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Polθ reverse transcribes RNA and promotes RNA-templated DNA repair

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Abstract

Genome-embedded ribonucleotides arrest replicative DNA polymerases (Pols) and cause



deoxyribonucleotide reveals that the enzyme undergoes a major structural transformation within the thumb subdomain to accommodate A-form DNA/RNA and forms multiple hydrogen bonds with template ribose 2'-hydroxyl groups like retroviral RTs. Last, we find that Polθ promotes RNA-templated DNA repair in mammalian cells. These findings suggest that Polθ was selected to accommodate template ribonucleotides during DNA repair.

INTRODUCTION

Polymerase θ (Polθ) is a unique DNA polymerase-helicase fusion protein in higher eukaryotes whose A-family polymerase domain evolved from Pol I enzymes (**Fig. 1A**) (1, 2). However, contrary to most Pol I enzymes, Polθ is highly error-prone and promiscuous (3–6), performs translesion synthesis (TLS) opposite DNA lesions (3, 7, 8), and facilitates microhomology-mediated end-joining (MMEJ) of double-strand breaks (DSBs) by extending partially base-paired 3' single-stranded DNA (ssDNA) overhangs at DSB repair junctions (5, 9–12). Polθ is not expressed in most tissues but is highly expressed in many cancer cells, which corresponds to a poor clinical outcome (13, 14). Furthermore, Polθ confers resistance to genotoxic cancer therapies and promotes the survival of cells deficient in DNA damage response pathways (11, 13–16). Thus, Polθ represents a promising cancer drug target.

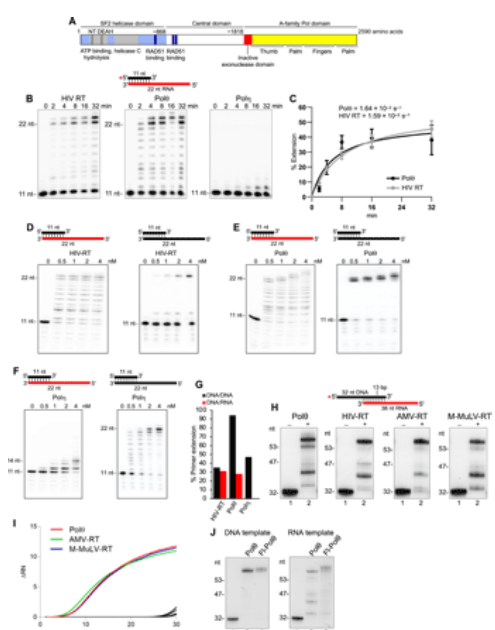


Fig. 1 Polθ exhibits reverse transcriptase activity.

(A) Schematic of full-length Polθ. (B) Denaturing gels showing a time course of DNA/RNA primer-template extension by the indicated polymerases. (C) Plot showing relative rate of DNA/RNA extension by Polθ and HIV RT. Data represent mean \pm SD; $n = 3$. (D to F) Denaturing gels showing DNA/RNA (left) and DNA/DNA (right) extension by the indicated polymerases. (G) Bar plot showing percent extension of DNA/RNA (red) and DNA/DNA (black) by the indicated polymerases (4 nM) [data from (D) to (F)]. (H) Denaturing gels showing DNA/RNA extension by the indicated polymerases. (I) Quantitative PCR chromatogram showing cDNA synthesis by the indicated polymerases. (J) Denaturing gels showing DNA/DNA (left) and DNA/RNA (right) extension by Polθ and FI-Polθ.

inactivating the 3'-5' proofreading function of closely related A-family bacterial Pol I Klenow fragment (KF) enables this polymerase to reverse transcribe RNA like retroviral reverse transcriptases (RTs), which lack proofreading activity (fig. S1A) (17, 18). Because Polθ is highly error-prone and promiscuous and contains an inactive proofreading domain, we hypothesized that it has RNA-dependent DNA synthesis activity. Given that ribonucleotides are the most frequently occurring nucleotide lesion in genomic DNA that arrest replicative Pols and cause DNA breaks (19, 20), we also envisaged that Polθ would tolerate template ribonucleotides during its DNA repair activities and thus promote RNA-templated DNA repair synthesis (RNA-DNA repair). Although RNA-DNA repair mechanisms have been demonstrated in genetically engineered yeast cells (21, 22), they remain obscure in mammalian cells.

RESULTS

Polθ exhibits RNA-dependent DNA synthesis activity

We tested whether the polymerase domain of Polθ (herein referred to as Polθ) reverse transcribes RNA like HIV RT using a DNA primer annealed to a RNA template (DNA/RNA). Polθ exhibits a similar rate of RT activity as HIV RT under identical conditions using substoichiometric amounts of enzyme relative to template (Fig. 1, B and C). Previous studies indicated that human Polη has RT activity at high micromolar concentrations (23). We find that Polη fails to perform reverse transcription beyond 3 nucleotides (nt) using conditions identical to those of Polθ and HIV RT at multiple concentrations (Fig. 1, B and C; compare Fig. 1F with Fig. 1, D and E; Fig. 1G). However, at significantly higher concentrations, Polη can further extend the DNA/RNA (figs. S1, B to E). Controls show that Polη is active on a DNA/DNA template like Polθ and HIV RT (compare Fig. 1F with Fig. 1, D and E; Fig. 1G). Overall, Polη exhibits increased stalling on RNA and requires higher enzyme concentration relative to Polθ for reverse transcription (figs. S1, B to E). Polθ RNA-dependent DNA synthesis activity is observed under various conditions and on different template constructs (figs. S1, F and G), and sequences (fig. S1H). Despite its robust activity on RNA, Polθ strongly discriminates against incorporating ribonucleotides (fig. S1I). Complementary DNA (cDNA) sequencing confirms Polθ's RNA-dependent DNA synthesis activity (fig. S2A) and reveals nucleotide misincorporations and indels, which is consistent with its low-fidelity DNA synthesis activity (fig. S2A) (3). HIV RT and other retrovirus RTs are also highly error-prone, demonstrating a shared characteristic between Polθ and retroviral RTs (18, 24–28). In addition to HIV RT, Polθ activity on RNA is nearly identical to RTs encoded by Moloney murine leukemia virus (M-MuLV) and avian myeloblastosis virus (AMV) (Fig. 1H). Polθ exhibits pausing events nearly identical to

The efficient RT activity of Polθ appears to be unique among human Polθs. Eight other human Polθs, representative of at least two enzymes from each polymerase family in humans (A, B, X, and Y), fail to reverse transcribe DNA beyond 2 to 3 nt under conditions identical to those of Polθ (Fig. 1F and fig. S3). Y-family polymerase κ (Polκ) exhibits limited RT activity under various conditions similar to Polη (Fig. S3H). Replicative Polθs δ and ε degrade the DNA primer on RNA due to exonuclease activity (fig. S3, F and G). All Polθs are active on DNA as expected (Fig. 1F and fig. S3). Full-length Polθ (Fl-Polθ) containing an N-terminal superfamily 2 and disordered central domain (Fig. 1A) (1) also has RT activity, suggesting that the endogenous protein performs RNA-DNA repair in cells (Fig. 1J). Because recombinant human Polθ and Fl-Polθ were purified from different organisms (*E. coli* and *Saccharomyces cerevisiae*, respectively) and by different methods (1, 9), the observed RT activity is not due to a protein contaminant. Consistent with this, human recombinant Polθ purified from *S. cerevisiae* also has robust RT activity (fig. S4A) (31).

Polθ exhibits higher velocity and fidelity of deoxyribonucleotide incorporation on RNA

We tested the relative velocity of Polθ deoxyribonucleoside monophosphate (dNMP) incorporation on RNA versus DNA. Single deoxyribonucleoside triphosphates (dNTPs) were incubated with Polθ during a time course on DNA/RNA versus DNA/DNA with identical sequence (Fig. 2A). Polθ exhibits a significantly higher velocity of incorporating deoxycytidine monophosphate (dCMP), deoxythymidine monophosphate (dTMP), and deoxyadenosine monophosphate (dAMP) on RNA (Fig. 2, B to D, and fig. S4B). The velocity of deoxyguanosine monophosphate (dGMP) incorporation was similar on RNA and DNA (Fig. 2E). Fl-Polθ also exhibits higher rates of dCMP, dTMP, and dAMP incorporation on RNA (fig. S5). Polθ exhibits a twofold higher affinity for DNA/RNA (fig. S4C), which may contribute to the higher velocity of nucleotide incorporation.

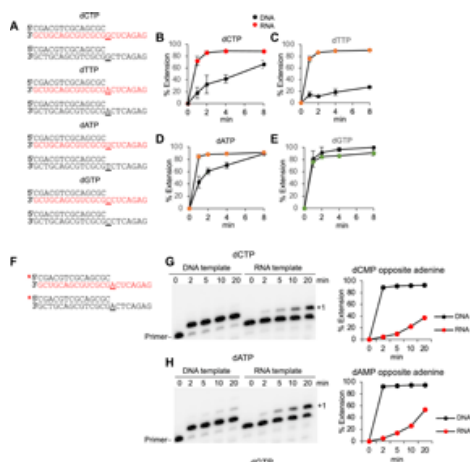


Fig. 2 Polθ exhibits a higher fidelity and velocity of deoxyribonucleotide incorporation on RNA.

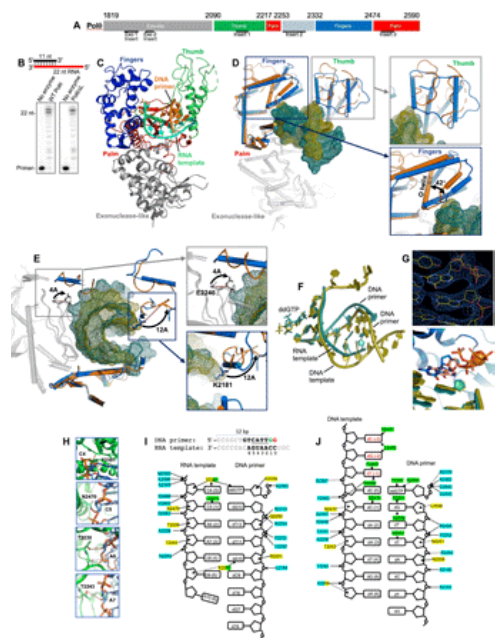
(A) Schematic of DNA/DNA and DNA/RNA templates used for the indicated dNTPs. Underlined base codes for the incoming nucleotide. Red, RNA; black, DNA. (B to E) Plots showing relative velocity of the indicated deoxyribonucleotide incorporation by Polθ on DNA/RNA and DNA/DNA primer-templates. Red, DNA/RNA; black, DNA/DNA. Data represent mean \pm SD; $n = 3$. (F) Schematic of DNA/RNA and DNA/DNA primer-templates. Underlined base codes for the correct

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To assess the relative fidelity of dNMP incorporation by Polθ on RNA versus DNA, we measured the relative velocity of nucleotide misincorporation. Remarkably, Polθ is significantly more accurate on RNA versus DNA as demonstrated by its severely limited ability to misincorporate nucleotides on RNA versus DNA (**Fig. 2, F to J**). For example, in **Fig. 2 (G and H)**, Polθ fully misincorporates dCMP and dAMP, respectively, on DNA in less than 2 min. Yet, on RNA, Polθ is unable to fully misincorporate dCMP or dAMP even after 20 min (**Fig. 2, G and H**). Polθ also fails to effectively misincorporate dTMP on RNA but efficiently misincorporates dTMP on DNA (**Fig. 2J**). Remarkably, Polθ rapidly misincorporates several consecutive dGMPs on DNA yet fails to fully misincorporate a single dGMP on RNA even after 20 min (**Fig. 2I**). Consecutive dGMP misincorporation events on DNA suggest that Polθ more easily misaligns the DNA/DNA primer-template (fig. S6A). The higher fidelity of Polθ on RNA versus DNA is also observed in a different sequence context (fig. S6, B to F). Hence, despite the enzyme's overall higher rate of correct dNMP incorporation on RNA, it exhibits a substantially slower rate of misincorporation on RNA. These data suggest that Polθ evolved to be more accurate on RNA similar to HIV RT (32).

Ternary structure of Polθ on a DNA/RNA primer-template

The higher fidelity of Polθ on RNA suggests that it binds the DNA/RNA and/or active site deoxyribonucleotide:ribonucleotide base pair in a distinct conformation. To investigate the molecular basis of Polθ RT activity, we solved a 3.2-Å crystal structure of Polθ on DNA/RNA with incoming 2',3'-dideoxyguanosine triphosphate (ddGTP) (**Fig. 3**). The construct used for x-ray crystallography (PolθΔL) was engineered to achieve higher *E. coli* expression by replacing five small disordered loops, which were not resolved in previous Polθ:DNA/DNA structures (2), with short glycine-serine inserts (**Fig. 3A** and fig. S7, A to C). The endogenous insert 2 loop promotes Polθ MMEJ of DNA/DNA with 3' ssDNA overhangs containing microhomology (9), and inserts 2 and 3 contribute to TLS (2, 7). WT (wild type) and PolθΔL exhibit similar DNA/DNA and DNA/RNA primer extension activities, demonstrating that the disordered loops do not substantially contribute to reverse transcription and canonical DNA/DNA extension (**Fig. 3B** and fig. S7D). In contrast to prior Polθ:DNA/DNA:ddNTP structures that were captured in the closed conformation



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Fig. 3 Ternary structure of Polθ on a DNA/RNA primer-template.

(A) Polθ polymerase. (B) DNA/RNA extension by Polθ and PolθΔL. (C) Structure of Polθ:DNA/RNA:ddGTP. (D) Superposition of Polθ:DNA/RNA (marine) and Polθ:DNA/DNA (orange, 4x0q). The fingers and thumb subdomains undergo reconfiguration. (E) Superposition of Polθ:DNA/RNA (marine) and Polθ:DNA/DNA (orange, 4x0q) highlighting a 12-Å shift of K2181 (blue box; thumb) and a 4.4-Å shift of E2246 (gray box; palm). (F) Superposition of nucleic acids and ddGTP from Polθ:DNA/RNA:ddGTP and Polθ:DNA/DNA:ddGTP structures. (G) Top: Electron density of ddGTP and 3' primer terminus in Polθ:DNA/RNA structure. Bottom: Zoomed-in image of the superposition of active sites, illustrating a different conformation of ddGTP in the Polθ:DNA/RNA (blue) and Polθ:DNA/DNA (salmon) complexes. (H) Interactions between ribose 2'-hydroxyl groups of the RNA template and residues in the Polθ:DNA/RNA structure. Red dashed lines, hydrogen bonds. (I) DNA/RNA used for cocrystallization with Polθ and ddGTP (top). Strong electron density is present for four base pairs [nucleotides located at positions 2 to 5 (underlined) of the DNA/RNA] and two base pairs resulting from an incorporated ddGMP (2',3' dideoxyguanosine monophosphate) (green; position 1) and a bound unincorporated ddGTP (red; position 0) in the active site (top). Interactions between Polθ and nucleic acids in Polθ:DNA/RNA:ddGTP (bottom). Interactions between residues and phosphate backbone, sugar oxygen, or nucleobase are shown in blue, yellow, and green, respectively. Hydrogen bonds between Polθ and ribose 2'-hydroxyl groups are indicated (boxed residues). (J) Interactions between Polθ and nucleic acids in Polθ:DNA/DNA:ddGTP (4x0q). Color scheme identical to (I).

Unexpectedly, the thumb subdomain undergoes a major reconfiguration. Fifty-seven percent of thumb subdomain residues refold from a helices to loops (Fig. 3D and fig. S7C, thumb domain), which may be necessary to accommodate the thicker A-form DNA/RNA. A loop shift in the palm subdomain involving E2246 is also observed on the opposite side of the DNA/RNA, suggesting a specific ribose 2'-hydroxyl interaction with its main-chain carbonyl likely mediated through a

mismatches) are observed near the upstream portion of the DNA/RNA (**Fig. 3I**). The incoming ddGTP and complementary cytidine on the RNA template also show a significant shift relative to the Polθ:DNA/DNA:ddGTP complex (**Fig. 3G**).

Polθ accommodation of DNA/RNA also involves many RNA template interactions (compare **Fig. 3, I and J**). Multiple hydrogen bonds with ribose 2'-hydroxyl groups along the RNA template are observed (**Fig. 3, H and I**). Polθ, HIV RT, and M-MuLV RT form a similar carbonyl hydrogen bond with the 2'-hydroxyl group of the active site template ribose, suggesting a conserved mechanism of active site RNA template binding (fig. S9, C and D) (**34, 35**). This interaction and additional Polθ:2'-hydroxyl ribose hydrogen bonds along the RNA template may suppress Polθ template misalignment errors and thus potentially contribute to its higher fidelity on RNA. Overall, the formation of multiple specific Polθ:RNA interactions combined with the major thumb subdomain reconfiguration and palm subdomain loop shift reveal how the polymerase becomes active on a DNA/RNA hybrid.

Polθ promotes RNA-templated DNA repair

To test whether Polθ promotes RNA-templated DNA repair synthesis (RNA-DNA repair) in a biological setting, we developed a cellular green fluorescent protein (GFP) reporter assay that can simultaneously quantitate Polθ MMEJ and RNA-DNA repair (**Fig. 4B**). Polθ is essential for MMEJ of DSBs, such as those caused by ionizing radiation (**1, 9–11**), and performs TLS (**7, 31**). Polθ therefore regularly uses aberrant template bases during its DNA repair activities. Polθ promotes MMEJ by using microhomology [≥ 2 base pairs (bp)] between 3' ssDNA overhangs generated by 5'-3' resection of DSBs and then extends the partially base-paired overhangs (**Fig. 4A**). The MMEJ GFP reporter assay is described as follows. Left and right DNA constructs respectively encoding the upstream and downstream portion of a GFP expression vector with 6 bp of overlapping sequence (microhomology) were conjugated with streptavidin at 5' DNA termini opposite from the microhomology tract to suppress 5'-3' exonuclease activity at these ends (**Fig. 4B**).

Introduction of 5 or 10 ribonucleoside monophosphates (NMPs) into the nontemplate transcription strand immediately upstream from the microhomology tract within the left construct enables analysis of RNA-templated DNA repair synthesis during MMEJ (**Fig. 4, B and C**). In the case of the left construct with five NMPs, base mutations within the transcription template strand directly opposite the RNA tract were added to engineer a stop codon in the GFP coding sequence (**Fig. 4C**, middle). Cotransfection of the right construct and left construct with or without NMP tracts into mouse induced pluripotent stem cells (iPSCs) activates GFP, demonstrating the capacity of MMEJ to use deoxyribose and ribose nucleobases as templates

containing 5 or 10 NMPs also results in significantly higher %GFP in *Polq*^{+/+} iPSCs, demonstrating that Polθ promotes RNA-DNA repair (**Fig. 4F**, middle and right). In the case of the left construct containing the stop codon opposite the 5-NMP tract (**Fig. 4C**, middle), RT activity is essential for generating a transcription template strand lacking the stop codon. Hence, these data confirm that Polθ promotes RNA-dependent DNA synthesis in cells.

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Fig. 4 Polθ promotes RNA-templated DNA repair synthesis in cells.

(**A**) Schematic of MMEJ. (**B**) Schematic of GFP MMEJ reporter assay. (**C**) Sequences of downstream end of left DNA GFP constructs with and without NMPs. (**D**) GFP FACS plots following no transfection (left) and cotransfection of the indicated right DNA constructs in *Polq*^{+/+} iPSCs. (**E**) Sequencing chromatograms showing MMEJ of the left and right constructs. Red, microhomology. (**F**) Bar plots showing normalized GFP cells following cotransfection of the indicated reporter constructs. Data pooled from three independent experiments performed at least in duplicate. ± SEM. *Statistical significance from paired *t* test: *P* = 0.04, 0 NMPs; *P* = 0.025, 5 NMPs; *P* = 0.04, 10 NMPs. (**G**) Δ7-GFP reporter integrated in U2OS cells for measuring CRISPR-Cas9 donor template [DNA (*phosphorothioate linkages) or DNA with two RNA bases (R2)]-mediated genome engineering. Ribonucleotides, rCrA; Green text, knock-in donor sequence. (**H**) Shown are frequencies of GFP⁺ cells (±SD) for the oligonucleotides shown, cotransfected with FLAG-POLQ, or empty vector (EV), normalized to transfection efficiency and to the DNA template (=1, left) and the R2 template in the parental cell line (=1, right). ***P* ≤ 0.0043, ****P* = 0.0001, and *****P* < 0.0001, one-way analysis of variance with Tukey's (left) and Dunnett's (right) posttests. (**I**) Immunoblot analysis of FLAG-POLQ and actin control. (**J**) Sequence of an amplification product from GFP⁺ cells isolated from *POLQ*^{e16m} cells with the Δ7 reporter assay using the R2 donor template and POLQ expression vector.

We further investigated Polθ RNA-DNA repair in cells using a previously published GFP knock-in reporter assay (Δ7-reporter) that is chromosomally integrated in human U2OS cells (**Fig. 4G**) (36). This assay has a GFP expression cassette disrupted by an insert sequence, along with a deletion of 7 bp of GFP sequence (Δ7). An oligonucleotide donor with these 7 nt flanked by 16-nt homology arms on each side can template restoration of active GFP⁺ (36). This event is induced via CRISPR-Cas9 to generate two DSBs that excise the insert. We compared the frequency of

line and to the mutant cells with expression of FLAG-POLQ (**Fig. 4I**). From either comparison, we found that Polθ promotes a significant increase in the frequency of GFP⁺ cells using the R2 donor template, which further confirms its ability to reverse transcribe template ribonucleotides during DNA repair in a biological setting (**Fig. 4H**, right).

DISCUSSION

Our study unexpectedly reveals that Polθ reverse transcribes RNA and undergoes a significant structural transformation to accommodate a DNA/RNA template. The structural transformation of Polθ's thumb subdomain is likely needed to maintain productive interactions on DNA/RNA, which adopts a significantly different conformation relative to DNA/DNA in the Polθ complex (**Fig. 3F** and fig. S11B). In contrast, structurally characterized retroviral RTs, such as HIV-RT, do not exhibit structural refolding of their thumb subdomain when acting on DNA/RNA (fig. S11A). The marked structural-functional switch within the thumb subdomain observed in Polθ has not been previously observed in other DNA polymerases or retroviral RTs and therefore may be unique to Polθ, which is an unusually promiscuous enzyme that is capable of acting on a variety of different templates including DNA/DNA, DNA/RNA, ssDNA, partial ssDNA, and single-stranded RNA (1, 5, 9, 37). Together, these structural studies reveal that Polθ has an extraordinary degree of structural plasticity that enables it to efficiently transcribe template ribonucleotides and accommodate a full RNA-DNA hybrid within its active site. Although future studies will be required to fully elucidate the physiological relevance of Polθ RT activity, our findings demonstrate that Polθ accommodates template ribonucleotides in an active configuration and promotes RNA-DNA repair, which may contribute to cellular tolerance of genome-embedded ribonucleotides.

MATERIALS AND METHODS

Primer-template extension assays

Relative velocity of RT activity (Fig. 1B). Polθ (0.5 nM), HIV RT, and Polη (catalytic core, residues 1 to 514) were incubated with 10 nM radiolabeled DNA/RNA template (RP559/RP493R) for the indicated times in buffer A [25 mM Tris-HCl (pH 7.8), 10 mM MgCl₂, 0.01% (v/v) NP-40, 1 mM dithiothreitol (DTT), bovine serum albumin (BSA; 0.1 mg/ml), and 10% (v/v) glycerol] with 100 μM dNTPs at 37°C. Percent extension was determined by dividing the intensity of the extended product by the intensity of the sum of extended and unextended products for each lane. All

Comparison of polymerase activities on DNA/DNA and DNA/RNA (Fig. 1, D to G). The primer extension assays on DNA/RNA (RP559/RP493R) and DNA/DNA (RP559/RP493D) templates were performed using the conditions in **Fig. 1B** with the following changes. The indicated concentrations of the indicated polymerases were used, and the reactions were performed for 32 min.

Relative RT activity (Fig. 1H). The indicated polymerases were incubated with 10 nM radiolabeled DNA/RNA template (SM98/SM44R) for 20 min in the presence of 10 μ M dNTPs at 37°C. Polθ and HIV RT reactions were performed in 25 mM Tris-HCl (pH 8.0), 10 mM KCl, 10 mM MgCl₂, 0.01% (v/v) NP-40, 1 mM DTT, BSA (0.1 mg/ml), and 10% (v/v) glycerol. AMV RT (20 units, New England Biolabs)-containing reactions were performed in buffer [50 mM Tris-acetate (pH 8.3), 75 mM potassium acetate, 8 mM magnesium acetate, and 10 mM DTT] and contained BSA (0.1 mg/ml). M-MuLV RT (400 units, New England Biolabs)-containing reactions were performed in buffer [50 mM Tris-HCl (pH 8.3), 75 mM KCl, 3 mM MgCl₂, and 10 mM DTT] and contained BSA (0.01 mg/ml).

Comparison of RT and DNA-dependent DNA synthesis activities by truncated and full-length Polθ (Fig. 1J). One hundred nanomolar of the indicated polymerases were incubated with 10 nM radiolabeled DNA/RNA (SM98/SM44R) or DNA/DNA (SM98/SM44) for 45 min with 50 μ M dNTPs and 25 mM Tris-HCl (pH 7.8), 2 mM MgCl₂, 4 mM KCl, 6 mM NaCl, 0.01% (v/v) NP-40, 1 mM DTT, BSA (0.1 mg/ml), 10% (v/v) glycerol, and 750 μ M adenosine triphosphate (ATP) at 37°C.

Relative velocity of single dNMP incorporation on radiolabeled DNA/RNA (Fig. 2, A to E). Polθ (2 nM) was incubated with 100 nM of the indicated radiolabeled DNA/RNA and DNA/DNA templates for the indicated times with 300 μ M of the indicated dNTP in buffer [25 mM Tris-HCl (pH 8.0), 10 mM MgCl₂, 4 mM KCl, 6 mM NaCl, 0.01% NP-40, 1 mM DTT, BSA (0.01 mg/ml), and 10% (v/v) glycerol] at 37°C. Percent extension was determined as described above.

Relative velocity of nucleotide (dNMP) misincorporation (Fig. 2, F to J). Polθ (20 nM) was incubated with 100 nM of the indicated radiolabeled DNA/RNA and DNA/DNA templates and 300 μ M of the indicated dNTP for the indicated times in buffer [25 mM Tris-HCl (pH 8.0), 10 mM MgCl₂, 10 mM KCl, 0.01% (v/v) NP-40, 1 mM DTT, BSA (0.1 mg/ml), and 10% (v/v) glycerol]. Percent extension was determined as described above. All oligonucleotides were radiolabeled using T4 polynucleotide kinase (New England Biolabs) and ³²P- γ -ATP (PerkinElmer) in recommended buffer for 37°C for at least 1 hour.

positive control to measure transfection efficiency, a wild-type linear DNA GFP expression construct was transfected simultaneously. GFP-positive cell frequencies were measured 3 days after transfection by flow cytometry using GUAVA easyCyte 5-HT (Luminex Corp.) in independent replicates and corrected for transfection efficiency and background events. Data are represented as the mean and SEM of three independent experiments, with at least duplicates per experiment. Statistical analysis was carried out by paired *t* test.

Preparation of GFP MMEJ reporter constructs

Some of the methods used here are similar to those in a previously published article (38). Polymerase chain reaction (PCR) preparation followed recommended conditions for the Phusion High-Fidelity DNA Polymerase (New England Biolabs M0530) using 10 ng of pCMV-GFP plasmid as template in 1× Phusion HF Buffer. PCR for the left-flank DNA was performed with primers RP500B and RP501. PCR for the right-flank DNA was performed with primers RP502 and RP503B. Following PCR, left- or right-flank DNA products were pooled together and digested with Dpn I (New England Biolabs) in 1× CutSmart buffer and then purified via Qiagen QIAquick PCR Purification Kit. PCR was then conjugated to streptavidin using PCR (110 ng/μl) and streptavidin (0.8 μg/μl) in 10 mM tris-HCl (pH 7.5) and 100 mM NaCl at 37°C for 1 hour. PCR for the left-flank DNA with five consecutive ribonucleotides was performed with primers RP500B and RP501. PCR was purified and then digested with Dpn I (New England Biolabs) and Sap I in 1× CutSmart buffer. PCR was purified again and then ligated to double-stranded DNA composed of annealed oligonucleotides RP550R-P and RP530a [oligos were annealed in the presence of a ribonuclease (RNase) inhibitor] for 16 hours at 16°C with T4 DNA Ligase (New England Biolabs) in 1× T4 DNA Ligase Buffer. Ligated PCR was purified and then conjugated to streptavidin as described above. PCR for the left-flank DNA with 10 consecutive ribonucleotides was prepared by the same methodology with ligation to the double-stranded DNA composed of oligos RP546P and RP530. Streptavidin conjugation and DNA amplification steps were confirmed in agarose gels stained with ethidium bromide.

CRISPR-Cas9 knock-in GFP reporter assay

The U2OS parental and *POLQ^{e16m}* cell lines with the Δ7-reporter and CAS9/single guide RNA (sgRNA) plasmids to target the DSBs in this reporter, DNA oligonucleotide template, and control oligonucleotide (LUC) were previously described (36). The R2 oligonucleotide has the same sequence as the DNA oligonucleotides, but with two RNA bases in the 7 nt missing from the Δ7-reporter (IDT). The cell lines were seeded at 1×10^5 on a 12-well dish and transfected the following day, and % GFP was analyzed 3 days after transfection using a CyAN-ADP (DAKO)

were performed with 4 μ l of Lipofectamine 2000 (Thermo Fisher Scientific) in 0.2 ml of Optimem (Thermo Fisher Scientific) and incubated with cells in 1 ml of antibiotic-free medium for 4 hours. Immunoblotting analysis for FLAG-POLQ involved extraction with ELB [250 mM NaCl, 5 mM EDTA, 50 mM Hepes, 0.1% (v/v) Ipegal, and Roche protease inhibitor] with sonication (Qsonica, Q800R) and using antibodies for FLAG (Sigma-Aldrich, A8592) or ACTIN (Sigma-Aldrich, A2066). Sequence analysis of reporter assay with R2 oligo: Cells were transfected as for the reporter assay with the R2 oligonucleotide template in the *POLQ^{et16m}* cells with the POLQ expression vector, GFP⁺ cells were isolated by fluorescence-activated cell sorting (FACS) (Becton Dickinson Aria Sorter), and the GFP repair product was amplified with CMVFWDFRT5 5' CGCAAATGGGCGGTAGGCGTG and BGHREVFRT5 5' TAGAAGGCACAGTCGAGG and sequenced with the CMVFWDFRT5 primer.

cDNA synthesis

cDNA synthesis reactions were performed by the indicated polymerase in the presence of 100 μ M dNTPs and optimal buffer for each enzyme: Polθ [25 mM tris-HCl (pH 8.0), 10 mM KCl, 10 mM MgCl₂, 0.01% (v/v) NP-40, 1 mM DTT, BSA (0.1 mg/ml), and 10% (v/v) glycerol]; AMV RT [50 mM tris-acetate (pH 8.3), 75 mM potassium acetate, 8 mM magnesium acetate, 10 mM DTT, and BSA (0.1 mg/ml)]; and M-MuLV RT [50 mM tris-HCl (pH 8.3), 75 mM KCl, 3 mM MgCl₂, and 10 mM DTT]. Reactions with synthetic DNA/RNA contained 10 ng of template. Polθ (100 nM), AMV-RT (20 units), or M-MuLV RT (400 units) were incubated at 37°C (Polθ) or 42°C (AMV-RT and M-MuLV RT) for 1 hour. Enzymes were then heat-inactivated at 85°C for 20 min. cDNA (0.1875 ng) was used for real-time PCR (Power SYBR Green Master Mix, Thermo Fisher Scientific) using primers SM246 and SM247.

Proteins

Polθ, Fl-Polθ, and Polδ were purified as previously described (1, 9). Pols β and λ were provided by S. Wilson. Polκ was purchased from Enzymax LLC. HIV RT RT52A optimized for x-ray crystallography was provided by Dr. E. Arnold. Polη was provided by Dr. S. Arora. Pols ε and α were provided by Dr. M. O'Donnell. Poly^{Exo-} was provided by Dr. W. Copeland. M-MuLV, AMV RTs, and KF Pols were purchased from New England Biolabs.

Protein purification for x-ray crystallography

The gene encoding Polθ (residues 1819 to 2590) was codon-optimized and cloned into the pSUMO vector to generate a sumo fusion that carries an N-terminal 6×His tag and a PreScission

resuspended in buffer L [50 mM Hepes (pH 8.0), 500 mM NaCl, 0.005% (v/v) Igepal CA 630, and 0.5 mM TCEP]; lysed by sonication or French Press in the presence of DNase I (20 µg/ml), RNase A (30 µg/ml), 5 mM MgCl₂, 2 mM CaCl₂, and 1 mM phenylmethylsulfonyl fluoride; and then centrifuged at 12,000g for 45 min. 6×His sumo fusion was captured by Ni-NTA agarose gravity-flow chromatography followed by a series of washes by buffer W [50 mM Hepes (pH 8.0), 500 mM NaCl, 0.005% (v/v) Igepal CA 630, 0.5 mM TCEP, and 10 mM imidazole]. Five milliliters of buffer L and Precision Protease was added to cleave Polθ from the 6×His tag by staying overnight at 4°C. The cleaved Polθ was eluted another two times by 5 ml of buffer L. The eluted protein was purified to homogeneity using a HiTrap Heparin column (GE Healthcare Life Sciences). The protein was concentrated to 5 mg/ml in a buffer of ammonium acetate (150 mM), KCl (150 mM), tris-HCl buffer (pH 8.0) (40 mM), TCEP (2.5 mM), and glycerol (1% v/v).

Crystallization and structure determination

The crystallization condition was identified by wide matrix screening. Sitting-drop crystallization screening plates were set at 18°C using ARI Crystal Gryphon Robot (ARI) and crystallization screening solutions (Qiagen and Hampton Research). Reacting Polθ (2.5 mg/ml) with a DNA/RNA hybrid (DNA primer: 5'-GCGGCTGTCATT and RNA template: 5'-CGUCCAAUGACAGCCGC) in the presence of ddGTP (1 mM), sucrose monolaurate (300 µM), MgCl₂ (1 mM), and spermine tetrahydrochloride (20 mM) prepared the sample for sitting drop vapor diffusion over a 50-µl reservoir containing 20% (w/v) ethanol by mixing 0.3 µl of the reservoir solution with an equal part of the reaction solution. Crystals of approximate final dimensions 20 µm by 20 µm by 20 µm grew over the next 2 weeks. Cryoprotection was achieved by looping the crystals into mother solution with additional 25% (v/v) glycerol before flash cooling into liquid nitrogen. Diffraction data were collected at beamline 23ID-D of the National Institute of General Medical Sciences and National Cancer Institute Structural Biology Facility at the Advanced Photon Source. A complete dataset for Polθ was collected, indexed, integrated, and scaled by the autoprocessing package in APS server (GMCAProc). The Phaser-MR program in the PHENIX package was used for molecular replacement, and COOT was used for model rebuilding and Phenix for simulated annealing and refinement. The structure of Polθ was determined by molecular replacement (MR) using Polθ [Protein Data Bank (PDB): 4x0q] as a search model. The full PDB structure of 4x0q could not yield a solution. An MR solution was only obtained by removing some of the loops and deleting the DNA/DNA duplex from the model. The initial phases of the MR protein model were improved by cyclic model building and refinement, which allow us to slowly build in the missing loops, refolded and reoriented finger and thumb domains, and the

polarity):

RP559 CGACGTCGCAG

RP493R rUrUrUrUrUrUrCrGrCrGrCrUrGrCrGrArCrGrUrCrG

RP493R TTTTTTTCGCGCTGCGACGTCG

RP500B/5Biosg/GCTAGCCAGTCAGTGGGCCCCGC

RP501 TCGGGCATGGCGGACTTGAAGAAGTCG

RP502 GCCCGAAGGCTACGTCCAGGAGCGCACC

RP503B/5Biosg/TGATTACGCCAAGTTAATTAAGGACGTCCTCCTGCTGG

RP550R-P/5Phos/TCTTCAAGTCCGCrCrArUrGrCCCCGA

RP530a TCGGGCTAGGCGGACTTGA

RP546P/5Phos/TCTTrCrArArGrUrCrCrGrCrCATGCCCGA

RP530 TCGGGCATGGCGGACTTGA

RP563 CCACAAGTTCAGCGTGTCCGGCGAGG

RP566 CCTTGATGCCGTTCTTCTGCTTGTGGCC

RP564 CGAGGGCGATGCCACCTACGGC

RP565 GTGGCTGTTGTAGTTGTACTCCAGCTTGTGCC

RP560 AGAACCTGTTGAA

RP468R rArCrArGrUrUrUrUrUrUrUrGrCrUrUrUrUrGrUrUrCrArArCrArGrGrUrUrCrU

SM98 ATTATGCGGCCGTGTACAACGGGTGTCGGGGC

SM44R GCUCUGAUGCCGCAUAGUUAAGCCAGCCCCGACACCCG

SM44 GCTCTGATGCCGCATAGTTAAGCCAGCCCCGACACCCG

SM248 AATTGAAGAGTTTGATCATGGC

SM270FAM/56-FAM/CGACGTCGCAGCGC

SM271 GAGACTCGGCGCTGCGACGTCG

SM271R GAGACUCGGCGCUGCGACGUCG

SM106 ATTATGCGGCCGTGTAC

SM107 GCTCTGATGCCGCATAG

RP561R2 G*G*TAGCGGCTGAAGCACTGrCrACGCCGTAGGTCAGGGT*G*G.

RP561 G*G*TAGCGGCTGAAGCACTGCACGCCGTAGGTCAGGGT*G*G.

SUPPLEMENTARY MATERIALS

Supplementary material for this article is available at <http://advances.sciencemag.org/cgi/content/full/7/24/eabf1771/DC1>

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