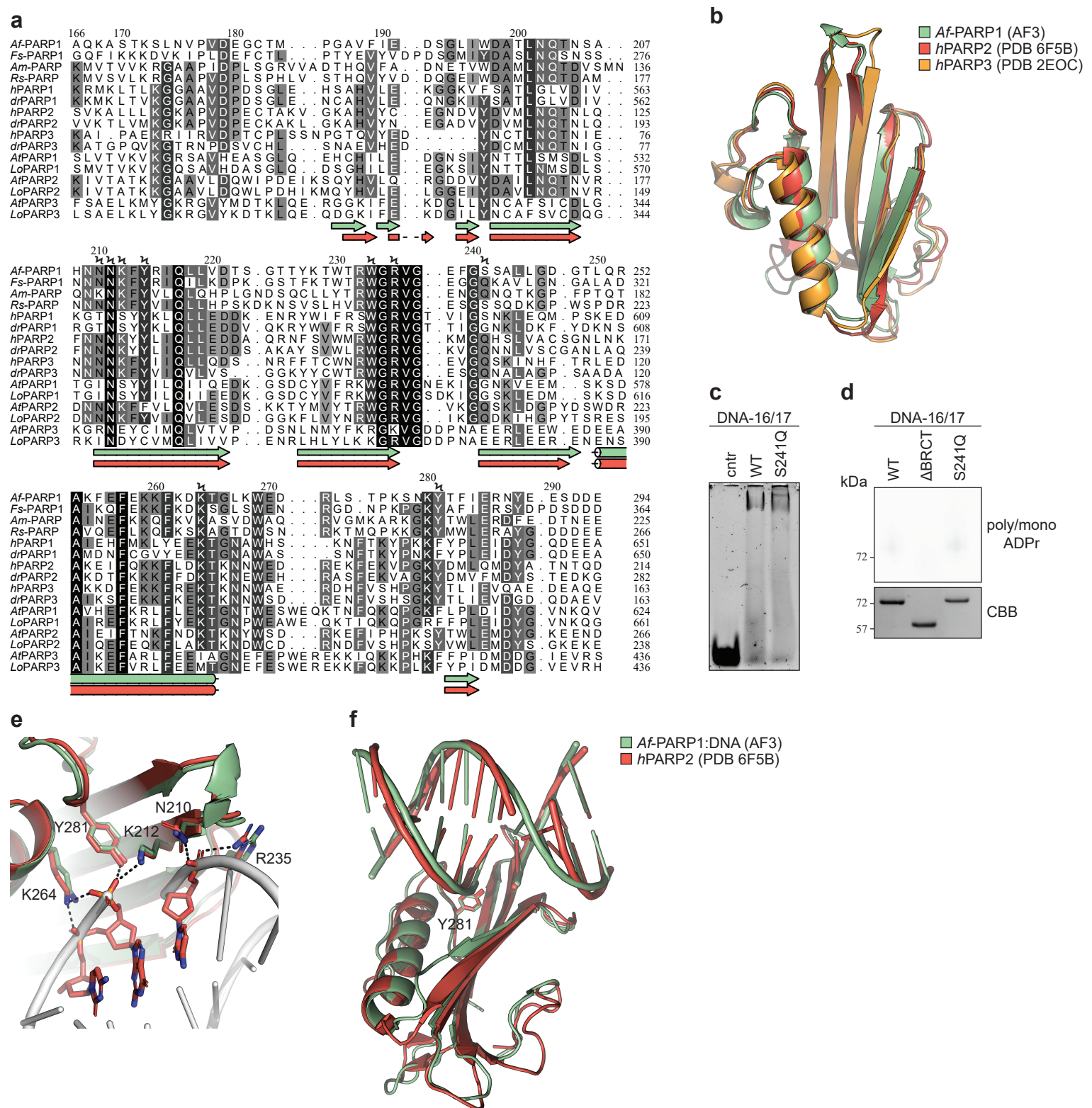




Extended Data Figure 2. DNA damage recognition by the Af-PARP1 WGR domain.

(a) Multiple sequence alignment of selected PARP-derived WGR domains from Fungi, Animalia, and Plantae. Residue numbers for Af-PARP1 and key DNA interaction residues (h) are indicated above the alignment. Secondary structures for Af-PARP1 (AlphaFold 3 model; green) and hPARP2 (PDB 6F5B; red) are given below the alignment.

(b) Ribbon representation of the structural alignment of Af-PARP1 WGR domain (AlphaFold 3 model) with the experimentally determined WGR domains from hPARP2 and -3.

(c) EMSA assay assessing the influence of Ser241 on the ability of the WGR domain to bind double-strand breaks. Af-PARP1 WT or S241Q (5 μ M) were incubated with DNA mimetic (1 μ M) for 1 h, separated via native PAGE and DNA visualised with SYBR safe staining.

(d) Assessment of the influence of Ser241 on the catalytic activity of Af-PARP1 in presence of 5'-phosphorylated DNA double-strand break mimetic. WT or S241Q Af-PARP1 (0.5 μ M) were incubated with DNA (0.6 μ M) and NAD⁺ (500 μ M) and formation of PAR detected using immunoblot.

(e) Ribbon-liquorice representation of the DNA coordination by hPARP2 (PDB 6F5B; red/white) overlaid with the Af-PARP1 WGR domain derived from the apo AlphaFold 3 model (green). Residues indicated represent Af-PARP1 numbering.

(f) Ribbon-liquorice representation of the structural alignment of Af-PARP1:DNA AlphaFold 3 model (only WGR domain shown) with the experimentally determined WGR:DNA complex from hPARP2. Af-PARP1 Tyr281 is indicated for orientation.