Characterization of serine hydroxymethyltransferase GlyA as a potential source of D-alanine in Chlamydia pneumoniae

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INTRODUCTION

Acute and chronic diseases caused by Chlamydiaceae are a global health problem. The Gram-negative obligate intracellular pathogens depend on eukaryotic host cells to maintain their unique biphasic developmental cycle. One elusive phenomenon of the chlamydial biology has fascinated researches for two decades: for endobacteria, such as Chlamydiaceae, there is no need to withstand osmotic challenges and a functional cell wall has not been detected in these pathogens so far. Nevertheless, penicillin inhibits cell division in Chlamydiaceae resulting in enlarged aberrant bodies, a phenomenon known as chlamydial anomaly. D-alanine is a unique and essential component in the biosynthesis of bacterial cell walls. In free-living bacteria like Escherichia coli, penicillin-binding proteins such as multiple transpeptidases PBPs and PBPs, the putative targets of penicillin in Chlamydiaceae, cross-link adjacent peptidoglycan strands via meso-diaminopimelic acid and D-Ala-D-Ala moieties of pentapeptide side chains. In the absence of genes coding for alanine racemase Alr and DadX homologs, the source of D-Ala and thus the presence of substrates for PBPs activity in Chlamydiaceae has puzzled researchers for years. Interestingly, Chlamydiaceae genomes encode GlyA, a serine hydroxymethyltransferase that has been shown to exhibit slow racemization of D- and L-alanine as a side reaction in E. coli. We show that GlyA from Chlamydia pneumoniae can serve as a source of D-Ala. GlyA partially reversed the D-Ala auxotrophic phenotype of an E. coli racemase double mutant. Moreover, purified chlamydial GlyA had racemase activity on L-Ala in vitro and was inhibited by D-cycloserine, identifying GlyA, besides D-Ala ligase MurC/Ddl, as an additional target of this competitive inhibitor in Chlamydiaceae. Proof of D-Ala biosynthesis in Chlamydiaceae helps to clarify the structure of cell wall precursor lipid II and the role of chlamydial penicillin-binding proteins in the development of non-dividing aberrant chlamydial bodies and persistence in the presence of penicillin.

Keywords: chlamydial anomaly, persistence, aberrant bodies, D-alanine, alanine racemase, GlyA, penicillin, D-cycloserine

For intracellular Chlamydiaceae, there is no need to withstand osmotic challenges, and a functional cell wall has not been detected in these pathogens so far. Nevertheless, penicillin inhibits cell division in Chlamydiaceae resulting in enlarged aberrant bodies, a phenomenon known as chlamydial anomaly. D-alanine is a unique and essential component in the biosynthesis of bacterial cell walls. In free-living bacteria like Escherichia coli, penicillin-binding proteins such as multiple transpeptidases PBPs and PBPs, the putative targets of penicillin in Chlamydiaceae, cross-link adjacent peptidoglycan strands via meso-diaminopimelic acid and D-Ala-D-Ala moieties of pentapeptide side chains. In the absence of genes coding for alanine racemase Alr and DadX homologs, the source of D-Ala and thus the presence of substrates for PBPs activity in Chlamydiaceae has puzzled researchers for years. Interestingly, Chlamydiaceae genomes encode GlyA, a serine hydroxymethyltransferase that has been shown to exhibit slow racemization of D- and L-alanine as a side reaction in E. coli. We show that GlyA from Chlamydia pneumoniae can serve as a source of D-Ala. GlyA partially reversed the D-Ala auxotrophic phenotype of an E. coli racemase double mutant. Moreover, purified chlamydial GlyA had racemase activity on L-Ala in vitro and was inhibited by D-cycloserine, identifying GlyA, besides D-Ala ligase MurC/Ddl, as an additional target of this competitive inhibitor in Chlamydiaceae. Proof of D-Ala biosynthesis in Chlamydiaceae helps to clarify the structure of cell wall precursor lipid II and the role of chlamydial penicillin-binding proteins in the development of non-dividing aberrant chlamydial bodies and persistence in the presence of penicillin.
FIGURE 1 | Proposed lipid II pathway in Chlamydiaceae. A complete cycle of lipid II biosynthesis, including translocation to the periplasm, processing and bactoprenol carrier recycling is required for coordinated function of the divisome machinery and modulation of Nod1 and Nod2 mediated host immune response to chlamydial muropeptides. Biosynthesis of lipid II takes place in the cytoplasm and at the inner leaflet of the cytoplasmic membrane. In two consecutive biosynthesis steps of L-Ala racemization and D-Ala ligation, catalyzed by GlyA and the MurC/Ddl fusion protein, the D-Ala-D-Ala dipeptide is produced. MurF adds the dipeptide to the nascent peptide chain to complete synthesis of the soluble precursor UDP-MurNAc-pentapeptide and to provide D-Ala-D-Ala moieties in the cell wall precursors for transpeptidation activity of the penicillin-binding proteins PBP2 and PBP3. Actin-ortholog MreB functionally organizes MurF, MraY, and MurG (Gaballah et al., 2011), the last three enzymes in lipid II biosynthesis, at the septum. The synthesized precursor is translocated to the periplasm and processed by the concerted activity of the PBP enzymes and amidase AmiA to allow for bactoprenol-P recycling. In the process, the rudimentary by-product found by Liechti et al. (2013), in which the peptide side chains are cross-linked by peptide bonds, might result (Ghuysen and Goffin, 1999) and released muropeptides might contribute to modulation of the host immune response (McCoy and Maurelli, 2006). Chlamydiaceae lack transglycosylases as well as endopeptidases and pyrophosphorylases described so far to link lipid II sugar units to form glycan chains, to cleave peptide bridges between cross-linked glycan chains and to dephosphorylate bactoprenol-PP, respectively. Moreover, the L-Glu racemase MurI is absent. Question marks and dashed arrows highlight steps of the pathway that remain to be clarified. (GlcNAc: N-acetylglucosamine; MurNAc: N-acetylmuramic acid).

essential component in the biosynthesis of bacterial cell walls. The non-proteinogenic amino acid is synthesized by alanine racemases Alr and DadX and ligated by Ddl to form D-Ala-D-Ala. The dipeptide is attached to the amino acid in position 3 by the action of MurF to complete the pentapeptide side chain. In the first membrane-linked step, MraY catalyzes the synthesis of lipid I by transferring UDP-MurNAc-pentapeptide to the lipid carrier bactoprenol-phosphate (undecaprenyl-P). With the addition of UDP-GlcNAc, MurG synthesizes lipid II, the completed peptidoglycan cell wall building block. Lipid II is then translocated by the flippase FtsW to the outside of the cell and incorporated into the peptidoglycan network by the action of penicillin-binding proteins (PBPs) which exhibit transglycosylase and DD-transpeptidase activities.

The human cytosolic Pattern Recognition Receptors, Nod1 and Nod2, sensing bacterial cell wall fragments, recognize intracellular C. pneumoniae and subsequently mediate activation of the transcription factor NFkB which plays a key role in regulating
the immune response to infection (McCoy and Maurelli, 2006). Nod1 and Nod2 receptor mediated recognition, together with the susceptibility to penicillin, suggests that cell wall precursors/peptidoglycan fragments are synthesized by Chlamydiaceae during infection.

A nearly complete lipid II biosynthesis pathway has been found in genomes of Chlamydiaceae (Figure 1) (McCoy and Maurelli, 2006), and functional conservation of enzymes catalyzing cytoplasmic steps (MurA, MurC/Ddl, CT390, DapF, MurE, MurF) and the two membrane-linked steps (MraY and MurG) of cell wall precursor biosynthesis has been demonstrated (McCoy and Maurelli, 2006; McCoy et al., 2006; Henrichfreise et al., 2009; Patin et al., 2009, 2012). Chlamydiaceae genomes code for only two PBPs that serve as DD-transpeptidases in free-living bacteria. PBP2 and PBP3 are the putative targets of penicillin in Chlamydiaceae and cross-link adjacent peptidoglycan strands via meso-diaminopimelic acid and D-Ala-D-Ala moieties in E. coli. AmiA is the only ortholog of septal peptidoglycan hydrolyzing amidases found in chlamydial genomes. Moreover, Chlamydiaceae harbor a rudimentary set of cell division proteins, lacking the central organizer FtsZ, but comprising FtsW, FtsI (PBP3) and FtsK, and possess, despite their spherical shape, the cytoskeletal protein MreB (Gaballah et al., 2011). Chlamydial MreB was shown to interact with key components in lipid II biosynthesis and FtsK (Gaballah et al., 2011; Ouellette et al., 2012).

We proposed that retaining biosynthesis of lipid II in cell wall-lacking “minimal bacteria,” like Chlamydiaceae, may reflect a vital role of the lipid II pathway in prokaryotic cell division (Henrichfreise et al., 2009). Moreover, we discussed a model for the role of the conserved lipid II pathway in maintaining a functional divisome and contributing to modulation of host response in Chlamydiaceae (Figure 1). A recent study revealed the presence of cell wall sacculi in Protochlamydia, a genus of evolutionary older amoeba symbionts with less reduced genomes as compared to Chlamydiaceae (Pilhofer et al., 2013). In the pathogenic Chlamydiaceae, however, no functional cell wall but ring-like shaped structures were found and supposed to contain peptidoglycan-like material and localize to the division plane (Liechti et al., 2013). These findings are consistent with our model described above which implicates a crucial function of the PBP2 and PBP3 DD-transpeptidase activity in lipid II processing and sustaining a complete cycle of lipid II biosynthesis and recycling. PBP catalyzed DD-transpeptidation depends on the presence of the D-Ala-D-Ala terminus in the pentapeptide side chain of cell wall building blocks. Genomes of Chlamydiaceae do not encode homologs of the pyridoxal-5′-phosphate (PLP) cofactor requiring alanine racemases Alr and DadX. Therefore, the source of D-Ala and thus the presence of substrates for PBP2 and PBP3 activity in Chlamydiaceae have remained unclear for years.

We searched chlamydial genomes for genes encoding other PLP dependent proteins and found serine hydroxymethyltransferase GlyA to be conserved in all chlamydial genera. Serine hydroxymethyltransferases are found in eu- and prokaryotes and are well known for their function in reversible interconversion of serine and glycine using tetrahydrofolate as the one-carbon carrier. In addition, the enzymes show a particularly broad reaction specificity and catalyze other side reactions typical for PLP dependent enzymes, such as decarboxylation, transamination and retroaldol cleavage (Contestabile et al., 2001). Moreover, an alanine racemase co-activity was proven in vitro for GlyA from E. coli (Shostak and Schirch, 1988).

The aim of this study was to analyze GlyA as a potential source of D-Ala in Chlamydiaceae.

Here, we demonstrate that GlyA from C. pneumoniae is capable of the racemization of alanine in vivo and in vitro implicating that the enzyme can substitute for the absent alanine racemases and that D-Ala is self-synthesized in Chlamydiaceae.

RESULTS

RACEMIZATION OF ALANINE IN CHLAMYDIAE

Using BLAST alignments, we searched Chlamydiaceae and environmental chlamydiae genomes to identify genes coding for orthologs of E. coli PLP cofactor-requiring enzymes known to confer alanine racemization activity (Table 1). In contrast to the Chlamydiaceae and Simkania, the three environmental chlamydiae genera Parachlamydia, Protochlamydia, and Waddlia harbored one ortholog of the Alr or DadX alanine racemases. GlyA

| Table 1 | PLP cofactor-requiring enzymes involved in biosynthesis of D-Ala. |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| **E. coli**     | **Cpn**         | **Ctr**         | **Pac**         | **Pam**         | **Wch**         | **Sne**         |
| Alr (alanine racemase) | –               | –               | –               | pc0631          | wcw_0679        | –               |
|                  | (2e–34)         | (3e–34)         |                 | (1e–32)         |                 |                 |
| DadX (alanine racemase 2) | –               | –               | PUV_23750       | –               | –               | –               |
|                  | (2e–28)         | (2e–28)         |                 | (2e–32)         |                 |                 |
| GlyA (serine hydroxymethyltransferase) | CPn0521 (2e–107) | CT432 (2e–108) | PUV_05830 (6e–118) | pc0444 (3e–107) | wcw_1457 (2e–117) | SNE_A20 270 (1e–114) |
| MetC (cystathionine beta-lyase) | –               | –               | PUV_18690 (2e–40) | –               | wcw_1145 (4e–40) | –               |

Locus tags of genes coding for enzymes involved in biosynthesis of D-Ala are shown for two exemplary Chlamydiaceae and for environmental chlamydiae species. The expected (E) values of BLAST P alignments are listed in brackets. E. coli, Escherichia coli W3110 (NC_007779.1); Cpn, Chlamydia pneumoniae CWL029 (NC_000922); Ctr, Chlamydia trachomatis D/UW-2/Cx (NC_000117); Pac, Parachlamydia acanthamoebae UV-7 (NC_015702.1); Pam, Protochlamydia amoebophila UWE25 (NC_005861.1); Wch, Waddlia chondrophila WSU 86-1044 (NC_014225.1); Sne, Simkania negevensis Z (NC_015713.1).
was the only enzyme to be encoded in *Chlamydiaceae* and in all environmental chlamydiae.

**IN VIVO ACTIVITY OF GlyA FROM C. pneumoniae**

In the absence of a tractable system to genetically manipulate *C. pneumoniae* we tested whether heterologously expressed *C. pneumoniae* GlyA (GlyACp) shows an effect on the D-Ala auxotrophic phenotype of an *E. coli ΔalrΔdadX* racemase double mutant strain. Our experiments in liquid and solid culture revealed that chlamydial GlyA did not completely reverse the need of exogenous D-Ala of the racemase mutant strain but favored its growth under D-Ala limited conditions (Figure 2). These findings suggest that GlyACp is a functional alanine racemase and capable of generating D-Ala in *E. coli*.

**RECOMBINANT GlyACp HAS L-ALANINE RACEMASE ACTIVITY**

To investigate the potential alanine racemase activity of the serine hydroxymethyltransferase GlyA from *C. pneumoniae*, we overexpressed recombinant GlyACp in *E. coli* and purified the Strep-tagged protein. *In vitro* activity of GlyACp was tested in a D-amino acid oxidase coupled enzymatic assay containing L-Ala and cofactor PLP. Alanine racemase from *Bacillus stearothermophilus* served as positive control. D-Ala that was produced by GlyA was converted to pyruvate by the activity of D-amino acid oxidase (DAAO) and colorimetrically quantified. The chlamydial GlyA converted L-Ala to D-Ala *in vitro* exhibiting weak racemase activity in comparison to the enzyme from *B. stearothermophilus* (Figure 3).

**GlyACp RACEMASE ACTIVITY IS SENSITIVE TO D-CYCLOSERINE**

D-cycloserine is a structural analog of D-Ala and competitively inhibits activity of alanine racemases and D-Ala ligases from free-living bacteria (Strominger et al., 1960; Lambert and Neuhaus, 1972). The inhibitor has anti-chlamydial activity in chick embryo yolk sac infection (Moulder et al., 1963) which can be reversed by the addition of D-Ala. For the *C. trachomatis* D-Ala ligase Ddl which is encoded as a fusion with MurC, as typical for *Chlamydiaceae*, sensitivity to D-cycloserine has been proven before (McCoy and Maurelli, 2005). We performed activity assays for GlyACp in the presence of D-cycloserine and identified the enzyme to be a second target of D-cycloserine in *Chlamydiaceae* (Figure 3).

**DISCUSSION**

The source of D-Ala in *Chlamydiaceae* and thus the presence of the transpeptidation substrates for PBPs is a crucial aspect of the chlamydial anomaly that will help to gain understanding of the penicillin induced persistence in these human pathogens. Penicillin and other beta-lactams structurally mimic the D-Ala-D-Ala terminus of the pentapeptide side chain of the lipid II cell wall building blocks and are recognized by the active sites of DD-transpeptidase PBPs. Beta-lactams are active against *Chlamydiaceae* in cell culture (McCoy and Maurelli, 2006; Skilton...
et al., 2009) and in vivo (Phillips Campbell et al., 2012) and the chlamydial monofunctional DD-transpeptidases PBP2 and PBP3, recovered in detergent-soluble fractions from whole cell preparations, bind [H\(^3\)]-penicillin (Barbour et al., 1982). Moreover, the recombinant MurC/Ddl protein from C. trachomatis has been shown to specifically ligate D-Ala but not the L-Ala enantiomer to form alanine dipeptidases and MurF was demonstrated to add these D-Ala-D-Ala dipeptides to the lipid II peptide side chain (McCoy and Maurelli, 2006; Patin et al., 2012). Moulder et al. (1963) proved anti-chlamydial activity of D-cycloserine in chick embryo yolk sac infection models and demonstrated the specific reversal of this effect by the addition of D-alanine (Moulder et al., 1963). D-cycloserine is a structural analog of D-Ala and well known as competitive inhibitor of alanine racemases and the D-Ala ligase Ddl in other bacteria (Strominger et al., 1960; Lambert and Neuhaus, 1972). Finally, feeding of replicating Chlamydia trachomatis with D-Ala-D-Ala probes revealed evidence for the incorporation of D-Ala into ring-like shaped peptidoglycan-like structures (Liechti et al., 2013). All in all, these data strongly indicate that D-Ala is present in the cells of Chlamydiaceae and plays an essential role in chlamydial cell biology. In the past, the mammalian host was discussed as a potential source of D-Ala (McCoy and Maurelli, 2006) and in Chlamydiaceae genomes a D-alanine permease (DagA_2) was annotated (Stephens et al., 1998) that could allow for the passive transport of D-Ala over the chlamydial cytoplasmic membrane. Though D-Ala is found almost exclusively in the microbial world and is not synthesized in mammalian cells, D-Ala was detected in trace quantities in mammalian urine and tissues, apparently due to breakdown products from intestinal and food bacteria (Guoyao, 2013). Nevertheless, experiments with Listeria monocytogenes and Shigella flexneri indicated that mammalian host cells cannot serve as source of D-Ala. Alanine racemase knockout mutants of both facultative intracellular species failed to survive within mammalian host cells unless exogenous D-Ala was added to the cell culture medium (McCoy and Maurelli, 2006). Based on these data, self-biosynthesis of D-Ala by alternative racemase activity is more likely the source of D-Ala in Chlamydiaceae.

For E. coli, besides the constitutive expressed Alr racemase and the catabolic DadX racemase, two other PLP dependent enzymes involved in the methionine pathway have been shown to confer L-Ala racemization as a side reaction. Overexpressed cystathionine beta-lyase MetC completely reversed the D-Ala auxotrophic phenotype of an E. coli racemase double mutant whereas racemase co-activity of serine hydroxymethyltransferase GlyA was not sufficient to allow for growth on D-Ala lacking medium (Kang et al., 2011). Recently, in vitro activity of MetC from Wolbachia indicated that the enzyme substitute for the absent alanine racemases in these endosymbionts of many arthropods and filarial nematodes. Wolbachia are excellent targets for anti-filarial therapy. Similar to Chlamydiaceae, these obligate intracellular bacteria lack a cell wall but treatment with lipid II biosynthesis blocking fosfomycin, results in enlarged Wolbachia cells (Vollmer et al., 2013). Among the Chlamydiaceae only the environmental Parachlamydia and Waddlia possess a MetC ortholog but all chlamydial genera harbor GlyA. The serine hydroxymethyltransferase is the only component of the methionine pathway that is encoded by Chlamydiaceae genomes and phylogenetic analysis indicated lateral transfer of the glyA gene from Actinobacteria to the common ancestor of Chlamydiaceae (Griffiths and Gupta, 2006). Moreover, transcription profiles revealed overlapping expression of the genes encoding GlyA, enzymes for lipid II biosynthesis (MurA to MurF, MraY, and MurG) as well as processing (PBP2, PBP3, and AmiA), the structural protein MreB, and the cell division proteins FtsW and FtsK (Belland et al., 2003). These data suggest an essential function of GlyA in chlamydial biology and correlation with the cellular processes of lipid II biosynthesis and cytokinesis.

We demonstrated L-Ala racemase activity for the GlyA enzyme from C. pneumoniae in vivo in an E. coli racemase double mutant and characterized the purified protein in vitro. Moreover, we identified GlyA as a second target of D-cycloserine besides MurC/Ddl in Chlamydiaceae.

Our results implicate that the enzyme can substitute for the absent alanine racemases and that D-Ala is present and self-synthesized in Chlamydiaceae. The observed weak alanine racemase activity of GlyACp cannot completely compensate D-Ala requirements of the E. coli racemase mutant to build a functional cell wall but might be sufficient to produce D-Ala in amounts that maintain the lipid II biosynthesis pathway in the cell wall-lacking Chlamydiaceae for the proposed functions of the cell wall precursor in co-ordination of cell division and modulation of the host immune response. Future research toward the elucidation of the chlamydial anomaly will include the isolation and structural characterization of lipid II building blocks and biochemical analysis of the penicillin binding proteins PBP2 and PBP3, the putative targets of penicillin whose activity depends on the presence of D-Ala-D-Ala in the pentapeptide side chain of lipid II.

Like other effectors, such as interferon-γ and tumor necrosis factor-α, penicillin can be used to induce persistence as an experimental tool to study chlamydial/host interactions. Knowledge of the underlying mechanisms of penicillin induced formation of ABs will help to assess these results on the pathogenicity of Chlamydiaceae.

Moreover, analysis of the molecular biology of penicillin induced persistence is important to improve understanding of long-term infection in patients in particular as to the role of chlamydial cell wall precursors in immune modulation and refractory of ABs to anti-chlamydial agents.

**MATERIALS AND METHODS**

**BACTERIAL STRAINS AND GROWTH CONDITIONS**

E. coli JM83 harboring the GlyACp expression vector was grown on Luria Bertani (LB) agar plates containing 30 μg/ml chloramphenicol and 100 μg/ml ampicillin, respectively. The temperature sensitive E. coli ΔalrΔdadX racemase double mutant TKL-10 was maintained on LB agar plates containing 25 μg/ml thymine.

**IN VIVO COMPLEMENTATION**

E. coli TKL-10 ΔalrΔdadX was transformed with pET21b-glyACp and grown in liquid or on solid LB medium supplemented with 5 mg/L D-Ala, 50 μM PLP, 25 μg/ml thymine, 50 μg/ml ampicillin and 0.1 mM IPTG at 42°C. For each experiment three independently generated transformants were used and controls with
E. coli TKL-10 ΔalrΔdadX harboring the empty pET21b vector were carried out.

**CLONING OF glyA**

The *glyA* gene from *C. pneumoniae* Gid was amplified by PCR using primer glyACp_f (5′-ATGGTAGGTCTCGGCCCTTGCTAAAAAGTTTGGAAATTTAAA-3′) and glyACp_r (5′-ATGGTAGGTCTCGGCCCTAACTAAAGCTTCTAATTCAATTTCCGG-3′) and cloned into pASK-IBA2c (IBA, Germany) using the BsaI restriction site to generate an N-terminal OmpA-leader peptide fused, C-terminal Strep-tagged protein for periplasmic overproduction. For cytoplasmic expression using NdeI and NotI restriction sites.

**OVERPRODUCTION AND PURIFICATION OF GlyACp**

*E. coli* JM83 cells, transformed with pASK-IBA2c-glyACp, were grown in no salt LB in presence of 30 μg/ml of chloramphenicol, 250 mM sucrose and 50 mM L-serine at 30°C. After induction at an OD600 of 1.2 with 200 ng/ml anhydrotetracycline (AHT), 50 μM PLP and 200 μM of folinic acid were added and the cells were incubated for 4 h at 25°C. The purification of GlyACp was performed using the protocol for cleared lysates recommended by the manufacturer (IBA, Germany) with small modifications: the buffers contained 2% N-lauroylsarcosine (or 0.1% N-lauroylsarcosine in the washing and elution buffer), 2 mM 1,4-dithiothreitol (DTT) and 50 μM PLP. Purity of the protein was controlled using SDS-PAGE.

**IN VITRO GlyACp ACTIVITY ASSAY**

 Racemase activity of GlyACp was determined in a DAOO coupled enzymatic assay as described previously with slight modifications (Francois and Kappock, 2007). Briefly, 1 μg GlyACp or DAOO were incubated with 50 mM L-alanine in a final volume of 60 μl for 16 h at 37°C in 50 mM KH2PO4, pH 8, 100 mM KCl, 80 μM PLP and 2 mM DTT. D-Ala that was derived from GlyACp racemization was deaminated into pyruvate by the DAOO from *E. coli* JM83 cells, transformed with pASK-IBA2c-glyACp, cloned into pET21b (Novagen, VWR, Germany) using NdeI and NotI restriction sites.

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