The TFIIH subunits p44/p62 act as a damage sensor during nucleotide excision repair

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ABSTRACT

Nucleotide excision repair (NER) in eukaryotes is orchestrated by the core form of the general transcription factor TFIIH, containing the helicases XPB, XPD and five ‘structural’ subunits, p62, p44, p34, p52 and p8. Recent cryo-EM structures show that p62 makes extensive contacts with p44 and in part occupies XPD’s DNA binding site. While p44 is known to regulate the helicase activity of XPD during NER, p62 is thought to be purely structural. Here, using helicase and adenosine triphosphatase assays we show that a complex containing p44 and p62 enhances XPD’s affinity for dsDNA 3-fold over p44 alone. Remarkably, the relative affinity is further increased to 60-fold by dsDNA damage. Direct binding studies show this preference derives from p44/p62’s high affinity (20 nM) for damaged ssDNA. Single molecule imaging of p44/p62 complexes without XPD reveals they bind to and randomly diffuse on DNA, however, in the presence of UV-induced DNA lesions these complexes stall. Combined with the analysis of a recent cryo-EM structure, we suggest that p44/p62 acts as a novel DNA-binding entity that enhances damage recognition in TFIIH. This revises our understanding of TFIIH and prompts investigation into the core subunits for an active role during DNA repair and/or transcription.

INTRODUCTION

Cell survival relies on accurate replication and transcription of genetic material. When DNA is damaged, rapid, efficient repair is essential to ensure genome integrity. Deficiencies in these repair processes have severe consequences as exemplified by defects in the nucleotide excision repair (NER) machinery leading to diseases such as xeroderma pigmentosum (XP), which is phenotypically characterized by UV hyper-sensitivity and increased incidence of skin cancers (1). Mutations found in many components of the NER pathway, including TFIIH can lead to XP. The core structure of TFIIH comprises two helicases; of these, XPB is enzymatically involved in both transcription and repair, whereas XPD’s activity is only essential for repair (2,3). In addition, TFIIH possesses five other core subunits: p44, p62, p34, p8 and p52 (4–13). While it has been shown that p44 and p52/p8 can stimulate XPD and XPB’s helicase activities respectively (8,14,15), most of the core subunits remain relatively unstudied. A recent cryo-EM structure of TFIIH reveals significant interactions between p44, p62 and XPD (16). Part of p62 is located in XPD’s DNA binding site, suggesting that p62 might assume a role in modulating XPD’s ability to interact with DNA. This would expand p62’s function beyond stabilizing TFIIH and acting as a hub for protein interactions in transcription and crucially with the initial damage sensor XPC-HR23B for NER (14,17–20).

Previous attempts to purify full-length p44 have been hampered by protein stability issues, however, a truncated form of p44 (N-p44) has been expressed and has been shown to activate the adenosine triphosphatase (ATPase) of XPD (21). In this study, we successfully purified full length p44 through co-expression with p62. We show that the complex formed by both proteins enhances XPD’s affinity for dsDNA. However, the presence of damaged dsDNA dramatically increases the relative affinity for the ternary XPD-p44/p62 complex 60-fold versus XPD-N-p44. Remarkably, in the absence of XPD, we observe that p44/p62 also preferentially binds to damaged single-stranded DNA compared to undamaged ssDNA, likely explaining the enhanced selectivity for damage in the presence of XPD-p44/p62. Using single molecule fluorescence microscopy, we show that p44/p62 alone binds and randomly diffuses on non-damaged dsDNA, but this complex is stalled in the presence of UV lesions, suggesting a direct response to dam-
Our data thus reveals that p44/p62 acts as a damage-specific DNA binding pocket when bound to single-stranded DNA (ssDNA). This hypothesis is supported by the in-depth analysis of a recent TFIIH cryo-EM structure (22), where we placed p62 fragments into unassigned regions of the EM maps to propose that p62 is located proximally to XPD’s DNA binding entity. The ternary p44/p62 complex was generated in 20 mM HEPES pH 7.5, 250 mM NaCl and 1 mM TCEP on a Superdex200 Increase 5/50 GL column (CYTIVA) containing 20 mM HEPES pH 7.5, 50 mM NaCl and 1 mM TCEP as loading buffer and the same buffer containing 1 M NaCl was used for elution. The final buffer after AEC was 20 mM HEPES pH 7.5, 250 mM NaCl and 1 mM TCEP. The proteins were concentrated to at least 5 mg/ml based on their calculated extinction coefficient using ProtParam (SwissProt) and then flash frozen for storage at −80°C.

**Figure 1.** SEC analysis of XPD, the dimeric p44/p62 and the ternary XPD/p44/p62 complexes. SEC chromatograms of XPD (92 kDa), p44/p62 (132 kDa) and the XPD/p44/p62 (224 kDa) complex were performed in 20 mM HEPES pH 7.5, 250 mM NaCl and 1 mM TCEP on a Superdex200 Increase 5/50 GL column (CYTIVA) containing 20 mM HEPES pH 7.5, 50 mM NaCl and 1 mM TCEP as loading buffer and the same buffer containing 1 M NaCl was used for elution. The final buffer after AEC was 20 mM HEPES pH 7.5, 250 mM NaCl and 1 mM TCEP. The proteins were concentrated to at least 5 mg/ml based on their calculated extinction coefficient using ProtParam (SwissProt) and then flash frozen for storage at −80°C.

**MATERIALS AND METHODS**

**Purification**

The genes encoding p44 and p62 were cloned from *Chaetomium thermophilum* cDNA, p62 was cloned into the pETM-11 vector (EMBL) without a tag. p44 was cloned into the pBADM-11 vector (EMBL) containing an N-terminal hexa-Histidine tag followed by a TEV cleavage site. p62 and p44 were co-expressed in *E. coli* BL21 CodonPlus (DE3)-RIL cells (Agilent) and were co-purified in liquid nitrogen for storage.

**DNA binding**

The ATPase assay used dsDNA substrates used:

- F26,50 contains a fluorescein moiety covalently attached to thymine (*);
- 5′GACATCGTACTGTTACGGCCTCCATCT*CTACCG CAATCAGGCGGATCTGC 3′
- The reverse complementary sequence to F26,50:
- 5′GAGATCTGGCGATTGGGTAGCGGAG TGGCACAGTACGGTC 3′
- F26,50 without the fluorescein moiety;
Nucleic Acids Research, 2020 3

Figure 2. Steady-state ATPase and helicase activity of XPD complexes in the presence of various DNA substrates and core TFIIH proteins. (A) XPD’s helicase activity is stimulated by N-p44 (hashed) and p44/p62 (white) on an open fork substrate with lengths indicated in nucleotides. XPD alone displays no helicase activity (21). Errors are shown as S.E.M from nine repeats. Statistical significance determined using a student’s t-test where * = P < 0.05, n.s = not statistically significant. (B) Summary table of the $k_{cat}$ values and $K_M$ values obtained from DNA substrate titrations. Errors are S.E.M fit values. (C) Summary of the effect of DNA substrates on XPD-p44/p62 (white) and XPD-N-p44 (hashed) affinities. XPD-p44/p62 shows a 60-fold enhancement in affinity versus XPD-N-p44. (D) XPD-p44/p62’s ATPase is stimulated by the presence of DNA substrates. Damaged dsDNA (squares) binds much more tightly than undamaged DNA (triangles), and approximately equivalent to ssDNA (circles). ATPase rates are normalized to the protein concentration used so that the asymptote reports the $k_{cat}$. Errors are shown as S.E.M from three or more repeats. In the absence of DNA, XPD possesses a slow ATPase that is unaffected by N-p44, p44/p62 or damaged DNA (Supplementary Table S1) indicating that p44/p62 displays no ATPase activity by itself.

$5'\text{GACTACGTACTGTACGGCTCCATCTCTACCGCAATCAGGCCAGATCTGC}3'$

The NADH-coupled ATPase assay was performed as described previously (23) in plate reader format. Imaging buffer containing the NADH-reaction components was supplemented with 1 mM fresh TCEP, protein (50 nM (equimolar concentrations for XPD N-p44 and XPD p44/p62)), and increasing amounts of DNA substrate (4–500 nM). The reaction was started with the addition of 2 mM ATP to each well, and the change in OD$_{340}$ (NADH) was monitored every 8 seconds/well over 30 min at room temperature in a Cariostar plate reader. The rates of NADH consumption were used to calculate $k_{cat}$. Reactions were repeated at least three times and include biological replicates, and S.E.M used as errors values.

$5'\text{AGCTACCATGCCCTCATCTCTACCGCAATCAGGCCAGATCTGC}3'$

Helicase activity assay

Helicase activity was analyzed utilizing a fluorescence-based assay. We used an open fork substrate with a Cy3 label at the 3’ end of the translocated strand where unwinding of the DNA substrate reduces quenching of the Cy3 fluorescence.

$5'\text{GCTATGACCATGATTACGAATTGCTTGGAATCCTGACGAACTGTAG}3'$ and a dabcyl modification on the 5’ end of the opposite strand

Assays were carried out in 20 mM HEPES pH 7.5, 50 mM KCl, 5 mM MgCl$_2$, and 1 mM TCEP. DNA was used at a concentration of 250 nM. Helicase activity was measured with equimolar concentrations of XPD in the presence of various DNA substrates and core TFIIH proteins.
RESULTS AND DISCUSSION

p44/p62 co-purify as a complex that stimulate XPD’s activity

Both p44 and p62 make extensive contacts with XPD in TFIIH (16). To investigate this interaction, recombinant p44, p62 and XPD from  C. thermophilum were utilized. All subunits share high sequence homology with the human TFIIH subunits and we have shown previously that they serve as an ideal model for the human proteins (3,29). Only the N-terminal region (residues 1–285) of p44, containing the von Willebrand domain (N-p44), has been successfully purified in isolation to date, nonetheless this is sufficient to activate XPD’s ATPase and helicase (3). By co-expressing full-length p44 with full-length p62, we purified a tightly associated complex of p44/p62 that also interacts with XPD (Figure 1), thus providing the first successful purification of full length p44 and suggesting that p44 requires p62 to fold correctly.

Recently Greber et al. showed by cryo-EM, that a portion of p62 is located in XPD’s DNA binding site within the DNA-free transcriptional apo-TFIIH structure (16). To understand how this might affect XPD, we measured the helicase activity of XPD-p44/p62 relative to XPD-N-p44. Surprisingly, XPD-p44/p62 possessed a significantly enhanced ability to unwind the DNA substrate compared to XPD-N-p44 (Figure 2A), indicating that p62 enhances rather than blocks DNA binding. This was confirmed using a dsDNA titration of XPD’s ATPase activity in complex with either N-p44 or p44/p62, (Figure 2B and C) revealing a substantial enhancement in the affinity of XPD-p44/p62 for dsDNA (57.8 ± 0.9 nM) versus XPD-N-p44 (149.6 ± 55 nM). Surprisingly, the ATPase $k_{cat}$ rate constant was only slightly affected by the presence of p62 (Figure 2B). Altogether these data suggest that p62 activates XPD by enhancing its affinity for DNA.

More remarkably, titrating different DNA substrates while measuring the ATPase of XPD-p44/p62 (Figure 2C and D) revealed a substantial enhancement in the affinity for fluorescein-modified (recognized as damage by NER (27)) dsDNA (2.7 ± 0.2 nM) versus undamaged dsDNA (57.8 ± 9.0 nM). In contrast, XPD-N-p44 showed near identical affinity for dsDNA regardless of the presence of DNA damage (Figure 2C). Both XPD-N-p44 and XPD-p44/p62 showed similar affinity for ssDNA (2.5 ± 0.5 versus 2.6 ± 0.2 nM), indicating p44/p62 is not simply displaying a higher affinity for exposed ssDNA (Figure 2B and D). These observations provide the first evidence that p62 in the ternary XPD-p44/p62 complex may enhance the affinity of the complex for DNA damage. Such behavior explains the sensitization to UV irradiation observed for truncations of the yeast p62 homolog (17,30,31).

The p44/p62 complex binds directly to DNA

How p62 enhances the affinity of the ternary complex for DNA is unknown; however, it is conceivable that p62 undergoes a conformational change and is thereby able to interact directly with the DNA. To investigate this hypothesis, we determined p44/p62’s affinity for different DNA substrates relevant to NER by fluorescence polarization (Figure 3). In support of our hypothesis, p44/p62 was found to bind ds-
DNA with a $K_D$ of 1.18 μM. Interestingly, DNA intermediates expected to form during NER showed even stronger binding by p44/p62 than dsDNA ($K_D$ (ssDNA) 0.84 μM and $K_D$ (open fork) 0.21 μM).

Since p44/p62 enhanced XPD’s affinity for damaged DNA, we also investigated the affinity of p44/p62 for different DNA damage containing substrates independent of XPD. p44/p62 bound dsDNA containing a fluorescein moiety with an affinity similar to that of undamaged DNA ($K_D$ 1.23 and 1.18 μM, respectively). In contrast, we observed a striking 40-fold increase in binding affinity to a ssDNA substrate containing a central fluorescein lesion ($K_D$ 0.02 μM), compared to the ssDNA substrate without damage. The tighter binding to these NER intermediates strengthens the hypothesis that p62 contacts DNA within TFIIH, and specifically locates to damage within an opened bubble.

The p44/p62 complex independently diffuses on DNA

To directly determine how p44/p62 interacts with DNA, we imaged individual fluorescently tagged proteins binding to DNA tightropes in real time using fluorescence microscopy (25). In this assay, p44/p62 is fluorescently labeled using a quantum dot (Qdot) conjugated to the protein via its poly-histidine purification tag and an anti-His IgG antibody (24). Individual protein molecules are then imaged interacting with single DNA molecules in real time (Figure 4A). We observed substantial binding of p44/p62 complexes to dsDNA, and of these ~80% diffused randomly ($n = 599$ total). To investigate the mechanism of p44/p62’s diffusion on the DNA, the diffusion constant and exponent (used to determine if motion is directionally biased) was calculated in high and low salt conditions (100 versus 10 mM KCl). At elevated KCl concentrations fewer p44/p62 complexes bound to DNA, and the calculated diffusion constant using mean-squared displacement analysis (25) showed no significant change ($P > 0.05$) between salt conditions (10 mM KCl 0.067 μm$^2$/s ± 0.006 versus 100 mM KCl 0.042 μm$^2$/s ± 0.010; Figure 4B), consistent with p44/p62 complexes sliding along the DNA helix (32). The diffusive exponent was found to be ~0.9 in either salt condition, which indicates that there is no directional bias to the diffusive motion (33). Based on the estimated size of a p44/p62 complex conjugated to a Qdot, the diffusion constant is consistent with rotation-coupled diffusion around the backbone of the DNA helix (25,34,35). Altogether, these data provide the first observation that the p44/p62 complex can bind dsDNA and randomly diffuse, independently of XPD.
The position of p62 in apo- and DNA bound TFIIH structures. (A) Representation of the Kokic et al. DNA-bound TFIIH structure (PDB ID: 6RO4) shown in cartoon mode with the EM-envelope displayed as gray mesh. (B) The unmodeled cryo-EM density around p44/p34 was fit using p62 from the Greber et al. apo-TFIIH structure (PDB ID: 6NMI) and the three helical bundle (residues 450–547) superimposed with an rmsd of 0.5 Å after rebuilding. Residues 397–450 differ in orientation due the concerted movement of XPD-p44 in 6RO4. The additional part of p62 interacting with XPD was modeled according to the cryo-EM density with no defined amino acid sequence. (C) Close-up view of XPD and p62 based on the apo-TFIIH structure, with the iron sulphur cluster in XPD shown as spheres. (D) Equivalent view of XPD from the DNA-bound TFIIH structure with the additional cryo-EM density modeled as p62. The modeled structure has been deposited in the worldwide protein databank, accession code: 7AD8.

p44/p62 complexes are stalled by the presence of damaged DNA

To further analyze these observations, we investigated the effect of DNA damage on the behavior of single p44/p62 complexes on DNA tightropes irradiated with 500 J/m² of 254 nm UV light, which generates cyclobutane pyrimidine dimers and 6–4 photoproducts (23). We observed a striking and significant decrease in the number of diffusing p44/p62 complexes from ∼80% to 50% (P < 0.05) (Figure 4C). These results clearly indicate that p44/p62 independently recognizes DNA damage and stalls at damaged sites. We also observed a change in the mechanism of p44/p62’s diffusion in the presence of UV damage; a population of p44/p62 complexes now appeared limited in their total excursion distance on DNA (Figure 4D), similar to the damage recognition protein XPC, which undergoes constrained diffusion around a lesion (36,37). Although p44/p62 cannot discriminate DNA damage in a dsDNA substrate without XPD (Figure 3), the observed stalling in the DNA tightrope assay suggests that p44/p62 may have located exposed ssDNA regions containing UV lesions.

In summary, we have shown an enhanced affinity of XPD-p44/p62 for damaged dsDNA (Figure 2), enhanced affinity of p44/p62 for damaged ssDNA but not to damaged dsDNA (Figure 3), and an ability to differentiate damage sites on dsDNA tightropes (Figure 4C). Altogether, these observations are consistent with a role for p44/p62 in recognizing DNA lesions in the opened bubble of TFIIH.

The structural basis of p62-DNA contacts within TFIIH

The recent cryo-EM structure of TFIIH bound to DNA shows the path of single-stranded DNA as it is passed from XPB to XPD (22). In this structure, p44 is located in the same position as in apo-TFIIH (16), but p62 is unassigned. However, additional cryo-EM densities can be observed in the structure at a position similar to p62 in apo-TFIIH (Figure 5A) (16). We modeled regions of p62 from the apo-TFIIH into the DNA-bound TFIIH structure and found excellent agreement with the position of p62 (Figure 5B). This model suggests that the cryo-EM density close to XPD could represent N-terminal regions of p62. Closer examination of the DNA binding groove within XPD shows that p62 occupies this site in the absence of DNA (Figure 5C), i.e. in the apo TFIIH structure. Analysis of the same region in the DNA-bound structure suggests that the p62 loop could undergo a conformational change to form a helical bundle that interacts with the Arch domain of XPD and the ssDNA bound to XPD (Figure 5D); however, we cannot entirely exclude the small possibility that another protein (e.g. XPG) also contributes to this density. How these interactions result in damage recognition is an open question since the precise mechanism of damage recognition/verification has not been established. However, it could be postulated that p62’s Arch interaction affects recognition at the FeS cluster pocket (38). Or even more speculative, given its position 5' from the FeS cluster, it is also conceivable that a second recognition site may exist, explaining the origin of...
differential damage detection (27). This would be in good agreement with damage recognition site postulated for the XPD homolog DinG that is located directly below the p62 Arch domain junction (39).

In conclusion, we present the first mechanistic characterization of the non-helicase TFIIH subunits p44/p62. Our biochemical data clearly show that the p44/p62 complex binds to DNA, but displays a profoundly tighter affinity for ssDNA containing damage (Figure 3). Modeling of non-assigned densities in a recent cryo-EM structure indicate p62 to be in a position where it could interact with XPD and DNA in a substrate dependent manner (Figure 5). Taken together, these data suggest that p44/p62 interacts with ssDNA bound to XPD and is capable of modulating its helicase/ATPase activity through interactions with the helicase and/or Arch domains. This provides a structural explanation for the observed DNA damage-dependent increase in XPD’s DNA substrate affinity in the presence of p44/p62 (Figure 2). Furthermore, we have used fluorescence polarization binding studies and single molecule imaging to directly demonstrate that p44/p62 complexes are capable of recognizing UV damage and consequently stalling on DNA (Figures 3 and 4). This behavior could be important for stalling XPD at a lesion and triggering damage verification. Nonetheless, our results clearly show that the p44/p62 complex plays an active role in TFIIH’s recognition of DNA damage and indicates p44/p62 can no longer be thought of simply as ‘structural subunits’.

**DATA AVAILABILITY**

All data will be made available through the University of Kent academic repository (https://kar.kent.ac.uk/). The structure modeled in Figure 5 has been deposited in the worldwide protein databank, accession code: 7AD8.

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

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**REFERENCES**

1. Bradford, P.T., Goldstein, A.M., Tamura, D., Khan, S.G., Ueda, T., Boyle, J., Oh, K.S., Imoto, K., Inui, H., Moriwaki, S. et al. (2011) Cancers and neurologic degeneration in xeroderma pigmentosum: long term follow-up characterizes the role of DNA repair. *J. Med. Genet.*, **48**, 168–176.

2. Coint, F., Oksenzh, V. and Egly, J.M. (2007) Distinct roles for the XBP/p52 and XPD/p44 subcomplexes of TFIIH in damaged DNA opening during nucleotide excision repair. *Mol. Cell.*, **26**, 245–256.

3. Kuper, J., Braun, C., Elias, A., Michels, G., Sauer, F., Schmitt, D.R., Poterszman, A., Egly, J.M. and Kisker, C. (2014) In TFIIH, XPD helicase is exclusively devoted to DNA repair. *PLoS Biol.*, **12**, e1001954.

4. Roy, R., Schaeffer, L., Humbert, S., Vermeulen, W., Weeda, G. and Egly, J.M. (1994) The DNA-dependent ATPase activity associated with the class II basic transcription factor BTF2/TFIIH. *J. Biol. Chem.*, **269**, 9826–9832.

5. Greber, B.J., Nguyen, T.H.D., Fang, J., Afonine, P.V., Adams, P.D. and Nogales, E. (2017) The cryo-electron microscopy structure of human transcription factor IIH. *Nature*, **549**, 414–417.

6. Sung, P., Bailly, Y., Weber, C., Thompson, L.H., Prakash, L. and Prakash, S. (1993) Human xeroderma pigmentosum group D gene encodes a DNA helicase. *Nature*, **365**, 852–855.

7. Schulz, A.F., Fribourg, S., Poterszman, A., Marlow, V., Moras, D. and Egly, J.M. (2000) Molecular structure of human TFIIH. *Cell*, **102**, 599–607.

8. Luo, J., Cimerman, P., Viswanathan, E., Beem, C.C., Kim, B., Dehecq, M., Raman, V., Greenberg, C.H., Pellarin, R., Sali, A. et al. (2015) Architecture of the human and yeast general transcription and DNA repair factor TFIIH. *Mol. Cell.*, **59**, 794–806.

9. Aroulo, S.J., Tirode, F., Coint, F., Pospiech, H., Sywosa, J.E., Stucki, M., Huber, U., Egly, J.M. and Wood, R.D. (2000) Nucleotide excision repair of DNA with recombinant human proteins: definition of the minimal set of factors, active forms of TFIIH, and modulation by CAK. *Genes Dev.*, **14**, 349–359.

10. Svejstrup, J.Q., Wang, Z., Feave, W.J., Wu, X., Bushnell, D.A., Donahue, T.F., Friedberg, E.C. and Kornberg, R.D. (1995) Different forms of TFIIH for transcription and DNA repair: Holo-TFIIH and a nucleotide excision repairosome. *Cell*, **80**, 21–28.

11. Habraken, Y., Sung, P., Prakash, S. and Prakash, L. (1996) Transcription factor TFIIH and DNA endonuclease Rad2 constitute yeast nucleotide excision repair factor 3: implications for nucleotide excision repair and Cockayne syndrome. *PNAS*, **93**, 10718–10722.

12. Sung, P., Prakash, L., Matson, S.W. and Prakash, S. (1987) RAD3 protein of Saccharomyces cerevisiae is a DNA helicase. *Proc. Natl. Acad. Sci. U.S.A.*, **84**, 8951–8955.

13. Sung, P., Prakash, L., Weber, S. and Prakash, S. (1987) The RAD3 gene product of Saccharomyces cerevisiae encodes a DNA-dependent ATPase. *PNAS*, **84**, 6045–6049.

14. Jawhari, A., Laine, J.P., Dubaele, S., Lamour, V., Poterszman, A., Coint, F., Moras, D. and Egly, J.M. (2002) p52 Mediates XPB function within the transcription/repair factor TFIIH. *J. Biol. Chem.*, **277**, 31761–31767.

15. Coint, F., Marini, J.C., Rodolfo, C., Fribourg, S., Pedrini, A.M. and Egly, J.M. (1998) Mutations in the XPD helicase gene result in XP and TTD phenotypes, preventing interaction between XPD and the p44 subunit of TFIIH. *Nat. Genet.*, **20**, 184–188.

16. Greber, B.J., Tos, D.B., Fang, J. and Nogales, E. (2019) The complete structure of the human TFIIH core complex. *eLife*, 8, e47711.

17. Gervais, V., Lamour, V., Jawhari, A., Frindel, F., Wasselewski, E., Dubaele, S., Egly, J.M., Thierry, J.C., Kieffer, B. and Poterszman, A. (2004) TFIIH contains a PH domain involved in DNA nucleotide excision repair. *Nat. Struct. Mol. Biol.*, **11**, 616–622.

18. Yoko, M., Masutani, C., Maekawa, T., Sugasawa, K., Ohkuma, Y. and Hanaoka, F. (2000) The xeroderma pigmentosum group C protein complex XPC-HR23B plays an important role in the recruitment of transcription factor IIH to damaged DNA. *J. Biol. Chem.*, **275**, 9870–9875.

19. Sugasawa, K., Masutani, C., Uchida, A., Maekawa, T., Van Der Spek, P., Bootsma, D., Hoeijmakers, J. and Hanaoka, F. (1996) HHR23B, a human Rad23 homolog, stimulates XPC protein in nucleotide excision repair in vitro. *Mol. Cell. Biol.*, **16**, 4852–4861.
20. Di Lello, P., Jenkins, L.M.M., Jones, T.N., Nguyen, B.D., Hará, T., Yamaguchi, H., Dikeakos, J.D., Appella, E., Legault, P. and Omichinski, J.G. (2006) Structure of the Tfb1/p53 complex: Insights into the interaction between the p62/Tfb1 subunit of TFIIH and the activation domain of p53. *Mol. Cell.*, 22, 731–740.

21. Kuper, J., Braun, C., Elias, A., Michels, G., Sauer, F., Schmitt, D.R., Poterszman, A., Egly, J.-M. and Kisker, C. (2014) In TFIIH, XPD helicase is exclusively devoted to DNA repair. *PLoS Biol.*, 12, e1001954.

22. Kokić, G., Chernev, A., Tegunov, D., Dienemann, C., Urlaub, H. and Cramer, P. (2019) Structural basis of TFIIH activation for nucleotide excision repair. *Nat. Commun.*, 10, 2885.

23. Burnett, J.T. and Kad, N.M. (2019) Understanding the coupling between DNA damage detection and UvrA’s ATPase using bulk and single molecule kinetics. *FASEB J.*, 33, 763–769.

24. Springall, L., Inchingolo, A.V. and Kad, N.M. (2016) DNA-protein interactions studied directly using single molecule fluorescence imaging of quantum dot labeled proteins moving on DNA tightropes. *Methods Mol. Biol.*, 1431, 141–150.

25. Kad, N.M., Wang, H., Kennedy, G.G., Warshaw, D.M. and Van Houten, B. (2010) Collaborative dynamic DNA scanning by nucleotide excision repair proteins investigated by single molecule imaging of quantum-dot labeled proteins. *Mol. Cell.*, 37, 702–713.

26. Jiang, Y., Ke, C., Mieczkowski, P.A. and Marszalek, P.E. (2007) Detecting ultraviolet damage in single DNA molecules by atomic force microscopy. *Biophys. J.*, 93, 1758–1767.

27. Buechner, C.N., Heil, K., Michels, G., Carell, T., Kisker, C. and Tessmer, I. (2014) Strand-specific recognition of DNA damages by XPD provides insights into nucleotide excision repair substrate versatility. *J. Biol. Chem.*, 289, 3613–3624.

28. Adams, P.D., Afonine, P.V., Bunkóczi, G., Chen, V.B., Davis, I.W., Echols, N., Headd, J.J., Hung, L.W., Kapral, G.J., Grosse-Kunstleve, R.W. et al. (2010) PHENIX: a comprehensive Python-based system for macromolecular structure solution. *Acta Crystallogr. D: Biol. Crystallogr.*, 66, 213–221.

29. Radu, L., Schoenwetter, E., Braun, C., Marcoux, J., Koelmel, W., Schmitt, D.R., Kuper, J., Cianferani, S., Egly, J.M., Poterszman, A., et al. (2017) The intricate network between the p34 and p44 subunits is central to the activity of the transcription/DNA repair factor TFIIH. *Nucleic Acids Res.*, 45, 10872–10883.

30. Gileadi, O., Feaver, W.J. and Kornberg, R.D. (1992) Cloning of a subunit of yeast RNA polymerase II transcription factor b and CTD kinase. *Science.*, 257, 1389–1392.

31. Matsu, P., DePaulo, J. and Buratowski, S. (1995) An interaction between the Tfb1 and Ssl1 subunits of yeast TFIIH correlates with DNA repair activity. *Nucleic Acids Res.*, 23, 767–772.

32. von Hippel, P.H. and Berg, O.G. (1989) Facilitated target location in biological systems. *J. Biol. Chem.*, 264, 675–678.

33. Saxton, M.J. (2001) Anomalous subdiffusion in fluorescence photobleaching recovery: a Monte Carlo study. *Biophys. J.*, 81, 2226–2240.

34. Schurr, J.M. (1979) The one-dimensional diffusion coefficient of proteins absorbed on DNA. *Biophys. Chem.*, 9, 413–414.

35. Hughes, C.D., Wang, H., Ghodke, H., Simons, M., Towheed, A., Peng, Y., Van Houten, B. and Kad, N.M. (2013) Real-time single-molecule imaging reveals a direct interaction between UvrC and UvrB on DNA tightropes. *Nucleic Acids Res.*, 41, 4901–4912.

36. Kong, M., Liu, L., Chen, X., Driscoll, K.I., Mao, P., Böhms, S., Kad, N.M., Watkins, S.C., Bernstein, K.A. and Wyrrick, J.J. (2016) Single-molecule imaging reveals that Rad4 employs a dynamic DNA damage recognition process. *Mol. Cell.*, 64, 376–387.

37. Cheon, N.Y., Kim, H.S., Yeo, J.E., Scherer, O.D. and Lee, J.Y. (2019) Single-molecule visualization reveals the damage search mechanism for the human NER protein XPC-RAD23B. *Nucleic Acids Res.*, 47, 8337–8347.

38. Wolski, S.C., Kuper, J., Hanzelmann, P., Truglio, J.J., Croteau, D.L., Van Houten, B. and Kisker, C. (2008) Crystal structure of the FeS cluster-containing nucleotide excision repair helicase XPD. *PLoS Biol.*, 6, e149.

39. Cheng, K. and Wigley, D.B. (2018) DNA translocation mechanism of an XPD family helicase. *eLife*, 7, e42400.