Molecular mechanism of poly(ADP-ribosyl)ation by PARP1 and identification of lysine residues as ADP-ribose acceptor sites

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ABSTRACT

Poly(ADP-ribose) polymerase 1 (PARP1) synthesizes poly(ADP-ribose) (PAR) using nicotinamide adenine dinucleotide (NAD) as a substrate. Despite intensive research on the cellular functions of PARP1, the molecular mechanism of PAR formation has not been comprehensively understood. In this study, we elucidate the molecular mechanisms of poly(ADP-ribosyl)ation and identify PAR acceptor sites. Generation of different chimera proteins revealed that the amino-terminal domains of PARP1, PARP2 and PARP3 cooperate tightly with their corresponding catalytic domains. The DNA-dependent interaction between the amino-terminal DNA-binding domain and the catalytic domain of PARP1 increased $V_{\text{max}}$ and decreased the $K_m$ for NAD. Furthermore, we show that glutamic acid residues in the auto-modification domain of PARP1 are not required for PAR formation. Instead, we identify individual lysine residues as acceptor sites for ADP-ribosylation. Together, our findings provide novel mechanistic insights into PAR synthesis with significant relevance for the different biological functions of PARP family members.

INTRODUCTION

Poly(ADP-ribose) polymerases (PARPs) use nicotinamide adenine dinucleotide (NAD) as substrate to synthesize poly(ADP-ribose) (PAR) (1). On the cellular level, PAR formation has been implicated in a wide range of processes, such as maintenance of genomic stability, transcriptional regulation, energy metabolism and cell death (2).

PARP1 was the first protein described to catalyze PAR formation in response to mitogenic stimuli or genotoxic stress (3–7). It contains three functionally distinct domains: an amino-terminal DNA-binding domain (DBD), an auto-modification domain (AD) and a carboxyl-terminal PARP homology domain that includes the catalytic domain (CAT) responsible for PAR formation (8). The DBD extends from the initiator methionine to threonine 373 in human PARP1. It contains two structurally and functionally unique zinc fingers (FI: aa, amino acid, 11–89; FII: aa 115–199) (2,9). Recently, a third and so far unrecognized zinc-binding motif was discovered (FIII: aa 233–373) (10,11). The DBD also contains a bipartite nuclear localization signal (NLS) of the form KRK-X(11)-KKKSKK (aa 207–226) that targets PARP1 to the nucleus (12). The PARP1 zinc fingers FI and FII are thought to recognize altered structures in DNA rather than particular sequences and have also been reported to be involved in protein–protein interactions (13). PARP1 strongly associates with DNA single and double strand breaks generated either directly by DNA damage or indirectly by the enzymatic excision of damaged bases during DNA repair processes (2,9). Several studies indicate that the first zinc finger is required for PARP1 activation by both DNA single and double strand breaks, whereas the second zinc finger may exclusively act as a DNA single strand break sensor (2,9).

The AD of PARP1 is located in the central region of the enzyme, between residues 373 and 525 of human PARP1 (14,15). It was identified as the domain containing acceptor amino acids for the covalent attachment of PAR (16). In addition, several recent studies identified a weak leucine-zipper motif in the amino-terminal region of the AD, which suggests that this motif might be involved in homo- and/or hetero-dimerization (9). The AD of PARP1 also comprises a breast cancer 1 protein (BRCA1) C-terminus (BRCT) domain (from aa 386 to 464 in...
human PARP1) as well as an unstructured loop that connects the AD with the PARP homology domain.

PARP1 contains an 80–90 amino acid long tryptophan-, glycine-, arginine-rich (WGR) domain carboxyl terminal of the AD. The WGR domain is named after the most conserved central motif of tryptophane (W), glycine (G), arginine (R) residues and may represent a nucleic-acid-binding domain (2). This region of PARP1 has not been extensively characterized and its function is still unknown. The CAT has been suggested to catalyze at least three different enzymatic reactions: the attachment of the first ADP-ribose moiety onto an acceptor amino acid (initiation reaction), the addition of further ADP-ribose units onto already existing ones (elongation reaction) and the generation of branching points (branching reaction) (8). The active site is formed by a phylogenetically well-conserved sequence of ~50 residues (aa 859–908 of hPARP1). This ‘PARP signature’ contains the NAD acceptor sites and critical residues involved in the initiation, elongation and branching of PAR.

Like PARP1, both PARP2 and PARP3 also contain a WGR as well as a CAT (16). PARP2 and PARP3 lack, however, most motifs present in the amino-terminal half of PARP1. Neither zinc-binding motifs nor leucine-zippers or BRCT domains have been described for PARP2 or PARP3. PARP2 contains an amino-terminal SAP/SAF motif/module [named after scaffold-associated protein/scaffold-associated factor SAF-A/B, Acinus and PIAS: (17)] and a eukaryotic module proposed to be involved in sequence- or structure-specific DNA and RNA binding (18). Furthermore, PARP2 contains an amino-terminal NLS which targets the protein to the nucleus. PARP3 is the least studied and smallest PARP identified so far (19). The protein domain structure of PARP3 is very similar to that of PARP2, featuring a small putative DBD consisting of only 54 residues and apparently containing a targeting motif that is sufficient to localize the enzyme to the centrosome (19,20).

Attempts to obtain structural information on the full-length proteins PARP1, PARP2 and PARP3 by X-ray crystallography or by nuclear magnetic resonance (NMR) have not been successful up to now. The 3D structures of single domains, however, have been solved and allow for a structure-based comparison of different PARP family members (8,21) (PDB: 1A26, 1GS0 and 2PA9). Although the amino acid identity between PARP1 and PARP2 or PARP3 is only moderate (40% and 32% in the CAT, respectively), the overall structure of the CATs of these three proteins is nearly identical. This conservation suggests, in general, similar capabilities to generate PAR. Both PARP1 and PARP2 have been shown to synthesize very complex branched polymers at least in vitro (2). The enzymatic activity of PARP3 and its isoforms has not yet been investigated in detail.

An unresolved issue regarding the mechanism of poly(ADP-ribose)ylation is how DNA binding in the amino-terminal DBD triggers enzyme activation in the carboxyl-terminal CAT and how the different domains of the different PARPs are coordinated during this process. Furthermore, earlier studies suggested that the auto-modification activity targets between 4 and 28 acceptor residues located in the AD and in the DBD of PARP1 (14,22,23). For histone H1, a major target for transpoly(ADP-ribosylation) by PARP1, glutamic acid residues have been described to function as acceptors for PAR (24). This, together with the reported chemical similarity between the ADP-ribose-PARP1 linkage and carboxyl esters in mono-ADP-ribosylated histones (23), led to the hypothesis that multiple glutamic acid residues present in the AD of PARP1 might function as acceptor sites for auto-poly(ADP-ribosylation) (16). However, despite intensive research during the last 40 years, the acceptor amino acids in PARP1 have not been confirmed by mutational studies.

Here, we comprehensively analyze PAR formation by PARP1, PARP2 and PARP3 and find a close cooperativity between the amino-terminal portions of the proteins and their corresponding CATs. We define the DBD (aa 1–373) and the WGR/CAT domain (aa 533–1014) as the minimal domains of PARP1 required for PAR formation. The DNA-dependent interaction between the DBD and the CAT increased $V_{\text{max}}$ and decreased the $K_m$ for NAD. Furthermore, by amino-acid substitutions, we establish that glutamic acid residues within the AD are not required for PAR formation and thus do not function as acceptor amino acids for PAR. Instead, we identify lysine residues within the AD of PARP1 as acceptor sites for ADP-riboseylation.

### MATERIALS AND METHODS

#### Chemicals and antibodies

$^3$H-NAD and protein A sepharose were purchased from GE Healthcare and $^{32}$P-NAD was from PerkinElmer. NAD was obtained from Sigma-Aldrich. Anti-PAR antibody LP96-10 was from Alexis Biochemicals or Becton Dickinson, anti-PARP1cat antibody H250 from Santa Cruz Biotechnology and anti-haemagglutinin (HA) antibody 16B12 from Covance.

#### Plasmids

The baculovirus expression vectors pQE-TriSystem (Qiagen) and BacPak8 (Clontech) were used for the expression of recombinant proteins in $S/21$ insect cells as described previously (25,26).

#### Cloning, expression and purification of recombinant proteins

Wild-type hPARP1 (NCBI ID: BC037545), hPARP2 (NCBI ID: NM_001042618) and hPARP3 (NCBI ID: BC014260) were cloned and expressed as carboxyl-terminal His-tagged proteins. PARP family chimera were generated by overlapping polymerase chain reaction (PCR) at the position corresponding to amino acid 533 in hPARP1 and expressed as carboxyl-terminal His-tagged proteins. Protein fragments and deletion mutants were generated by PCR and expressed as carboxyl-terminal His-tagged proteins as described before (25,26). Amino-acid substitutions were introduced by site-directed PCR-based mutagenesis and mutant proteins.
were expressed as described before (25,26). All recombinant proteins were purified by one-step affinity chromatography using ProBond resin according to the manufacturer’s recommendations (Invitrogen). Expression and purification of all recombinant proteins was analyzed by SDS–PAGE followed by coomassie staining. For the stacking gel a 4.5% acrylamide-bis solution [37.5:1, 40% (w/v), Serva] and for the separating gel a 10–12.5% acrylamide-bis solution was used.

PAR formation assays

$^3$H-NAD time course experiments. One hundred picomoles recombinant purified enzyme and 5 μg of protein fragments in PAR reaction buffer (50 mM Tris–HCl pH 8.0, 4 mM MgCl$_2$, 250 μM DTT, 1 μg/ml pepstatin, 1 μg/ml bestatin, 1 μg/ml leupeptin) in the presence of 50 pmol annealed double-stranded oligomer (5’-GGGAATTCC-3’) were supplemented with $^3$H-NAD to a final concentration of 400 μM. PAR formation was allowed for 1, 3, 5, 15 and 60 min at 30°C. Reactions were stopped by addition of ice-cold 10% TCA/2% Na$_2$P$_2$O$_7$. Polymers were precipitated for 10 min on ice and then applied onto filter papers. Counts per minute were obtained by liquid scintillation counting. For the determination of $V_{\text{max}}$ and $K_m$, initial reaction velocities ($V_0$) were obtained by measuring PAR levels generated after 0, 1, 3 and 5 min incubation at different $^3$H-NAD concentrations and using the GraphPad Prism software for nonlinear regression analysis assuming a one-site binding model. $V_{\text{max}}$ and $K_m$ were calculated from $V_0$ according to Michaelis–Menten.

Anti-PAR western blot. Unless otherwise stated, 10 pmol recombinant purified enzyme and 0.5 μg of protein fragments in PAR reaction buffer in the presence of 5 pmol annealed double-stranded oligomer (5’-GGGAATTCC-3’) were supplemented with NAD to a final concentration of 400 μM. PAR formation was allowed for 5 min at 30°C. Reactions were stopped by addition of SDS–PAGE loading buffer and boiling for 5 min at 95°C. Samples were subjected to SDS–PAGE followed by anti-PAR western blot.

$^{32}$P-NAD auto-modification. Unless otherwise stated, 10 pmol recombinant purified enzyme and 0.5 μg of protein fragments in PAR reaction buffer in the presence of 5 pmol annealed double-stranded oligomer (5’-GGGAATTCC-3’) were supplemented with $^{32}$P-NAD to a final concentration of 100 nM. Auto-modification was allowed for 10 s at 30°C. Reactions were stopped by addition of SDS–PAGE loading buffer and boiling for 5 min at 95°C. Samples were subjected to SDS–PAGE followed by detection of auto-modification by autoradiography.

PAR detection by silver staining. Following synthesis of PAR as for western blot analysis, PAR chains were purified and separated by modified DNA sequencing gel electrophoresis as described by Fahren et al. (27).

In vitro co-immunoprecipitation. Ten picomoles recombinant purified enzyme and 0.5 μg of protein fragments were incubated for 5 min at 30°C in Co-IP buffer (50 mM Tris–HCl pH 8.0, 100 mM NaCl, 4 mM MgCl$_2$, 0.2% NP-40, 250 μM DTT, 1 μg/ml pepstatin, 1 μg/ml bestatin, 1 μg/ml leupeptin) in the absence or presence of 5 pmol annealed double-stranded oligomer (5’-GGGAATTCC-3’). The CAT of PARP1 was allowed to bind to the anti-PARP1 antibody for 1 h at 4°C. Protein A sepharose was added and samples were incubated for another 2h at 4°C. Samples were washed three times for 5 min in Co-IP buffer containing 300 mM NaCl before being subjected to SDS–PAGE followed by western blot.

RESULTS

Purified full-length human PARP1 and PARP2 are enzymatically active

In order to gain detailed insights into the mechanism of PAR formation by different PARP family members, we expressed and purified full-length human PARP1, PARP2 and PARP3 using the baculovirus expression system (Figure 1A and B). PARP3 showed a slower migration velocity than predicted in SDS–PAGE, possibly due to the high content of hydrophobic amino acids in the CAT. To measure PAR formation, the purified proteins were incubated for different time periods with 400 μM tritium-labeled NAD in the presence of double strand break mimicking DNA. Reaction products were precipitated by trichloroacetic acid (TCA) before they were analyzed using a beta counter. PARP1 generated PAR in a time-dependent manner (Figure 1C, left panel). PARP2 also synthesized PAR in a time-dependent manner, however, not as efficiently as PARP1 (Figure 1C, middle panel). The reduced amount of product formed by PARP2 most probably represents a quantitative rather than a qualitative difference, since the length distribution of PAR chains formed by PARP2 was comparable to the length distribution of PAR formed by PARP1 (Supplementary Figure 4B). Human PARP3 did not produce detectable amounts of PAR under the tested conditions (Figure 1C, right panel).

Assuming that mono(ADP-ribosyl)ation precedes PAR formation, we assessed the auto-modification of the three proteins after 10s incubation with 100 nM radiolabeled NAD (Figure 1D). The short incubation period and the low concentration of NAD were chosen to prevent polymer formation. The discrete bands observed using this approach indeed suggest that under these conditions mostly mono(ADP-ribosyl)ation occurred. In line with the time-course experiments, PARP1 and PARP2 were able to auto-modify themselves in an NAD- and DNA-dependent manner while PARP3 was not (Figure 1D). PARP1 synthesized increasing PAR levels in a time- and DNA-dependent manner detected also by western blot and vacuum slot blot using 400 μM NAD (Supplementary Figure 1A and B). PAR formation after 5 min incubation with 400 μM NAD caused a pronounced shift of the coomassie blue-stained proteins in the denaturing gel due to a severely reduced migration velocity of the poly(ADP-ribosyl)ated proteins when...
compared to unmodified proteins (Supplementary Figure 1C). The observed basal activity of PARP1 in the absence of DNA can be explained either by a contamination with DNA or by the intrinsic DNA-independent activity of the CAT as described by Simonin et al. (28). Analysis of PAR formation by silver staining after modified DNA sequencing gel electrophoresis confirmed PAR formation by PARP1 and PARP2 and no PAR formation by PARP3 (Supplementary Figure 4B).
The carboxyl-terminal domains of PARP1, PARP2 and PARP3 cannot compensate for each other

Next, we investigated the crosstalk between the different amino-terminal domains of PARP1, PARP2 and PARP3 with their carboxyl-terminal domains (i.e. WGR/CAT). Therefore, we generated chimera proteins by replacing the WGR/CAT domain of PARP1 with the WGR/CAT domain of PARP2 or PARP3 [named chimera PARP1-2 (aa 1–532 of PARP1 fused to aa 81–570 of PARP2) or chimera PARP1-3 (aa 1–532 of PARP1 fused to aa 48–533 of PARP3)], respectively (Figure 2A and B). We analyzed PAR formation by these proteins and found that replacing the WGR/CAT domain of PARP1 by the one of PARP2 (i.e. chimera PARP1-2) resulted in an active enzyme that showed roughly similar PAR formation in time course experiments as PARP2 (Figure 2C, middle panel). Replacement of the WGR/CAT domain of PARP1 by that of PARP3 (i.e. chimera PARP1-3) resulted in an enzyme that did not produce detectable amounts of PAR under the tested conditions (Figure 2C, right panel and Supplementary Figure 1D). In line with these findings, chimera PARP1-2 was able to auto-modify itself whereas chimera PARP1-3 was not (Figure 2D). Together these results suggest that the WGR/CAT domains of the investigated PARP proteins cannot compensate for each other. The WGR/CAT domains cooperate tightly with their corresponding amino-terminal domains and limit poly(ADP-ribosyl)ation capacity and the ability for auto-modification, despite high levels of structural similarity between the CATs (see Supplementary Figure 2A).

The carboxyl-terminal domain of PARP1 is not activated by the amino-terminal domains of PARP2 or PARP3

In a second set of chimera proteins we fused the WGR/CAT domain of PARP1 to the amino-terminal domains of PARP2 or PARP3, or deleted the amino-terminal domain
of PARP1 [named chimera PARP2-1 (aa 1–80 of PARP2 fused to aa 533–1014 of PARP1), chimera PARP3-1 (aa 1–47 of PARP3 fused to aa 533–1014 of PARP1) or PARPs-1 (aa 533–1014 of PARP1)], respectively (Figure 3A and B). Analysis of these proteins revealed that chimera PARP2-1, chimera PARP3-1 and PARPs-1 did not generate detectable levels of PAR (Figure 3C and Supplementary Figure 1E). Furthermore, no auto-modification of the three proteins was observed under the tested conditions (Figure 3D). These results indicate that the WGR/CAT domain of PARP1 is only stimulated by its corresponding amino-terminal domain, but not by the amino-terminal domains of PARP2 or PARP3.

The DBD of PARP1 is sufficient to stimulate its WGR/CAT domain

To further investigate the cooperativity between the amino-terminal domain of PARP1 and its WGR/CAT
domain, the inactive chimera PARP2-1 was co-incubated with the PARP1 E988K mutant, which lacks the ability to generate PAR. Surprisingly, co-incubation of chimera PARP2-1 with PARP1 E988K strongly induced PAR synthesis, suggesting that PARP1 E988K was able to stimulate the WGR/CAT domain of chimera PARP2-1 (Figure 4A). To map the minimal domain of PARP1, which was able to stimulate the WGR/CAT domain of chimera PARP2-1, we expressed and purified different fragments of PARP1 covering all domains from A to F (see Figure 1A). Analysis of PAR synthesis by western blot upon co-incubation of chimera PARP2-1 with these fragments revealed that the DBD of PARP1 comprising amino acid 1–373 was the only fragment able to stimulate chimera PARP2-1 (Figure 4B). Further dissection of the DBD revealed that only the complete and undisrupted DBD from amino acid 1 to 373 containing FI, FII and FIII was able to stimulate chimera PARP2-1 (Figure 4C). The stimulation of chimera PARP2-1 by the DBD was salt resistant up to 300 mM NaCl (Supplementary Figure 3B, left panel).

To further assess the specificity of the observed stimulation, the DBD was incubated with different proteins (chimera PARP2-1, chimera PARP3-1, PARP-1 and PARP1 656–1014) and the time course of PAR formation was analyzed using tritium-labeled NAD. Of note, besides chimera PARP2-1, only chimera PARP3-1 and PARP-1, but neither the CAT of PARP1 nor full-length PARP2, PARP3, chimera PARP1-2 or chimera PARP1-3, were stimulated by the DBD of PARP1 (Figure 4D and Supplementary Figure 4A and C). Analysis of PAR formation by silver staining after polymer separation using modified DNA sequencing gel electrophoresis confirmed that the observed stimulation in the time course experiments correlated with the synthesis of PAR containing 1 to more than at least 50 ADP-ribose units (Supplementary Figure 4B).

The stimulation of chimera PARP2-1 by the DBD was strongly dependent on DNA (Figure 4E, left panel), which suggests that DNA tightly regulates the interaction necessary for the activation of the CAT. Furthermore, our observation that PARPs-1 but not PARP1 656–1014 together with the DBD was able to generate PAR indicates that the WGR domain of PARP1 is absolutely essential for enzymatic activity. Since chimera PARP2-1 does not exist physiologically, PAR synthesis by a PARP1 DBD deletion mutant (aa 373–1014) co-incubated with the DBD was analyzed. Interestingly, the DBD was able to stimulate PARP1 373–1014 in a DNA-dependent manner and comparable to PARP2-1 (Figure 4E, right panel and Supplementary Figure 3B, right panel), suggesting that the observed stimulation of PARP2-1 by the DBD represents a physiological regulatory mechanism in the PARP1 full-length context.

The DBD of PARP1 interacts with its CAT domain

The results described above suggest that the DBD of PARP1 interacts with the CAT and/or the WGR domain to stimulate PAR synthesis by the CAT. To test this hypothesis experimentally, in vitro co-immunoprecipitation assays were performed with purified proteins and fragments. The complete DBD (aa 1–373), but not aa 1–214 alone, specifically bound to chimera PARP2-1 in a manner that was stabilized by DNA (Figure 4F). Similarly, the DBD also bound to PARP1 373–1014, and this interaction was enhanced by DNA (Figure 4G, left panel). Interestingly, the CAT domain of PARP1 without the WGR (aa 656–1014) was sufficient for the DNA-dependent interaction with the DBD (Figure 4G, right panel). Since PAR formation was only observed when combining the DBD with PARPs-1 (expressing WGR/CAT) but not with the CAT domain of PARP1 alone (Figure 4D and Supplementary Figure 4A), we conclude that an intact DBD (aa 1–373) interacts with the CAT in a DNA-dependent manner and that the WGR domain is additionally required to allow PAR formation.

The DBD bound to DNA activates the CAT by increasing $V_{\text{max}}$ and decreasing $K_m$

Next, we determined the enzymatic parameters of chimera PARP2-1 stimulated by the PARP1 DBD in the absence or presence of DNA. We measured the incorporation of tritium-labeled NAD into TCA-precipitable polymers at early reaction time points and obtained initial reaction velocities ($V_0$) for different substrate concentrations by nonlinear regression analysis assuming one substrate-binding site. In the absence of the DBD, chimera PARP2-1 did not generate detectable levels of PAR independent of the addition of DNA (Table 1, second and third column), thus confirming our previous results. In the presence of the DBD, PAR generation was strongly dependent on DNA. Without DNA, the obtained PAR levels were low, but still allowed for curve fitting and calculation of $V_{\text{max}}$ and $K_m$ values (Table 1, fourth column). Addition of DNA increased the maximum reaction velocity $V_{\text{max}}$ about 4-fold and reduced $K_m$ 8-fold (Table 1, compare fifth to fourth column). The reaction efficiency $K_{\text{cat}}/K_m$ was thereby increased by more than 30-fold. DNA could thus be considered a $V^++K$-type activator, affecting both turnover rate and substrate affinity. Remarkably, the enzymatic parameters obtained for chimera PARP2-1 together with the PARP1 DBD closely match the values reported for full-length PARP1 (Table 1, compare fifth to first column). Together, these results provide evidence that DNA containing double strand breaks is recognized and bound by the DBD of PARP1, which subsequently binds to the CAT domain to induce structural changes within the catalytic cleft in order to increase the affinity for NAD and stabilize reaction intermediates.

PARP1 forms a catalytic dimer which requires at least one functional FI and FIII domain for activity

The CAT of PARP1 was previously described to dimerize (29). To investigate whether our purified proteins were also able to form dimers, the enzymatic activity of full-length PARP1 was assessed by western blot analysis after co-incubation with different molar ratios of two catalytically inactive PARP1 mutants (E988K or M890V/D899N,
Figure 4. The DBD of PARP1 interacts with and is sufficient to stimulate its WGR/CAT domain. (A) PAR formation by chimera PARP2-1 co-incubated with catalytically inactive PARP1 E988K. PAR was detected by western blot using anti-PAR antibody LP96-10. Substrate concentration: 400 μM NAD. (B) PAR formation by chimera PARP2-1 co-incubated with the indicated fragments of PARP1 or with PARP1 E988K. (C) PAR formation of chimera PARP2-1 co-incubated with the indicated fragments or combination of fragments of PARP1. (D) Time course of PAR formation by chimera PARP2-1, chimera PARP3-1, PARPs-1 and PARP1 656–1014 in the absence or presence of fragment 1–373 as in Figure 1C. Black without fragment 1–373 and grey with fragment 1–373. (E) PAR formation of chimera PARP2-1 or PARP1 373–1014 co-incubated with fragment 1–373 in the absence or presence of DNA. (F) *In vitro* interaction between chimera PARP2-1 and 1–373. Chimera PARP2-1 was bound to protein A sepharose using an antibody against the CAT of PARP1 (a-PARP1 cat) and was then incubated with HA-tagged fragment 1–373 or 1–214 in the absence or presence of DNA. HA-tagged fragments were detected by western blot. PARP1 cat antibody coupled to beads without chimera PARP2-1 served as control (ctr.). (G) *In vitro* interaction between PARP1 373–1014 or 656–1014 with 1–373. Experiments were performed as described in (F). Molecular size markers in kilo Daltons and the border between stacking and separating gel (asterisk) are indicated.
VNC: not calculable (product levels below detection limit).

K/C1 of the regions containing zinc finger FI (inactive due to lost activation of the CAT. Deletion one of the DBD sub-domains would render the protein indeed able to form dimers and to regulate each other. (Figure 4). This finding suggests that disruption of the DBD of PARP1 by deleting the WGR domain is required for the catalytic activity of the enzyme and function as acceptors for PAR. First, we deleted the BRCT domain as part of the AD (Figure 6A). A PARP1 ΔBRCT mutant was as active as its wild-type counterpart with regard to auto-modification (Figure 6B, first four lanes) and PAR formation (Supplementary Figure 5A, left panel). Next, in the context of the PARP1 ΔBRCT mutant, we additionally mutated all eight glutamic acid residues in the remaining auto-modification loop between amino acids 484 and 557 to glutamine (ΔBRCT/E) (Figure 6A). Surprisingly, these substitutions also did not reduce auto-modification (Figure 6B, last two lanes) or PAR formation (Supplementary Figure 5A, right panel). These results strongly indicate that glutamic acid residues within the AD of PARP1 are not required for enzymatic activity and are unlikely to serve as acceptors for PAR.

**Lysine residues are acceptor sites in PARP1**

In contrast to the deletion of the BRCT domain, deletion of the remaining amino acids in the AD of PARP1 (ΔAc, aa 466–525) (Figure 6A), a region previously reported to be acetylated (26), resulted in severely impaired auto-modification (Figure 6C) and reduced PAR formation (Supplementary Figure 5B), suggesting that acceptor sites are localized in this region of PARP1. As PAR levels generated by PARP1 ΔAc were decreased but did not drop completely, additional PAR acceptor sites are likely to exist in other domains of PARP1. Trans-poly(ADP-ribosylation) of different fragments of PARP1 by wild-type PARP1 indeed confirmed that not only the AD but also a fragment containing amino acid 1–214 is modified (Supplementary Figure 5C).

ADP-ribose has earlier been described to be a potent histone glycation and glycoxidation agent *in vitro*, leading to the formation of ketoamine glycation conjugates (30).

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**Table 1. Kinetic parameters of chimera PARP2-1**

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<th>hPARP1&lt;sup&gt;a&lt;/sup&gt;</th>
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NC: not calculable (product levels below detection limit).

<sup>a</sup>Values as reported in the literature.
Figure 5. PARP1 forms a catalytic dimer which requires at least one functional FI and FIII domain for activity. (A) Domain organization of the PARP1 deletion mutants used for this figure. (B) PAR formation by PARP1 when co-incubated with the indicated inactive proteins or fragments at a molar ratio of 1:1 or 1:5. According to the manufacturer, the anti-PAR antibody LP96-10 cross reacts with bovine serum albumin (BSA) (band at
When we analyzed the chemical linkage stability of automodified PARP1, we found that it was stable up to pH 10 (but not at pH 13.5) and that incubation with 1 M hydroxylamine at pH 7 for 30 min at 30°C did not release the modification (Supplementary Figure 5D). This observation suggests that the protein-ADP-ribose linkage indeed might occur on lysines. To exclude that the investigated auto-modification of PARP1 was due to traces of ADP-ribose within the provided NAD, the inactive PARP1 mutant M890V/D899N was incubated with radioactive NAD. Only upon long exposure a faint labeling of PARP1 M890V/D899N was observed while PARP1 E988K was able, as described earlier, to modify itself (Supplementary Figure 5E), confirming that the observed auto-modification of PARP1 was due to its enzymatic activity.

Next, we analyzed the trans-poly(ADP-ribosyl)ation of a fragment comprising the AD (aa 373–525) of PARP1 by full-length PARP1. Although auto-modification of PARP1 was much more efficient than trans-poly(ADP-ribosyl)ation, specific labeling of the AD fragment was observed (Figure 6D, second lane). In order to identify individual lysine residues within the AD which serve as acceptor sites for PAR, we analyzed trans-poly(ADP-ribosyl)ation of the AD fragment containing three lysine to arginine substitutions (K498, K521 and K524, called KTR). These sites were previously reported to be targets for acetylation (26). Modification of the 373–525 KTR fragment by full-length PARP1 was reduced as compared to 373–525 wt (Figure 6D, third lane). Since trans-poly(ADP-ribosyl)ation of a protein fragment might lead, due to structural constrains, to unspecific modification of amino acids and might thus not be comparable to modification of the full-length protein, we generated a full-length PARP1 mutant which contains the three lysine to arginine substitutions at position 498, 521 and 524.

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Importantly, this mutant showed strongly reduced auto-
modification, very much comparable to the levels
observed for the PARP1 ΔAc mutant (Figure 6E).
Overall, these experiments provide evidence that not glut-
amic acid residues but instead at least three lysine res-
dues within the auto-modification loop (aa 466–525)
and additional residues within the first 214 amino acids
of PARP1 are target sites for enzymatic auto-ADP-
ribosylation.

DISCUSSION

In this study we analyzed the poly(ADP-ribosyl)ation
capacity of PARP1 and the closely related proteins
PARP2 and PARP3 under standardized reaction condi-
tions and investigated the molecular mechanism of PAR
formation. Human PARP1 and PARP2 were able to
auto-modify themselves and generate PAR, although to
different levels. Neither polymer formation nor auto-
modification was observed for PARP3 under the tested
conditions.

PARP1 deletion mutants and fusion proteins had been
successfully employed before to study different aspects of
poly(ADP-ribosyl)ation (31,32). Here, we have generated
PARP family chimeras to analyze the molecular mech-
anism of PAR formation. The PARP chimera revealed
that the WGR/CAT domains of PARP1, PARP2 and
PARP3 tightly cooperate with their corresponding
amo-no-terminal domains. Closer examination of PARP1
revealed that FI, FIII and the WGR/CAT domain of
PARP1 are required and sufficient for PAR formation.
FII and the BRCT domain, however, were not essential
for the enzymatic activity. The DBD interacted directly
with the CAT domain of PARP1. DBD bound to DNA
increased \( V_{\text{max}} \) and reduced the \( K_m \) of the CAT for NAD.
We also provide evidence that PARP1 forms a catalytic
dimer in which lack of either FI or FIII could be func-
tionally complemented by a protein containing these
domains. Finally, we identified three lysine residues
within the AD and additionally the first 214 amino acids
of the DBD as target sites for enzymatic covalent auto-
poly(ADP-ribosyl)ation by PARP1.

We employed three different methods to assess
poly(ADP-ribosyl)ation. First, \(^3\)H-NAD at a concentra-
tion of 400 \( \mu \)M was used to measure TCA-precipitable
polymer formation in time course experiments. Second,
\(^32\)P-NAD at a concentration of only 100 nM was used to
measure auto-modification after short incubation periods
(10 s). This approach resulted in distinct bands corre-
sponding to the modified protein and most likely repre-
senting mono(ADP-ribosyl)ation or short oligomers of
ADP-ribose attached to the labeled protein. Third, unlabeled
NAD at a concentration of 400 \( \mu \)M was used to measure
PAR formation detected by western blot. The anti-PAR
antibody typically detected high molecular weight polymers, most of which remained as a smear at
the top of the separating gel or even in the stacking gel.
This approach was not very suitable to make quantitative
statements but could be readily applied to analyze whether
a protein was active or not.

Human PARP3 was previously described by Augustin
et al. to be an active enzyme, as detected by autoradiogra-
phy after 15 min incubation with 10 \( \mu \)M \(^32\)P-NAD (19).
Augustin and co-workers did not, however, compare the
activity of PARP3 to that of PARP1 or any other PARP
family member under these conditions. We analyzed
PARP3 in comparison to PARP1 and PARP2 under Stan-
dardized reaction conditions and could not observe
any activity for this protein. However, when we applied
the conditions provided by Augustin et al. to measure
PARP3 activity by autoradiography, we could also
observe PARP3 auto-modification (data not shown), sug-
gesting that the protein possesses some degree of activity
under certain well-defined conditions. Further investiga-
tions are needed to analyze the extent of PAR formation
by PARP3 as well as its physiological relevance.

The DBD of PARP1 interacted in a coordinated
and DNA-dependent manner with the CAT domain of
PARP1, but not with that of PARP2 or PARP3. Thus,
deep the high level of structural similarity between the
CATs of PARP1, PARP2 and PARP3, these domains
cannot compensate for each other and may possess unan-
ticipated intrinsic regulatory functions. Since the PARP1
DBD is not or only partially present in other PARP family
members, the newly identified intra-molecular interaction
might provide a promising surface for the development of
PARP1 specific inhibitors.

In our study, several enzymatic dead mutants with
deletions in the DBD could be functionally complemented
by another inactive PARP1 mutant containing the missing
domain, thus implicating that PARP1 is forming a dimer
for poly(ADP-ribosyl)ation. The existence of catalytically
active protein dimers in which each monomer is lacking
a domain required for enzymatic activity was surprising
and suggests that PARP1 is a highly flexible molecule
with rather loose domain architecture.

Consistent with earlier reports (33,34) our results
showed that zinc finger FI is absolutely required for the
DNA-dependent activation of the protein, whereas zinc
finger FII is dispensable. Zinc finger FII may, however,
determine the binding specificity for DNA single strand
breaks as suggested previously by Gradwohl et al. (35).
Our data revealed that the recently discovered zinc-
binding motif FIII is essential for the interaction of the
DBD with the CAT and thus also for the activation
of the enzyme. Furthermore, the so far uncharacterized
WGR domain is an indispensable prerequisite for PAR
formation, although this domain is not necessary for the
interaction between the DBD and the CAT.

The interaction between the DBD bound to DNA and
the CAT domain increased the maximum reaction velocity
\( V_{\text{max}} \) by a factor of four and reduced the \( K_m \) for NAD
roughly from 1 mM to 140 \( \mu \)M (Table 1). The reaction
efficiency \( \text{Kcat/} K_m \) was thereby increased by a factor of
more than 30. The total cellular NAD concentration was
previously estimated to be around 350 \( \mu \)M (36). Zhang
et al. argued that NAD cofactors should readily pass
through nuclear pores, which would suggest that cytoplas-
ic NAD levels reflect nuclear NAD concentrations (37).
The same group estimated the free nuclear NAD concen-
tration to be around 70 \( \mu \)M (38). Despite this uncertainty
in the estimation of free nuclear NAD concentrations, we believe that increasing the affinity of PARP1 for NAD by binding to DNA double strand breaks might be an important regulatory step to allow PAR formation at physiological NAD concentrations. Release of PARP1 from DNA would consequently reduce the affinity of PARP1 for NAD and terminate PAR formation. Importantly, the nuclear concentration of NAD can be modulated by NMN adenylyl transferase 1 (NMNAT-1), which catalyzes the final step of NAD biosynthesis. A recent study revealed that NMNAT-1 is able to interact with and stimulate PARP1. It is thus tempting to speculate that PARP1 activation by its binding to DNA strand breaks is supported by the localized action of NMNAT-1.

Our results suggest that activation of PARP1 occurs in defined sequential steps (Figure 7A). First, the DBD binds to certain damages within the DNA. This enhances the interaction between the DBD and the WGR/CAT domain. As a consequence, minor structural rearrangements within the catalytic cleft occur, resulting in an increased affinity for NAD. Increasing substrate affinity and additionally substrate turnover rates then allows for high reaction efficiency and very rapid auto-modification at distinct lysine residues followed by PAR chain elongation. An analogous model can be envisioned for the protein chimera PARP2-1, which is activated by the PARP1 DBD in the presence of double strand breaks mimicking DNA (Supplementary Figure 6A).
The assumption that PARP1 is modified at glutamic acid residues was based mainly on the chemical stability of the ADP-ribose-PARP1 linkage, which was very heterogeneous but in part of a similar type as carboxyl esters in mono-ADP-ribosylated histones isolated from cells (23). The presented mutation analysis studies revealed that neither deletion of all glutamic acid residues in the BRCT domain (aa 385–476, containing nine glutamic acid residues) nor additional mutation of the remaining glutamic acid residues to glutamine in the AD (aa 477–557, containing eight glutamic acid residues) affected auto-modification or PAR formation and thus provide strong evidence that these amino acids in the AD are not the acceptor sites for poly(ADP-ribosyl)ation. Interestingly, mutation of the three lysines K498, K521 and K524 in the AD of PARP1 to arginines strongly reduced the auto-modification of the enzyme, suggesting that these residues in fact are acceptors for PAR. A longer exposure of the gel revealed a weak labeling of PARP1 KTR (data not shown) and it may well be that additional lysine residues serve as acceptor sites in this domain. Furthermore, acceptor sites can also be expected in the DBD of PARP1. Whether these sites are also lysine residues or whether outside the AD other amino acids serve as PAR acceptors is currently not known.

Modification of proteins by ADP-ribose can be characterized according to their chemical properties. ADP-ribosylated lysine residues were described to be stable in the presence of 1 M hydroxylamine at pH 7, while chemically modified glutamic and aspartic acid residues would rapidly release the ADP-ribose moiety (40,41). Our chemical analysis of modified PARP1 revealed that the observed linkage most likely corresponds to the glycation linkage described above. Thus we propose that ADP-ribosylation of PARP1 is catalyzed by its NADase activity, which subsequently allows modification of lysine residues positioned close to the catalytic active site to Lys-ADP-ribose ketamine (Figure 7B, C and D). This moiety could then serve as acceptor for the elongation reaction, which is catalyzed by glutamic acid residue E988 in human PARP1. We are currently investigating whether other ADP-ribose acceptor proteins are modified by PARP1 in the same manner.

To date two enzymes, poly(ADP-ribose) glycohydrolase (PARG) and ADP-ribosyl protein Lyase, have been described to be involved in PAR catabolism (42,43). While PARG possesses both exon- and endoglycosidic activities, the Lyase was described to cleave the bond between proteins and mono(ADP-ribose). ADP-ribosylation of lysines creates a chemical bond, which is not a substrate for PARG, which cleaves the ribose-ribose bonds. Breaking a lysine-ADP-ribose linkage would instead require the activity of a Lyase (Figure 7D). Alternatively, the last ADP-ribose moieties might remain on PARP1 to serve as elongation sites for the next round of PAR formation or to mark the chromatin to memorize the location of previous DNA damage repair.

Lysine residues K498, K521 and K524 were previously identified as targets for acetylation by p300 and P300/CBP-associated factor (PCAF) in a stimulus-dependent manner (26). Remarkably, simple addition of PCAF reduced poly(ADP-ribosyl)ation by PARP1 (unpublished observation), suggesting that the interaction domain of PARP1 with PCAF is overlapping with the AD-ribose acceptor sites. Furthermore, we recently showed that acetylation of PARP-2 strongly reduced the enzymatic activity (44). Already more than 20 years ago, a possible interrelation between poly(ADP-ribosyl)ation reactions and post-translational protein acetylation had been discussed (45,46). Our finding that acetylation of lysine residues interferes with ADP-ribosylation supports this idea and points at an interesting crosstalk between acetylation of and AD-ribosylation by PARP family members. This crosstalk hypothesis is further strengthened by the finding that the enzymatic activity of PARP1 is not required for the function as transcriptional co-activator of NF-κB, a role which requires acetylation of PARP1 (47).

During apoptosis, PARP-1 is cleaved by different caspases to generate 89-kDa and 24-kDa fragments, a well-characterized hallmark of apoptosis. The data shown provide a functional explanation for the observed inactivation of PARP1 upon caspase cleavage, as this cleavage is separating FI and FII (aa 1–214) from FIII (aa 214–373), thus no longer allowing the DBD to interact as an intact polypeptide with the CAT domain for subsequent activation.

Our chemical and mutational analyses provide evidence that lysine residues are acceptor sites for auto-modification by PARP1 in vitro (Figure 6B). As PARP1 is the main acceptor protein for poly(ADP-ribosyl)ation in vivo (48,49), our findings are most likely also relevant in vivo. Confirming acceptor sites in vivo, however, is very difficult for different reasons. First, mutations within the DBD to eliminate the acceptor sites in this region and to allow only the analysis of the three lysine residues in the auto-modification domain would affect the activation of PARP1 by DNA. Second, PARP1 is known to be modified by PARP2 and possibly by other PARP family members. Whether these proteins are modifying PARP1 also at the auto-modification sites or at other residues is currently not known. In any case, however, this crosstalk would interfere with in vivo analysis of PARP1 auto-modification.

In conclusion, we propose that PARP1 forms a catalytic dimer that allows the interaction of the DNA-binding domain with the CAT to modify distinct lysine residues as ADP-ribose acceptor sites in the AD as well as additional acceptor sites in the DNA-binding domain. These insights will allow further investigations to elucidate the biological functions of PARP1 and its enzymatic activity.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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