BSA.
In the reciprocal experiment, MglA-His\(_6\) was preloaded with GDP, then mantGTP was added followed by addition of His\(_6\)-RomX and/or MalE-RomR or BSA, and the increase in fluorescence intensity associated with release of GDP and binding of mantGTP by MglA-His\(_6\) was followed. In the presence of MglA alone or together with RomR or BSA no increase in fluorescence intensity was observed indicating that GDP was not being exchanged for mantGTP (Figure 42, B). After addition of only RomX, a transient increase in fluorescence intensity was observed. However, in the presence of RomX together with RomR a significant increase in fluorescence intensity was observed indicating GDP for mantGTP exchange by MglA (Figure 42, B). We speculate that the reason why no spontaneous exchange of GDP for mantGTP by MglA was observed in this experiment is due to the 20-fold higher concentration of GDP compared to mantGTP.

Note than in the experiment in Figure 42, A in which spontaneous exchange of mantGDP for GTP by MglA was observed, GTP was added in 5-fold excess of mantGDP. Based on the *in vitro* experiments, we conclude that the RomR/RomX complex has MglA GEF activity and also interacts directly with MglA-GTP. *In vivo* these two activities would bring about polar localization of MglA-GTP by increasing the concentration of MglA-GTP via the GEF activity and by RomX/RomR binding and recruiting MglA-GTP directly to a pole.

Binding of effectors or a GAP to a small GTPase containing a bound mant-nucleotide can cause a change in mant fluorescence (Kuhlmann *et al.*, 1997, Leonard *et al.*, 1998, Hemsath *et al.*, 2005). To exclude the possibility that the increase in fluorescence intensity observed in the experiment in Figure 42, B in the presence of the RomR/RomX complex was caused by simple binding of RomR/RomX to the MglA/mantGTP complex a control experiment was performed. In this experiment, we tested the change in fluorescence of mantGTP bound to MglA following addition of RomX and/or RomR. As a positive control, we used MglA preloaded with GDP and then added mantGTP with RomX and RomR as described in the previous experiment. We did not observe an increase in fluorescence for MglA/mantGTP alone or with RomX, RomR or BSA. After addition of the RomX/RomR complex to MglA/mantGTP, we detected an increase in fluorescence. However, the increase was lower than in the mantGTP exchange experiment with RomX and RomR (Figure 43). Thus, the data support that the increase in mant fluorescence for the previous experiment is mainly due to the binding of mantGTP by MglA and not the RomR/RomX complex interacting with MglA/mantGTP. Furthermore, because we are so far not able to purify nucleotide free MglA (data not
shown) we cannot exclude that the increase in fluorescence observed upon addition of the RomR/RomX complex to MglA/mantGTP is a consequence of increased binding of mantGTP to MglA. Regardless, the data is consistent and support the conclusion that RomX/RomR complex is a GEF and stimulates MglA nucleotide exchange.

Figure 43. RomX/RomR complex binds to MglA-GTP.
MglA was loaded with GDP and mGTP binding was measured in real time in the presence of RomX/RomR complex (as shown in the scheme above; orange line on the graph) or MglA was loaded with mGDP and change of the fluorescence intensity due to possible interaction with MglA was measured in real time (as shown in the scheme above) in the absence and in the presence of RomX, RomR, RomX/RomR or BSA. Arrow indicates start of the reaction by adding MglA-GDP and RomX/RomR or MglA with or without RomX, RomR, RomX/RomR or BSA.

2.3. Characterization of RomY in Myxococcus xanthus

Alignment of all 28 identified RomY homologs revealed a conserved N-terminal region, a proline rich low complexity region of variable length, and a conserved C-terminal motif (Figure 44, A). None of these domains matches any currently characterized domain models.

The genetic neighborhood of romY is conserved in closely related species; however, the flanking genes did not suggest any function in regulation of motility. In M. xanthus, the ABC transporter FtsEX involved in cell division is encoded downstream of romY and immediately upstream of romY is the gene encoding the carotenoid synthesis regulator, carF (Figure 44, B).
Figure 44. Bioinformatics analysis of RomY

(A) Alignment of identified RomY homologs. N-terminal conserved region and C-terminal conserved motif are indicated.

(B) Conservation of romY neighbourhood in selected myxobacteria. Genes are depicted as arrows. Arrow orientation indicates coding direction. Homologous proteins are indicated in same colours. Numbers in arrows in M. xanthus display gene number. Numbers in arrows (outside of M. xanthus) indicate identity and similarity of their encoded protein to their M. xanthus homolog. Numbers were obtained from Pairwise Sequence Alignment of protein using EMBOSS Needle. Right part shows predicted proteins functions based on the NCBI BLAST and conserved domain analysis (Marchler-Bauer et al., 2017)

2.3.1 RomY is important for both motility systems

In order to test whether RomY is involved in motility and/or its regulation we generated an in-frame romY deletion. The ΔromY mutant showed very short flares on
0.5% agar in comparison to the WT strain. In the case of gliding motility, we observed fewer single cells for the ΔromY mutant in comparison to WT on 1.5% agar (Figure 45). To confirm that the motility defects in the ΔromY mutant was caused by lack of RomY, we generated a complementation strain in which a RomY-YFP fusion protein was expressed ectopically under the control of the native promoter. This strain did not show motility defects (Figure 45). Based on these experiments we conclude that RomY is important for both T4P-dependent and gliding motility.

**Figure 45. romY mutant shows defect in T4P-dependent and gliding motility.**
Motility assay as described in Figure 21 compares WT to the ΔromY mutant. Scale bars, 500 µm (T4P-dependent motility), 500 µm (gliding motility – left panel) and 50 µm (gliding motility – right panel).

### 2.3.2 RomY acts in the same pathway as MglA, MglB, RomR and RomX

Next, we tested whether RomY acts in the same pathway as MglA, MglB, RomR and RomX. To this end, genetic epistasis experiments were performed (Figure 46). Both ΔmglA and ΔmglAΔromY mutants had a flat colony edge on 0.5% as well as on 1.5% agar. This suggests that MglA acts downstream of RomY. The ΔmglB mutant was indistinguishable from the ΔromY mutant under both conditions. Both strains showed very short flares on 0.5% agar and only few single cells at the colony edges on 1.5% agar. Furthermore, the same phenotype was observed for the double deletion ΔmglBΔromY
mutant under both conditions. Moreover, measured expansions of the colonies after 24 hours of the ΔmglB, ΔromY and ΔmglBΔromY mutants were very similar and the differences in colony expansion were not statistically significant (t-test p>0.01). Based on this observation we conclude that lack of the RomY caused the same defect as ΔmglB and the double deletion caused no additive effect. The ΔromR mutant showed misformed flares on 0.5% agar and the ΔromRΔromY mutant had a similar phenotype. On 1.5% agar, the double ΔromRΔromY mutant showed more single cells at the colony edge than the ΔromR mutant and a colony expansion after 24 hours that was comparable to that of the ΔromY mutant. Both ΔromX and ΔromXΔromY mutants had short misformed flares in contrast to the very short flares of the ΔromY mutant. On 1.5% agar, the double ΔromXΔromY mutant showed a larger colony expansion after 24 hours in comparison to the single ΔromX mutant and somewhat similarly to the ΔromY mutant. Furthermore, the differences in colony expansion after 24 hours were statistically significant (t-test p<0.01) (Figure 46, A).
Figure 46. RomY acts in the same pathway as MglA, MglB, RomR and RomX.  
(A) Motility assay as described in Figure 21 compares single and double mutants of romY, mglA, mglB, romX and romR. The numbers indicate the increase in colony diameter in mm ± standard deviation from three technical replicates after 24h. Scale bars, 1000 μm (T4P-dependent motility), 500μm (gliding motility – left panel) and 50 μm (gliding motility – right panel).  
(B) Reversal frequency of WT and selected mutants on 1.5% TPM agar. Shown are boxplots of the measured reversals of isolated cells for 15 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. Student’s t-test was employed for statistical analysis and * indicates p< 0.001. n>50.
RESULTS

The epistasis experiment suggested that RomY acts in the same pathway as MglA, MglB, RomR and RomX. Moreover, comparison of the ΔmglB and ΔromY mutants suggested that lack of MglB and RomY cause the same motility defects. Because lack of MglB causes cells to hyper-reverse, we quantified the reversal frequency on 1.5% agar of the ΔromY mutant. Under the tested conditions, WT reversed on average 1.40 times in 15 min while the ΔmglB mutant reversed on average 3.41 times in 15 min. Importantly, the ΔromY mutant also hyper-reversed and with an average of 3.5 reversals in 15 min. In the double ΔmglBΔromY mutant, we did not observe additive effects of the single deletions. (Figure 46, B). Thus, the ΔromY mutant is hyper-reversing and RomY acts in the same pathway as MglA, MglB, RomR and RomX.

2.3.3 RomY localization studies

MglA, MglB, RomR and RomX are polarly localized and mutually depend on each other for polar asymmetry and localization (chapter 2.2.4). We decided to test whether RomY is polarly localized and whether localization of RomY depends on MglA, MglB, RomR and RomX.

First, we tested accumulation and stability of RomY-YFP fusion protein in the ΔromY mutant, as well as in absence of MglA, MglB, RomR and RomX. The RomY protein in WT was detected by α-RomY antibodies. Moreover, RomY-YFP accumulated in the complementation strain at a slightly higher level than the native protein in the WT. RomY-YFP accumulated at similar level as in the complementation strain in the absence of MglA, MglB, RomR and RomX (Figure 47).

Figure 47. RomY-YFP is stable in investigated mutants.
Immunoblot of RomY-YFP. Cells were grown in liquid culture, harvested, and total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using α-RomY. RomY and RomY-YFP with calculated molecular masses are indicated. Immunoblot with α-PiC serves as loading control.
RomY-YFP localized predominantly unipolarly with the cluster at the lagging cell pole. Moreover, when cells reversed, RomY-YFP changed localization to the new lagging cell pole (Figure 48, A). RomY-YFP localized predominantly in a unipolar pattern in the absence of MglA. RomY-YFP localized mostly diffusely throughout the cytoplasm in the absence of MglB. In the absence of RomR, RomY-YFP localization shifted towards more diffuse throughout the cytoplasm. However, more than 50% of cells still showed a polar RomY-YFP cluster. In the absence of RomX, RomY-YFP localized predominantly diffusely throughout the cytoplasm (Figure 48, B). Nevertheless, the effect of the ΔromX mutation on RomY localization was not as strong as in the case of the ΔmglB mutation. Therefore, polar localization of RomY depends strongly on MglB and less on RomR and RomX.
Figure 48. RomY localizes predominantly at the lagging pole.

(A) Localization of RomY-YFP. Upper left panel shows snapshot, lower left panel shows time-lapse fluorescence microscopy at 30 s intervals. Cells were treated as in Figure 26. Percent of fluorescence signal for each pole and cytoplasm were calculated and plotted on the upper right graph. Cyan dot represents the mean. The localization patterns observed are indicated in the schematics. The ratios between polar signals were calculated to distinguish between unipolar, asymmetric bipolar and symmetric bipolar localization as in figure 26. Scale bar: 5 µm. n>200.
Localization of RomY-YFP in indicated strains. Left panels show fluorescent microscopy pictures. Cells were treated as in Figure 26. Percent of fluorescence signal for each pole and cytoplasm were calculated and plotted on the left graph. Cyan dot represents the mean. The localization patterns observed are indicated in the schematics. The ratios between polar signals were calculated to distinguish between unipolar, asymmetric bipolar and symmetric bipolar localization as in figure 26. n>200. Scale bar: 5 µm.

Localization of MglA-mVenus, MglB-mCherry and RomR-mCherry in indicated strains. Cells were treated as in Figure 26. Left panel shows fluorescent microscopy picture. Percent of fluorescence signal for each pole and cytoplasm were calculated and plotted on the upper graph. Cyan dot represents the mean. The localization patterns observed are indicated in the schematics. The ratios between polar signals were calculated to distinguish between unipolar, asymmetric bipolar and symmetric bipolar localization as in figure 26. n>200. Scale bar: 5 µm.

In the reciprocal experiments, we observed that MglA predominantly localized in a unipolar pattern in WT and changed localization towards bipolar asymmetric and symmetric in the absence of RomY. Thus, polar asymmetry of MglA depends on the RomY. MglB, in the absence of RomY, localized in a bipolar asymmetric pattern as in WT. RomR, in the absence of RomY, localized predominantly in the bipolar asymmetric pattern as in WT. However, in the absence of RomY less RomR was polarly bound. Thus, RomY helps RomR to bind to the poles. RomX localized mostly in the bipolar asymmetric pattern in WT and in the absence of RomY, we observed a shift towards a more bipolar symmetric pattern. Also, more RomX was polarly bound in the absence of RomY in comparison to WT. Thus, polar asymmetry of RomX depends on RomY (Figure 48, C).

2.3.4 RomY interacts with RomX and MglA in bacterial two hybrid system

To test whether RomY interacts directly with MglA, MglB, RomR or RomX, we performed bacterial two hybrid experiments. In these experiments, we found that RomY self-interacts and also interacts with RomX and MglA (Figure 49, A).
2.4 Characterization of the importance of RomR for reversals

2.4.1 ΔromR and ΔromX mutants respond to IAA

The RomR response regulator was suggested to be the central output of the Frz system because its substitution of the phosphorylatable Asp residue in the receiver domain to Glu, which has been reported to mimic the phosphorylated state in many response regulators (Domian et al., 1997), and Gln, which cannot be phosphorylated and mimics the non-phosphorylated form, caused a hyper-reversal and hypo-reversal phenotype, respectively (Keilberg et al., 2012, Leonardy et al., 2007). Thus, phosphorylation of RomR could link signaling by the Frz system to the polarity switch (Leonardy et al., 2007). Additionally, it has been shown that a ΔromR mutant does not
reverse while cells are moving by T4P-dependent motility (Guzzo et al., 2015). As reported here, cells that lack RomX also do not reverse while moving by T4P-dependent motility. To test whether RomX and/or RomR are essential to generate an output of the Frz system, we artificially increased Frz signaling in ΔromX and ΔromR mutants by adding isoamyl alcohol (IAA) to cells. The exact target of IAA is not known; however, its action strictly depends on the Frz system and IAA activates the Frz system (McBride et al., 1992, Bustamante et al., 2004).

We tested T4P-dependent motility on 0.5% agar in the presence of gradually increasing IAA concentrations (Figure 50). In the absence of IAA, WT formed flares, the ΔpilA mutant that served as a negative control did not, and the ΔfrzE mutant showed misformed flares. The ΔromX and ΔromR mutants formed short misformed flares. Importantly, increasing concentrations of IAA caused the formation of shorter flares in all strains except the ΔfrzE mutant that was not affected by IAA (Figure 50). We conclude that IAA in the absence of RomX or RomR still induces reversals.

We also tested the effect of IAA on gliding motility on 1.5% agar (Figure 50). Without IAA, WT showed single cells at the colony edge while the ΔaglQ mutant did not. ΔfrzE mutant showed single cells at the colony edge comparable to WT. For the ΔromX and ΔromR mutants, only movement of groups of cells was observed. Also, for gliding motility increasing concentrations of IAA caused fewer visible single cells at the colony edges in all strains except the ΔfrzE mutant that was not affected by IAA (Figure 50). We conclude that IAA in the absence of RomX or RomR still induces reversals. In total, these data suggests that neither RomX nor RomR are essential for the output of the Frz system.
Figure 50. The ΔromX and ΔromR strains are sensitive to IAA.
Motility assays as in Figure 21; right part show colony after 24h on the 0.5% CTT, 0.5% agar plates without and with indicated IAA concentrations. Light part show colony after 24h on the 0.5% CTT, 1.5% agar plates without and with indicated IAA concentrations. Bars, 1000 μm (T4P-dependent motility – left part), 50 μm (gliding motility – right part).

To confirm the effect of IAA on motility observed in the population-based assay, we analysed single cells moving by T4P-dependent motility in the absence and presence of IAA. As shown in Figure 51, the ΔaglQ mutant reversed on average 0.62 times in 10 minutes while the ΔaglQΔromR and ΔaglQΔromX mutants did not reverse. As reported previously (Blackhart & Zusman, 1985), the ΔaglQΔfrzE mutant did not reverse. However, in the presence of 0.3% IAA, the reversal frequency of the ΔaglQ, ΔaglQΔromX and ΔaglQΔromR mutants increased while the ΔaglQΔfrzE mutant was not affected by IAA (Figure 51). Therefore, artificial activation of Frz signaling increases the reversal frequency of the normally non-reversing ΔromX and ΔromR mutants, supporting that RomR and RomX are not essential to generate the output of the Frz system.
Figure 51. The ΔromX and ΔromR mutants are IAA sensitive.
Reversal frequency of WT and selected mutants on 1% methylcellulose with or without 0.3% IAA. Shown are boxplots of the measured reversals of isolated cells for 10 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. n>50.

2.4.2 Mimicking and blocking of possible RomR phosphorylation site has no effect on the motility

To further investigate how RomR is important for reversals induced by the Frz system, we ectopically expressed RomR$^{D53E}$, which has been suggested to mimic the phosphorylated state of RomR (Leonardy et al., 2007, Keilberg et al., 2012), and RomR$^{D53N}$, which cannot be phosphorylated and mimics the non-phosphorylated form (Leonardy et al., 2007, Keilberg et al., 2012), under the control of the strong, constitutively active pilA promoter. Unfortunately, both variants of RomR did not accumulate (data not shown). Next, romR, romR$^{D53E}$ and romR$^{D53N}$ were expressed ectopically under control of a vanillate inducible promoter.
In the absence of inducer, the strains expressing $romR$, $romR^{D53N}$ or $romR^{D53E}$ showed the $\Delta romR$ phenotype. On plates with and without 50 µM vanillate, WT showed flares on 0.5% agar and single cells at the colony edge on 1.5% agar (Figure 52). Therefore, vanillate has no effect on motility. Furthermore, strains that expressed $romR$, $romR^{D53N}$ or $romR^{D53E}$ under the control of the inducible promotor were indistinguishable from WT in the presence of 50 µM vanillate (Figure 52). Thus, blocking and mimicking of possible phosphorylation site of RomR has no effect on the motility.

To further investigate the effect of possible RomR phosphorylation on reversals, we examined single cells expressing $romR$, $romR^{D53N}$ or $romR^{D53E}$. Synthesis of the RomR variants was induced in suspension with 50 µM vanillate and samples were withdrawn at different time points to determine the accumulation level of RomR variants. In the absence of vanillate, none of the RomR variants accumulated (Figure 53, A). After
RESULTS

2 hours, the accumulation level of the RomR variants had reached the WT level. After 3 hours, the level of the RomR variants were higher than in WT (Figure 53, A). At all time points, the strains expressing romR, romRD53N and romRD53E behaved similarly with respect to reversals (Figure 53, B).

**Figure 53.** Mimicking or blocking phosphorylation site of RomR has no effect on the reversal frequency when cell are moving by gliding motility.

(A) Immunoblots of RomR. Details of experiment are explained in text. Total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using α-RomR. Immunoblot with α-PilC serves as a loading control.

(B) Reversal frequency of WT and selected mutants on 1.5% TPM agar. Shown are boxplots of the measured reversals of isolated cells for 15 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. ΔromR was not moving as single cells in the tested conditions.

In previous experiment, we induced synthesis of the RomR variants over time. Next, we analysed the effect of the RomR variants on the reversal frequency in steady state after 8 generation of induction in the presence of 15 µM vanillate (Figure 54, A). As expected, romR, romRD53N and romRD53E mutants that were growing for 8 generations without inducer showed no RomR accumulation (Figure 54, A) and no sufficient movement to calculate reversals. In the presence of 15 µM vanillate, RomR, RomRD53N
and RomR^{D53E} accumulated to WT level (Figure 54, A) and romR, romR^{D53N} and romR^{D53E} mutants had the same reversal frequency as WT (Figure 54, B). Therefore, blocking and mimicking of possible RomR phosphorylation have no effect on the reversal frequency.

Figure 54. Mimicking or blocking phosphorylation site of RomR has no effect on the reversal frequency.

(A) Immunoblots of RomR. Details of experiment explained in text. Total protein (1 mg per lane) was separated by SDS–PAGE and analyzed by immunoblot using α-RomR. Immunoblot with α-PilC serves as a loading control.

(B) Reversal frequency of WT and selected mutants on 1.5% TPM agar. Shown are boxplots of the measured reversals of isolated cells for 15 min, boxes enclose the 25th and 75th percentile, whiskers represent the 10th and 90th percentile, and dots outliers. ΔromR was not moving as single cells in the tested conditions.
3. Discussion

To identify additional proteins that may regulate motility in *M. xanthus* we sought in 1611 prokaryotic genomes for genes with the same genomic distribution as *romR*. We identified two genes that encode uncharacterized proteins without any conserved domains that could indicate their function. These two proteins, named RomX and RomY, have not been identified before in screens for genes involved in gliding, T4P-dependent motility or regulation of reversals. We speculate that the small size of *romX* (264 bp) and *romY* (651 bp) decreases the possibility of random insertion during transposon mutagenesis. Therefore, bioinformatics analysis serves as a powerful tool to find new proteins potentially involved in different cellular processes.

3.1 RomX regulates cell polarity together with MglA, MglB and RomR


Generally, *romX* is found in genomes that contain *mglA, mglB* and *romR* suggesting that RomX is functionally connected to MglA, MglB and RomR. In this study, we have shown that RomX is important for both motility systems. Furthermore, more detailed studies show that RomX is necessary for gliding motility and regulates reversals in the T4P-dependent motility system. Furthermore, epistasis experiments showed that RomX acts in the same pathway as MglA, MglB and RomR. Δ*romX* and Δ*romR* mutants share the same motility phenotype and the double deletion Δ*romRΔromX* mutant did not show an additive phenotype. Strikingly, deletion of *mglB* partially restored gliding motility in the Δ*romX* and Δ*romR* mutants, suggesting that MglB acts downstream of RomX and RomR. Furthermore, epistasis experiments suggest that RomX and RomR have opposite function to MglB. Finally, strains that lack RomX or RomR and have MglA locked in the
GTP bound form show completely restored gliding motility and reversals. In conclusion, we suggest that RomX and RomR act upstream of MglB and MglA-GTP (Figure 55).

![Figure 55. RomX together with RomR act upstream of MglB and MglA-GTP in the regulation of polarity in motility of M. xanthus. Arrow shows stimulation while T-bar shows inhibition.]

RomX localizes in a bipolar asymmetric pattern similarly to RomR. RomX polar localization depends on RomR, and MglA polar localization depends on RomX and RomR. However, RomR polar localization is RomX independent. MglA only binds to the cell pole in its active GTP bound form (Leonardy et al., 2010, Zhang et al., 2010, Keilberg et al., 2012, Zhang et al., 2012). MglA locked in the GTP-bound form (MglA_{Q82A}) localizes in a bipolar pattern; however, polar localization is strongly impaired in the absence of RomX and RomR. These data suggest that at the leading cell pole RomR binds RomX that, in turn, recruits MglA-GTP. To further investigate the possibility that RomX and RomR form a functional protein complex at the cell pole, we compared localization of RomR and RomX in the absence of MglA and MglB. Both, RomR and RomX localize mainly unipolar in the absence of MglA. This is a common pattern of localization of the motility regulatory proteins in the absence of MglA and was previously described for FrzS, MglB, RomR and MglC (Zhang et al., 2012, Keilberg et al., 2012, McLoon et al., 2015). By contrast, in the absence of MglB, MglA, RomR and RomX change localization toward bipolar symmetric. These localization patterns support the idea that RomR and RomX form a complex at the cell pole. Furthermore, in pull-down experiments with purified proteins, RomR interacts with RomX and RomX interacts with MglA-GTP. Furthermore, the RomR/RomX complex interacts with MglA-GTP. Moreover, RomR does not interact with MglA-GTP and MglA-GDP. Together with the observations from bacterial two hybrid system, where direct interactions RomX-RomR and RomX-MglA were observed, we propose a model in which RomR binds to the leading pole and, in turn, recruits RomX. Next, the RomR/RomX complex binds MglA-GTP to form a RomR/RomX/MglA-GTP complex.
3.2 RomR/RomX complex acts as an MglA GEF

Epistasis analysis has shown that MglA-GTP acts downstream of RomX and RomR. Moreover, quantitative analysis of motility parameters (velocity and reversal frequency) of gliding cells showed no differences between \textit{mglA}^{Q82A}, \textit{mglA}^{Q82A}\textit{\Delta}romX and \textit{mglA}^{Q82A}\textit{\Delta}romR mutants. Thus, the gliding motility defects caused by \textit{\Delta}romX and \textit{\Delta}romR mutations can be bypassed by high MglA-GTP concentration. Similarly, the defects in gliding motility in the absence of RomR or RomX are corrected in the absence of MglB. Based on these observations we hypothesized, that the RomR/RomX complex not only functions as a polar binding determinant of MglA-GTP but also plays a role in the regulation of the nucleotide-bound state of MglA. \textit{In vitro} analyses with purified proteins demonstrated that neither RomR alone, RomX alone nor the RomR/RomX complex inhibit MglA GTP hydrolysis. Similarly, RomR and/or RomX do not inhibit MglB GAP activity. However, the RomR/RomX complex stimulates MglA nucleotide exchange. Thus, we conclude that the RomR/RomX complex is an MglA GEF. In total, the RomR/RomX complex has two functions. First, it has MglA GEF activity. Second, it binds MglA-GTP. \textit{In vivo} both activities contribute to a high local concentration of MglA-GTP at the leading pole.

The interaction between GTPases and their cognate GEFs have been extensively studied in eukaryotes. In the current model, a GEF associates with the GDP-bound GTPase forming a low affinity complex. Upon nucleotide release, the GEF forms a high affinity complex with nucleotide free GTPase. Upon binding of GTP, a low affinity complex is formed again and, finally, the GEF is released (Bos \textit{et al.}, 2007). The mechanism of the RomR/RomX GEF activity remains to be investigated. In particular, it will be interesting to investigate whether the catalytic part of the complex is formed by both proteins or whether one of the two proteins undergoes structural changes in the RomR/RomX complex in that way allowing it to function as a GEF. Of note, the active site of the TRAPP complex, which has Rab1 GEF activity, is assembled from several subunits (Cai \textit{et al.}, 2008). Activation of a GEF by an interaction partner was described for the Ras GEF Sos that is activated by the interaction with the adaptor Grb2 (Aronheim \textit{et al.}, 1994) and p115RhoGEF in which GEF activity is activated by interaction with subunits of heterotrimeric G proteins (Fukuhara \textit{et al.}, 2001).

We have shown that RomR/RomX complex has two distinct activities with respect to MglA: GEF and binding of MglA-GTP at the leading cell pole. GEF proteins that also bind to their cognate small Ras GTPase in their GTP-bound form were described for some
eukaryotic GEFs. Detailed biochemical studies revealed that the interaction is part of a positive feedback loop and activates the GEF, in turn, leading to accumulation of GTPase-GTP (Margarit et al., 2003, Menetrey et al., 2007, Richardson et al., 2012, Cohen et al., 2007, Chen et al., 2010, Lin et al., 2006). So far, we do not have any evidence that RomR/RomX and MglA localization involves a similar positive feedback loop.

In eukaryotes, the GEFs of the five subfamilies of small GTPases are structurally unrelated. Interestingly, neither RomX nor RomR show sequence similarity to known GEF proteins. Therefore, the RomR/RomX complex may represent a new family of GEFs. Previously, it has been shown that the GAP of MglA, MglB, belongs to a new group of GAPs proteins with a unique mechanism of stimulating GTP hydrolysis (Miertzschke et al., 2011). Interestingly, MglA and MglB are widespread in prokaryotes (Keilberg et al., 2012, Wuichet & Søgaard-Andersen, 2014) whereas RomR and RomX have a more narrow distribution, suggesting the existence of other GEFs for MglA-like proteins.

3.3 Formation of focal adhesion complexes depends on RomX and RomR

MglA-GTP is an integral part of focal adhesion complexes. MglA-GTP is incorporated into the focal adhesion complexes at the leading cell pole and focal adhesion complexes are disassembled at the lagging cell pole due to MglB GAP activity there (Zhang et al., 2010, Patryn et al., 2010, Keilberg et al., 2012, Treuner-Lange et al., 2015). Moreover, experiments with inactive MglA-GDP provided evidence that only active, MglA-GTP stimulates gliding motility (Leonardy et al., 2010). In the non-gliding ΔromX and ΔromR mutants, MglA does not localize to focal adhesion complexes. Moreover, when we used AglZ-YFP as a readout for formation of the focal adhesion complexes, we confirmed lack of focal adhesion complex formation in the absence of RomX and RomR. We speculate that lack of gliding motility and focal adhesion complexes could be a result of non-sufficient MglA-GTP concentration. To test this hypothesis, we generated an in frame deletion of mglB in the backgrounds of mglA-mVenusΔromX, mglA-mVenusΔromR and mglA-mVenusΔromXΔromR. In the absence of MglB and RomX and/or RomR, we observed MglA-mVenus in the focal adhesion complexes. Moreover, we confirmed these observations by detecting AglZ-YFP in the focal adhesion complexes. Thus, RomX and RomR are necessary for focal adhesion complex formation only when the MglA nucleotide-bound state is tightly regulated by its cognate GAP MglB. Moreover, these data are in agreement with the analysis of gliding motility in single cells in which we observed partial recovery of the gliding motility in the double ΔmglBΔromX and
ΔmglBΔromR mutants. Also, these observations are consistent with our model in which RomR/RomX complex acts as an MgIA GEF and stimulates MgIA-GTP concentration at the leading cell pole and in focal adhesion complexes. These observations also suggest that a sufficient level of MgIA-GTP accumulates in the absence of MgIB, RomR and RomX to stimulate assembly of the gliding motility complexes. Along the same lines, even in the presence of MgIB, a sufficiently high level of MgIA-GTP accumulates to stimulate T4P-dependent motility.

3.4 RomX and RomR localize to the focal adhesion complexes

Quantification of the fluorescence signal of the MglB-mCherry fusion, which is only detected as a full-length protein in immunoblots, showed that more than 90% of MglB is localized diffusely in the cytoplasm. Based on this observation, we hypothesized that RomR/RomX complex may localize to the focal adhesion complexes to replenish MgIA-GTP in case of MglB stimulated GTP hydrolysis by MgIA. To this end, we observed that the motor protein AglQ colocalizes with RomX and AglZ colocalizes with RomR. All proteins are incorporated into the focal adhesion complexes at the leading cell pole and disassemble near lagging cell pole. Interestingly, in the double ΔmglBΔromR mutant we did not observe RomX in the focal adhesion complexes. By contrast, RomR localizes to focal adhesion complexes in the ΔmglBΔromX mutant. Furthermore, RomR localizes to focal adhesion complexes in the absence of RomX in the strain with MgIA locked in the GTP bound form. Therefore, we speculate that RomR interacts directly with one or more of the cytoplasmic components of the gliding machinery. Additionally, this data strongly suggest that RomR is a binding scaffold for RomX in the focal adhesion complexes. We conclude that RomX and RomR are new cytoplasmic components of the focal adhesion complexes; however, as opposed to all other proteins in these complexes, RomR and RomX are only essential for formation of these complexes in the presence of MgIB.

3.5 RomX and RomR are important for attachment, stability and directionality of the focal adhesion complexes

Previously, the focal adhesion complexes were described to be assembled at the leading cell pole, stay stationary in the respect to the substratum in moving cell and disassemble near the lagging cell pole (Mignot et al., 2007, Jakobczak et al., 2015).
Recent studies also revealed clusters moving along the cell body. In moving cells, these clusters are assembled at the leading cell pole and move towards the lagging cell pole. In non-moving cells multiple dynamic focal adhesion complexes assemble at one cell pole and move to the opposite cell pole. Dynamic focal adhesion complexes are thought to be unattached motility complexes (Faure et al., 2016).

Quantitative analysis of the movement of focal adhesion complexes within cells showed that in the absence of RomX and RomR in strains with high MglA-GTP concentration (ΔmglB mutants or with MglA locked in the GTP-bound form) these complexes are assembled and more motile than in WT. We would like to propose that these motile focal adhesion complexes do not produce or produce less force to propel a cell. Therefore, we speculate that RomX and RomR are important for the correct assembly of functional focal adhesion complexes. Moreover, in ΔmglBΔromX, ΔmglBΔromR and ΔmglBΔromXΔromR mutants we observe a shorter lifetime of the focal adhesion complexes. Therefore, RomX and RomR are also important for their stability. However, shorter lifetime can be explained by lower MglA-GTP concentration than in WT, as there is no effect on the stability of focal adhesion complexes in mglA<sup>Q82A</sup>-mVenusΔromX and mglA<sup>Q82A</sup>-mVenusΔromR mutants. We observe only partially restored gliding motility in the ΔmglBΔromX and ΔmglBΔromR mutants and fully restored gliding in mglA<sup>Q82A</sup>ΔromX and mglA<sup>Q82A</sup>ΔromR mutants, thus, we speculate that the main motility defect is coming from instability of the focal adhesion complexes. However, we cannot exclude the additive effect of the instability and intracellular mobility of the focal adhesion complexes.

Qualitative analysis of the focal adhesion complexes in the absence of romX and romR in the strains with elevated MglA-GTP concentration shows loss of the directionality. For some cells, we observe focal adhesion complexes that are assembled at the opposite cell poles and are moving in opposite directions. Loss of the directionality is observed in the strains with increased mobility of the focal adhesion complexes. However, it remains unclear, if increase of mobility leads to loss of directionality.

Taken together, we conclude that RomX and RomR by localizing to the focal adhesion complexes help to maintain their attachment, stability and directionality. It remains unclear whether the only role of RomR is to bring RomX to the focal adhesion complexes. Lack of attachment and loss of directionality of focal adhesion complexes are observed when RomX is absent and RomR is present in the focal adhesion complexes. It remains unknown how RomR/RomX complex binds to the focal adhesion complexes and
what the mechanism of action is to stimulate focal adhesion complexes attachment and directionality.

### 3.5 RomX/RomR establishes front-rear polarity, cell asymmetry and gliding motility by its triple function

We propose the following three functions of RomR and RomX: First, RomR binds RomX at the leading cell pole. Second, the RomR/RomX complex by its GEF activity stimulates the formation of a local pool MglA-GTP at this pole. RomR/RomX binds MglA-GTP forming a RomR/RomX/MglA-GTP complex at the leading cell pole. Third, the RomR/RomX/MglA-GTP complex is loaded onto the focal adhesion complexes. The RomR/RomX complex stimulates the stability of focal adhesion complexes by replenishing MglA-GTP in these complexes in case of MglB stimulated MglA GTPase activity. RomX or the RomX/RomR complex by a so far unknown mechanism is giving directionality and increases adhesion of the focal adhesion complexes. At the lagging cell pole, GTPase activity of MglA-GTP is stimulated by MglB, GTP is hydrolysed to GDP and focal adhesion complex is disassembled (Figure 56, A and B).

![Figure 56. Model for RomX/RomR complex actions in the cell.](image)

**A** Spatial organization of the motility regulatory proteins. Description in text. Arrow from GDP to GTP indicates RomR/RomX GEF activity on MglA. Arrow from GTP to GDP indicates hydrolysis of GTP by MglA stimulated by MglB.

**B** Model for MglA GTP/GDP cycle.
In eukaryotes, small Ras-like GTPases play an important role in regulation of cell polarity and motility and function in spatially organized manners (Jaffe & Hall, 2005, Charest & Firtel, 2007, Kortholt & van Haastert, 2008, Chiou et al., 2017). Small GTPases may be spatially regulated by the colocalization with their cognate GEF. This has been described for Cdc42 and Cdc24 from Saccharomyces cerevisiae at the bud site (Chiou et al. 2017) and for Rab5 and its cognate GEF Rabex-5 in the early endosomes (Blumer et al., 2013). Alternatively, the activated GTPase colocalizes with its GEF(s) and is spatially separated from its cognate GAP, as is the case during invagination of epithelial cells during embryogenesis in Drosophila where the two Rho1 GEFs colocalize with Rho1 at the apical membrane while the single Rho1 GAP localizes to the basolateral membrane (Simões et al., 2006). Thus, colocalization of MglA with the RomR/RomX complex, separation from MglB and spatial control of MglA activation are similar to mechanisms observed in eukaryotes. Particularly challenging question is how MglA, MglB, RomR and RomX become asymmetrically localized. For some of above described eukaryotic systems, positive feedback plays a crucial role in acquiring asymmetry. However, as mentioned before, there is no evidences for such a positive feedback loop for RomR/RomX and MglA. More detailed biochemical studies are required to get better insight into how asymmetry is established.

3.6 RomY regulates cell polarity together with MglA, MglB, RomR and RomX

Generally, romY occurs in genomes together with mglA, mglB and romR suggesting a functional connection to regulation of the cell motility. We confirmed our predictions by analyzing motility of an in-frame romY deletion strain. Lack of RomY causes formation of shorter flares on 0.5% agar and less single cells movement on 1.5% agar in comparison to WT. Thus, RomY seems to be important for regulation of motility rather than movement per se. Moreover, the finding that the ΔromY mutant hyper-reverses supports the hypothesis that RomY plays a role in regulation of the reversal frequency in both motility system. Epistasis experiment shows that RomY acts in the same pathways as MglA, MglB, RomR and RomX. Moreover, ΔmglB and ΔromY mutants share the same motility phenotype and the double ΔmglBΔromY mutant does not show additive phenotype. Furthermore, quantitative reversal frequency analysis shows no statistically relevant difference between ΔmglB, ΔromY and ΔmglBΔromY mutants. For ΔromRΔromY and ΔromXΔromY mutants, we observe partially restored gliding motility. Based on our previous studies on ΔromXΔmglB and ΔromRΔmglB mutants and
observations of the partial complementation of the gliding motility, we speculate that lack of RomY, similarly to lack of MglB, increases MglA-GTP concentration.

To further understand RomY function in the regulation of polarity we determined RomY localization dependencies. RomY localizes mainly at the lagging cell pole. Surprisingly, in the absence of RomY, MglA localizes mainly in the bipolar pattern. Furthermore, RomX shows strong shift towards bipolar symmetric localization in the absence of RomY. RomR, in the absence of RomY, does not show shift towards more symmetric localization, however we observe strongly increased signal from the cell body. Interestingly, similar localization changes for MglA, RomX and RomR we observe in the absence of MglB. Based on the data we thought of two possible mechanisms of RomY action: RomY stimulates MglB GAP activity or RomY has GAP activity on its own. However, we did not observe direct interaction between RomY and MglB. Of note, we are unable to exclude direct interactions between MglB and RomY based on the negative results from the bacterial two hybrid system. Nevertheless, we observe that polar localization of RomY strongly depends on MglB suggesting a functional connection between these two proteins. Interestingly, we have found that RomY directly interacts with MglA and RomX. The function of the MglA-RomY interaction is unknown. RomY may stimulate GTPase activity directly or by interacting with MglA may stimulate MglB GAP activity. Further studies are necessary to identify the function of RomY in regulation of polarity. GTPase assay could also directly answer whether RomY is a GAP, directly stimulates MglA GTPase activity or stimulates MglB GAP activity.

In two (out of 28 species that romY was originally found) species romY is present without mglB arguing against the idea that RomY stimulates MglB GAP activity. Well described example of organism with mglA, romR and romY but without mglB is Bdellovibrio bacteriovorus. B. bacteriovorus is small predatory deltaproteobacterium, which invades other Gram-negative bacteria wherein it replicates. Bdellovibrio can encounter their prey by fast motility, driven by flagellum in liquid environment (Iida et al., 2009), or by twitching motility driven by T4P on a solid surface. B. bacteriovorus does not show social T4P-dependent motility as in M. xanthus. B. bacteriovorus requires T4P for invasion into the prey cell periplasm. The ΔmglA B. bacteriovorus strain is hypopiliated and unable to invade prey (Lambert et al., 2011). B. bacteriovorus MglA (MglABb) shares significant sequence similarity to MglA from M. xanthus (MglAMx) with 64% protein identity and 82% similarity (Milner et al., 2014). Major difference is in a P-loop region that is important for GTP hydrolysis (Miertschke et al., 2011). The P-loop
region of \( \text{MglA}_{\text{Bb}} \) contains a serine at residue 21; the corresponding G21S substitution in eukaryotic RAS proteins locks them in the GTP-bound form (Sei et al., 2009). Moreover, \( \text{MglA}_{\text{Mx}}^{G21V} \) is insensitive to \( \text{MglB} \) and thus locked in the GTP bound form (Miertzschke et al., 2011). Taken all together, \( \text{MglA}_{\text{Bb}} \) is most probably locked in the GTP bound state. Of course, we are not able to exclude that \( \text{RomY}_{\text{Bb}} \) has GAP activity and by a mechanism that is different from that of \( \text{MglB} \), stimulates \( \text{MglA}_{\text{BB}} \) GTPase activity and thus \( \text{MglA}_{\text{Bb}} \) cycles between GTP and GDP bound state. However, up to date the function of \( \text{RomY} \) in \( B. \text{bacteriovorus} \) is unknown.

![Figure 57. Model of polar localization of proteins involved in the regulation of motility in \( M. \text{xanthus} \).](image)

Proteins colour on the model correspond to the colours in the legend.

In this study, we identified two new proteins that are important for the polarity of motility in \( M. \text{xanthus} \), \( \text{RomX} \) and \( \text{RomY} \). Furthermore, we have found that \( \text{RomX} \) and \( \text{RomY} \) show dynamic polar localization similarly to other proteins that are involved in the regulation of motility – \( \text{RomR} \), \( \text{MglA} \) and \( \text{MglB} \) (Figure 57). All these proteins mutually depend on each other for the correct localization. We have shown that, two proteins localize to the cell pole independently – \( \text{RomR} \) and \( \text{MglB} \). We hypothesize that \( \text{MglB} \) form directly or indirectly a localization landmark for \( \text{RomY} \). We have shown that \( \text{RomR} \) recruits \( \text{RomX} \), which, in turn, recruits \( \text{MglA} \)-GTP. So far, the exact mechanism how \( M. \text{xanthus} \) cells establish correct polarity and protein asymmetry remains unclear. Further protein interaction analysis together with extensive localization studies could be helpful in gaining detailed insight into regulation of polarity.

3.7 \( \text{RomX} \) and \( \text{RomR} \) are not essential for generating an output from the Frz system

\( \text{RomR} \) was previously thought to be the central output of the Frz system. However, direct connection between \( \text{RomR} \) and the Frz system is not known up to date (Leonardy et al., 2007, Keilberg et al., 2012, Guzzo et al., 2015). Our detailed studies show that
ΔromX and ΔromR mutants do not glide but still show T4P-dependent movement on WT level, however they do not reverse. To test whether RomX or/and RomR may be essential for generating an output from the Frz system we took advantage of IAA that simulates Frz dependent reversals. On the population level, using motility assay as a read out, we have found that both ΔromX and ΔromR mutants respond to the IAA in the same way as WT. Moreover, analysis of single cells shows that reversals of the ΔromX and ΔromR mutants can be artificially increased by the addition of IAA. However, stimulation of reversals does not reach the WT level. Previously, it has been shown that ΔromR mutant is insensitive to the IAA. This observation was done by tracking oscillations of FrzS-YFP, protein important for T4P-dependent motility that localize mainly at the leading cell pole (Guzzo et al., 2015). We suggest that lack of RomR decreases dynamic oscillations of FrzS and it cannot be correlated with cellular reversal. Moreover, our studies in which cells were growing for 24 hours on medium with the IAA clearly show that ΔromR and ΔromX mutants are IAA sensitive. The data does not definitively rule out that RomX and RomR are not a part of signal transduction from the Frz system. It has been shown that reversals of the ΔfrzZ mutant, that lacks the output protein of the Frz system, could be induced by the IAA and this stimulation is at a lower level than for WT (Guzzo et al., 2015). MglA-GTP at the leading cell pole is thought to play a regulatory role for Frz dependent reversals. In the current model of the Frz system dependent reversals, phosphorylated output of the Frz system (FrzZ) interacts at the leading pole with MglA-GTP and releases it from this cell pole (Kaimer & Zusman, 2013). We hypothesize that artificially increased activity of the Frz system and thus, high level of phosphorylated output(s) of this system, can bypass the absence of MglA-GTP at the leading cell pole in the ΔromR and ΔromX mutants.

RomR contains response regulator domain with putatively phosphorylatable conserved aspartic acid residue (Leonardy et al., 2007, Keilberg et al., 2012). However, phosphorylation of RomR has not been shown and a kinase that could phosphorylate RomR is not known. Previously, it was hypothesized that the Frz system could phosphorylate RomR by an unknown mechanism. For many response regulators it was shown that phosphorylation of a conserved aspartic acid residue is required for activity (Stock et al., 2000). Blocking phosphorylatable residue by a mutation of aspartic acid (D) to the asparagine (N) results in loss of the ability of phosphorylation and thus, loss of activity. Mutation of aspartic acid (D) to glutamic acid (E) was shown to partially mimic phosphorylation state for some response regulators (Domian et al., 1997). For romR
insertional mutant expressing $\text{romR}^{D53N}$-gfp under the control of the strong, constitutively active pilA promoter, no reversals were observed for 50 cells for 10 minutes. Phosphorylation mimicking mutant ($\text{romR}::\text{nptII P}_{\text{pilA}}\text{romR}^{D53E}$-gfp) has shown 1.5 fold more reversals than strain expressing $\text{romR}$-gfp (Leonardy et al., 2007). Moreover, later studies with RomR-GFP variants expressed under the pilA promoter in the in frame romR deletion mutant have shown similar patterns. Blocking RomR phosphorylation resulted in hypo-reversing strain, however, some reversals were still observed. Mimicking of the RomR phosphorylation generated hyper-reversing strain but this effect was less prominent than in the previous studies (Keilberg et al., 2012). We revisited function of potential RomR phosphorylation by testing motility phenotypes of mutants that express RomR variant that mimics phosphorylation or is blocked in the phosphorylation site.

Our population-based experiments show no differences between wild type version of RomR, RomR with blocked phosphorylation and mimicking phosphorylation for the T4P-dependent and gliding motility. In our experimental conditions, all three tested variants of RomR behave like WT. We confirmed our observations in two independent experiments in which we analyzed reversals of single gliding cells. We do not observe statistically significant differences in reversal frequency between WT and strains that express RomR, RomR$^{D53N}$ and RomR$^{D53E}$ as an only copy of RomR. Thus, RomR phosphorylation is not regulating reversal frequency under the conditions tested. Based on the data, we concluded that RomR is not essential for generating an output from the Frz system. In the experiment in which we induced $\text{romR}$, $\text{romR}^{D53N}$ and $\text{romR}^{D53E}$ over the time, we found that a low level of RomR causes cells to hypo-reverse. We hypothesize that strong overexpression of RomR may lead to the hyper-reversing cells. However, so far we do not have any experimental evidence to confirm this hypothesis, future studies with RomR overexpression could be performed to give directly answer whether RomR overexpression will lead to hyper-reversing cells. Nevertheless, we would like to propose that RomR concentration effect on the reversal frequency could explain previously published observations (Leonardy et al., 2007, Keilberg et al., 2012).
4. Materials and Methods

4.1 Chemicals, equipment and software

All chemicals, enzymes and kits are listed together with their suppliers in the Table 1. All devices together with their manufacturers and applications are listed in the Table 2. Specific softwares together with their manufacturers are listed in the Table 3.

<table>
<thead>
<tr>
<th>Reagents, enzymes, antibiotics and kits</th>
<th>Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reagents</td>
<td>Supplier</td>
</tr>
<tr>
<td>Chemicals</td>
<td>Roth (Karlsruhe), Merck (Darmstadt), Sigma-Aldrich (Taufkirchen)</td>
</tr>
<tr>
<td>Media components, agar</td>
<td>Roth (Karlsruhe), Merck (Darmstadt), Difco (Heidelberg), Invitrogen (Darmstadt)</td>
</tr>
<tr>
<td>Rabbit antisera</td>
<td>Eurogentec (Seraing, Belgium)</td>
</tr>
<tr>
<td>Goat anti-rabbit IgG, goat anti-rabbit IgG</td>
<td>Pierce/Thermo Scientific (Dreieich)</td>
</tr>
<tr>
<td>DyLight 549</td>
<td>Pierce/Thermo Scientific (Dreieich)</td>
</tr>
<tr>
<td>Anti-GFP monoclonal antibody</td>
<td>Roche Diagnostics GmbH (Mannheim)</td>
</tr>
<tr>
<td>Anti-mCherry</td>
<td>BioVision</td>
</tr>
<tr>
<td>Anti-mouse sheep IgG antibody, horseradish peroxidase linked</td>
<td>GE Healthcare Europe GmbH (Freiburg)</td>
</tr>
<tr>
<td>Nitrocellulose membrane</td>
<td>GE Healthcare Europe GmbH (Freiburg)</td>
</tr>
<tr>
<td>SDS gel electrophoresis size standards</td>
<td>Pierce™ Thermo Scientific™ (Darmstadt)</td>
</tr>
<tr>
<td>Pageruler™ Plus Prestained Protein Ladder</td>
<td>Millipore Merck Chemicals GmbH (Schwalbach)</td>
</tr>
<tr>
<td>Luminata Western HRP Substrate</td>
<td>Millipore Merck Chemicals GmbH (Schwalbach)</td>
</tr>
<tr>
<td>Oligonucleotides</td>
<td>Eurofins MWG Operon (Ebersberg)</td>
</tr>
<tr>
<td></td>
<td>Invitrogen™ life technologies (Karlsruhe)</td>
</tr>
<tr>
<td>2-log DNA Ladder</td>
<td>New England Biolabs (NEB) (Frankfurt a. M.)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Enzymes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antarctic Phosphatase</td>
</tr>
<tr>
<td>T4 DNA Ligase</td>
</tr>
<tr>
<td>restriction enzymes</td>
</tr>
<tr>
<td>5 PRIME MasterMix</td>
</tr>
<tr>
<td>Phusion High-Fidelity DNA Polymerase</td>
</tr>
<tr>
<td>Q5 High-Fidelity DNA Polymerase</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Antibiotics</th>
</tr>
</thead>
<tbody>
<tr>
<td>kanamycin sulfate,</td>
</tr>
</tbody>
</table>
ampicillin sodium sulfate,
gentamycin sulfate,
oxytetracycline dehydrate,
tetracycline hydrochloride

Streptomycin sulfate

**Kits**

| DNA purification (plasmid DNA), | Zymo Research (Freiburg), |
| PCR purification, | Qiagen (Hilden), |
| Gel purification | Macherey-Nagel (Düren) |
| DNA purification (chromosomal DNA) | Epicentre Biotechnologies (Wisconsin, USA) |

**Table 2. Equipment**

<table>
<thead>
<tr>
<th>Application</th>
<th>Device</th>
<th>Manufacturer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cell disruption</td>
<td>Branson Sonifier 250, French pressure cell press</td>
<td>G. Heinemann (Schwäbisch Gmünd), SLM instruments (Urbana, IL)</td>
</tr>
<tr>
<td>Centrifugation</td>
<td>Centrifuge 5424, Centrifuge 5424 R, Multifuge X1 R, Mega Star 1,6R</td>
<td>Eppendorf (Hamburg), Thermo Scientific (Dreieich), VWR International GmbH (Darmstadt)</td>
</tr>
<tr>
<td>PCR</td>
<td>Mastercycler personal, Mastercycler epgradient</td>
<td>Eppendorf (Hamburg)</td>
</tr>
<tr>
<td>Thermomixer</td>
<td>Thermomixer compact</td>
<td>Eppendorf (Hamburg)</td>
</tr>
<tr>
<td>DNA illumination</td>
<td>UVT_20 LE</td>
<td>Herolab (Wiesloch)</td>
</tr>
<tr>
<td>DNA illumination and documentation</td>
<td>E-BOX VX2 imaging system</td>
<td>Bio-Rad (München)</td>
</tr>
<tr>
<td>Protein electrophoresis</td>
<td>Mini-PROTEAN® 3 cell</td>
<td>Bio-Rad (München)</td>
</tr>
<tr>
<td>Western blotting</td>
<td>TransBlot®Turbo™ Transfer System, Hoefer™ TE77</td>
<td>Bio-Rad (München), Amersham (Freiburg)</td>
</tr>
<tr>
<td>Chemiluminescence detection</td>
<td>Luminescent image analyzer LAS-4000</td>
<td>Fujifilm (Düsseldorf)</td>
</tr>
<tr>
<td>Microscopes</td>
<td>MZ75 Stereomicroscope, M205FA Stereomicroscope, DM IRE2 Inverted microscope, DMi8 Inverted microscope, DMi6000B inverted microscope, DM6000B</td>
<td>Leica (Wetzlar)</td>
</tr>
<tr>
<td>Determination of optical densities,</td>
<td>Ultrospec 2100 pro Spectrophotometer,</td>
<td>GE Healthcare Europe GmbH (Freiburg),</td>
</tr>
</tbody>
</table>
### MATERIALS AND METHODS

- **nucleic acids absorption**: Nanodrop ND-1000 UV-Vis spectrophotometer, DS11+ (Nanodrop (Wilmington), DeNovix Inc. (Wilmington))

- **Multifunktionsreared Infinite M200 pro with Monochromatoroptics**: Detection of absorption changes during enzymatic assay (Tecan Deutschnland GmbH (Crailsheim))

- **AKTA pure with Fraction collector F9-C**
  - 5 ml Strep Traap HP
  - 5 ml HiTrap Q HP
  - 5 ml MBPTrapHP
  - 5 ml HITrap Chelating HP
  - HiLoad 16/600 Superdex 200 pg
  - HiLoad 16/600 Superdex 75 pg
  - Size, affinity purification of recombinant proteins, anion exchange chromatography, exclusion chromatography (GE Healthcare Europe GmbH (Freiburg))

- **Measuring of mant fluorescence**: ISS PC1 spectrofluorometer with a cooled photomultiplier (ISS (USA))

<table>
<thead>
<tr>
<th>Table 3. Software Application</th>
<th>Software</th>
<th>Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Data analysis of microscopy pictures</td>
<td>Metamorph® v 7.7.5.0, ImageJ 1.51s</td>
<td>Molecular Devices (Union City, CA), Wayne Rasband (National Institutes of Health, USA)</td>
</tr>
<tr>
<td>Automatic detection of cells on the microscopy pictures</td>
<td>Oufi</td>
<td>Jacobs-Wagner Lab (Paintdakhi et al., 2016)</td>
</tr>
<tr>
<td>Automatic analysis of fluorescence signals</td>
<td>MATLAB R2016b</td>
<td>The MathWorks, Inc (Natick, USA)</td>
</tr>
<tr>
<td>Checking of DNA and proteins sequences, in silico cloning of plasmids and data management of DNA, protein and plasmid sequences.</td>
<td>Vector NTI advance software, suite 11, DNASTAR</td>
<td>Invitrogen™ life technologies (Karlsruhe), DNASTAR, Inc (Madison, USA)</td>
</tr>
<tr>
<td>Preparation of kymograph and its analysis</td>
<td>Icy</td>
<td>Icy - copyright 2017 - Institut Pasteur (de Chaumont et al., 2012)</td>
</tr>
</tbody>
</table>

### 4.2 Media

*E. coli* cells were cultivated in Luria-Bertani (LB) liquid media or on LB agar plates with 1.5% agar concentration. *M. xanthus* cells were cultivated in CTT media or on CTT
agar plates with 1.5% agar concentration. Motility assays of *M. xanthus* were performed on A- or S-motility plates. Media composition is described in Table 4.

**Table 4. Growth media for *E. coli* and *M. xanthus***

<table>
<thead>
<tr>
<th>Media</th>
<th>Composition</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>E. coli</strong></td>
<td></td>
</tr>
<tr>
<td>LB medium</td>
<td>1% (w/v) tryptone,</td>
</tr>
<tr>
<td></td>
<td>0.5% (w/v) yeast extract,</td>
</tr>
<tr>
<td></td>
<td>1% (w/v) NaCl</td>
</tr>
<tr>
<td>2 x YT</td>
<td>16 g Bacto tryptone,</td>
</tr>
<tr>
<td></td>
<td>5 g yeast extract,</td>
</tr>
<tr>
<td></td>
<td>5 g NaCl, pH 7.2</td>
</tr>
<tr>
<td>LB agar plates</td>
<td>LB medium, 1.5% (w/v) agar</td>
</tr>
<tr>
<td><strong>M. xanthus</strong></td>
<td></td>
</tr>
<tr>
<td>CTT</td>
<td>1% (w/v) Bacto casitone,</td>
</tr>
<tr>
<td></td>
<td>10 mM Tris-HCl pH 8.0,</td>
</tr>
<tr>
<td></td>
<td>1 mM potassium phosphate buffer pH 7.6,</td>
</tr>
<tr>
<td></td>
<td>8 mM MgSO4</td>
</tr>
<tr>
<td>CTT agar plates</td>
<td>CTT medium, 1.5% agar</td>
</tr>
<tr>
<td>CTT soft agar</td>
<td>CTT medium, 0.5% agar</td>
</tr>
<tr>
<td><strong>Motility assays</strong></td>
<td></td>
</tr>
<tr>
<td>A-motility plates</td>
<td>0.5% CTT, 1.5% agar</td>
</tr>
<tr>
<td>(Hodgkin &amp; Kaiser, 1977)</td>
<td></td>
</tr>
<tr>
<td>S-motility plates</td>
<td>0.5% CTT, 0.5% agar</td>
</tr>
<tr>
<td>(Hodgkin &amp; Kaiser, 1977)</td>
<td></td>
</tr>
<tr>
<td><strong>Microscopy media</strong></td>
<td></td>
</tr>
<tr>
<td>TPM agar</td>
<td>10 mM Tris-HCl pH 7.6,</td>
</tr>
<tr>
<td></td>
<td>1 mM KH2PO4 pH 7.6,</td>
</tr>
<tr>
<td></td>
<td>8 mM MgSO4, 1.5% agar</td>
</tr>
<tr>
<td>MC7</td>
<td>10 mM MOPS, pH 7.0,</td>
</tr>
<tr>
<td></td>
<td>1 mM CaCl2</td>
</tr>
<tr>
<td>Methylcellulose solution</td>
<td>MMC, 1% methylcellulose</td>
</tr>
<tr>
<td>Chitosan 100x solution</td>
<td>2M acetic acid, 15mg/ml chitosan</td>
</tr>
</tbody>
</table>
The adequate antibiotics were added to cultures when needed. For the protein induction in *E. coli* IPTG was added, for *M. xanthus* vanilate and for selection X-GAL or galactose were added (Table 5)

**Table 5. Additives used for *E. coli* and *M. xanthus***

<table>
<thead>
<tr>
<th>Additive</th>
<th>Final concentration</th>
<th>Dissolved in</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>E. coli</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ampicillin sodium sulfate</td>
<td>100 μg/ml</td>
<td>H₂O</td>
</tr>
<tr>
<td>Tetracyclin</td>
<td>15 μg/ml</td>
<td>99.99% ethanol</td>
</tr>
<tr>
<td>Kanamycin sulfate</td>
<td>50 μg/ml</td>
<td>H₂O</td>
</tr>
<tr>
<td>5-Brom-4-chlor-3-indoxyl-β-D-galactopyranosid (X-gal)</td>
<td>40 μg/ml</td>
<td>Dimethylformamide</td>
</tr>
<tr>
<td>IPTG</td>
<td>1mM</td>
<td>H₂O</td>
</tr>
<tr>
<td><strong>M. xanthus</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Kanamycin sulfate</td>
<td>50 μg/ml</td>
<td>H₂O</td>
</tr>
<tr>
<td>Oxytetracycline</td>
<td>10 μg/ml</td>
<td>0.1M HCl</td>
</tr>
<tr>
<td>Galactose</td>
<td>2.5%</td>
<td>H₂O</td>
</tr>
<tr>
<td>Isoamyl alcohol</td>
<td>0.03%-0.3%</td>
<td></td>
</tr>
<tr>
<td>Vanilate</td>
<td>15μM-150μM</td>
<td>H₂O (adjusted to pH 7.6 with KOH)</td>
</tr>
</tbody>
</table>

**4.3 Microbial methods**

**4.3.1 *E. coli* strains used in this study**

**Table 6. *E. coli* strains used in this study**

<table>
<thead>
<tr>
<th>Strain</th>
<th>Relevant characteristics</th>
<th>Source or reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mach1</td>
<td>ΔrecA1398 endA1 tonA ΔlacM15 ΔlacX74 hsdR(rK-mK+)</td>
<td>Invitrogen (Darmstadt)</td>
</tr>
<tr>
<td>TOP10</td>
<td>F- mcrA Δ(mrr-hsdRMS-mcrBC) ΔlacZΔM15 ΔlacX74 recA1 araD139 Δ(aara leu) 7697 galU galK rpsL (StrR) endA1 nupG</td>
<td>Invitrogen™ life technologies (Karlsruhe)</td>
</tr>
<tr>
<td>BTH101</td>
<td>F- cya-99 araD139 galE15 galK16 rpsL1 (StrR) hsdR2 mcrA1 mcrB1</td>
<td>Euromedex, (Souffelweyersheim, France)</td>
</tr>
<tr>
<td>Rosetta 2 (DE3)</td>
<td>F-ompT hsdSB(rB-mB-) gal dcm(DE3) pRARE2(CmR)</td>
<td>Novagen/Merck (Darmstadt)</td>
</tr>
</tbody>
</table>
4.3.2 *M. xanthus* strains used in this study

For strains containing plasmid integrated at the Mx8 *attB* site, the gene expressed including promoter driving the expression is indicated in brackets.

Table 7. *M. xanthus* strains used in this study

<table>
<thead>
<tr>
<th>Strain</th>
<th>Genotype</th>
<th>Source or reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>DK1622</td>
<td>Wild type</td>
<td>(Kaiser, 1979)</td>
</tr>
<tr>
<td>DK10410</td>
<td>Δ<em>pilA</em></td>
<td>(Wu &amp; Kaiser, 1996)</td>
</tr>
<tr>
<td>SA5293</td>
<td>Δ<em>algQ</em></td>
<td>(Jakobczak <em>et al.</em>, 2015)</td>
</tr>
<tr>
<td>SA3683</td>
<td>Δ<em>romX</em> Δ<em>pilA</em></td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA6974</td>
<td>Δ<em>romX</em>, Δ<em>pilA</em></td>
<td>This work</td>
</tr>
<tr>
<td>SA7123</td>
<td>Δ<em>romX</em>, Δ<em>algQ</em></td>
<td>Anna Potapova</td>
</tr>
<tr>
<td>SA5990</td>
<td>Δ<em>romX</em>, <em>attB</em>::pDK131 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;-yfp)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA6997</td>
<td>Δ<em>romX</em>, <em>attB</em>::pDK157 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3300</td>
<td>Δ<em>romR</em></td>
<td>(Keilberg <em>et al.</em>, 2012)</td>
</tr>
<tr>
<td>SA9676</td>
<td>Δ<em>romR</em>, Δ<em>pilA</em></td>
<td>This work</td>
</tr>
<tr>
<td>SA7110</td>
<td>Δ<em>romR</em>, Δ<em>algQ</em></td>
<td>Anna Potapova</td>
</tr>
<tr>
<td>SA8802</td>
<td>Δ<em>frzE</em></td>
<td>Dorota Skotnicka</td>
</tr>
<tr>
<td>SA7135</td>
<td>Δ<em>frzE</em>, Δ<em>algQ</em></td>
<td>Anna Potapova</td>
</tr>
<tr>
<td>SA4420</td>
<td>Δ<em>mgla</em></td>
<td>(Miertzschke <em>et al.</em>, 2011)</td>
</tr>
<tr>
<td>SA3387</td>
<td>Δ<em>mglB</em></td>
<td>(Leonardy <em>et al.</em>, 2010)</td>
</tr>
<tr>
<td>SA6300</td>
<td>Δ<em>mgla</em>, Δ<em>romX</em></td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3615</td>
<td>Δ<em>mglB</em>, Δ<em>romX</em></td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3619</td>
<td>Δ<em>romR</em>, Δ<em>romX</em></td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3936</td>
<td>Δ<em>mglB</em>, Δ<em>romR</em></td>
<td>(Keilberg <em>et al.</em>, 2012)</td>
</tr>
<tr>
<td>SA3631</td>
<td>Δ<em>mgla</em>, Δ<em>romX</em>, <em>attB</em>::pDK131 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;-yfp)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3635</td>
<td>Δ<em>mglB</em>, Δ<em>romX</em>, <em>attB</em>::pDK131 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;-yfp)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3633</td>
<td>Δ<em>romR</em>, Δ<em>romX</em>, <em>attB</em>::pDK131 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;-yfp)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA3636</td>
<td>Δ<em>romY</em>, Δ<em>romX</em>, <em>attB</em>::pDK131 (P&lt;nsub&gt;<em>nat</em>&lt;sup&gt;romX&lt;/sup&gt;-yfp)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>SA8185</td>
<td><em>mgla</em>-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA7593</td>
<td>Δ<em>mglB</em>, <em>mgla</em>-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8369</td>
<td>Δ<em>romR</em>, <em>mgla</em>-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA7580</td>
<td>Δ<em>romX</em>, <em>mgla</em>-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA7577</td>
<td>Δ<em>romY</em>, <em>mgla</em>-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>Strain</td>
<td>Description</td>
<td>Reference</td>
</tr>
<tr>
<td>--------</td>
<td>-------------</td>
<td>-----------</td>
</tr>
<tr>
<td>SA6963</td>
<td>mglB-mCherry</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>SA3971</td>
<td>ΔmglA, mglB-mCherry</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>SA3966</td>
<td>ΔromR, mglB-mCherry</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>SA6935</td>
<td>ΔromX, mglB-mCherry</td>
<td>This work</td>
</tr>
<tr>
<td>SA6911</td>
<td>ΔromY, mglB-mCherry</td>
<td>This work</td>
</tr>
<tr>
<td>SA7507</td>
<td>romR-mCherry</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA7579</td>
<td>ΔmglA, romR-mCherry</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8308</td>
<td>ΔmglB, romR-mCherry</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8172</td>
<td>ΔromX, romR-mCherry</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8306</td>
<td>ΔromY, romR-mCherry</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA3833</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>SA8183</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8191</td>
<td>ΔromR, mglA&lt;sup&gt;Q82A&lt;/sup&gt;-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA8190</td>
<td>ΔromX, mglA&lt;sup&gt;Q82A&lt;/sup&gt;-mVenus</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>SA7108</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromR</td>
<td>Anna Potapova</td>
</tr>
<tr>
<td>SA6962</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromX</td>
<td>This work</td>
</tr>
<tr>
<td>SA6960</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromX; attB::pDK131&lt;sup&gt;(P&lt;sub&gt;natromX-yfp&lt;/sub&gt;)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6957</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromR, ΔromX; attB::pDK131&lt;sup&gt;(P&lt;sub&gt;natromX-yfp&lt;/sub&gt;)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA7113</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromR; attB::pSH700&lt;sup&gt;(P&lt;sub&gt;pilAromR-gfp&lt;/sub&gt;)&lt;/sup&gt;</td>
<td>Anna Potapova</td>
</tr>
<tr>
<td>SA6958</td>
<td>mglA&lt;sup&gt;Q82A&lt;/sup&gt;, ΔromX, ΔromR; attB::pSH700&lt;sup&gt;(P&lt;sub&gt;pilAromR-gfp&lt;/sub&gt;)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6932</td>
<td>ΔromR; attB::pSH700&lt;sup&gt;(P&lt;sub&gt;pilAromR-gfp&lt;/sub&gt;)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6984</td>
<td>ΔromX, ΔromR, mglA-mVenus</td>
<td>This work</td>
</tr>
<tr>
<td>SA6985</td>
<td>ΔmglB, ΔromX, mglA-mVenus</td>
<td>This work</td>
</tr>
<tr>
<td>SA6996</td>
<td>ΔmglB, ΔromR, mglA-mVenus</td>
<td>This work</td>
</tr>
<tr>
<td>SA6986</td>
<td>ΔmglB, ΔromX, ΔromR, mglA-mVenus</td>
<td>This work</td>
</tr>
<tr>
<td>MxH2262</td>
<td>ΔaglZ</td>
<td>(Zhang et al., 2010)</td>
</tr>
<tr>
<td>SA6987</td>
<td>aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA9100</td>
<td>ΔmglB, aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6990</td>
<td>ΔromR, aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6988</td>
<td>ΔromX, aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6994</td>
<td>ΔmglB, ΔromR, aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
<tr>
<td>SA6993</td>
<td>ΔmglB, ΔromX, aglZ::aglZ-yfp&lt;sup&gt;(pSL65)&lt;/sup&gt;</td>
<td>This work</td>
</tr>
</tbody>
</table>
4.3.3 Cultivation and storage of *E. coli* and *M. xanthus*

*E. coli* strains were grown in LB liquid media with 230 rpm horizontally shaking at 37 °C or on LB agar plates at 37 °C, strain BTH 101 was grown on LB agar plates at 30 °C. The optical densities of cultures were determined photometrically at 600 nm. Glycerol
MATERIALS AND METHODS

stocks for long storage were made with overnight culture by adding glycerol to the final concentration of 10%, freezing in liquid nitrogen and stored at -80 °C.

*M. xanthus* cells were grown on CTT agar plates at 32 °C in dark with appropriate antibiotics when necessary. For the liquid cultures, cells were harvested from the plate, resuspended in 1 ml of CTT and then transferred to the bigger volume of media. Liquid cultures were incubated with horizontal shaking 220 rpm at 32 °C. The optical density of *M. xanthus* cultures were determined photometrically at 550 nm. Glycerol stocks for long storage were made with exponentially growing culture of *M. xanthus* by adding the glycerol to final concentration 4%, freezing in liquid nitrogen and stored at -80 °C.

4.3.4 Bacterial Two Hybrid Assay (BACTH)

The Bacterial Two Hybrid system (Karimova *et al.*, 2005) was used in this studies to detect direct interactions between two proteins of interest in heterologous system in *E. coli*. Plasmids containing the T18 (pUT18 and pUT18C) and the T25 (pKT25 and pKNT25) fragment of the *Bordetella pertussis* adenylate cyclase gene were provided by the manufacturer (Euromedex, Souffelwewersheim, France). Prepared plasmids contain N-terminal or C-terminal fusions of genes of interest to T18 or T25 fragment of the adenylate cyclase. For the assay chemi-competent cell of BTH101 strain, lacking the *cyaA* gene, encoding the catalytic domain of the adenylate cyclase, were transformed with two plasmids as described by the manufacturer. Cells of reporter strain BTH101 do not produce cyclic adenosine monophosphate (cAMP). *In vivo* protein-protein interaction expressed from transformed plasmids can restore the activity of adenylate cyclase from *Bordetella pertussis* in *E. coli* BTH 101 reporter strain and produces cAMP. cAMP production leads to activity of the lac-operon and production of β-galactosidase. The enzyme cleaves present in growth medium X-gal, which allows for blue-white screening of colonies. To test for interactions, plasmids containing gene of interest fused with T18 fragment were used as a bait and co-transformed with a plasmid containing the second gene of interest fused to the T25 fragment. For transformation 10 ng to 25 ng of plasmid DNA were used. Co-transformed cells were plated on LB agar plates containing 100 µg/ml ampicillin, 50 µm/ml kanamycin and 1 mM IPTG. For comparison, for each screen the plasmids pUT18C-Zip and pKNT25-Zip were co-transformed as a positive control. Additionally each plasmid used used in the screen was co-transformed with pUTC-Zip or pKNT25-Zip as a negative control. Plates were incubated on 30 °C for 48 h. For direct comparison 3 corresponding colonies of each co-transformation plate were inoculated in 0.5 ml LB medium containing 100 µg/ml ampicillin, 50 µm/ml kanamycin and 1 mM IPTG.
IPTG and incubated for 3 h shaking at 37 °C. After incubation 5 μl of each culture were spot with all the controls on the same LB agar plate containing 100 μg/ml ampicillin, 50 μm/ml kanamycin, 1 mM IPTG and 40 μg/ml X-gal. Plate were incubated on 30 °C for 48 h. Pictures were acquired before and after additional incubation of plates at 4 °C for 24 h and 48 h to enhance the blue color of positive colonies.

4.3.5 Motility assays for *M. xanthus*

For motility assay, *M. xanthus* cells from exponentially growing cultures were harvested at 4700 rpm for 10 min and resuspended in 1% CTT to density of $7 \times 10^9$ cells/ml. 5 μl aliquots of the resuspension were spotted on 0.5% CTT supplemented with 0.5% for T4P-dependent motility (S-motility) and 1.5% agar for gliding motility (A-motility) and incubated in dark at 32 °C. After 24h, colony morphology and colony edges were observed using a Leica MZ75 Stereomicroscope or Leica M205FA Stereomicroscope and visualized using Leica DFC280 and Hamamatsu ORCA-flash V2 Digital CMOS cameras, respectively. Additionally, on 1.5% agar colonies edges were observed using Leica DM IRE2 Inverted microscope or Leica DM6000B microscope and visualized by Leica DFC280 and Photometrics Cascade II 1024 EMCCD cameras, respectively.

4.3.6 Reversal frequency assay for *M. xanthus* on 1.5% agar, 0.5% CTT

For reversal frequency assay, 5 μl of the exponentially growing overnight culture was spotted on 0.5% CTT supplemented with 1.5% agar, covered by cover slide and incubated in dark at 32 °C. After 4 to 6h, cells were observed using Leica DMi 6000B inverted microscope or Leica DM 6000B or Leica DMi8 microscope microscope and visualized using Hamamatsu Flash 4.0 sCMOS, Photometrics Cascade II 1024 EMCCD and Leica DFC9000 GT cameras, respectively. Cells were recorded for 15 min at 30 s intervals. Analysis was done in Metamorph (Molecular Devices) and ImageJ (Wayne Rasband).

4.3.7 Reversal frequency assay for *M. xanthus* moving by the T4P-dependent motility

For T4P-dependent reversal frequency assay, 5 μl of the exponentially growing overnight culture was spotted into the 24 polystyrene well-plate (Falcon; Sarsted). After 10 minutes incubation in the dark at room temperature cells were covered with 500 μl of 1% methylcellulose in the MMC buffer (10 mM MOPS pH 7.6, 4 mM MgSO₄, 2 mM CaCl₂). Next, plates were incubated in the dark at room temperature for 30 min. Cells
were observed using Leica DM IRE2 Inverted microscope or Leica DMi8 microscope and visualized by Leica DFC280 and Leica DFC9000 GT. Cells were recorded for 10 min at 20 s intervals. Analysis was done in Metamorph (Molecular Devices) and ImageJ (Wayne Rasband).

4.3.8 Trypan blue and congo red dyes binding assay

To determine ability of M. xanthus to bind tryphan blue and congo red dyes plate assay was carried out. Cells from exponentially growing cultures were harvested at 4700 rpm for 10 min and resuspended in 1% CTT to density of $7 \times 10^9$ cells/ml. 10 µl aliquots of resuspension were spotted on 0.5% CTT supplemented with 0.5% agar and 20 µg/ml trypan blue or 40 µg/ml congo red. Plates were incubated at 32 ºC for 24h.

4.3.9 Epi fluorescence microscopy

For epifluorescence microscopy, M. xanthus cells were placed on a thin 1.5 % agar pad buffered with TPM buffer (10 mM Tris-HCl pH 8.0, 1 mM potassium phosphate buffer pH 7.6, 8 mM MgSO4) on a glass slide and immediately covered with a coverslip. After 30 min incubation in the dart at 32 ºC cells were observed using temperature controlled (32 ºC) Leica DMi8 and visualized with Hamamatsu ORCA-flash V2 Digital CMOS camera. Cells on the phase contrast images were automatically detected in Oufti (Jacobs-Wagner Lab). Next, fluorescence signal for segmented cells were analyzed by custom Matlab (The MathWorks) script. Script detects polar clusters only if they have average fluorescence higher than mean of the cytoplasmic fluorescence plus two standard deviations of the cytoplasmic fluorescence and it is bigger than three pixels. Total fluorescence of the polar clusters and cytoplasm was recalculated as a percent of the total cell fluorescence. The data was plotted as a fluorescence at the pole 1 in the function of fluorescence at the pole 2 while pole 1 was always the polar cluster with higher fluorescence. Additionally, omega value that represents asymmetry between the polar clusters was calculated from the equation:

$$\omega = \frac{\text{total fluorescence at pole 1} - \text{total fluorescence at pole 2}}{\text{total fluorescence at pole 1} + \text{total fluorescence at pole 2}}$$

Omega value is between 0 (bipolar symmetric) and 1 (unipolar localization). The localization patterns were determined from omega value: unipolar ($\omega > 0.9$), bipolar asymmetric ($0.9 > \omega > 0.2$) and bipolar symmetric ($\omega<0.2$). Diffuse localization was determined when script was not able to detect any polar cluster that fulfilled mentioned before criteria.
For the time-lapse epifluorescence microscopy, cells were prepared as for snapshot analysis. Time-lapses were taken for 15 min with time resolution 30 s. Data was processed with Metamorph (Molecular Devices) and ImageJ (Wayne Rasband).

4.3.10 TIRF microscopy

For TIRF microscopy, 50 – 150 µl of *M. xanthus* overnight, exponentially growing culture was diluted in 1 ml of the MC7 buffer (10 mM MOPS pH 7.0, 1 mM CaCl2) and spotted on the chitosan coated glass and visualized after 10 min of incubation at room temperature in the dark. Chitosan coated glass was prepared as described in (Ducret et al., 2013) with further modifications. Freshly prepared chitosan 100x counting solution (15 mg/ml chitosan in 2 M acetic acid) diluted 100 fold with deionized water was used for coating the µDish (IBIDI GMBH, Martinsried). 1 ml solution was incubated in the µDish for 30 min. Then, chitosan solution was removed, µDish washed with 1 ml of deionized water and 1 ml of the MC7 buffer. Cells were observed with Leica DMi8 inverted microscope with a 100x flat field apochromatic oil-immersion objective (NA=1.47) and dual color laser Leica AM TIRF MC (488 nm solid state laser used for YFP and mVenus and 561 diode laser used for mCherry imaging) and visualized with Hamamatsu ORCA-flash V2 Digital CMOS camera. TIRF images and time-lapses were taken with penetration depth of 110 nm. For the time lapses, cells were observed for 10 min with the time resolution of 20 s. Active autofocus was used to correct any changes in the objective – sample distance. Obtained data was further processed with Metamorph (Molecular Devices) and ImageJ (Wayne Rasband) and analyzed in ICE (Institut Pasteur). ICE was used to create and analyze kymographs.

4.4 Molecular biology methods

4.4.1 Plasmids and oligonucleotides

All oligonucleotides that were used in this study are listed in Table 8. Underlined sequences display restriction sites used for cloning. All plasmids used in this study are listed in Table 9.

<table>
<thead>
<tr>
<th>Name</th>
<th>Sequence (5’-3’)</th>
<th>Purpose</th>
</tr>
</thead>
</table>
| RomR fw| ATCGAAGCTTATGCCCCAAGAATCGCTGG        | Amplification of *romr-gfp* to clone to pSW105  
<p>| pSW105 |                                      |                                   |
| GFP rv | ATCGGAATTCCTTACTTGACGCTGTCAT         |                                   |</p>
<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence</th>
<th>Function</th>
</tr>
</thead>
<tbody>
<tr>
<td>RomR fw</td>
<td>ATCGCATATGCCCCAGAGATCTGCTGGTCGC</td>
<td>Amplification of the full length romR</td>
</tr>
<tr>
<td>RomR rv stop</td>
<td>ATCGGGAATTGATACGTGGGTCTGGTTTCTTGA</td>
<td></td>
</tr>
<tr>
<td>BTHRomRfw</td>
<td>TCGGTCTAAGATGCCCAAGAATCTGCTGGTCGC</td>
<td>Amplification of the receiver domain of romR</td>
</tr>
<tr>
<td>BTHRecrv</td>
<td>ATCGGGGAATTGAGCACCTTTGCTGGAGCACCTTGG</td>
<td></td>
</tr>
<tr>
<td>BTHGlufw</td>
<td>ATCGGTCTAAGATGGCCCGGGAATGGGGGC</td>
<td>Amplification of the C-terminal domain of romR</td>
</tr>
<tr>
<td>BTHRomR rv stop</td>
<td>ATCGGGAATTGATACGTGGGTCTGGTTTCTTGA</td>
<td></td>
</tr>
<tr>
<td>romX E</td>
<td>GAGGCTCCGTCCGGAGCCGGG</td>
<td>Primers used for verification of romX deletion</td>
</tr>
<tr>
<td>romX F</td>
<td>CTTCTGGAGGCGGCCACCGGCCGC</td>
<td></td>
</tr>
<tr>
<td>RomY E</td>
<td>GGGCGGATGAGCCTTCCTGCCCGAGC</td>
<td>Primers used for verification of romY deletion</td>
</tr>
<tr>
<td>RomY F</td>
<td>TCTCGCGCGCCCTCCGGCGGGG</td>
<td></td>
</tr>
<tr>
<td>romR E</td>
<td>GAGAGGCCCTGGCCCAGCACC</td>
<td>Primers used for verification of romR deletion</td>
</tr>
<tr>
<td>RomR F</td>
<td>GGCACCGTACATCAGGCC</td>
<td></td>
</tr>
<tr>
<td>pilaF2</td>
<td>CAGCAGTCCGGTAGACCTGGGC</td>
<td>Primers used for verification of pilA deletion</td>
</tr>
<tr>
<td>PilaE</td>
<td>CGCTTCCGCCGCCGAGCACCGG</td>
<td></td>
</tr>
<tr>
<td>pilaH</td>
<td>CGATACCCAGTCATCGAA</td>
<td></td>
</tr>
<tr>
<td>pilaG</td>
<td>CCTGCCGCCATCGGCCATCC</td>
<td></td>
</tr>
<tr>
<td>MglBfwsur</td>
<td>ATCGGAAGCTTTCGCTGAGGCCCTCATTCAGGTGAGC</td>
<td>Primers used for verification of mglb deletion or integration of mglB-mcherry</td>
</tr>
<tr>
<td>MglB sur rv</td>
<td>ATCGGGAATTCTCGCCTGGTTTGTACTGGA</td>
<td></td>
</tr>
<tr>
<td>int18-19C forw</td>
<td>CCCACGGAGAGCTCGCTGGA</td>
<td>Primers used to verify integration at 18-19 site</td>
</tr>
<tr>
<td>int18-19C rev</td>
<td>GAGAAGGGGTGCGGGACCTGTC</td>
<td></td>
</tr>
<tr>
<td>int18-19P forw</td>
<td>CGGAAGGGCAGACAGGTGCAGG</td>
<td></td>
</tr>
<tr>
<td>int18-19P rev</td>
<td>CCCTGGCGGAGATTCATGGTAAA</td>
<td></td>
</tr>
<tr>
<td>attB right</td>
<td>GGAATGATCGGGACCAGCTGGA</td>
<td>Primers user to verify integration at Mx8 phage attachment site</td>
</tr>
<tr>
<td>attB left</td>
<td>CGGCACACCTGAGGGCCACATA</td>
<td></td>
</tr>
<tr>
<td>attP right</td>
<td>GCTTTCCGGGACATGGAGGA</td>
<td></td>
</tr>
<tr>
<td>attP left</td>
<td>GGGAAAGCTCTGGGTACGAA</td>
<td></td>
</tr>
<tr>
<td>M13 uni (-43)</td>
<td>AGGGTTTTTCCCACTCAGCGAGG</td>
<td></td>
</tr>
<tr>
<td>M13 rev (-49)</td>
<td>GACCGGATAACCATATTTCACACAGG</td>
<td></td>
</tr>
<tr>
<td>KA254</td>
<td>GTGGCGACACCTGGGTGGCAGG</td>
<td></td>
</tr>
<tr>
<td>pKT25fw</td>
<td>GGGCATTCGCTGGAGCAGG</td>
<td></td>
</tr>
<tr>
<td>pUT18fwrow</td>
<td>TCCGGCTCGATGTGGTTGTGGG</td>
<td></td>
</tr>
<tr>
<td>pUT18V-fw</td>
<td>CTTGGAACCGGTGCGGCGCCG</td>
<td></td>
</tr>
</tbody>
</table>

Table 9. Plasmids used in this study
<table>
<thead>
<tr>
<th>Plasmid</th>
<th>Description</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>pSW105</td>
<td>P_pilA, kanR</td>
<td>S. Weiss (MPI Marburg)</td>
</tr>
<tr>
<td>pSWU30</td>
<td>terR</td>
<td>(Wu et al., 1997)</td>
</tr>
<tr>
<td>pMR3691</td>
<td>P_van, tetR</td>
<td>(Iniesta et al., 2012)</td>
</tr>
<tr>
<td>pBJ114</td>
<td>KanR, galK</td>
<td>(Julien et al., 2000)</td>
</tr>
<tr>
<td>pDK95</td>
<td>pBJ114; in frame deletion of romY, kanR</td>
<td>Daniela Keilberg, PhD thesis, 2013</td>
</tr>
<tr>
<td>pSL37</td>
<td>pBJ114; in frame deletion of romR, kanR</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>pES2</td>
<td>pBJ114; in frame deletion of mglB, kanR</td>
<td>(Leonardy et al., 2010)</td>
</tr>
<tr>
<td>pDK145</td>
<td>pBJ114; in frame integration of mglB-mCherry, kanR</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>pSL65</td>
<td>pBJ114; in frame integration of aglZ-yfp, kanR</td>
<td>(Leonardy et al., 2010)</td>
</tr>
<tr>
<td>pLC32</td>
<td>pBJ114; in frame integration of romR-mCherry, kanR</td>
<td>Luis Carreira</td>
</tr>
<tr>
<td>pMAT162</td>
<td>pBJ114; in frame deletion of pilA, kanR</td>
<td>Anke Treuner-Lange</td>
</tr>
<tr>
<td>pDK131</td>
<td>pSW105; P_natromX-yfp</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pDK132</td>
<td>pSW105; P_natromY-yfp</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pDK157</td>
<td>pSWU30; P_natromX</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pSH700</td>
<td>pSW105; P_pilAromR-gfp</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>pDSZ10</td>
<td>pSW105; P_pilRomR^{D53E}-GFP</td>
<td>This work</td>
</tr>
<tr>
<td>pDSZ11</td>
<td>pSW105; P_pilRomR^{D53E}-GFP</td>
<td>This work</td>
</tr>
<tr>
<td>pDSZ16</td>
<td>pMR3691; P_vanromR</td>
<td>This work</td>
</tr>
<tr>
<td>pDSZ17</td>
<td>pMR3691; P_vanromR^{D53E}</td>
<td>This work</td>
</tr>
<tr>
<td>pDSZ18</td>
<td>pMR3691; P_vanromR^{D53E}</td>
<td>This work</td>
</tr>
<tr>
<td>pTM1</td>
<td>Overexpression MglA-His6</td>
<td>(Zhang et al., 2010)</td>
</tr>
<tr>
<td>pTM2</td>
<td>Overexpression His6-MglB</td>
<td>(Zhang et al., 2010)</td>
</tr>
<tr>
<td>pTB005</td>
<td>Overexpression His6-RomX</td>
<td>Tobias Bender</td>
</tr>
<tr>
<td>pDK28</td>
<td>Overexpression MalE-RomR</td>
<td>(Keilberg et al., 2012)</td>
</tr>
<tr>
<td>pAH159</td>
<td>Overexpression of Strep-RomX</td>
<td>Andrea Harms</td>
</tr>
<tr>
<td>pUT18</td>
<td>ampR</td>
<td>Euromedex (France)</td>
</tr>
<tr>
<td>pUT18C</td>
<td>ampR</td>
<td>Euromedex (France)</td>
</tr>
<tr>
<td>pKT25</td>
<td>kanR</td>
<td>Euromedex (France)</td>
</tr>
<tr>
<td>pKNT25</td>
<td>kanR</td>
<td>Euromedex (France)</td>
</tr>
<tr>
<td>pDK70</td>
<td>MglA (pKNT25)</td>
<td>(McLoon et al., 2015)</td>
</tr>
<tr>
<td>pDK143</td>
<td>MglA (pKT25)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pDK76</td>
<td>MglA (pUT18)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pDK75</td>
<td>MglA (pUT18C)</td>
<td>(McLoon et al., 2015)</td>
</tr>
<tr>
<td>pDK71</td>
<td>MglB (pKNT25)</td>
<td>(McLoon et al., 2015)</td>
</tr>
<tr>
<td>pDK142</td>
<td>MglB (pKT25)</td>
<td>Daniela Keilberg</td>
</tr>
<tr>
<td>pDK77</td>
<td>MglB (pUT18)</td>
<td>Daniela Keilberg</td>
</tr>
</tbody>
</table>
4.4.2 Plasmids construction

Genomic DNA of *M. xanthus* DK1622, SA3980 or SA3981 was used to amplify DNA fragments. Plasmid construct were transformed to *E. coli* Mach1 or TOP10 cells. Purified plasmids were sequenced by the Eurofins MWG Operon (Eldersber) company to verify the correct sequences. Sequencing results were analyzed using ContigExpress from the VectorNTI advance suite 11 software (Invitrogen) or with SeqMan Pro from DNASTAR (DNASTAR) software package.

The plasmids pDSZ10 and pDSZ11 are derivatives of pSW105 and were generated for the expression of *romR<sup>D53N</sup>-gfp* and *romR<sup>D53E</sup>-gfp* under the control of the strong, constitutively active promotor. To amplify *romR<sup>D53N</sup>-gfp* genomic DNA of SA3980 and primers “RomR fw pSW105” and “GFP rv” were used. To amplify *romR<sup>D53E</sup>-gfp* genomic DNA of SA3981 and primers “RomR fw pSW105” and “GFP rv” were used. The products were cloned with HindIII/EcoRI sites to pSW105.

The plasmids pDSZ16, pDSZ17 and pDSZ18 are derivatives pf pMR3691 and were generated for the expression of *romR*, *romR<sup>D53N</sup>* and *romR<sup>D53E</sup>* under control of the inducible promotor. Genomic DNA of DK1622, SA3980 and SA3981 were used
respectively. To amplify romR, romR^{D3N} and romR^{D53E} “RomR fw pMR” and “RomR rv stop” primers were used. The products were cloned at the Ndel/EcoRI sites to pMR3691.

The plasmid pDSZ12 and pDSZ13 are derivatives of pUT18 and pKT25 respectively and were generated for testing protein-protein interaction in the Bacterial two hybrid system. pDSZ12 contains C-terminal RomR domain, DNA fragment was amplified with primers “BTHGlufw” and “BTHRomRrv stop”. The product was cloned with XbaI/EcoRI sites to pKT25. pDSZ13 contains RomR N-terminal receiver domain, DNA fragment was amplified with primers “BTHRomRfw” and “BTHrecrv”. Product was cloned with Xhol/EcoRI sites to pUT18.

4.4.3 Generation of in-frame deletion mutants

In-frame deletion mutants were generated by two-step homologous recombination as described (Shi et al., 2008). Briefly, the upstream and downstream flanking region of approximately 500bp were amplified using AB and CD primer pairs. B and C primers contain overlapping compatibilie ends essential for the fusion of AB and CD fragments in a PCR reaction using A and D primers. AB and CD fragments serve as a template to generate the in-frame deletion fragment AD. Next, AD fragment was cloned into pBJ114 vector. The correct pBJ114AD plasmid was transformed into M. xanthus DK1622. The plasmid integration was checked by PCR reaction with E (binds upstream of A primers) and F (binds downstream to D primer), E and M13forward (binds to pBJ114), F and M13reverse (binds to pBJ144) primer pairs. One clone from up- and downstream plasmid integration was used for the second step of homologous recombination. Plasmid pBJ114 contains the selection marker galK (galaktokinase) gene. The gene product GalK converts galactose into galactose-1-phosphate which cannot be metabolize by M. xanthus and accumulates up to toxic levels when cells are grown on media supplemented with galactose. Thus, only clones that undergo second homologous recombination lost plasmid are able to grown on media with galactose. For the second homologous recombination event cells were grown in CTT liquid shaking culture to exponential growth phase. Series of dilutions were plated on CTT agar plates supplemented with 10 µg/ml gentamycin and 2.5% galactose. Galactose resistant and kanamycin sensitive clones were checked with PCR reaction using E and F, G(binds downstream of B primer) and H ( binds upstream of C primer) primer pairs. The EF prime pair PCR reaction product was longer for the WT then for the deletion mutant, while GH fragment was amplified only in the WT.
4.4.4 DNA isolation from *E. coli* and *M. xanthus*

Plasmid DNA from *E. coli* was isolated using the QIAprep Spin Miniprep Kit (Qiagen) or the NucleoSpin Plasmid QuickPure kit (Macherey-Nagel) in accordance to the instructions provided by the manufacturer. Genomic DNA of *M. xanthus* was isolated using MasterPure DNA preparation Kit (Epicentre) according to the instructions provided by the manufacturer. Concentration and purity of DNA was determined with the Nanodrop ND-1000 spectrophotometer (Nanodrop, Wilmington) or with DS11+ spectrophotometer (DeNovix Inc., Wilmington). Crude genomic DNA of *M. xanthus* for colony PCR was prepared by resuspending in 50 µl H2O cells grown on CTT agar plates and boiling for 5 min at 96°C and centrifuging the sample for one minute at 13 000 rpm. 3 µl of resulting supernatant was used for the PCR reaction.

4.4.5 Polymerase Chain Reaction (PCR)

For the amplification of specific DNA fragments, the Phusion High-Fidelity DNA Polymerase (Thermo Scientific™, Darmstadt) or Q5® Hot Start High-Fidelity DNA Polymerase (New England Biolabs, Frankfurt a. M.) was used in a total reaction volume of 50 µl. The colony PCR was performed used 5 PRIME MasterMix in total volume of 20 µl. The composition of the PCR reaction mix is described in Table 10.

**Table 10. PCR reaction mix**

<table>
<thead>
<tr>
<th>Component</th>
<th>Volume</th>
<th>Final concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>PCR with Phusion High-Fidelity DNA Polymerase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Template DNA</td>
<td>1 µl</td>
<td>~50 ng</td>
</tr>
<tr>
<td>10 µM primer (each)</td>
<td>2.5 µl</td>
<td>1.25 µM</td>
</tr>
<tr>
<td>10 mM dNTPs</td>
<td>1 µl</td>
<td>0.2 mM</td>
</tr>
<tr>
<td>5 x Phusion GC buffer</td>
<td>10 µl</td>
<td>1x</td>
</tr>
<tr>
<td>5 x enhancer</td>
<td>10 µl</td>
<td>1x</td>
</tr>
<tr>
<td>DMSO</td>
<td>2.5 µl</td>
<td>5% (v/v)</td>
</tr>
<tr>
<td>Phusion DNA polymerase</td>
<td>0.5 µl</td>
<td>0.02 unit/µl</td>
</tr>
<tr>
<td>ddH2O</td>
<td>To 50 µl</td>
<td></td>
</tr>
<tr>
<td>PCR with Q5® Hot Start High-Fidelity DNA Polymerase</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Template DNA</td>
<td>1 µl</td>
<td>~50 ng</td>
</tr>
<tr>
<td>10 µM primer (each)</td>
<td>2.5 µl</td>
<td>1.25 µM</td>
</tr>
<tr>
<td>10 mM dNTPs</td>
<td>1 µl</td>
<td>0.2 mM</td>
</tr>
<tr>
<td>5 x Q5 Reaction Buffer</td>
<td>10 µl</td>
<td>1x</td>
</tr>
<tr>
<td>5 x Q5 High GC Enhancer</td>
<td>10 µl</td>
<td>1x</td>
</tr>
<tr>
<td>DMSO</td>
<td>2.5 µl</td>
<td>5% (v/v)</td>
</tr>
<tr>
<td>Q5 Hot Start High-Fidelity DNA</td>
<td>0.5 µl</td>
<td>0.02 unit/µl</td>
</tr>
<tr>
<td>ddH2O</td>
<td>To 50 µl</td>
<td></td>
</tr>
</tbody>
</table>
Colony PCR

<table>
<thead>
<tr>
<th>Component</th>
<th>Volume</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crude genomic DNA</td>
<td>3 µl</td>
<td>~ 200 ng</td>
</tr>
<tr>
<td>10 µM primer (each)</td>
<td>1 µl</td>
<td>0.5 µM</td>
</tr>
<tr>
<td>5 PRIME MasterMix</td>
<td>10 µl</td>
<td></td>
</tr>
<tr>
<td>DMSO</td>
<td>2 µl</td>
<td>10% (v/v)</td>
</tr>
<tr>
<td>ddH₂O</td>
<td>To 20 µl</td>
<td></td>
</tr>
</tbody>
</table>

The PCR programs used in this study are represented in Table 11. PCR conditions were modified depending on the primer annealing temperature and expected product size.

### Table 11. PCR programs

<table>
<thead>
<tr>
<th>Step</th>
<th>Temperature</th>
<th>Time</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Standard/check PCR</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Initial denaturation</td>
<td>98 °C</td>
<td>30 min</td>
</tr>
<tr>
<td>denaturation</td>
<td>98 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>Primer annealing</td>
<td>5 °C below predicted melting temperature</td>
<td>30 sec</td>
</tr>
<tr>
<td>elongation</td>
<td>72 °C</td>
<td>1 min/kb</td>
</tr>
<tr>
<td>final elongation</td>
<td>72 °C</td>
<td>3 min</td>
</tr>
<tr>
<td>hold</td>
<td>4 °C</td>
<td>∞</td>
</tr>
<tr>
<td><strong>Touch down PCR</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Initial denaturation</td>
<td>98 °C</td>
<td>30 min</td>
</tr>
<tr>
<td>denaturation</td>
<td>98 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>Primer annealing</td>
<td>65 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>elongation</td>
<td>72 °C</td>
<td>1 min/kb</td>
</tr>
<tr>
<td>denaturation</td>
<td>98 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>Primer annealing</td>
<td>62 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>elongation</td>
<td>72 °C</td>
<td>1 min/kb</td>
</tr>
<tr>
<td>denaturation</td>
<td>98 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>Primer annealing</td>
<td>58 °C</td>
<td>30 sec</td>
</tr>
<tr>
<td>elongation</td>
<td>72 °C</td>
<td>1 min/kb</td>
</tr>
<tr>
<td>final elongation</td>
<td>72 °C</td>
<td>3 min</td>
</tr>
<tr>
<td>hold</td>
<td>4 °C</td>
<td>∞</td>
</tr>
</tbody>
</table>

### 4.4.6 Agarose gel electrophoresis

Nucleic acid fragments were separated by size on 1% agarose gels within 0.01% (v/v) ethidium bromide in TBE buffer (Invitrogen) at 120 V. DNA samples were mixed with 5 x DNA loading buffer (32.5 % sacharose, 5 mM EDTA, 5 mM Tris-HCl pH 7.5, 0.15% bromophenol blue). As a DNA marker the 2-log DNA ladder (NEB) was used. Agarose gels were imaged using E-BOX VX2 imaging system (PeqLab).
4.4.7 DNA restriction and ligation

Restriction of DNA fragment (0.5-2 μg) was performed with restriction endonucleases at 37 °C for 2 h. Restricted DNA was purified using the DNA Clean & Concentration kit (Zymo Research) according to manufacturer protocol.

Ligation reactions were performed with T4 DNA ligase (NEB). DNA fragments were ligated into vector with 3- to 5-fold molar excess of insert DNA. After 1 h at room temperature ligation mixtures were used for transformation into *E. coli* Mach1 or TOP10.

4.4.8 Preparation and transformation of chemically competent *E. coli* cells

To prepare chemical competent *E. coli* cells, the overnight culture was used to inoculate 200 ml of LB media. Cultures were grown with shaking at 230 rpm at 37 °C to an OD₆₀₀ of 0.5 – 0.7. The cells were harvested by centrifugation at 4700 rpm for 20 min 4 °C and resuspended in 50 ml ice-cold sterile 50 mM CaCl₂ solution. The cells were pelleted again at the same conditions and washed again. The cells were centrifuged again in the same conditions and resuspended in 2 ml ice-cold sterile 50 mM CaCl₂ with 10% (v/v) glycerol solution. 50 μl aliquots of cells were frozen in liquid nitrogen and kept at -80 °C until used.

For transformation one 50 μl aliquote was thawed on ice and 10 μl of ligation mixture was added to cells and mixed carefully. After 30 min incubation on ice to perform heat shock cells were transferred to 42 °C for 1 min 30 sec. After 5 min incubation on ice, 1 ml LB-medium was added and cells were incubated for 60 min shaking at 37 °C. After harvesting, cells pellet were resuspended in 50 μl LB medium and plated on LB agar plates supplemented with appropriate antibiotics. Plates were incubated at 37 °C overnight. Grown colonies were checked for the presence of the plasmid containing the insert by colony PCR reaction.

4.4.9 Preparation and transformation of electrocompetent *M. xanthus* cells

*M. xanthus* cells were grown overnight with shaking at 230 rpm at 32 °C to an OD₅₅₀ of 0.5 – 0.9. Cells were harvested at 13 000 rpm for 1 min and the pellet was washed twice in 1 ml sterile ddH₂O and resuspended in 30 μl H₂O. The cell resuspension was immediately used for electroporation. 1 μg plasmid DNA was added to 30 μl cells and the mixture was transferred into an electroporation cuvette (Bio-Rad, Munchen) and pulsed with 0.65 kV, 25 μF and 400 Ω. 1 ml of CTT media was added and mixed with cells, the cell suspension was transferred to a 25 ml Erlenmyer flask containing 1 ml of CTT media and incubated with shaking at 230 rpm at 32 °C for 6 h. Then, 2 ml (for
plasmid integrating at the endogenous site) and 100 µl (for plasmids integrating at the Mx8 site and Mxan 18-19 site) were mixed with 3.5 ml of soft agar and plated on CTT agar plates supplemented with appropriate antibiotics. The plates were incubated for 5 – 10 days at 32 ºC in the dark. Grown colonies were transferred to fresh agar plates. Plasmid integration was verified by colony PCR.

4.5 Biochemical methods

4.5.1 SDS-polyacrylamide gel electrophoresis (SDS-PAGE)

To separate proteins by size under denaturing conditions SDS polyacrylamide gel electrophoresis (Laemmli, 1970) was performed with SDS gels with 10% to 16% polyacrylamide concentration. To denature proteins, samples were mixed with loading buffer (10% (v/v) glycerol, 60 mM Tris-HCl pH 6.8, 2% (w/v) SDS, 100 mM DTT, 3 mM EDTA, 0.005% (w/v) bromophenol blue) and boiled at 95 ºC for 10 min before loading on the gel. Gel electrophoresis was made in Bio-Rad electrophoresis chamber (Bio-Rad, München) at 80-140 V in 1x Tris Glycin SDS (TGS) running buffer (Bio-Rad, München). To determine size of the proteins, the PageRuler Prestained Protein Ladder (Fermentas) was used for comparison.

4.5.2 Determination of total protein concentration in cell extracts

To determine protein concentration in a lysate of *M. xanthus* the Bio-Rad proteins assay kit was used accordingly to the manufacturer’s manual. In order to make standard curve of protein concentration Pre-Diluted Protein Assay Standards: Bovine Serum Albumin (BSA) set (Thermo Scientific, Dreieich) was used. Standards were diluted in three volumes of H2O and one volume of 5 x SDS sample buffer to get 10 µg/µl, 5 µg/µl, 2.5 µg/µl, 1.25 µg/µl and 0.625 µg/µl concentrations. These resulting dilutions were diluted again 25 times in H2O and 20 µl of these samples were mixed with 980 µl 1:5 diluted Bradford reagent. After 5 min incubation at room temperature and in the dark, absorbance was measured at 595 nm with Ultrospec 2100 pro spectrophotometer (GE Healthcare Europe GmbH, Freiburg). To prepare *M. xanthus* cell lysate a pellet of cells were dissolved in 1 x SDS sample buffer and heated at 95 ºC for 10 min. Lysate were diluted 1:25 in H2O and 20 µl was mixed with 980 µl of 1:5 diluted Bradford reagent and incubated 5 minutes in the dark. Absorbance was measured ass described before. Protein concentration was calculated from the standard curve.

To determine protein concentration of the purified proteins the Bradford reagent (Bio-Rad) was used accordingly to yhe manufacturer’s manual. The protein standard
curve was generated using Pre-Diluted Protein Assay Standards: Bovine Serum Albumin (BSA). The reaction samples were prepared in duplicates 1 ml reaction volume. After 5 min incubation at room temperature and in the dark, absorbane was measured at 595 nm with Ultrospec 2100 pro spectrophotometer (GE Healthcare Europe GmbH, Freiburg). Protein concentrations were determined based on the linear slope of the standard curve.

4.5.3 Immunoblot analysis

Proteins from cell extracts were separated in the gel by SDS-PAGE and transferred to a nitrocellulose membrane using “TransBlot® TurboTM Transfer System” from Bio-RaD at 1.3 A (0.0208/cm²), 25 V for 7 min with transfer buffer (300 mM Tris, 300 mM Glycin, 0.05% SDS, pH 9.0) or using Hoefer™ TE77 from Amersham at 50 mA (0.8 mA/cm²) for 1h with anode transfer buffer (25 mM Tris, 192 mM Glycin, 0.01% SDS, 25% Methanol) and cathode transfer buffer (50 mM Tris, 384 mM Glycin, 0.2% SDS, 10% Methanol). After transfer, the membrane was blocked in 5% non-fat milk powder (w/v) in TBST (20 mM Tris-HCl, 137 mM NaCl, 0.05% (v/v) Tween 20, pH 7.0) for 1 h or overnight at 4 °C shaking. After blocking, the membrane was washing with TBST buffer. After washing, the primary antibody was added at the dilution listed in Table 12 in TBST buffer supplemented with 2% non-fat milk powder and incubated overnight at 4 °C shaking. After washing with TBST buffer, the membrane was incubated with secondary anti-rabbit immunoglobulin G peroxidase conjugate (Sigma) in a dilution 1:15 000 or with secondary anti-mouse immunoglobulin G, horseradish peroxidase lined whole antibody (GE Healthcare) in a dilution 1:2 000 for 1h at 4 °C shaking. After washing with TBST buffer the membrane was developed with the luminescent image analyzer LAS-4000 (Fujifilm).

### Table 12. Dilutions of primary antibodies used for immunoblots analysis

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Dilution</th>
</tr>
</thead>
<tbody>
<tr>
<td>α-MglA</td>
<td>1:2000</td>
</tr>
<tr>
<td>α-MglB</td>
<td>1:2000</td>
</tr>
<tr>
<td>α-RomR</td>
<td>1:5000</td>
</tr>
<tr>
<td>α-RomX</td>
<td>1:2000</td>
</tr>
<tr>
<td>α-RomY</td>
<td>1:5000</td>
</tr>
<tr>
<td>α-PilC</td>
<td>1:5000</td>
</tr>
<tr>
<td>α-GFP</td>
<td>1:2000</td>
</tr>
<tr>
<td>α-mCherry</td>
<td>1:1000</td>
</tr>
</tbody>
</table>

4.5.4 Proteins purification

Purification of MalE-RomR
To purify the proteins, \textit{E. coli} Rosetta 2 (DE3)/pLysS strain (Novagen) was transformed with relevant plasmids. The culture were grown in 1.0 l or 2.0 l of LB with 0.5 % glucose and supplemented with appropriate antibiotics and 0.5% glucose when necessary at 30 °C to an \textit{OD}_{600} of 0.5-0.7. The protein expression was induced by addition of IPTG to a final concentration of 0.5 mM. Then, cells were growing overnight at 18 °C.

The cells were harvested by centrifugation at 6 000 x g, 20 min at 4 °C and washed in wash buffer (50 mM Tris pH 7.5, 150 mM NaCl, 1 mM EDTA, 1mM DTT) and resuspended in 50ml Lysis buffer (50 ml of the wash buffer supplemented with PMSF 100 µg/ml, DNase 1 10U/ml and protease inhibitors – Complete Protease Inhibitor Cocktail Tablet from Roche). Next, cells were sonicated and centrifuged at 4 700 x rpm for 30 min at 4 °C to pellet the cellular debris. Cleared supernatant was loaded onto 5 ml MBPTrapHP (GE Healthcare) column equilibrated with wash buffer. Proteins were eluted with elution buffer (50 mM Tris pH 7.5, 150 NaCl, 1mM EDTA, 1mM DTT, 5% glycerol, 10 mM maltose). Eluted fractions containing MalE-RomR were loaded onto 5 ml HiTrap Q HP column (GE Healthcare) equilibrated with buffer A (50 mM Tris pH 7.5, 50 mM NaCl, 5 mM MgCl$_2$, 1 mM DTT, 5% glycerol. Proteins were eluted with buffer A along a gradient of 20 columns volume (CV) from 50 mM to 500 mM NaCl. Next, fractions containing MalE-RomR were loaded onto HiLoad 16/600 Superdex 200 pg (GE Healthcare) column that was equilibrated with Gefi buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5 mM MgCl$_2$, 1 mM DTT, 5 % glycerol). Fractions with MalE-RomR were pooled, span-frozen in liquid nitrogen and stored at -80 °C.

**Purification of MglA-His$_6$**

To purify the proteins, \textit{E. coli} Rosetta 2 (DE3)/pLysS strain (Novagen) was transformed with relevant plasmid. The culture were grown in 1.0 l or 2.0 l of LB with 0.5% glucose and supplemented with appropriate antibiotics and 0.5% glucose when necessary at 30 °C to an \textit{OD}_{600} of 0.5-0.7. The protein expression was induced by addition of IPTG to a final concentration of 0.5 mM. Then, cells were growing overnight at 18 °C.

The cells were harvested by centrifugation at 6 000 x g, 20 min at 4 °C and washed in wash buffer (50 mM Tris pH 7.5, 150 mM NaCl, 10 mM imidazole, 5% glycerol, 5mM MgCl$_2$) and resuspended in Lysis buffer (50 ml of wash buffer supplemented with 1 mM DTT, 100 µg/ml PMSF, 10 U/ml DNase 1 and protease inhibitors – Complete Protease Inhibitor Cocktail Tablet from Roche). Next, cells were sonicated and centrifuged at 4 700 x rpm for 30 min at 4 °C to pellet the cellular debris. Cleared supernatant was filter with 0.45 µm sterile filter (Millipore Merek, Schwalbach). To purify MglA-His$_6$ only soluble fraction was used. Clear soluble fraction was loaded onto 5 ml HITrap Chelating
MATERIALS AND METHODS

HP (GE Healthcare) column, preloaded with NiSO₄ and equilibrated with Wash buffer. Proteins was eluted with Elution buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 5 mM MgCl₂, 500 mM imidazole) along a gradient of 16 CV. Elution fractions containing MglA-His₆ were loaded onto HiLoad 16/600 Superdex 75 pg (GE Healthcare) column that was equilibrated with Gefi buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5 mM MgCl₂, 1 mM DTT, 5% glycerol). Fractions with MglA-His₆ were pooled, span-frozen in liquid nitrogen and stored at -80 °C.

Purification of His₆-MglB

To purify the proteins, E. coli Rosseta 2 (DE3)/pLysS strain (Novagen) was transformed with relevant plasmid. The culture were grown in 1.0 l or 2.0 l of LB with 0.5% glucose and supplemented with appropriate antibiotics and 0.5% glucose when necessary at 30 °C to an OD₆₀₀ of 0.5-0.7. The protein expression was induced by addition of IPTG to a final concentration of 0.5 mM. Then, cells were growing overnight at 18 °C.

The cells were harvested by centrifugation at 6 000 x g, 20 min at 4 °C and washed in wash buffer (50 mM Tris pH 7.5, 150 mM NaCl, 10 mM imidazole, 5% glycerol, 5M MgCl₂) and resuspended in Lysis buffer (50 ml of wash buffer supplemented with 1 mM DTT, 100 µg/ml PMSF, 10 U/ml DNase 1 and protease inhibitors – Complete Protease Inhibitor Cocktail Tablet from Roche). Next, cells were sonicated and centrifuged at 4 700 x rpm for 30 min at 4 °C to pellet the cellular debris. Cleared supernatant was filter with 0.45µ sterile filter (Millipore Merck, Schwalbach). To purify His₆-MglB only soluble fraction was used. Clear soluble fraction was loaded onto 5 ml HITrap Chelating HP (GE Healthcare) column, preloaded with NiSO₄ and equilibrated with Wash buffer. Proteins was eluted with Elution buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 5 mM MgCl₂, 500 mM imidazole) along a gradient of 16 CV. Elution fractions containing His₆-MglB were loaded onto HiLoad 16/600 Superdex 75 pg (GE Healthcare) column that was equilibrated with Gefi buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5 mM MgCl₂, 1 mM DTT, 5% glycerol). Fractions with His₆-MglB were pooled, span-frozen in liquid nitrogen and stored at -80 °C.

Purification of His₆-RomX

To purify the proteins, E. coli Rosseta 2 (DE3)/pLysS strain (Novagen) was transformed with relevant plasmid. The culture were grown in 1.0 l or 2.0 l of LB with 0.5% glucose and supplemented with appropriate antibiotics and 0.5% glucose when necessary at 37 °C to an OD₆₀₀ of 0.5-0.7. The protein expression was induced by addition of IPTG to a final concentration of 1.0 mM. Then, cells were growing for 3 hours at 37 °C.
The cells were harvested by centrifugation at 6000 x g, 20 min at 4 °C and washed in wash buffer (50 mM Tris pH 7.5, 150 mM NaCl, 10 mM imidazole, 5% glycerol, 5mM MgCl₂) and resuspended in Lysis buffer (50 ml of wash buffer supplemented with 1 mM DTT, 100 μg/ml PMSF, 10 U/ml DNase 1 and protease inhibitors – Complete Protease Inhibitor Cocktail Tablet from Roche). Next, cells were sonicated and centrifuged at 4700 x rpm for 30 min at 4 °C to pellet the cellular debris. Cleared supernatant was filter with 0.45µ sterile filter (Millipore Merck, Schwalbach). To purify His₆-RomX only soluble fraction was used. Clear soluble fraction was loaded onto 5 ml HITrap Chelating HP (GE Healthcare) column, preloaded with NiSO₄ and equilibrated with Wash buffer. Proteins was eluted with Elution buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 5 mM MgCl₂, 500 mM imidazole) along a gradient of 16 CV. Elution fractions containing His₆-RomX were loaded onto HiLoad 16/600 Superdex 75 pg (GE Healthcare) column that was equilibrated with Gefi buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5 mM MgCl₂, 1 mM DTT, 5 % glycerol). Fractions with His₆-RomX were pooled, span-frozen in liquid nitrogen and stored at -80 °C.

Purification of Strep-RomX

To purify the proteins, E. coli Rosseta 2 (DE3)/pLysS strain (Novagen) was transformed with relevant plasmid. The culture were grown in 1.0 l or 2.0 l of 2xYT with 0.5% glucose and supplemented with appropriate antibiotics and 0.5% glucose when necessary at 37 °C to an OD₆₀₀ of 0.5-0.7. The protein expression was induced by addition of anhydrotetracycline to a final concentration of 200 µg/l. Then, cells were growing for ON at 18 °C.

The cells were harvested by centrifugation at 6000 x g, 20 min at 4 °C and washed in wash buffer (100 mM Tris pH 8.0, 150 mM NaCl, 1 mM EDTA, 1 mM DTT) and resuspended in Lysis buffer (50 ml of wash buffer supplemented with 100 µg/ml PMSF, 10 U/ml DNase 1 and protease inhibitors – Complete Protease Inhibitor Cocktail Tablet from Roche). Next, cells were sonicated and centrifuged at 10000 x rpm for 20 min at 4 °C to pellet the cellular debris. Cleared supernatant was filter with 0.45 µm sterile filter (Millipore Merck, Schwalbach). To purify Strep-RomX only soluble fraction was used. Clear soluble fraction was loaded onto 5 ml Strep Traap HP (GE Healthcare) column, equilibrated with wash buffer. Proteins was eluted with Elution buffer (150 mM Tris pH 8.0, 150 mM NaCl, 1 mM EDTA, 2.5 mM Desthiobiotin). Elution fractions containing Strep-RomX were loaded onto HiLoad 16/600 Superdex 75 pg (GE Healthcare) column that was equilibrated with Gefi buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5 mM MgCl₂,
1 mM DTT, 5% glycerol). Fractions with Strep-RomX were pooled, span-frozen in liquid nitrogen and stored at -80°C.

4.5.5 GTPase assay

GTP-hydrolysis by MglA-His6 was measured in an enzyme coupled spectrophotometric assay as reported in (Ingerman & Nunnari, 2005). For the assay, MglA-His6 was preincubated for 30 min with 100 times molar excess of GTP in the room temperature. In parallel, equimolar combinations of His6-MglB, RomX-His6 and MalE-RomR were preincubated for 10 min before starting reaction. Eventually, RomX-His6 and MalE-RomR were preincubated for 10 min with MglA-His6-GTP in a molar ratio 2:1. Reactions were started by adding His6-MglB, His6-RomX and/or MalE-RomR to the MglA-His6/GTP or to MglA-His6/GTP/His6-RomX and/or MalE-RomR mixtures. Final concentration in the reactions: MglA-His6 3 µM, His6-MglB 6 µM, His6-RomX 6 µM and MalE-RomR 6 µM. GTPase assay was performed in the reaction buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 1 mM DTT, 5 mM MgCl2). Absorption at 340 nm was followed for 60 min in one minute intervals at 37°C. Absorbance versus time was plotted. In order to decrease noise, we did not utilize data from early time points. Only absorbance data that is linear with respect to time was used. By using the “add trendline” function of Microsoft Excel, data from each experiment was fitted. The slope of this line was multiplied by 60 and divided by the molar absorbivity of NADH (6220 M⁻¹cm⁻¹) and by previously determined path length (0.23 cm) and by concentration of MglA-His6. That processed data showed amount of hydrolyzed GTP per hour per molecule of MglA.

4.5.6 Pull down experiment

To test direct protein-protein interactions, 128 µg MalE-RomR, 58 µg His6-RomX, 64 µg MalE, 48 µg MglA-His6 were used. For the experiments to test affinity of MalE-RomR and His6-RomX to the amylose resin, interactions between MalE-RomR and His6-RomX and between MalE and His6-RomX proteins were applied to 200 µL Amylose Resin previously equilibrated with buffer 1 (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 1 mM DTT, 5 mM MgCl2). Then, samples were incubated for 30 min at the room temperature. Next, resin was loaded into the column and washed 3 times 1 ml with buffer 1. Proteins were eluted with 2 times 200 µl of buffer 2 (buffer 1 supplemented with 10 mM maltose). Fractions were analyzed by the SDS-PAGE.

To test interactions between MglA-His6 and Strep-RomX 27 µg of Strep-RomX (final concentration 10 µM) was mixed with 46 µg MglA-His6 (final concentration 10 µM). MglA-His6 was preincubated with GTP or GDP (final concentration 40 µM) for 30
min at RT. After 30 min incubation of Strep-RomX with MglA-His6-GDP/GTP at RT proteins were incubated with 10 µl of strep-tactin XT MagStrep ‘type3’ XT Beads (IBA GmbH) for 30 min at RT. Than beads were washed 10x with 1 ml wash buffer (50 mM Tris pH 7.5, 150 mM NaCl, 5% glycerol, 1 mM DTT, 5 mM MgCl2) supplemented with GDP or GTP, respectively. Proteins were eluted with 2x 100µl elution buffer (100 mM Tris pH 8.0, 150 mM NaCl, 1 mM EDTA, 50 mM Biotin). Fractions were analyzed by the SDS-PAGE.

To test interactions between MglA-His6 and MalE-RomR and MalE-RomR/His6-RomX complex, MglA-His6 was preincubated with 4 times molar excess of GTP or GDP for 30 min at the room temperature. Next, MglA-His6 GTP/GDP was applied to an amylose resin with prebound MalE-RomR or MalE-RomR/His6-RomX for 30 min incubation at the room temperature. Proteins were prebounded to the resin and washed as described before. Next, resin was loaded into the column and washed with 3 times 1 ml of buffer 1 supplemented with 40 µM GTP or GDP respectively. Proteins were eluted from the column with 2 times 200 µl of buffer 2. Fractions were analyzed by the SDS-PAGE.

4.5.7 Nucleotide exchange experiments

Experiment of the release of mGDP from MglA was done as described in (Lenzen et al., 1995) with modifications. Briefly, MglA-His6 (2.016 µM) was preloaded with mGDP (403.226 nM) for 300 s at 25 ºC in the presence or absence of His6-RomX (4.032 µM) and/or MalE-RomR (4.032 µM) or BSA (8.064 µM). Loading was observed real time with monitoring fluorescence of mant (excitation 336, emission 450) on a temperature controlled ISS PC1 spectrofluorometer. Next, GTP was added to the final concentration of 2 µM. Reaction was performed in the exchange buffer (20 mM Tris pH 7.5, 200 mM NaCl, 10% glycerol, 1 mM DTT) at 25 ºC and fluorescence was measured every second. Reaction was followed from this point for 800 s. Obtained data was standardized, time point after addition of GTP, when fluorescence signal was stabilized, was set up as time point zero and fluorescence arbitrary units 1.

For the experiment with binding of mGTP to MglA, MglA-His6 (26.61 µM) was preloaded with GDP (106.48 µM) for 30 min at the room temperature. Subsequently, the MglA-His6/GDP mix was added to a reaction cuvette with mGTP (524.62 nM). After signal stabilization, His6-RomX and/or MalE-RomR or BSA were added. Final concentrations: MglA-His6 2 µM, 8 µM GDP, mGTP 400 nM, His6-RomX 8 µM, MalE-RomR 8 µM, BSA 16 µM. Reaction was performed in the exchange buffer (20 mM Tris
pH 7.5, 200 mM NaCl, 10% glycerol, 1 mM DTT) at 25 °C and mant fluorescence (exc 336, emi 450) was measure every second and followed for 1500s on a temperature controlled ISS PC1 spectrofluorometer. Obtained data was standardized, time point after addition of His6RomX and/or MalE-RomR or BSA, when fluorescence signal was stabilized, was set up as time point zero and fluorescence arbitrary units at zero. For the experiments that tested concentration dependences of His6RomX/MalE-RomR complex, conditions and concentration of MglA-His6, GDP and mGTP were the same as previously described.

In the experiment that tested binding of His6-RomX/MalE-RomR to MglA-His6 GTP, MglA-His6 was loaded with mGTP for the 150s, monitored real life with mant fluorescence (exc 336, emi 450) at 25 °C on a temperature controlled ISS PC1 spectrofluorometer. Subsequently, His6-RomX and/or MalE-RomR or BSA were added. Final concentrations in the reactions: MglA-His6 2 µM, mGTP 400 nM, His6-RomX 8 µM, MalE-RomR 8 µM, BSA 16 µM. Reaction was performed in the exchange buffer. After proteins addition, reaction was followed for 350 s with fluorescence read out every second. Data was standardized, time point after addition of His6RomX and/or MalE-RomR or BSA, when fluorescence signal was stabilized was set up as time point zero and fluorescence arbitrary units at zero.
5. References


Cherfils, J. & M. Zeghouf, (2013) Regulation of small GTPases by GEFs, GAPs, and GDIs. *Physiological reviews* **93**: 269-309.


REFERENCES


REFERENCES


Acknowledgements

First of all I would like to thank my mentor, Prof. MD, PhD Lotte Søgaard-Andersen for giving me opportunity to work with fascinating world of regulation of motility in *M. xanthus*, for advices and ideas. Thanks to this challenging project and under her supervision, I could develop scientific skills and critical thinking.

Furthermore, I would like to thank my IMPRS and thesis committee, specifically: Prof. Dr. Martin Thanbichler, Prof. Dr. Gert Bange, Prof. Dr. Victor Sourjik and Dr. Simon Ringgaard. I would also like to thank Dr. Dorota Skotnicka and Dr. Magdalena Polatynska for all the help in writing and proof reading of this work.

I am very thankful to the collaborators, Prof. Dr. Ulrich Gerland and Manon Wigbers for great help with developing tools and procedures for fluorescent images analysis. Furthermore, I would like to thank Andrea Harms for all the work with proteins purification and biochemical analysis, without her help this work would lack conclusions. For founding, I would like to thank International Max Planck Research School for Environmental, Cellular and Molecular Microbiology and the German-Israeli Project Cooperation “Spatial and Temporal Regulation of Macromolecular Complex Formation in Bacteria”.

I would also like to thank members of “motility lab”, Anna and Luis for great working atmosphere. I could always ask them for help and resources and could count on a good word on the bad day. It was a great pleasure to work with you.

I am happy that I meet so many nice people in the lab. I really appreciate time spent with former and current members of the lab: Bene, Monika, Tobi, Dorotka, Magdziszka, Anna, Soficzka, Maria, Nuria, Luis, Deepak and all the others. I would also like to thank Steffi for all the help. Starting a life in Germany is not easy and I really appreciate her support in everyday life.

Finally, I would never accomplish all this without love and support of my family. They have never lost the faith that I can finish this work and thanks to this, I stayed strong and kept thinking positively.
Curriculum Vitae

Personal data

Name: Dobromir Szadkowski
Date of birth: 05.10.1989
Place: Nowy Dwor Mazowiecki, Poland

Education:

09/2008 - 06/2011 Secondary School No.1
Lowicz, Poland

10/2008 – 07/2011 Bachelor of Science,
Univerity of Wroclaw, Poland
Supervisor: Dr. Anna Krasowska

10/2011 - 07/2013 Master of Science
Univerity of Wroclaw, Poland
Supervisor: Prof. Dr. Jolanta Zakrzewska-Czerwinska

From 08/2013 PhD program (Dr. rer. nat) Philipps-Universität Marburg
Max-Planck-Institute for Terrestrial Microbiology, Marburg
Supervisor: Prof. Dr. Lotte Søgaard-Andersen
List of publications

MglC, a Paralog of *Myxococcus xanthus* GTPase-Activating Protein MglB, Plays a Divergent Role in Motility Regulation,

The RomX/RomR MglA GEF complex establishes front-rear polarity for efficient motility in *M. xanthus*

Erklärung


Die Dissertation wurde in der jetzigen oder einer ähnlichen Form noch bei keiner anderen Hochschule eingereicht und hat noch keinen sonstigen Prüfungszwecken gedient.

Marburg, den ____.___.____

________________________

Dobromir Szadkowski
Einverständniserklärung

Hiermit erkläre ich mich einverstanden, dass die vorliegende Dissertation mit dem Titel

"Identification and characterization of RomX and RomY, two novel motility regulators in Myxococcus xanthus"

in Bibliotheken zugänglich gemacht wird. Dazu gehört, dass sie
von der Bibliothek der Einrichtung, in der meine Dissertation angefertigt wurde, zur Benutzung in ihren Räumen bereitgehalten wird;
in konventionellen und maschinen lesbaren Katalogen, Verzeichnissen und Datenbanken verzeichnet wird;
im Rahmen der urheberrechtlichen Bestimmungen für Kopierzwecke benutzt werden kann.

Marburg, den____.____.____

_______________________
Dobromir Szadkowski

_______________________
Prof. Dr. MD Lotte Søgaard-Andersen