Strand-specific Recognition of DNA Damages by XPD Provides Insights into Nucleotide Excision Repair Substrate Versatility

Recognition and removal of DNA damages is essential for cellular and organismal viability. Nucleotide excision repair (NER) is the sole mechanism in humans for the repair of carcinogenic UV irradiation-induced photoproducts in the DNA, such as cyclobutane pyrimidine dimers. The broad substrate versatility of NER further includes, among others, various bulky DNA adducts. It has been proposed that the 5’-3’ helicase XPD (xeroderma pigmentosum group D) protein plays a decisive role in damage verification. However, despite recent advances such as the identification of a DNA-binding channel and central pore in the protein, through which the DNA is threaded, as well as a dedicated lesion recognition pocket near the pore, the exact process of target site recognition and verification in eukaryotic NER still remained elusive. Our single molecule analysis by atomic force microscopy reveals for the first time that XPD utilizes different recognition strategies to verify structurally diverse lesions. Bulky fluorescein damage is preferentially detected on the translocated strand, whereas the opposite strand preference is observed for a cyclobutane pyrimidine dimer lesion. Both states, however, lead to similar conformational changes in the resulting specific complexes, indicating a merge to a “final” verification state, which may then trigger the recruitment of further NER proteins.

Maintenance of genomic integrity is one of the most important cellular tasks and is largely achieved by a number of different DNA repair systems targeting diverse types of DNA lesions, such as erroneous alterations in the genetic code, chemical base modifications, or bulky adducts (1, 2). Nucleotide excision repair (NER) is an essential DNA repair mechanism with an exceptionally large range of chemically and structurally unrelated targets. In humans, it is furthermore the only repair system for the removal of UV irradiation-induced damages, and dysfunctional NER is responsible for severe diseases including xeroderma pigmentosum (3, 4). Eukaryotic NER encompasses a total of ~30 proteins, including the xeroderma pigmentosum group proteins (XPA–XPG). In the current model of NER, repair can either be initiated by a stalled RNA polymerase in transcription coupled NER or via global genome NER through high affinity binding of the XPC-HR23B heterotrimer to short distorted and destabilized DNA structures containing ss/ds-DNA junctions (1–4). The ATPase/helicase XPB, which is part of the 10 subunit transcription factor IIG (TFIIH) complex, directly interacts with XPC (5), and ATP-dependent conformational rearrangements of XPB likely further enhance the size of the nascent DNA bubble (6). XPD, the second helicase within TFIIH, is a functional 5’-3’ helicase, and its helicase activity is exploited to further increase the size of the unpaired region (7) to permit the binding of additional NER factors. More importantly, however, XPD has been proposed to assume a central role in damage verification (8–12). Once the damage has been verified, the NER cascade proceeds with the recruitment of additional proteins including the endonucleases XPG and XPF-ERCC1, resulting in the excision of a 24–32-nt oligonucleotide containing the lesion (13–15).

Within a DNA repair mechanism, the process of verifying a target site is of paramount importance, because this step affords a mechanism the high specificity that grants efficient processing of cytotoxic or carcinogenic DNA lesions while preventing futile repair. Crystal structures of XPD from different archaeal organisms have provided valuable insight into the general architecture of this enzyme (8, 9, 16). Archaeal XPDs share high sequence homology with the human XPD protein and are exploited as model systems for analyses of the structure and function of their human counterpart (8–12, 16, 17). In our studies, we used XPD from the archaeal organism Thermoplasma acidophilum, a single-stranded DNA: CPD, cyclobutane pyrimidine dimer; TFIIH, transcription factor IIG; taXPD, XPD from T. acidophilum; ATPγS, adenosine 5’-O-(thiotriphosphate).
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TABLE 1

DNA substrates for AFM and BLI experiments

The sequences that form the DNA bubbles are underlined. F, fluorescein adducted thymine; CPD, cyclobutane pyrimidine dimer; [CPD]).

| Substrate | DNA sequence |
|-----------|--------------|
| A Bottom strand | GGT CGA CTC TAG AGG ATC AGT CTT GCT GCT AGT |
| B Top −/− bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| C Top −/ bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| D Top F/bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| E Top F/bubble (14 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| F Top F’/5 bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| G Top F’/3 bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| H Top F−/− | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| I Top CPD/bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| J Top CPD’/5 bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| K Top CPD’/3 bubble (8 nt) | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |
| L Top CPD−/− | GCA TGC CTC GAG TCT AGA GCT ACC GAG ACC GCT |

plasma acidophi lum (tXPD). The enzyme consists of four domains: two RecA-like helicase domains, a domain coordinating an iron-sulfur cluster, and an arch domain. The iron-sulfur and arch domains together with helicase domain 1 comprise a narrow pore with ~1-nm diameter (8, 9). In addition, the crystal structure of tXPD in complex with a short stretch of ssDNA, as well as reverse footprinting analysis, have led to a model of the possible path of the DNA across the enzyme (11, 18). In this model, the DNA threads through the protein pore and is in close proximity to the iron-sulfur cluster, consistent with a proposed role of such clusters in DNA damage investigation (19–23) and the recent identification of a dedicated lesion recognition pocket near the pore (12). However, the exact mechanism of lesion verification by and in particular the impressive substrate versatility of XPD remained elusive so far.

We used the single molecule technique of atomic force microscopy (AFM) to directly visualize individual XPD-DNA complexes at nanometer resolution. By introducing a specific lesion at a known position in long DNA fragments (916 base pairs), we created substrates that more closely resemble the natural occurring in vivo substrates than the short DNA oligonucleotides utilized with other methods for the analysis of protein-DNA interactions. Importantly, the exact knowledge of the lesion position within the DNA substrate allows us to distinguish between specifically bound protein complexes (bound at the lesion site) and nonspecifically bound complexes (bound elsewhere on homoduplex DNA). We exploited this approach to investigate the ability of XPD to recognize and verify two different types of lesions and to directly visualize conformational responses of the complexes to damage verification. The lesions are representatives of two distinct classes of damage repaired by NER, a fluorescein as a representative for bulky DNA adducts (24–26), and a cyclobutane pyrimidine dimer (CPD) as the major species of DNA damage resulting from UV radiation (27, 28). Our data clearly demonstrate specific stalling of tXPD at these target sites upon ATP-driven translocation on long DNA substrates. Most notably, however, our AFM data unambiguously show different DNA strand selectivity for the two lesions, indicating that tXPD utilizes distinct verification strategies for structurally diverse types of DNA damage.

EXPERIMENTAL PROCEDURES

Protein Expression and Purification—XPD from T. acidophilum (wild-type and K170A variant) was expressed and purified as described previously (8). Briefly, expression at 14 °C for 18 h of the N-terminally His-tagged protein in Escherichia coli BL21-CodonPlus (DE3)-RIL cells (Stratagene) was induced with 0.1 mM isopropyl-β-D-thiogalactoside. XPD was purified by metal affinity (nickel-nitrilotriacetic acid; Invitrogen) followed by size exclusion chromatography (HiLoad 26/60 Superdex 200 prep grade; GE Healthcare) in 50 mM Tris, pH 8.0, 300 mM NaCl, and 1 mM DTT.

DNA Substrates—DNA substrates for AFM and biolayer interferometry are listed in Table 1. The plasmid pUC19N (2,729 bp), kindly provided by S. Wilson’s laboratory (NIEHS, National Institutes of Health), served as circular DNA for the AFM experiments and as the basis for the 916-bp linear specific DNA substrates. The linear DNA substrates for AFM experiments were prepared as described previously (29). Details are given in the supplemental materials. Briefly, the modified DNA plasmid pUC19N contains closely spaced restriction sites of the nickase Nt.BstNBI (New England Biolabs). Incubation with Nt.BstNBI followed by heating in the presence of an excess of complementary oligonucleotide results in the removal of the ssDNA stretch between the nick positions, which can subsequently be replaced by a substrate containing a particular specific target site (listed in Table 1). Each step of the DNA substrate preparation was confirmed using restriction enzyme assay testing (supplemental Fig. S1). For AFM experiments, linear DNA fragments of 916-bp length containing either a fluorescein or a CPD lesion at 28–33% of the fragment length (depending on the presence and position of a DNA bubble directly surrounding the lesion or 5’ or 3’ of the lesion; see Table 1) were produced by digestion of the DNA with the restriction enzymes SspI and BspQI (NEB). All DNA fragments were purified by gel extraction (NucleoSpin Extract II kit; Macherey-Nagel) prior to the experiments. For biolayer interferometry, DNA oligomer A (Table 1) was obtained with a 3’ biotin modification (Integrated DNA Technologies) and annealed with complementary oligomers (Table 1, substrates B–L) at equimolar amounts.

Biolayer Interferometry DNA Binding Assay—DNA binding affinities were measured by biolayer interferometry (BLI) on an
TABLE 2
DNA binding affinities

| substrate       | $K_d$ [nM] | SD [nM] | n   | P         |
|-----------------|------------|---------|-----|-----------|
| ssDNA           | 155        | 61      | 13  | -         |
| dsDNA nsp       | 541        | 234     | 12  | $6.4 \times 10^4$ |
| - / bubble (8nt) | 155        | 75      | 16  | 0.48      |
| F / bubble (8nt)| 163        | 69      | 14  | 0.36      |
| F / -           | 490        | 210     | 12  | $7.9 \times 10^5$ |
| CPD / bubble (8nt)* | 138   | 52      | 8   | 0.28      |
| CPD / -         | 340        | 94      | 11  | $5.5 \times 10^5$ |

Octet RED system (Fortebio, Menlo Park, CA) using 50 nM DNA and 100–300 nM XPD for the DNA substrates containing ssDNA regions and 100 nM DNA and 400–500 nM XPD for fully dsDNA substrates. The binding assays were carried out in XPD incubation buffer (20 mM Tris, pH 8.5, 10 mM KCl, 5 mM MgCl$_2$, 1 mM EDTA, 5 mM DTT) supplemented with 0.1 mg/ml BSA and were performed in at least triplicate and with three different protein batches. In BLI, the thickness of protein molecular layers bound to DNA immobilized on the surface of a streptavidin-coated fiber optic sensor (ForteBio) is determined from the interference of light reflected from the sensor surface. DNA oligomers (Table 1) were purchased with a biotin group attached to the 3’ end of the bottom strand and coupled to the sensor surface via streptavidin-biotin interaction. DNA loading concentrations were optimized in a concentration series for the individual DNA substrates. Assuming a 1:1 binding model, dissociation ($K_{\text{dis}}$) and association rate constants ($k_{\text{ass}}$) were determined with the Octet data analysis software from increased and decreased attached protein layer thicknesses. Dissociation constants ($K_d$, Table 2) were calculated as the ratio of dissociation and association rate constants for applied protein concentration $c$.

$$K_d = \frac{k_{\text{dis}}}{k_{\text{ass}}} = \frac{k_{\text{dis}}}{k_{\text{obs}} \times (c^{-1})}$$  \hspace{1cm} (Eq. 1)

Atomic Force Microscopy Experiments—Incubations for AFM experiments were carried out at 350 nM XPD (wild-type or K170A) and 15 nM DNA substrate (see above and Table 1) for 30 min at 37°C in XPD incubation buffer (see “Biolayer Interferometry DNA Binding Assay” above) ± 2 mM ATP or ATPγS. DNA substrates were heated to 65°C for 10 min and slowly cooled down to ambient temperature prior to incubation to remove salt microcrystals that may form during storage. For sample deposition, the incubations were diluted 8-fold in AFM deposition buffer (25 mM HEPES, pH 7.5, 25 mM sodium acetate, 10 mM magnesium acetate) to a final volume of 20 μl and immediately pipetted onto freshly cleaved mica (Grade V; SPI Supplies), rinsed with ultra-pure deionized water and dried in a gentle stream of nitrogen. In the rinsing step, the negatively charged DNA polymers (± bound proteins) are stably chelated to the negative charges on the mica surface (at pH 7.5) by Mg$^{2+}$ ions in the applied buffer (29). Free protein molecules in the incubation also deposit and are fixed by the drying procedure on the substrate surface. For the visualization of minute protein-DNA complexes, mica is superior as an AFM substrate to all other currently known materials (29). This layered silicate provides extremely clean, flat, and smooth surface properties with typical surface roughness of 0.05 nm (root mean square) and the further advantage of rapid and easy experimental sample preparation (29). AFM images were captured with a molecular force probe 3D-Bio AFM (Asylum Research, Santa Barbara, CA) in tapping mode using OMCL-AC240TS (Olympus) noncontact/tapping mode silicon probes with spring constants of ~2 N/m and resonance frequencies of ~75 kHz. Images were collected at a scan speed of 2.5 μm/s with scan sizes of 2 μm × 2 μm, 4 μm × 4 μm, or 8 μm × 8 μm and pixel resolution of ~2 nm.

AFM Data Analysis—AFM images were third order plane-fitted and flattened using Asylum Research software on Igor Pro and subsequently analyzed using ImageJ (National Institutes of Health) similarly as described previously (30, 31). Example images for several representative conditions are shown in Fig. 1 and supplemental Figs. S2 and S3. DNA aggregates and fragments that were cut off by the image margins were excluded from analysis. For quantification of DNA coverage with protein (Fig. 1B), experiments were carried out at least in triplicate for each substrate and condition, with $n = 340–876$ DNA bound protein peaks for each substrate type (representative images in supplemental Fig. S2). DNA fragments and XPD-DNA complexes in the AFM images were counted to obtain binding densities per bp as shown in Equation 2.

$$\text{Binding density} = \frac{\#\text{protein peaks}}{(\#\text{DNA fragments} \times \# \text{basepairs})}$$  \hspace{1cm} (Eq. 2)

To exclude the possibility that protein molecules located on DNA in the images were merely accidentally co-localized upon sample deposition instead of being bound to DNA, experiments were also repeated with a protein (of similar size) involved in the regulation of neuronal synapses. This protein is not expected to display any significant degree of DNA binding, consistent with a significantly reduced degree of DNA coverage ((0.6 ± 0.1)/1,813-bp DNA compared with (1 ± 0.2) and (1.8 ± 0.5) for XPD in the presence and absence of ATP, respectively ($p = 0.000019$ and 0.00015, respectively), from analyses of three to seven independent experiments; supplemental Fig. S4).

Binding preferences for a target site are expressed as specificities (S) and were determined from Gaussian fits to the position distributions of protein peaks on DNA as described previously (30, 31) using the software Origin Pro. Briefly, DNA fragment lengths and positions of protein complexes on the
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DNA were measured as the length of a line along the entire DNA backbone or from the complex location to the DNA fragment ends, respectively. Position distributions were then plotted as normalized to the full length of the DNA fragment. Enhanced binding of a protein to a specific target site within a DNA fragment results in a peak in the distribution at the corresponding position: 0% for complexes bound at DNA fragment ends, 50% for complexes bound at a DNA fragment center, ~31% for complexes bound at the lesion, and/or DNA bubble for specific DNA substrates in these experiments. Many proteins involved in DNA repair show a propensity to also bind to DNA fragment ends, likely as a result of local helix destabilization (29, 32). XPD also shows a slight preference for DNA ends (bars at 0% DNA length in position distributions of Fig. 1D). Preferential localization of XPD at DNA fragment ends, at an unpaired DNA bubble region, or at a lesion site are, however, distinct and separate effects. Because here we are interested in recognition by XPD of specific target sites in the DNA (at ~31% of DNA length), we hence excluded complexes bound at DNA ends from the distribution histograms for statistical analysis (starting at 5% DNA length). Importantly, because of the vast excess of nonspecific sites over the one specific lesion site, we also observe many complexes at nonspecific DNA sites in the images (for example supplemental Figs. S2 and S3). However, the imbalance between the frequencies of these different types of DNA sites (>900 nonspecific sites versus 1 specific site) can still result in a considerable specificity value (site preference) as calculated from statistical analyses. Such preference is visible as a Gaussian shaped peak in the data at the specific site (~31% DNA length). A Gaussian fit to these distributions provides the maximum position of the peak and its surface area (specific area, $A_{sp}$), which reflects the fraction of specific complexes. The area of the background, which represents the fraction of complexes bound nonspecifically to the DNA fragment ($A_{nsp}$), is given by the product of DNA length as a percentage and the height of the background from the Gaussian fit to the specific site. Target site specificity ($S$) is then given by Equation 3 (30),

$$S = N \times \frac{A_{sp}}{A_{nsp}} + 1 \quad \text{(Eq. 3)}$$

where $N$ is the number of available binding sites on the DNA (here $N$ = 914, excluding DNA fragment ends). For better comparison of specificities for different targets, Gaussian widths were all fixed at 2.6%, the width of the Gaussian fit to the substrate with highest specificity (DNA with fluorescein directly in the context of an 8-nt DNA bubble).

Statistical DNA fragment length distributions displayed a distinct maximum at ~292 ± 11 nm, corresponding to a deviation of ~6% from the theoretical length of a 916-bp DNA fragment of 311 nm (assuming 0.34 nm/bp). Such underestimation of DNA lengths in AFM by up to 10% is typical and likely due to limited resolution of smaller DNA backbone convolutions in the images (29). Only DNA fragments within two standard deviations from the center of a Gaussian fit to the DNA length distributions were included in the analyses to ensure correct allocation of the lesion position at 28–33% of the DNA fragment length (see “DNA Substrates” above). We distinguish between specific protein-DNA complexes (bound at the site of the lesion and/or DNA bubble at ~31% of DNA length ± 2 S.D.) and nonspecific complexes, based on complex position. As in previous studies on other protein systems (31, 32), we avoided labeling of DNA fragment ends to allow for loading of protein at the fragment ends. As a consequence, the two fragment ends are indistinguishable, and the statistical position distributions were plotted only to 50% of DNA lengths, with the center of the fragment corresponding to 50% DNA length and the fragment ends to 0% DNA length. This approach results in a low nonspecific background at the position of the specific site in the distribution. However, because this background is present at all positions along the DNA substrate, it merely contributes noise to the distribution but does not affect the ratio of specific and nonspecific areas and hence does not change the value of $S$. The resolution of this specificity determination is typically limited to $S > 10$ (31). Specificity results in Table 3 are given as the averages from at least three individual experiments (except for batch 2 protein experiments, which were mostly performed in duplicate) with errors derived from the standard deviations between experiments. For all DNA substrates and conditions, analysis with the pooled distributions from the individual experiments led to similar results and analytical errors comparable to the variations between the individual experiments. The position distributions shown in supplemental Fig. S5 are from pooled data from repeated experiments. The results shown in Fig. 2 are all from the same protein batch for better comparison between different substrates; results from a second batch are shown for comparison in Table 3 (in gray), clearly confirming the observed strand selectivity.

**TABLE 3**

**DNA binding specificities**

The results are given as averages from 2–4 experiments with standard deviations between experiments. In gray below, results obtained with a second protein batch for some of the DNA substrates indicate good reproducibility. Examples of position distributions of XPD on the different DNA substrates, from which these specificities were calculated, are shown for several conditions in supplement Fig. S5.

| substrate | + ATP | + ATPys | - ATP |
|-----------|-------|--------|-------|
| F / bubble (8nt) | 131 ± 51 | n.d. | 213 ± 21 |
| F / bubble (8nt) | 851 ± 280 | 231 ± 67 | 169 ± 168 |
| F / bubble (14nt) | 468 ± 157 | n.d. | n.d. |
| F / 5’ bubble (8nt) | 445 ± 167 | 102 ± 60 | 101 ± 53 |
| F / 3’ bubble (8nt) | 95 ± 24 | n.d. | n.d. |
| F / - | 113 ± 89 | n.d. | n.d. |
| CPD / bubble (8nt) | 522 ± 112 | 374 ± 105 | 66 ± 51 |
| CPD / 5’ bubble (8nt) | 117 ± 39 | 96 ± 48 | n.d. |
| CPD / 3’ bubble (8nt) | 283 ± 99 | 82 ± 70 | n.d. |
| CPD / - | 137 ± 105 | n.d. | n.d. |
DNA bend angles are defined as deviation from a straight DNA backbone and were measured by manual tangent overlay as described previously (32), using ImageJ. Briefly, lines a and b were placed along the DNA backbone on each side of a protein complex. (180° − α) is the DNA bend angle, with α being the angle between a and b. For analysis using Origin Pro, complexes bound at positions within two standard deviations (based on Gaussian widths) from the center of the fit to the position distribution (at 30 ± 1% of DNA fragment length) were classed as specific, whereas complexes bound between 0 and 23% or between 37 and 50% of the DNA fragment lengths were considered nonspecific complexes. Bend angle analysis was carried out separately for specific and nonspecific XPD-DNA complexes. Bend angle distributions (for example Fig. 3) are from pooled data from repeated experiments. Significance of changes in bend angles was determined using the center positions of Gaussian fits to the data and standard deviations as given by the Gaussian fit widths (see “Statistical Analysis of Significance” below). In control experiments of intrinsic nonspecific DNA bending (supplemental Fig. S6), a mask comparable to the size of the protein was moved along the DNA fragments, and at regular intervals the bend angle was determined as described above. Intrinsic bending at lesion or bubble sites in the absence of bound protein was similarly determined using the protein sized mask at 30% of the DNA length from both of the DNA ends (producing a 50% nonspecific background population).

Statistical Analysis of Significance—Affinities to DNA, specificities for DNA sites, and DNA bend angle analyses were quantitatively compared for different DNA substrates and conditions. Significance was calculated based on a one-tailed Student’s t test and is classed as p < 0.05 (*), p < 0.01 (**), and p < 0.005 (***)..

RESULTS

XPD Loads Preferentially on Unpaired Regions within dsDNA—XPD has been previously shown to bind ssDNA with high affinity in the nanomolar range (10, 11, 17, 19, 33). We used BLI to analyze binding affinities of taXPD to different DNA structures, namely to dsDNA containing an unpaired region 8 nt in length, uninterrupted dsDNA, or ssDNA (see “Experimental Procedures”). To focus specifically on DNA affinity effects, BLI experiments were carried out in the absence of ATP. The results can be grouped into high affinity and low affinity binding, with dissociation constants of ∼150 and ∼500 nM, respectively, depending on the presence (high affinity) or absence (low affinity) of a single-stranded region (Table 2). Importantly, our data show similar affinities of taXPD for purely ssDNA and for dsDNA of the same length containing an 8-nt-long unpaired region, suggesting that the size of an 8-nt DNA bubble is sufficient to support loading of taXPD at this site. Consistent with the BLI results, our AFM analyses also demonstrate that taXPD can bind to fully base paired dsDNA (Fig. 1) but binds preferentially at ssDNA regions within a DNA bubble, as well as to DNA fragment ends (Fig. 1D).

Lesion Recognition and Stalling at Lesion Sites Requires ATP Hydrolysis—When adding ATP, we notice a significantly more pronounced decrease in protein coverage for linear dsDNA compared with circular dsDNA substrate (Fig. 1B, compare circu-
due to impaired loading onto the DNA, consistent with the higher affinity of XPD for ssDNA compared with dsDNA.

To separate effects of lesion recognition and of XPD loading onto DNA at an unpaired region, we further analyzed position distributions of XPD on DNA substrates containing an access site at a defined distance from the lesion site (Fig. 2, supplemental Fig. S5, and Table 3). We produced DNA substrates containing an 8-nt DNA bubble at a distance of 27 bp 5’ to the fluorescein lesion (see “Experimental Procedures”), a distance that can be overcome by the helicase activity of XPD (10). Although this separation is too close for us to distinguish in the AFM images between protein complexes localized at the DNA bubble and complexes at the lesion, these studies provide distinct information on the effects of separating loading and target recognition. When XPD was loaded at the bubble 5’ to the fluorescein lesion, the protein clearly accumulated at the target with high specificity, indicating 5’-3’ translocation on the DNA and recognition of the fluorescein lesion on the translocated DNA strand (Fig. 2). Our data also show that the process of DNA translocation that leads to lesion recognition is dependent on ATP hydrolysis because lesion localization specificity was reduced to background levels for XPD loaded at a bubble 5’ to the lesion in the presence of the nonhydrolyzable or only slowly hydrolyzable ATP analog ATP\(_{\text{S}}\) (S\(_{\text{S}}\)) comparable to the specificity for (−/bubble); Fig. 2). Lesion localization specificity in the presence of ATP was approximately half for this substrate compared with the fluorescein lesion within the context of a bubble (S\(_{\text{S}}\) compared with S\(_{\text{S}}\) of XPD molecules bound at the bubble moving in 5’ to 3’ direction on the DNA strand that does not contain the lesion, thus never encountering the lesion. In contrast, when loaded on the bubble directly surrounding the lesion, a stable XPD-lesion complex can form (10), leading to considerably lower losses from total protein binding.

Interestingly, our data suggest two modes of XPD translocation on our DNA substrates. Although both modes of translocation depend on ATP, they differ significantly. Nonspecific sliding of XPD on homoduplex DNA apparently can proceed over distances of hundreds of base pairs with eventual sliding off at the DNA fragment ends. In contrast, loading of XPD at
However, when we used a DNA substrate containing a DNA lesion at the bubble site rather than by stalling at the lesion. In contrast to the CPD lesion, XPD specificity was low for fluorescein when located on the nontranslocated strand of the protein occurred at a bubble 5’ to a lesion (with S.D. of 3°) independent of the substrate used and independent of the presence or absence of ATP or ATPγS (Table 4 and Fig. 3). Variation between substrates and experiments was small, with a S.D. of <3°. In striking contrast, the bend angles of complexes located at the specific (lesion) site strongly depended on the presence of ATP (black bars in Fig. 3, A–C). In the absence of ATP (Fig. 3A), we measured an average DNA bend angle of 48° with small variation between the different substrates and experiments (S.D. of ±5°), similar to the bending of XPD complexes observed at nonspecific sites (see above). In the presence of ATP, bend angle distributions of XPD bound at the specific site displayed a maximum at ~65° for all substrates containing a lesion (with S.D. of 3°; Fig. 3 and Table 4). This constitutes a highly significant shift in DNA bend angles as compared with the average bend angle observed for nonspecific complexes (p < 10^{-7}; Table 4 and “Experimental Procedures”), indicative of a conformational change in the specific XPD complexes at a lesion site, which clearly is not observed in the absence of ATP. Interestingly, however, a shift to an average specific site bend angle of ~64° (S.D. of ±2°) also occurred in the presence of the nonhydrolyzable (or only slowly hydrolyzable) ATP analog ATPγS (Fig. 3C and Table 4). These experiments thus clearly demonstrate that ATP (re)binding but not hydrolysis upon reaching and recognizing the target site is absolutely required for processing of the lesion, likely by conformational rearrangements. In addition, bend angle distributions of specific (lesion-bound) complexes in the presence of ATP or ATPγS were described better by a double Gaussian than by a single Gaussian fit (Fig. 3). According to the relative surface areas of the double Gaussians, typically ~65% of the specific complexes display DNA bend angles of ~65°, and ~35% of the complexes show bending by a smaller angle of ~30°. The second, smaller bend angle population will be further discussed below.

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FIGURE 2. DNA translocation and lesion recognition by XPD. Localization specificities of XPD in the presence (black bars) or absence (gray bars) of ATP or in the presence of ATPγS (white bars) were obtained from Gaussian fits (supplemental Fig. S5) to the statistical AFM position distributions of XPD on different DNA substrates (schematically indicated below the plot). Numbers for DNA substrates are also given, consistent with those in Table 3, which lists specificities for all targets and ATP conditions. Striped bars show specificities of the taXPD K170A variant for the fluorescein/5’ bubble DNA and CPD/3’ bubble DNA substrate, as indicated. The red circle represents a fluorescein, and the blue rectangle represents a CPD lesion. Significance (calculated as p < 0.05 (*) and p < 0.01 (**) was calculated for differences in specificity compared with a DNA bubble without a lesion in the presence of ATP (first bar).
Lesion Recognition Is Coupled to Helicase Stalling—To further analyze the significance of the ability of XPD to recognize a lesion, we repeated the experiments using a helicase mutant of taXPD, K170A. This XPD variant harbors a mutation within the iron-sulfur cluster domain, which affects and significantly enhances its helicase activity compared with the wild-type protein, as described previously (11). Lesion specificities of taXPD-K170A obtained from AFM imaging on DNA substrates containing a DNA bubble 5’/H11032 or 3’/H11032 to a fluorescein or CPD lesion (Fig. 2, striped bars) are consistent with (and even slightly lower than) the background level resulting from a slight preference for loading at an unpaired DNA bubble site over homoduplex DNA. These data strongly argue that DNA translocation by this XPD variant is not stalled by the lesion, likely because of its increased helicase activity resulting in its scanning past potential lesion sites. We also measured DNA bend angles induced by taXPD-K170A bound at the position of the fluorescein and CPD lesions in DNA (Fig. 3, F and I). The complete absence of a shift to the specific bend angle state in these distributions strongly supports the inability of this protein variant to investigate and process target sites.

DISCUSSION

Damage recognition and verification are critical processes in DNA repair, which have to ensure the speedy detection and processing of DNA lesions yet avoid futile incisions. The heli-
case XPD has been proposed to play a critical role in NER damage verification. Previous studies further indicated that XPD translocation may be stalled by DNA lesions, suggesting that the protein may be involved in damage recognition (10, 35). Single molecule imaging by AFM allowed us to investigate the contributions of the XPD enzyme to the NER lesion recognition and verification process and to analyze this function in the presence of long DNA substrates (>900 bp), which mimic more closely the in vivo situation in the cell as compared with the short substrates required for other in vitro studies.

We analyzed the ability of taXPD to interact with various DNA substrates. ssDNA or ss/dsDNA junctions were bound with moderately high affinity (~150 nM). Interestingly, the affinity to dsDNA (not containing any ssDNA regions) was only less than 1 order of magnitude weaker than to ssDNA (Table 2). Furthermore, our AFM data indicate that taXPD was not only able to bind to but also to translocate along dsDNA in the presence of ATP (Fig. 1). However, in the absence of a ssDNA region in the DNA substrate, lesions in the DNA were only poorly recognized by the enzyme (Fig. 2). It is therefore tempting to speculate that taXPD adopts at least two different binding modes upon DNA binding, in which only the second binding mode, which is induced by an initial interaction with ssDNA, is competent of dsDNA unwinding and supports successful lesion verification.

| Bend angle (degrees) | + ATP | + ATPγs | - ATP |
|----------------------|-------|---------|-------|
| Substrate            |       |         |       |
| - / bubble (8nt)     |       |         |       |
| 1                    | 44.1 ± 25.1 n = 164 | 50.8 ± 22.4 n = 123 | n.d. | n.d. | 46.6 ± 18.8 n = 246 | 47.7 ± 22.1 n = 196 |
| F / bubble (8nt)     | 51.9 ± 13.9 n = 94 | 27.2 ± 12.9 (27%) 65.0 ± 12.9 (73%) n = 137 | 49.4 ± 19.3 n = 242 | 28.0 ± 12.9 (26%) 65.0 ± 12.9 (72%) n = 224 | 48.5 ± 21.3 n = 174 | 51.2 ± 22.5 n = 113 |
| F / bubble (14nt)    | 49.7 ± 13.1 n = 212 | 40.1 ± 8.6 (36%) 65.9 ± 8.6 (64%) n = 223 | n.d. | n.d. | n.d. | n.d. |
| F / 5’ bubble (8nt)  | 52.9 ± 17.5 n = 184 | 32.9 ± 10.2 (37%) 63.3 ± 10.2 (66%) n = 188 | 50.7 ± 17.1 n = 201 | 27.4 ± 11.5 (35%) 62.7 ± 11.5 (65%) n = 166 | n.d. | n.d. |
| F / 3’ bubble (8nt)  | 47.4 ± 19.4 n = 253 | 32.1 ± 13.3 (38%) 67.7 ± 13.3 (62%) n = 118 | n.d. | n.d. | n.d. | n.d. |
| F / -                | 49.7 ± 23.0 n = 205 | 40.4 ± 10.0 (44%) 68.3 ± 10.0 (56%) n = 98 | n.d. | n.d. | n.d. | n.d. |
| CPD / bubble (8nt)   | 51.9 ± 17.9 n = 156 | 36.1 ± 11.5 (31%) 66.4 ± 11.5 (69%) n = 195 | 47.8 ± 19.4 n = 103 | 29.5 ± 11.7 (27%) 65.1 ± 11.7 (73%) n = 103 | 48.5 ± 19.6 n = 195 | 44.0 ± 22.4 n = 97 |
| CPD / 5’ bubble (8nt) | 51.5 ± 23.4 n = 313 | 26 ± 10.0 (34%) 59.8 ± 10.0 (66%) n = 212 | n.d. | n.d. | n.d. | n.d. |
| CPD / 3’ bubble (8nt) | 49.3 ± 18.0 n = 184 | 31.5 ± 10.8 (27%) 65.5 ± 10.8 (73%) n = 150 | n.d. | n.d. | n.d. | n.d. |

**TABLE 4**

DNA bend angles measured from AFM images are given for XPD-DNA complexes bound specifically at a target site (spec) or at non-specific (nsp) positions within different DNA substrates (as indicated schematically). The given values were derived as the maxima of single or double Gaussian fits to pooled bend angle distributions from 2–4 experiments. The total number of data points is given as n. The significances of the shift in average bend angle between nonspecific and specific complexes (in the presence of ATP) were $1.6 \times 10^{-11}$, $4.3 \times 10^{-11}$, $9.8 \times 10^{-11}$, $4.3 \times 10^{-10}$, $5.6 \times 10^{-10}$, $7.4 \times 10^{-10}$, $7.4 \times 10^{-9}$, and $1.4 \times 10^{-8}$ for the lesion containing DNA substrates 2–8, respectively. Significances of shifts in bend angles between the absence and presence of ATP (or ATPγs) for specific complexes were $2.2 \times 10^{-8}$ ($4.8 \times 10^{-8}$) and $9.5 \times 10^{-10}$ ($5.3 \times 10^{-10}$) for DNA substrates 2 (fluorescein) and 6 (CPD), respectively. Exemplary bend angle distributions are shown in Fig. 3.
Lesion Recognition by XPD

Most importantly, our analysis compared directly for the first time NER damage recognition and verification for different lesions. Introduction of a lesion into a DNA substrate within the context of an unpaired DNA region led to complex formation with high specificity both for a CPD and for a fluoroscein lesion, which differs significantly in their structure. However, when the lesion was removed from the unpaired region and positioned 3’ or 5’ to this region, a clear distinction in damage recognition became apparent (model shown in Fig. 4). A bulky fluoroscein adduct leads to a stalled taXPD-DNA complex when the lesion is located on the translocating strand. In contrast, a CPD lesion is preferentially recognized when it is located on the opposite strand, i.e., the nontranslocating strand.

Differences in NER mechanistic details may be related to the strong observed dependence of DNA repair efficiencies on the degree of DNA helix destabilization by different lesions (36, 37). A possible explanation for the strand selectivity observed in our studies is therefore a different recognition mechanism based on the diverse structural prerequisites for different types of lesions. In our studies, the bulky fluoroscein adduct may result in direct mechanical blocking of XPD translocation, but only when it is encountered on the actual strand that the protein "holds on to." Importantly, under the conditions used in our experiments, the fluoroscein adduct is most likely negatively charged and would therefore not destabilize the negatively charged DNA duplex via intercalation. Loading of XPD onto DNA carrying the type of lesion represented by a fluoroscein hence requires the presence of a DNA bubble (Fig. 2), which is provided in vivo by the concerted action of XPC and XPB. A CPD lesion also does not lead to a major destabilization of the DNA double helix (38–40). However, distortion of the CPD containing ssDNA strand by the thymine dimer may be sufficient to provide an access site for XPD loading, resulting in slightly enhanced XPD localization to the lesion site in the absence of a DNA bubble (Fig. 2). When loaded at an access site (DNA bubble) at a distance from the lesion, translocation of taXPD along the lesion-containing strand appears to be feasible and is not strongly hindered by the presence of the lesion. Therefore a different, so far unidentified verification process could be envisioned. It is tempting to speculate that, when the lesion is located on the nontranslocated strand, it may be in close proximity to the iron-sulfur cluster (Fig. 4A), thus supporting the hypothesis that the iron-sulfur cluster may act as a damage detector, as has been shown for other protein systems containing such clusters (20, 22). However, the exact location of the DNA strand that XPD does not directly bind to (the nontranslocated strand) is not resolved in the crystal structure and is hence so far not known with certainty (11). Further structural studies are clearly required to elucidate the (different) mechanism(s) of lesion verification by XPD.

It should be noted that the strand selectivity observed in our AFM experiments for recognition of a CPD lesion is in contrast to a recent publication by Naegeli and co-workers (10), who reported stalling of the archaeal *Ferroplasma acidarmanus* XPD helicase by a CPD lesion located both in the nontranslocated and in the translocated DNA strand. Their studies provide support for the formation of a stable complex between *F. acidarmanus* XPD helicase and a CPD lesion in the translocated DNA strand, with interactions that are strong enough to withstand incision by the CPD processing glycosylase T4 Endo V. Both mechanistic deviations between XPD from different species and/or variations caused by different experimental approaches are conceivable and will be worth investigating in future studies. Notably, compared with the short oligonucleotides employed in these biochemical experiments, the long DNA substrates in our AFM studies provide better stability of the DNA duplex and, importantly, more closely resemble physiological conditions. Our data do not argue against such stable complex formation with CPD lesions in the translocated strand, but report a strong preference for detection of CPD lesions in the nontranslocated strand versus in the translocated strand. Importantly, once verified, lesion processing by taXPD appears to be similar for all substrates in our experiments, as suggested by vast differences in lesion specificities (Fig. 2) but comparable bend angle distributions for complexes engaged at specific lesion sites (Fig. 3).

In the context of DNA damage search, taXPD clearly requires a ssDNA region for successful stalling at a lesion site. In eukaryotic NER, XPD is part of the TFIIH complex, which is initially recruited to the damaged DNA by XPC. The presence of a destabilized DNA region (as for instance in a 3-nt DNA bubble) has been shown to be essential for XPC-induced loading of NER factors and the subsequent excision of a CPD lesion (34). Although Sugasawa et al. (34) show convincing evidence for recognition of CPD damages in the same DNA strand that the TFIIH complex is loaded on, these studies do not exclude incision competent recognition of CPD on the nontranslocated strand after XPC loading on a symmetrical DNA bubble 3’ to the lesion. Importantly, their data further corroborate a two-step (bipartite) model and the importance of TFIIH orientation for correct lesion recognition and processing.

Once taXPD has verified the presence of an NER target, processing of the lesion involves conformational changes in the...
stalled complex at the lesion visible by a significant shift in the maximum of the distribution of induced DNA bend angles at the site of the bound protein from ~50° to ~65° (Fig. 3). These transitions were independent of the lesion type and of the details of the preceding lesion recognition strategy. In the context of the eukaryotic repair cascade, this conformational change may be the prerequisite for the recruitment of the remaining NER machinery, including the endonucleases XPG and XPF for damage removal. This conformational shift is completely absent in samples of the taXPD variant K170A, which is incapable of detecting NER target sites. These results further underline the significance of stalled XPD translocation for concomitant lesion recognition as a prerequisite for lesion-specific processing by XPD in NER. Fluorescein- and CPD-DNA structures have previously been shown to display intrinsic bending by 15–30° at the lesion (41, 42). However, AFM bend angle distributions obtained on the lesion sites in the absence of protein (supplemental Fig. S6) contain no major population displaying these bend angles and hence argue against an innate preformed DNA conformation that XPD binds to. It is conceivable that the ~30° bend angle conformation observed in all specific site bend angle distributions may represent a complex conformation sampled by the protein on the path to the specific lesion repair signaling complex. However, it is the larger bend angle state (~65°) that is dominant in the lesion-specific complexes (~70% of all complexes) and comparable in population to the significantly less bent state (~50°) in the nonspecific complexes at homoduplex DNA sites. We therefore interpret this bend angle conformation as the specific, lesion associated state competent for induction of subsequent DNA repair. Interestingly, the conformational changes occurred in the presence of either ATP or ATPγS, indicating that the lesion-dependent rearrangement of the taXPD-DNA complex involves ATP binding but not hydrolysis. The requirement of ATP rebinding for lesion-dependent conformational changes is strongly reminiscent of the prokaryotic NER mechanism. In the prokaryotic NER damage search and recognition complex, UvrB is thought to undergo initial conformational changes upon ATP hydrolysis. This process leads to its localization at the lesion site, followed by ATP rebinding and concomitant formation of a stable, specific preincision complex at the lesion (43), which is required for the recruitment of the endonuclease UvrC (44). General conservation of the mechanistic NER approach between the prokaryotic UvrABC system and the eukaryotic xeroderma pigmentosum system has often been described (45–47), despite a complete lack of sequence and structural homology between the involved enzymes. This is the first example showing that an individual step within the verification process in the NER cascade may be strikingly similar between prokaryotic and eukaryotic NER, corroborating the conservation of this biologically essential DNA repair system.

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