Development of ADPribosyl Ubiquitin Analogues to Study Enzymes Involved in Legionella Infection

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Abstract: Legionnaires’ disease is caused by infection with the intracellularly replicating Gram-negative bacterium Legionella pneumophila. This pathogen uses an unconventional way of ubiquitinating host proteins by generating a phosphoribosyl linkage between substrate proteins and ubiquitin by making use of an ADPribosylated ubiquitin (UbADPr) intermediate. The family of SidE effector enzymes that catalyze this reaction is counteracted by Legionella hydrolases, which are called Dups. This unusual ubiquitination process is important for Legionella proliferation and understanding these processes on a molecular level might prove invaluable in finding new treatments. Herein, a modular approach is used for the synthesis of triazole-linked UbADPr, and analogues thereof, and their affinity towards the hydrolyse DupA is determined and hydrolysis rates are compared to natively linked UbADPr. The inhibitory effects of modified Ub on the canonical eukaryotic E1-enzyme Uba1 are investigated and rationalized in the context of a high-resolution crystal structure reported herein. Finally, it is shown that synthetic UbADPr analogues can be used to effectively pull-down overexpressed DupA from cell lysate.

Introduction

The dogma in post-translational modification by ubiquitin (Ub) is that Ub-activating enzymes (E1), Ub-conjugating enzymes (E2), and Ub ligases (E3) are required to work together to activate the C-terminal carboxylate of Ub, in an adenosine triphosphate (ATP)-dependent manner, and subsequently ligate it to the N-terminal glycine of the substrate. DupA and DupB were identified on the basis of their structural homology to the SidE effector enzymes, which are able to ADP-ribosylate the δ-guanidinium group of arginine 42 (Arg42) of Ub at the expense of nicotinamide adenine dinucleotide (NAD⁺) by using their mono-ADP-transf erase (mART) domain in the first step, followed by the action of their phosphodiesterase (PDE) domain, which catalyzes the transfer of phosphoribosyl-Ub (UbPr) to the serine of a substrate protein, while expelling adenosine monophosphate (AMP; Figure 1). Legionella has its own regulatory mechanism in place to control the temporal activity of these SidE ligases by blocking their active-site glutamate using the glutamylase SidJ. The recently identified debiquitinases for phosphoribosyl ubiquitination (Dups), DupA and DupB, also known as LaiE and LaIF, counteract the SidE-mediated attachment of phosphoribosyl-linked Ub to substrates. DupA and DupB were identified on the basis of their structural homology to the SidE PDE domains, but lack the ability to Pr-ubiquitinate the substrate protein Rab33B upon incubation with ADPribosylated Ub. These DUPS, however, were shown to release proteins that were Pr-
ubiquitinated by SidE ligases by cleaving the phosphodiester bond between the substrate serine and Arg42UbPr.[4a] Although SidE ligases and Dups have opposite functions, they are structurally very similar and, even more so, the ligase SdeA is shown to mediate hydrolysis of the pyrophosphate bond in UbADPr if no suitable substrate protein is present. The ligase effectively mediates transfer of a water molecule instead of a serine residue to the activated pyrophosphate bond, thereby expelling AMP.[1b]

By using a catalytically inactive version of DupA to enrich for Pr-ubiquitinated substrates in HEK293T cells infected with Legionella, 180 host proteins were identified based on their affinity for DupA.[4a] Most of these proteins are involved in endoplasmic reticulum membrane recruitment to Legionella-containing vacuoles (LCVs). This highlights the importance of Pr-ubiquitination upon Legionella infection because maintaining LCV integrity is essential for Legionella proliferation and the onset of Legionnaires’ disease.

In the canonical ubiquitination pathway, the use of chemically prepared tools, such as substrate reagents and activity-based probes, has been a widely applied and successful approach to allow the study of kinetic parameters, as well as capturing and identifying both ligases and proteases.[5] The recent development of fluorescent polarization based assay reagents and inhibitors to study enzymes involved in the Pr-ubiquitination pathway highlights the applicability of chemically synthesized tools to study Pr-ubiquitination.[6] Hence, the construction of probes targeting the ADPr-mediated ubiquitination machinery will be a similarly useful asset in studying the enzymes involved. We set out to prepare α-O-propargyl ADPr (1; Figure 2) and its stabilized methylene bisphosphonate analogue, α-O-propargyl me-ADPr (2), in which oxygen in the pyrophosphate linkage is replaced with a methylene group.[7] Facile copper-catalyzed Huisgen azide-to-alkyne 1,3-dipolar cycloaddition (CuAAC, or click reaction) of these propargyl-containing ADPr analogues to azide-modified Ub allowed the gen-

![Figure 1. Schematic representation of substrate ubiquitination by noncanonical Legionella SidE enzymes and substrate release by DupA.](image1)

![Figure 2. A) Modular approach of using click chemistry to construct triazoleUb analogues. B) Schematic structure of native ArgUbADPr.](image2)
eration of probes 4 and 5 to investigate Legionella enzyme ac-

tivity. The rationale behind the oxygen-to-carbon substitution in 5 is that the PDE activity in SidE enzymes relies on expelling AMP. Replacing the diphosphate with a methylene bisphosphonate prevents this step from occurring, thereby blocking SidE-mediated conjugation to substrate proteins. This stabilized Ubme-ADPr conjugate 5 would thus be able to capture the Legionella enzyme and function as a suitable nonhydroyzable probe to target such enzymes. Additionally, little is known about the role of the phosphoribosyl residue that remains on the Ub moiety after DupA-mediated hydrolysis of the targeted substrate protein, and we envision Ub-based tools, such as 6, to be essential to decipher the role of Ub.

Results and Discussion

The inherent incompatibility of ADPr and other nucleotide-based structures with strongly acidic conditions routinely used in fluoroenzymeoxycarbonyl (Fmoc) tert-butyloxycarbonyl (Boc) solid-phase peptide synthesis (SPPS) prohibits the total chemical synthesis of large ADPr peptides or proteins and only allows for the construction of relatively short ADPr peptides by adapting protecting-group schemes. This has triggered the development of modular synthetic approaches towards such structures, in which the polypeptide can be treated with a strong acid to remove protecting groups and be released from a peptide synthesis resin followed by HPLC purification, before it is attached to the delicate ADPr moiety. To allow this final conjugation step to be executed under mild conditions, we envision click chemistry to be the most effective strategy. Upon substituting Arg42 of Ub with azidothomoalanine through SPPS, conjugation can be achieved at physiological pH with a minimum of chemical additives (3 mM CuSO4, 20 mM sodium ascorbate, and 3 mM tris[1-benzyl-4-triazolyl]-methyl]amine (TBTA) ligand) to the α-oriented propargyl ether on the anomic position of the riboside in ADPr (1), me-ADPr (2), or Pr (3) (Figure 2A). The Ub60Cu conjugate formed in such a CuAAC reaction carries a triazole linkage between the ribose and peptide part, from now on referred to as triazoleUbADPr, thus slightly deviating from the native arginine guanidinium linkage (Figure 2B).

After the successful CuAAC reactions of 1, 2, and 3 to Ub carrying an azidothomoalanine mutation on position 42, triazole-linked triazole42UbADPr (4), triazole42Ubme-ADPr (5), and triazole42UbPr (6) were obtained. We set out to compare these triazole-linked conjugates, and natively linked Arg42UbADPr, which was prepared enzymatically by using a SdeA mutant, for their affinity towards DupA.

To this end, we used bio-layer interferometry (BLI), and repeated the assay that was described earlier, by immobilizing the different Ub analogues on streptavidin (SA) biosensor tips through the biotin handle attached on the N terminus of Ub, and using glutathione S-transferase (GST)-tagged DupA-H67A as the analyte. With this setup, conjugates 4 and 5 show very high affinities of 11.2 and 10.6 nM, respectively, which are comparable to the Kd value of 5.7 nM observed for native Arg42UbADPr (Figure S3A in the Supporting Information). However, the observed nanomolar affinity for unmodified Ub (54.5 nM) would render all DupA inside a human cell bound to unmodified Ub (product-like) and unavailable for catalysis. We repeated the experiment with the catalytically inactive mutant, DupA-H67A, lacking the GST tag (Figure 3A). The results obtained show biologically plausible Kd values of 2.2 and 1.2 μM for 4 and 5, respectively, whereas 6 and unmodified Ub have at least a 15-fold reduced affinity (Figure 3B and Figure S3B in the Supporting Information). The discrepancy between the two assays could potentially be attributed to an artefact arising from dimerization of the GST-tagged analyte that we cannot fully explain at this point (see Figure S3D in the Supporting Information).

The resulting Kd values in the absence of the GST tag make biological sense and would fit with the mechanism of the hydrolase, which accepts substrates linked through a phosphodiester bond to ribosylated Ub, with micromolar affinity, and releases the phosphomoenoester Ub product due to the lower affinity of the latter.

Next, we wondered whether DupA could hydrolyze 4 to form triazoleUbPr, as reported previously for native Arg42UbADPr. We indeed observed robust hydrolysis of natively linked Arg42UbADPr (1 μM) by 300 nM DupA after incubation for 1 h at 37 °C (Figure 4A). Upon applying the same conditions to triazole-linked 4, we observed a similar hydrolysis reaction and formation of phosphoribosyl Ub 6, as monitored by means of mass spectrometry (Figure 4B), whereas DupA was not able to mediate hydrolysis of stabilized 5 (see Figure S4 in the Supporting Information). In control experiments on both native Arg42UbADPr and triazole-linked 4 in the absence of DupA, only a minor amount of hydrolysis of the pyrophosphate bond is observed, which is most likely due to the acidic conditions employed during mass spectrometry. DupA-mediated hydrolysis can be attributed to the catalytic specificity of the enzyme because control experiments with triazole-linked triazole54UbADPr, triazole74UbADPr, and triazole74UbPr showed neither hydrolysis nor formation of the corresponding Ub products. To investigate this further, we assessed these control compounds for DupA affinity using our BLI setup (Figure S3C in the Supporting Information). We could not detect significant binding of triazole54UbADPr or triazole74UbADPr to DupA H67A, giving a clue to why they are

![Figure 3. Results of BLI analysis. A) Concentration-dependent response curves of 4 to DupA-H67A. B) A comparison of the binding affinities of 4, 5, 6, and Ub.](image-url)
not processed by DupA. For triazole72UbADPr, however, we could detect binding to DupA H67A with a $K_d$ of 9.3 $\mu$m, suggesting that the adenosine moiety could be positioned in a manner resembling the configuration present in 4, but so that the di-phosphate linkage is not oriented appropriately for hydrolysis towards triazole72UbPr. To investigate any differences in catalysis of DupA on 4 or native Arg42UbADPr, we followed DupA-mediated UbPr formation over time by mass spectrometry using a lower enzyme concentration of DupA (30 n$m$) on $3 \mu$m of both hydrolyzable substrates. We observed a clear reduction in velocity (3.5-fold), when comparing relative $V_{max}$ for DupA-mediated hydrolysis of triazole-linked 4 to that of native Arg42UbADPr (Figure 4 C). It is apparent that, although accepted by DupA, triazole-linked 4 is hydrolyzed at a reduced rate relative to that of native Arg42UbADPr. Most likely, this reduced cleavage rate is caused by the more sterically demanding and rigid triazole linkage.

ADPribosylation or phosphoribosylation of Arg42 in Ub impairs the conventional ubiquitination machinery because activation by E1, trans-thioesterification to E2, and E3-mediated discharge from the E2 were shown to be compromised upon the introduction of the modification by Legionella ligase SdeA.[1b] From the crystal structure of Arg42UbADPr, it becomes apparent that any modification of Arg42 or Arg72 will interfere with Ub binding to E1, which could explain the inability of E1 to activate the Ub$^\beta$ molecule.[1b] These two arginine residues are reported to be critical in the interaction with the E1 enzyme Uba1, as in a previous study mutations of Arg42 or Arg72 to leucine were shown to result in a dramatically lower affinity between the E1 enzyme and Ub adenylate.[11] In addition, residue 72 is crucial for determining Ub-like specific recognition by E1, where for Ub this residue is an arginine, for Nedd8 it is an alanine, and for SUMO-family members it is either a glutamate or glutamine residue.[12] We managed to improve the resolution of our previously reported X-ray structure of Saccharomyces cerevisiae Uba1 in complex with Ub from crystals diffracting anisotropically to 2.03 $\AA$ (Figure 5A), which shows the C-terminal tail of Ub reaching towards the adenylation site of Uba1. This yeast homolog of Uba1 has a conserved overall structure with high sequence identity (68%) in the active adenylation domain compared to human Uba1.[13] Figure 5A shows the crossover loop connecting the adenylation domain to the catalytic cysteine domain encompassing the C-terminal tail of Ub just above the Arg42 and Arg72 guanidinium groups of Ub. The close spatial positioning of these residues could explain our observation that Ub ADPribosylated at Arg72 can still bind to DupA.

Furthermore, we observe a weak electron density for the guanidinium groups of Arg54 and Arg74 in this structure, indicating flexibility and the possibility for these residues to adopt ognition by E1, where for Ub this residue is an arginine, for Nedd8 it is an alanine, and for SUMO-family members it is either a glutamate or glutamine residue.[12] We managed to improve the resolution of our previously reported X-ray structure of Saccharomyces cerevisiae Uba1 in complex with Ub from crystals diffracting anisotropically to 2.03 $\AA$ (Figure 5A), which shows the C-terminal tail of Ub reaching towards the adenyla-
multiple conformations. Both the $\sigma$-weighted 2$F_o$—$F_c$ electron density map and the B factors of the guanidinium groups of the arginine residues suggest that Arg42 and Arg72 remain in a more rigid conformation, as part of the binding interface with the Ub1 adenylation domain, compared to Arg74 and Arg54. The B factors of the CZ atom of the guanidinium groups of Arg42, Arg54, Arg72, and Arg74 of Ub are 38.5, 63.7, 30, and 55.9 Å$^2$, respectively. The guanidinium groups of Arg42 and Arg72 show well-defined electron densities, indicative of their fixed placement in a single conformation, necessary for binding to the adenylation domain of Ub1. To validate whether the triazole-linked UbADP analogues would interfere with Ub1-mediated activation of Ub, we incubated triazole$^{54}$Ub$^{\text{me-ADP}}$, triazole$^{72}$Ub$^{\text{me-ADP}}$, triazole$^{74}$Ub$^{\text{me-ADP}}$, triazole$^{42}$Ub$^{\text{Pr}}$, triazole$^{72}$Ub$^{\text{Pr}}$, and triazole$^{74}$Ub$^{\text{Pr}}$ with human Ub1 (E1) in the presence of sodium 2-sulfanylthanesulfonate (MESNa) and ATP, and monitored thioester formation using mass spectrometry (Figure 5B). It became apparent that both me-ADP$^\text{Pr}$ and Pr modification of positions 72 and 42 completely abolished formation of the Ub-Gly76-MESNa thioester, whereas the same modifications at positions 54 and 74 had no effect since efficient thioester formation was observed. When using Arg-to-azidohomoalanine Ub mutants, the precursors used for click chemistry, all azido-containing mutants were accepted and processed by the E1 enzyme to form Ub-MESNa thioesters (bars labeled N 3 in Figure 5C). Notably, the Arg72-Aha mutant was significantly slower and Arg42-Aha was moderately slower than WT could be pulled down from cell lysates using biotinylated probes. For this purpose, we decided to test whether DupA-WT was able to bind and enrich DupA-WT and Ub ligase TRIM28 for 5, as well as the deubiquitinating enzyme OTUD4 for 5. A decreased interaction with the deubiquitinating enzyme USP5 is observed for both sites of modifications at positions 54 and 74, whereas controls with either mCherry (lane 3) or a non-specific interaction with the SA beads (lane 2) showed no or only minimal DupA recovery, respectively. Similarly, pull-down with biotin-Ub only showed marginal enrichment for DupA (Figure 6, lane 6) to a comparable extent to that in the beads-only control. We repeated this experiment with a slight excess of biotin-Ub and quantified these results using densitometry, showing >10-fold enrichment of mCherry-DupA recovery by biotinylated 5 compared with biotinylated wild-type Ub or beads (Figure 6). We then performed pull-downs from HEK293T cell lysate using nonhydrolyzable probes 5 and triazole$^{54}$Ub$^{\text{me-ADP}}$ and subjected the interacting proteins to trypsin digestion and MS/MS analysis to compare their interactome versus native Ub (Figure S7 in the Supporting Information).

Intriguingly, both sites of ADPribosylation lead to increased interaction with distinct proteins compared with that of unmodified Ub, such as the Ub ligase MYCBP for triazole$^{72}$Ub$^{\text{me-ADP}}$ and Ub ligase TRIM28 for 5, as well as the deubiquitinating enzyme OTUD4 for 5. A decreased interaction with the deubiquitinating enzyme USP5 is observed for both sites of modification in comparison with unmodified Ub. The change of interaction partners for Ub$^{\text{me-ADP}}$ contains, among others, deubiquitinating enzymes, Ub ligases, and proteins involved in intracellular (endosomal) trafficking or endoplasmic reticulum–Golgi
maintenance. These initial results need further validation and are a worthy subject of further research, to define the underlying cellular pathways wherein Ub ADPribosylation plays a role.

Conclusion

The preparation of ADP-ribose, adenosine methylenebisphosphonate ribose, and phosphoribose carrying an α-oriented alkyne on the anomeric position allowed us to conjugate a site-specific hydrolysis at positions 54, 72, and 74. These are not the final page numbers!

Conflict of interest

The authors declare no conflict of interest.

Keywords: ADPribosylation · click chemistry · Legionella · post-translational modifications · ubiquitin

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FULL PAPER

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Modular modifications: Analogues of ADPribosylated ubiquitin, including a nonhydrolyzable methylenebisphosphonate version, are constructed and subsequently applied to investigate (de-)ubiquitinating enzymes involved in Legionella infection. These probes are used in the study of the bacterial effector DupA and the human ubiquitin activating E1 enzyme.

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