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Evidence for interplay among yeast replicative DNA polymerases alpha, delta and epsilon from studies of exonuclease and polymerase active site mutations

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Abstract

Background: DNA polymerase ϵ (Pol ϵ) is essential for S-phase replication, DNA damage repair and checkpoint control in yeast. A *pol2-Y831A* mutation leading to a tyrosine to alanine change in the Pol ϵ active site does not cause growth defects and confers a mutator phenotype that is normally subtle but strong in a mismatch repair-deficient strain. Here we investigate the mechanism responsible for the mutator effect.

Results: Purified four-subunit Y831A Pol ϵ turns over more deoxynucleoside triphosphates to deoxynucleoside monophosphates than does wild-type Pol ϵ , suggesting altered coordination between the polymerase and exonuclease active sites. The *pol2-Y831A* mutation suppresses the mutator effect of the *pol2-4* mutation in the exonuclease active site that abolishes proofreading by Pol ϵ , as measured in haploid strain with the *pol2-Y831A,4* double mutation. Analysis of mutation rates in diploid strains reveals that the *pol2-Y831A* allele is recessive to *pol2-4*. In addition, the mutation rates of strains with the *pol2-4* mutation in combination with active site mutator mutations in Pol δ and Pol α suggest that Pol ϵ may proofread certain errors made by Pol α and Pol δ during replication *in vivo*.

Conclusions: Our data suggest that Y831A replacement in Pol ϵ reduces replication fidelity and its participation in chromosomal replication, but without eliminating an additional function that is essential for viability. This suggests that other polymerases can substitute for certain functions of polymerase ϵ .

Background

Multiple DNA polymerases are thought to be present at the eukaryotic replication fork [1-4]. Some of their functions could be unique while others could be overlapping. Different polymerases may compete for certain DNA sub-

strates and several polymerases may sometimes act in concert [5-9]. Under normal circumstances, chromosomal replication requires at least three DNA polymerases, Pol α , Pol ϵ and Pol δ . All of these polymerases are multi-subunit complexes [1,4] and all subunits are required for their

proper function (see recent papers [10-12], and references therein). Pol α is not very processive and lacks an intrinsic proofreading exonuclease. It has a tightly associated activity for the synthesis of RNA primers at replication origins and on the lagging DNA strand. Pol α extends these RNA primers by synthesizing short stretches of DNA, and then a switch occurs to processive synthesis by Pol ϵ and/or Pol δ .

The exact roles of Pol ϵ and Pol δ in replication are not yet fully understood. Among several possible models, it has been proposed that Pol ϵ is primarily responsible for copying the leading strand DNA template and Pol δ is responsible for lagging strand replication [13]. Another model has proposed the opposite [14]. Either model is consistent with the fact that Pol ϵ and Pol δ both possess intrinsic 3' to 5' exonuclease activity, and with genetic data in yeast suggesting that these nucleases proofread replication errors on opposite DNA strands during chromosomal [15] or plasmid [16] DNA replication. However, the replication functions of Pol δ and Pol ϵ are not equivalent [17]. When proofreading or base selectivity is impaired by homologous active site point mutations in *POL3* (encoding Pol δ) and *POL2* (encoding Pol ϵ), the mutator effects are much stronger for *pol3* mutants than for *pol2* mutants [7,18,19]. A yeast strain with an amino-terminal deletion of the polymerase domain of the *POL2* gene but retaining the carboxyl-terminal domain grows slowly but is nonetheless viable [20,21], and references therein), indicating that another polymerase can substitute for the polymerization function of Pol ϵ . Thus, it is possible that Pol δ may perform the bulk of chain elongation during chromosomal replication [17], while Pol ϵ serves more specialized roles. One possibility is a role in the S phase checkpoint control when replication fork progression is impeded [22], perhaps by sensing single-stranded DNA [23]. Pol ϵ also interacts with Dpb11 and, thus, may function during initiation of DNA replication at origins [24,25]. It has also been suggested that Pol ϵ may participate in replication during late, but not early, S phase [26], as well as in the establishment of sister chromatid cohesion [27].

The fidelity of nuclear DNA replication in eukaryotic cells relies on three steps that are thought to operate in series: the base selectivity of DNA polymerases that operate at the fork, proofreading by the exonucleases of DNA polymerases ϵ and δ or possibly by extrinsic exonucleases, and DNA mismatch repair. This is supported genetically by the fact that the double mutants that are deficient in proofreading by either DNA polymerase and mismatch repair are hypermutable, suggesting a sequential action of proofreading and mismatch repair [19,28]. As one attempt to understand the complex enzymology that determines fidelity and influences eukaryotic genome stability, we recently described the effects on viability and

mutagenesis in *Saccharomyces cerevisiae* resulting from replacing a conserved tyrosine in the active sites of three replicative polymerases with alanine [7]. Important for our understanding of the role of Pol ϵ , a haploid yeast strain with this *pol2-Y831A* mutation grew normally, suggesting no major defects in replication. However, we observed a modest, spontaneous base substitution mutator effect and a strong, spontaneous frameshift mutator effect when DNA mismatch repair was nonfunctional. These mutator effects suggested a defect in replication fidelity at a step preceding mismatch repair (for example, reduced polymerase base selectivity and/or proofreading) [7]. The *pol2-Y831A* mutation was semi-dominant in heterozygous diploid cells, suggesting that the mutator effects reflected an aberrant function of the altered Pol ϵ that is not masked by the presence of wild-type Pol ϵ . Here we further investigate the mutator effects conferred by the *pol2-Y831A* mutation using biochemical and genetic approaches. The results suggest decreased participation of Y831A Pol ϵ in chromosomal replication. We also investigate the possibility that wild-type Pol ϵ proofreads replication errors generated by altered variants of DNA polymerases α and δ .

Results

Biochemical characterization of Y831A Pol ϵ

We purified the four-subunit DNA polymerase ϵ from a yeast strain with a chromosomal *pol2-Y831A* mutation (Figure 1). The quality of the preparation, in terms of purity and specific activity, was comparable to the wild-type polymerase ϵ that we purified previously by the same method; exonuclease activity co-eluted with polymerase activity [23,29]. We compared the polymerase and exonuclease activity of the current preparation with DNA polymerase ϵ purified earlier from a wild-type strain. We first compared turnover of deoxythymidine triphosphate (dTTP) to deoxythymidine monophosphate (dTMP) during DNA synthesis by wild-type and mutant polymerases on poly(dA) template by thin-layer chromatography (TLC) (Figure 2A,2B). This method enables simultaneous measurement and comparison of exonuclease and DNA polymerase activity. dTMP turnover rates, $((\text{pmols dTMP excised})/(\text{pmols dTMP incorporated} + \text{excised}) \times 100)$, for wild-type and the mutant polymerases were 22% and 50%, respectively (Figure 2C,2D). The value for the mutant Pol ϵ was unchanged, even after further purification of the enzyme preparation using a gel-filtration column chromatography, suggesting that the high turnover number feature is intrinsic to the mutant Pol ϵ . In further experiments, when the exonuclease activity was assayed separately with single-stranded DNA as a substrate (see Methods), both enzyme preparations had the same ratio of 3' to 5' exonuclease activity to polymerase activity, 3.7 for wild-type and 3.5 for Pol ϵ Y831A. This suggests that the increased rate of dTMP turnover by the mutant

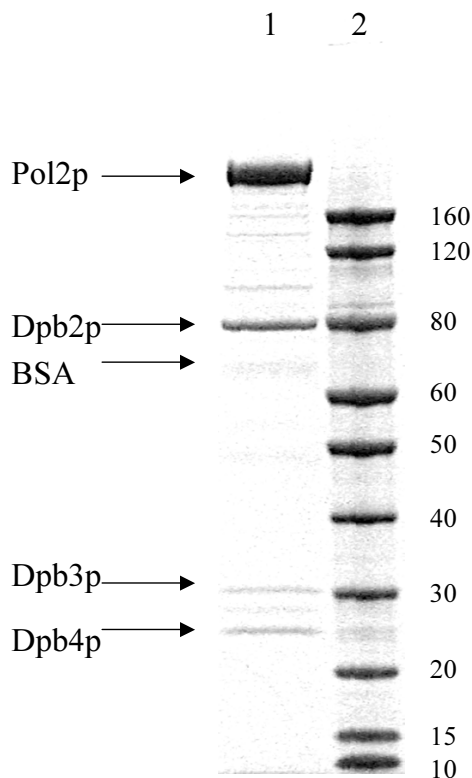


Figure 1

Purified four-subunit Y831A Pol ϵ . SDS-PAGE analysis of peak fraction from final SMART MonoS column (see Methods). 1 μ l of collected fraction 15 was mixed with Invitrogen loading buffer and run on denaturing 4–12% NUPAGE Bis-Tris polyacrylamide gel for 50 minutes at 200 V in MOPS buffer. The gel was stained with SimpleStain colloidal blue stain as recommended by the vendor (Invitrogen). Lane 1, Pol ϵ Y831A; lane 2, the Benchmark His-tagged protein standards (Invitrogen).

polymerase is not due to an alteration in the catalytic efficiencies of polymerase and proofreading exonuclease activities, but rather that the tyrosine to alanine substitution in the polymerase active site alters communication between the polymerase and exonuclease active sites.

Mutation rates in haploids with double *pol2*-Y831,4 mutation

Strains with the *pol2*-Y831A mutation are mutators in a mismatch repair defective background [7], suggesting that they are defective in controlling replication fidelity at a step prior to mismatch repair (MMR). If this defect is due to an altered base selectivity of Y831A Pol ϵ , the resulting

mismatches could be proofread by the intrinsic 3' exonuclease activity of Pol ϵ . In this case, mutational inactivation of the exonuclease should elevate mutation rates. Such a synergistic increase in mutation rate was previously observed in a strain with *pol2-4, C1089* double mutation that inactivates proofreading and alters the polymerase domain [30]. To test this possibility with the *pol2*-Y831A mutant, we constructed haploid yeast strains with either single or double mutations in the *POL2* gene, one in the polymerase active site (*pol2*-Y831A) and another in the exonuclease active site (*pol2-4*). These are derivatives of the previously developed strain 8C-YUNI101, which allows concomitant measurement of mutation rates at

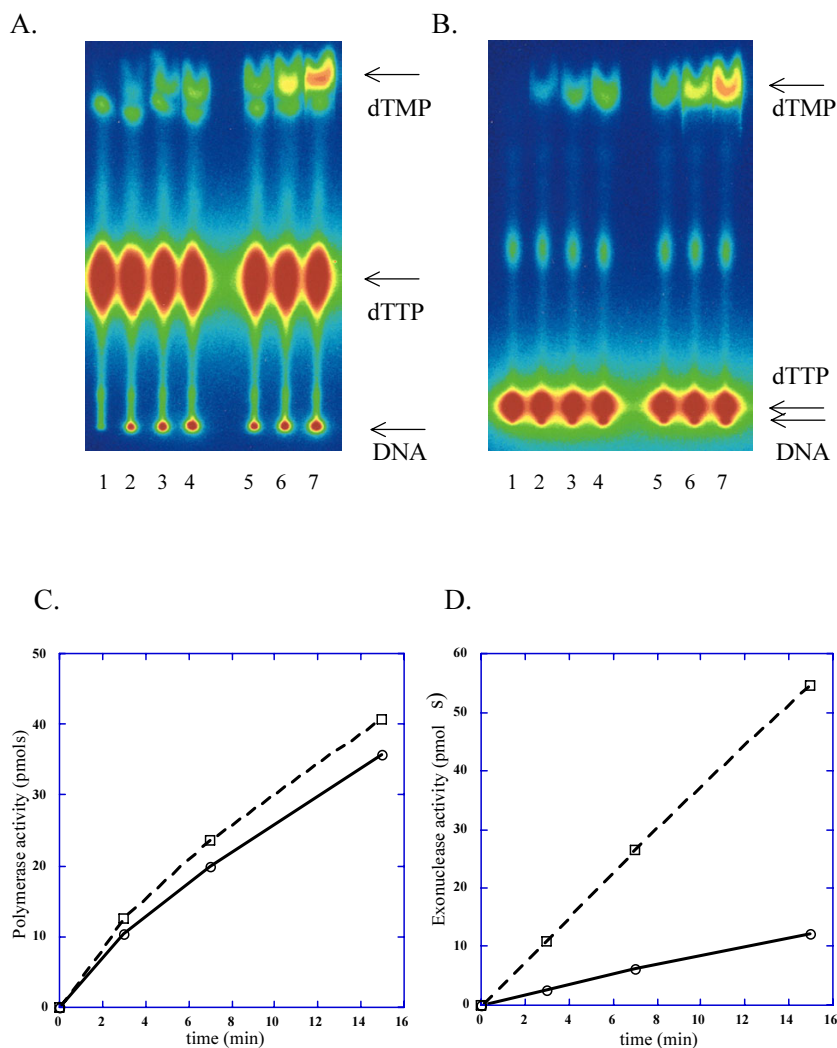


Figure 2

Elevated nucleotide turnover by Y831A Pol ε. DNA synthesis and dTTP turnover by Pol ε was measured on poly(dA)/oligo(dT) substrate. Reactions were performed as described in Methods, using 0.13 U of each enzyme for 20 μl reactions. **(A)** Analysis of polymerase reaction by TLC in 1 M LiCl running buffer. Lanes 1, 2, 3, 4: wild-type Pol ε at 0, 3, 7 and 15 minutes of reaction, respectively. Lanes 5, 6, 7: reactions with Y831A Pol ε at 3, 7 and 15 minutes, respectively. Positions of unincorporated label, label in DNA and dTMP are shown by arrows. **(B)** Analysis of polymerase reaction by TLC in 0.4 M LiCl running buffer. Lane assignment is the same as in (A). **(C)** Plot of time-course of DNA synthesis and dTTP turnover by wild-type Pol ε. Open circles connected by solid line represent dTTP retained into DNA; open rectangles connected by a dashed line represent excised dTTP. **(D)** Plot of time-course of DNA synthesis and dTTP turnover by Y831A Pol ε. Symbols are the same as in (C).

Table 1: Effect of combining the exonuclease and active site mutations in Pol ε gene on mutation rates in haploids in 8C-YUN1101 genetic background

POL2 allele	Mutation rates for different markers*		
	Forward mutation	Reversion	
		Base substitutions	Frameshifts (+1)
	Can ^r (× 10 ⁻⁷)	Ura ⁺ × (10 ⁻⁸)	His ⁺ × (10 ⁻⁸)
Mismatch repair proficient strains			
wild-type	3.3 2.5–4.9	1.3 1.1–1.7	1.1 0.9–3.5
<i>pol2-Y83 1A</i>	5.1 3.5–12.1	1.2 0.8–2.8	1.6 0.5–2.7
<i>pol2-4</i>	27 15–42	3.4 ¹ 2.4–5.7	45 ² 35–78
<i>pol2-4,Y83 1A</i>	9 7–15	1.5 1.1–1.9	3.9 ³ 2.3–5.4
Mismatch repair deficient strains (<i>pms1</i>)			
wild-type	90 72–150	2.8 2.4–4.1	87 54–150
<i>pol2-Y83 1A</i>	340 230–420	12 ⁴ 11–19	470 ⁵ 270–790
<i>pol2-4</i>	760 120–1060	160 ⁶ 90–420	1600 ⁷ 950–2800
<i>pol2-4,Y83 1A</i>	270 190–440	15 ⁸ 8–55	410 ⁹ 260–440

*Here and thereafter, median for 9–27 independent cultures with 95% confidence limits below. ¹14/16 of revertants sequenced were due to C->A transversions ²20/20 revertants sequenced contained a +1A frameshift event ³16/16 revertants sequenced were due to +1A frameshift ⁴12/14 revertants were due to C->A change ⁵15/15 revertants sequenced were due to +1A frameshift ⁶12/12 revertants sequenced were due to C->A mutation ⁷16/16 revertants sequenced were due to +1A frameshift ⁸12/15 revertants sequenced were due to C->A mutation ⁹14/14 revertants were due to +1A frameshift

Table 2: Effect of combining the exonuclease and active site mutations in DNA polymerase ε on mutation rates in diploids. Mismatch repair proficient strains

Relevant mutation	His ⁺ reversion rate (× 10 ⁻⁸)	Fold increase
+/+	0.5 0.3–1.2	1
+/ <i>pol2-Y83 1A</i>	1.1 0.7–1.7	2.2
<i>pol2-Y83 1A</i> / <i>pol2-Y83 1A</i>	2.2 1.8–5.7	4.4
+/ <i>pol2-4</i>	3.8 2.6–4.7	7.6
<i>pol2-4/pol2-4</i>	26 16–45	52
<i>pol2-4/pol2-Y83 1A</i>	28 16–53	56
<i>pol2-4/pol2-4,Y83 1A</i>	21 18–34	42
<i>pol2-Y83 1A/pol2-4,Y83 1A</i>	1.8 1.2–3.1	3.6
<i>pol2-4,Y83 1A/pol2-4,Y83 1A</i>	1.8 1.1–2.3	3.6

Table 3: Effect of combining the exonuclease and active site mutations in DNA polymerase ϵ on mutation rates in diploids. Mismatch repair deficient strains (*pms1/pms1*)

Relevant mutation	His ⁺ reversion rate ($\times 10^{-8}$)	Fold increase
+/+	110 90–140	1
+/ <i>pol2-Y831A</i>	400 270–600	4
<i>pol2-Y831A/pol2-Y831A</i>	820 750–950	7
+/ <i>pol2-4</i>	1,500 930–3600	14
<i>pol2-4/pol2-4</i>	9,400 5,900–14,000	85
<i>pol2-4/pol2-Y831A</i>	8,400 3,500–17,000	76
+/ <i>pol2-4,Y831A</i>	210 180–270	2
<i>pol2-4/pol2-4,Y831A</i>	11,000 8,100–22,000	100
<i>pol2-Y831A/pol2-4,Y831A</i>	1,200 1,030–1,460	11
<i>pol2-4,Y831A/pol2-4,Y831A</i>	1,020 840–2,300	9

several loci. These include the forward mutation rate at the *CAN1* locus, where mutations reflect a variety of substitution, frameshift and more complex events; the rate of reversion of the *ura3-29* missense mutation which reverts via three specific base pair substitutions at a single G→C base pair in a TCT codon encoding for serine [7]; and reversion of the *his7-2* mutant allele, which occurs mainly via + 1 frameshifts in a homopolymeric A→T run [7,31]. In a MMR-proficient background (Table 1, top), the *pol2-Y831A* mutation results in a slight or no mutator effect, while loss of proofreading by Pol ϵ elevates the mutation rate at *CAN1*, *ura3-29* and *his7-2* by eight-fold, 2.6-fold and 41-fold, respectively. In an MMR-deficient background (Table 1, bottom), the *pol2-Y831A* mutation and the *pol2-4* mutation each elevates the mutation rate at all three loci. Surprisingly, when both mutations are present in the same gene, the *pol2-Y831A* mutation strongly suppresses the mutator effects of the *pol2-4* mutation for all three markers tested in both MMR-proficient and MMR-deficient strains. Thus, despite generating a mutator effect itself, *pol2-Y831A* actually suppresses mutations resulting from loss of proofreading by Pol ϵ . This suppression suggests that Y831A Pol ϵ may not participate in replication to the same extent as does wild-type Pol ϵ .

To further test this hypothesis, we measured His⁺ reversion rates in a series of diploid strains. In MMR-proficient diploid strains (Table 2), the *pol2-Y831A* allele appeared to be semi-dominant, since the mutator effect of the het-

erozygote was half that of the homozygote. This may be due to either effective competition of error-prone mutant polymerase with the wild-type enzyme or due to perturbations in DNA synthesis caused by Y831A Pol ϵ . The heterozygous +/*pol2-4* and homozygous *pol2-4/pol2-4* diploid strains had a reversion rate elevated by 7.6-fold and 52-fold, respectively, suggesting that wild-type Pol ϵ may efficiently compete with the exonuclease-deficient enzyme and proofread replication errors. When one chromosome possessed the *pol2-4* allele while the other chromosome had the either *pol2-Y831A* alone or *pol2-4,Y831A* double mutation, the reversion rate was similar to the homozygous *pol2-4/pol2-4* diploid strain. However, homozygosity for *pol2-Y831A* suppressed the mutator effect of the *pol2-4* mutation, slightly in the +/*pol2-4* diploid (two-fold, Table 2, penultimate line) and strongly in the *pol2-4/pol2-4* diploid strain (14-fold, Table 2, last line). Similar effects were obtained in diploid strains in an MMR-deficient background (Table 3). These results are consistent with the results in the haploid strains, and further support the hypothesis that Y831A Pol ϵ does not compete with wild-type exonuclease-deficient Pol ϵ .

Lack of genetic interaction between *pol2-Y831A* and *pol3-01* mutation in exonuclease domain of Pol δ

The idea of altered communication between the polymerase and exonuclease active sites (see Figure 2) and the putative inability of Y831A Pol ϵ to compete with exonuclease-deficient Pol ϵ (Tables 1, 2 and 3) suggest that

Table 4: Effect of combining the mutation in the exonuclease domain of the Pol δ gene with the active site mutation in Pol ϵ genes on mutation rates in mismatch repair-proficient derivatives of 8C-YUN1101 strain

Relevant genotype	Mutation rates for different markers		
	Forward mutation	Reversion	
	Can ^r \times (10 ⁻⁷)	Base substitutions Ura ⁺ (\times 10 ⁻⁸)	Frameshifts (+1) His ⁺ (\times 10 ⁻⁸)
wild-type	2.9	1.5	1.5
	1.6–4.9	1.1–2.3	1.2–2.1
+/ <i>pol3-01</i> *	28	n/a	33
	22–28		31–39
+/ <i>pol3-01</i> * <i>pol2-Y831A</i>	31	n/a	27
	25–39		24–34
<i>pol3-01</i>	190	19	110
	140–260	6–35	80–160
<i>pol3-01 pol2-Y831A</i>	130	14	34
	90–210	13–28	18–43

n/a, non applicable, since these strains are Ura⁺. *Haploids with duplication of the *POL3* gene.

Y831A Pol ϵ may not edit its errors using its intrinsic proofreading exonuclease. In this case, a different exonuclease-proficient DNA polymerase may proofread errors by Y831A Pol ϵ . The precedent for this hypothesis comes from both *in vitro* [32] and *in vivo* [6] experiments indicating that the exonuclease activity of Pol δ can proofread errors by other replicative polymerases. To test if there is an effect of proofreading by Pol δ on the mutation rate in the *pol2-Y831A* mutant strain, we constructed strains with a combination of the *pol2-Y831A* mutation and the *pol3-01* mutation that inactivates proofreading by Pol δ . If the exonuclease activity of Pol δ is involved, a synergistic increase in mutation rate is expected in the double mutant. In a haploid strain with duplication of the *POL3* gene, encoding a catalytic subunit of Pol δ , when one allele is the *pol3-01* and another allele is the wild-type, there is 9.6-fold increase in Can^r forward mutation rate and a 22-fold increase in frameshift mutation rate (Table 4). A haploid strain with the *pol3-01* mutation showed a 72-fold increase in the Can^r mutation rate, a 15-fold increase in the rate of *ura3-29* reversion and a 73-fold increase in frameshift mutation rate. These results are consistent with the published data that the *pol3-01* is a strong mutator allele that is semi-dominant in respect to the wild-type allele [6,19]. The mutation rates at these loci are no higher in the same strains containing additional *pol2-Y831A* mutation (Table 4). According to our previous data, the synergistic interactions are readily detected with the *pol2-Y831A* allele, which is a weak mutator, as exemplified by an increase of mutation rates in the *pol2-Y831A pms1* double mutants [7]. We conclude that errors made by Y831A Pol ϵ are not proofread by Pol δ .

Genetic interaction between active site mutations in genes encoding Pol α and Pol δ and proofreading defective mutations in *POL2* and *POL3*

Although errors made by Y831A Pol ϵ do not appear to be proofread by Pol δ , it is possible that Pol ϵ might proofread errors by other polymerases. To investigate this and to determine whether the effects of combining exonuclease-defective alleles of *POL2* (Table 2) and *POL3* (Table 4) with a polymerase active site defect were specific to Y831A Pol ϵ , we looked for genetic interaction between proofreading and mutator alleles of the *POL1* (encoding Pol α) and the *POL3* (encoding Pol δ). For this we chose the *pol1-Y869A* and *pol3-Y708A* mutations that lead to conserved tyrosine in the active site of Pol α (in region II) and Pol δ (in region III, in the same place as Tyr 831 in Pol ϵ) to change to alanine. The *pol1-Y869A* mutation results in a weak mutator phenotype in a haploid yeast strain (Table 5, line 2). However, in marked contrast to decreased mutation rates with the *pol2-Y831A* mutation in the *POL2*, when the *pol1-Y869A* mutation is combined with the *pol2-4* mutation, the mutation rates at the *CAN1* and *his7-2* loci are increased. The increase at the *CAN1* locus is multiplicative rather than additive (Table 5, compare line 4 to lines 2 plus 3), and the increase at the *his7-2* locus is intermediate between multiplicative and additive. The interaction between the *pol1-Y869A* mutation and the *pol3-01* mutation that inactivates proofreading by Pol δ is somewhat different. In that case, the increases in mutation rates at the *CAN1* and *his7-2* loci are additive rather than multiplicative (Table 5, compare line 6 to lines 2 plus 5). This suggests that errors arising in the *pol1-Y869A* mutant strain may be proofread by the intrinsic exonuclease activity of Pol ϵ , but not by the exonuclease activity of Pol δ . Finally, when the Pol δ mutator allele

Table 5: Effect of combining the mutation in exonuclease domains of DNA polymerases δ and ϵ genes with active site mutation in DNA polymerases α and δ genes on mutation rates in mismatch repair proficient haploid derivatives of 8C-YUN1101

Relevant mutation	Mutation rates					
	Forward Can ^r mutation		Ura ⁺ reversion (base substitutions)		His ⁺ reversion (+ 1 frameshifts)	
	Rate ($\times 10^{-7}$)	Fold increase	Rate ($\times 10^{-8}$)	Fold increase	Rate ($\times 10^{-8}$)	Fold increase
no	4.4	1	1.3	1	1.9	1
	2.9–5.8		1.2–1.7		1.4–3.6	
<i>pol1-Y869A</i>	12	2.7	1.1	0.8	6.4	3.3
	6.2–28		0.8–2.2		4.1–16	
<i>pol2-4</i>	17	3.8	5.1	3.9	51	27
	12–24		3.7–5.5		45–71	
<i>pol2-4 pol1-Y869A</i>	46	10	8.8	6.7	120	63
	41–58		4.6–11		77–160	
<i>pol3-01</i>	190	43	19	15	110	59
	136–260		6–35		80–160	
<i>pol3-01 pol1-Y869A</i>	220	50	34	26	114	60
	140–240		27–48		87–210	
<i>pol3-Y708A</i>	28	6.3	6.1	4.7	6.0	3.1
	14–32		5.0–7.7		4.8–11	
<i>pol2-4 pol3-Y708A</i>	92	21	15	12	73	38
	73–120		14–18		61–104	

pol3-Y708A was combined with the *pol2-4*, the double *pol3-Y708A pol2-4* mutant showed clear multiplicity for forward mutations at the CAN1 locus – a 21-fold increase for the *pol2-4 pol3-Y708A* double mutant compared to a 3.8-fold increase for *pol2-4* and a 6.3-fold increase for *pol3-Y708A* (Table 5). However, the relationship at the two reversion loci was closer to additivity. This suggests that some but not all errors arising in the *pol3-Y708A* mutant strain may be proofread by the intrinsic exonuclease activity of Pol ϵ . Other explanations are also considered below.

Discussion

Investigation of the mechanism of the mutator phenotype of the yeast Y831A DNA Pol ϵ mutant began with the observation that, for an equivalent amount of polymerization activity, the purified Y831A enzyme has an increased ability to convert dNTPs to dNMPs relative to wild-type Pol ϵ (Figure 2). The replacement of alanine for a tyrosine in the polymerase active site that is conserved in the B family of DNA polymerases alters the relative activities of the polymerase and exonuclease active sites of Pol ϵ . The mutator effect of such mutations is unexpected, since in classical studies with T4 DNA polymerase, mutants that lead to increased nucleotide turnover were antimutators (see [33]).

As the Y831A amino acid replacement is in the polymerase active site rather than the exonuclease active site, it

seems likely that the altered protein is somehow compromised in its ability to extend primers, perhaps reflecting reduced catalytic efficiency or increased dissociation from DNA. Any polymerization defect resulting from the alanine substitution is apparently not severe enough to affect the essential role of Pol ϵ in cell viability, because a haploid yeast strain containing the *pol2-Y831A* mutation has normal growth characteristics [7]. This viability provided the opportunity, and the increased dNTP turnover provided the motivation, to investigate further the role of Pol ϵ and its proofreading activity *in vivo* by looking at the combined effects of polymerase and exonuclease active site mutations on mutation rates. These experiments lead to several interpretations.

The relative increase in dNTP turnover suggests altered communication between the polymerase and exonuclease. This may be relevant to the frameshift mutagenesis promoted by the *pol2-Y831A* mutation (Tables 1,2,3,4 and [7]). For example, misaligned intermediates may form as the primer partitions between double-stranded DNA in the polymerase active site and single-stranded DNA in the exonuclease active site (for example, see [34–36]).

The fact that *pol2-Y831A* suppresses mutations resulting from inactivation of proofreading by Pol ϵ by the *pol2-4* mutation (Tables 1, 2 and 3) suggests limited involvement of Y831A Pol ϵ in DNA synthesis *in vivo*. This is con-

sistent with the turnover data and the idea that the Y831A change compromises the ability of Pol ϵ to extend primers. On the other hand, the *pol2-Y831A* mutation does not lead to any obvious effects on growth. In this respect, our results are different from the study of a *pol3-t* mutation in the *POL3* gene encoding catalytic subunit of Pol δ . The *pol3-t* also suppresses the mutator effect of the *pol3-01* exonuclease defective allele in a strain with a double mutation but causes a temperature-sensitive growth phenotype [37]. One possibility that explains our data is increased polymerase dissociation from the primer-template. Increased dissociation has been proposed earlier to explain observations with mutant forms of two other replicative polymerases, *Escherichia coli* DNA polymerase III [38] and bacteriophage RB69 DNA polymerase [39]. For example, when wild-type Pol ϵ misinserts a nucleotide, the major pathway of error avoidance is exonucleolytic proofreading (Figure 3A, thick black arrow), whereas the mutation is generated if the mismatch is extended (central arrow). Inactivation of the intrinsic exonuclease of Pol ϵ , as with the *pol2-4* allele, yields substantial mutator effects, suggesting that under normal circumstances, wild-type Pol ϵ dissociation to allow mismatch excision by a different exonuclease may be rare (as indicated by the thin arrow). However, it may be that for Y831A Pol ϵ (Figure 3A, bottom), the dissociation mechanism prevails (thick arrow) over the proofreading mechanism, such that the mutator effect of a Pol ϵ proofreading deficiency is small, as observed here.

An alternative model is based on the hypothesis that Pol ϵ has three distinct roles in replication (Figure 3B). We propose that wild-type Pol ϵ has a limited DNA synthetic role during S-phase (Figure 3B, left). Indeed, its *in vitro* fidelity is distinct from all replicative DNA polymerases studied so far [40]. Since the mutator effect of the *pol2-4* is approximately 10-fold lower than that of the *pol3-01*, we can tentatively estimate that Pol ϵ would catalyze less than 10% of the bulk DNA synthesis during replication (indicated by thin arrow). This calculation is made under the assumption that Pol ϵ and Pol δ have similar base selectivity (currently under investigation). However, Pol ϵ may have more substantial roles in replicating certain regions/sites in the genome (arrow in the center), perhaps where proofreading is less active (for example, long homopolymeric runs, see [41]), and in performing limited reactions that are important for vegetative growth, for example, initiation of DNA replication [24]). This hypothesis is consistent with the genetic interactions of the *pol2-4* and *pol2-Y831A* combinations in *cis* and *trans* (Tables 1,2,3). In this model, the mutator effect of the *pol2-4* allele results from Pol ϵ involvement in general S-phase replication (Figure 3B, left). The Y831A amino acid change alters the enzyme as discussed above, reducing its involvement in general S-phase replication (indicated by a dashed thin arrow), but

permits its functions in the two other pathways, albeit with reduced fidelity (Figure 3B, right). The mutator effects of the *pol2-Y831A* might then reflect inaccurate replication at specific sites, or general perturbation of replication, rationalizing the lack of synergistic increases in mutation rates at some loci in strains with the *pol2-4* (Table 1) or *pol3-01* (Table 4) alleles and, on the other hand, semi-dominant mutator effect of *pol2-Y831A* in diploids when another allele is wild-type *POL2*. Alternatively or additionally, altered replication efficiency or fidelity in strains harboring mutant polymerase alleles may lead to activation of a checkpoint response that results in the accumulation of mutations [42].

The model in Figure 3B can also explain the increases in mutagenesis in the *pol2-4 pol1-Y869A* and *pol2-4 pol3-Y708A* mutant strains (Table 5), by invoking increased DNA synthesized by Pol ϵ during S-phase when Pol α or Pol δ is impaired. Especially interesting here are the synergistic increases in mutation rates at the *CAN1* locus. This synergy suggests that under certain circumstances, at least some errors generated by Pol α and Pol δ may be proofread by the intrinsic exonuclease activity of Pol ϵ . In the future, it will be interesting to test this hypothesis further, to determine under what circumstances, for example, translesion DNA synthesis [43] or base excision repair [44] errors made by one DNA polymerase may be proofread by an exonuclease intrinsic to a second DNA polymerase or even another protein [45].

Conclusions

Y831A replacement reduces replication fidelity and participation of Pol ϵ in chromosomal replication, but without eliminating an additional function of Pol ϵ that is essential for viability. This suggests that other polymerases can substitute for certain functions of Pol ϵ . Conversely, Pol ϵ can proofread errors made by polymerases α and δ .

Methods

Strains

For mutant construction we used strains 8C-YUNI101 and E134 described earlier [7]. Y831A DNA polymerase ϵ holoenzyme was purified from the protease-deficient strain ep831-T334 (*MAT α pol2-Y831A pep4-3 prb1-1122 reg1-501 gal1 ura3-52 leu2-3,112 trp1::hisG can1 LYS2*). To construct the strain, we introduced the *pol2-Y831A* allele into the strain T334 [46], a *Trp⁻* derivative of 334 described in [47], by a method described earlier [7].

Construction of mutants with DNA polymerase mutations

All mutations were introduced into chromosomal DNA polymerase genes by the integration-excision method described earlier [7,18]. Double mutants were constructed sequentially. To construct a double mutation in the *POL2* gene, we transformed a strain already possessing *pol2-*

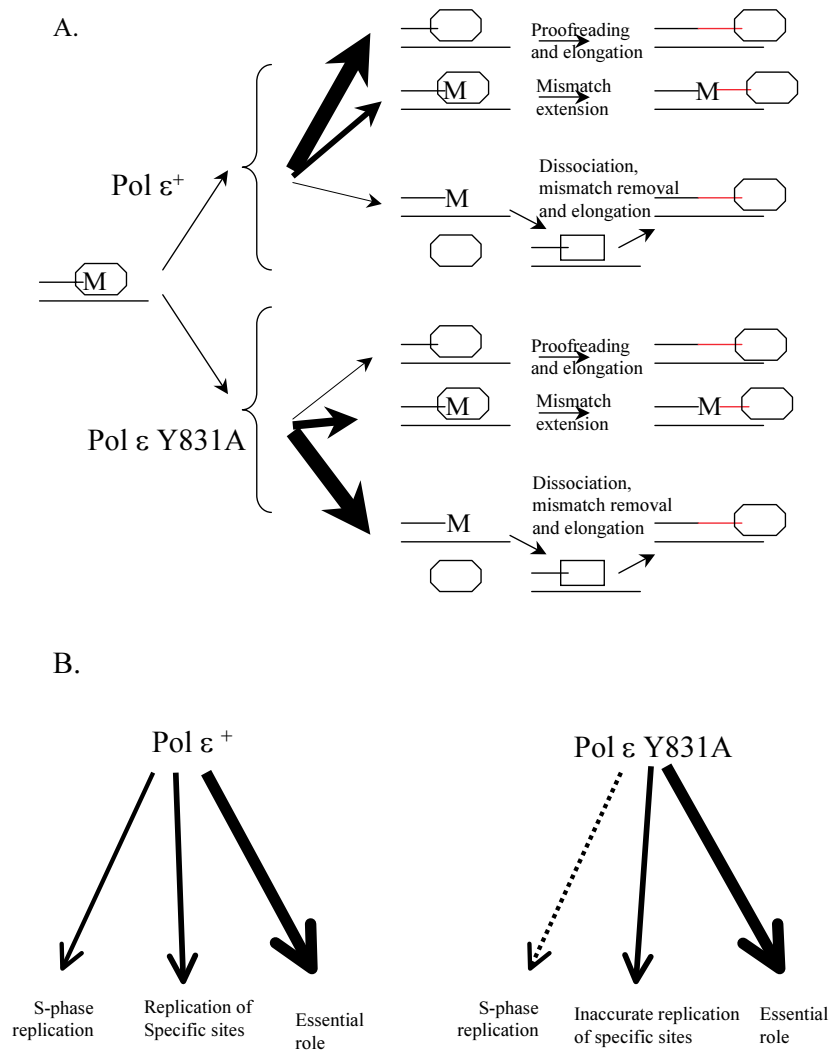


Figure 3

Models explaining results on the genetic interaction of mutants with defects in the active site and proofreading of DNA polymerase ϵ . **Thickness of arrows indicates the relative probability of a depicted pathway. See text for explanations. (A)** Dissociation-proofreading. The newly synthesized DNA is in red. 'M' stands for incorrectly inserted nucleotide. DNA polymerase is drawn as octagonal oval; an exonuclease is drawn as rectangle. **(B)** Three distinct roles of Pol ϵ in replication.

Y831A to Ura⁺ with the plasmid pJB1 (carrying the *pol2-4* mutation) cut by *Bam*HI. Transformants were plated on a medium with 5-fluoroorotic acid to select for Ura⁻ clones resulting from the loss of the plasmid. Resulting clones were tested by PCR for the presence of both mutations. Detection of *pol2-Y831A* was described previously [7]. To detect the *pol2-4* mutation, we amplified a 481 bp region of the *POL2* gene using primers M32-m, 5'TCCGAGTATC-TATAGACAAGGA and M36, 5'CTCACCTTCAG-CATCTGG. The resulting fragment was digested with *Sfc*I. The presence of the *pol2-4* mutation creates a new *Sfc*I site; the PCR fragment is cut into 119 and 362 base pair fragments. The same mutations were created in basic haploid strains 8C-YUNI101 and E134. The resulting mutants were then crossed to generate diploids homozygous or heterozygous for polymerase mutations.

Mutation rate measurements

Measurements of mutation rates were performed as described [7].

Purification of DNA polymerase ϵ

The four-subunit DNA polymerase ϵ was purified from 1.2 kg of the protease deficient ep831-T334 strain as described in [23,29] with modifications. Briefly, after an ammonium sulfate precipitation step and SP-Sepharose chromatography [23,29], the dialyzed sample was loaded onto a Q Sepharose HP60/100 column (Amersham Biosciences Corp., Piscataway, NJ, USA) and eluted with nine column volumes of a linear gradient of NaCl from 50 to 500 mM. Four peaks with DNA polymerase activity were obtained, and the peak containing the four-subunit Pol ϵ was identified by immunoblotting with polyclonal antibodies [29]. Further fractionation of pooled fractions from this peak by MonoS H/R and HiTrap heparin columns (Amersham Biosciences) was performed as described [23]. Finally, fractions with DNA polymerase ϵ from the Heparin column were loaded onto a MonoS 1.6/5 column in a SMART system (Amersham Biosciences). The polymerase was eluted by a linear NaCl gradient from 50 to 1000 mM. Fractions (30 μ l) were collected and assayed for activity. One UV absorbance peak was observed and it coincided with one peak of polymerase activity. The purity of the preparation was approximately 90% as judged from SDS-PAGE electrophoresis and SimpleStain (Invitrogen, Carlsbad, California, USA) colloidal coumassie staining (Figure 1). Peak fractions with specific polymerase activity around 4,000 U/mg were pooled and stored at -80°C.

Enzyme activity determination

Polymerase activity was determined using a poly(dA)₃₀₀/oligo(dT)₁₀ substrate as described [23]. Reaction mixtures (20 μ l) were incubated at 30°C, and 5 μ l aliquots were withdrawn at the indicated time intervals and placed in

tubes with 1 μ l of 300 mM EDTA. One μ l of these mixtures was spotted onto each of the two PEI (Merck, Whitehouse Station, NJ, USA) plates. One was developed in 1 M LiCl, 1 M formic acid, 50 mM NaH₂PO₄, thus separating the unincorporated label from label incorporated into DNA. A second plate was developed with essentially the same solution but containing only 0.4 M LiCl. This permits quantification of the amount of labeled nucleoside monophosphate generated (that is, turnover of dNTP by the DNA polymerase-associated proofreading exonuclease, see Figure 2A,2B). The intensity of the spots was quantified by BAS2500 Bio Image Analyzer (Fuji Photo Film Company, Tokyo, Japan).

Exonuclease assays employed a 3'-radiolabelled poly(dA)₃₀₀ single-stranded substrate. Three μ g of poly(dA)₃₀₀ were labeled by terminal transferase with 40 μ Ci of α -³²P deoxyadenosine triphosphate (dATP) in 1 mM dATP and 1x One-for-All Plus buffer (Promega, Madison, WI, USA). Reaction products were purified using a Qiagen nucleotide removal kit (Qiagen, Valencia, CA, USA) and then examined by TLC. An average of nine dATP molecules were added per each poly(dA)₃₀₀ molecule. Exonuclease reactions (20 μ l) with 37 ng (0.14 μ Ci) of labeled substrate and 200 ng of unlabeled poly(dA)₃₀₀ were performed in the same buffer as the polymerase reaction, also at 30°C. At each time point, one μ l of reaction mixture was spotted onto a PEI plate and developed with 0.4 M LiCl as described above. The polymerase activity estimate was compensated by dividing the apparent value of incorporation (acid insoluble nucleotides) with 0.78 for the wild-type enzyme (turnover rate = 22%) and 0.50 for the mutant enzyme (turnover rate = 50%).

Authors' contributions

YIP and SM purified and did initial characterization of polymerase ϵ Y831A. SM purified wild-type enzyme and performed most of the biochemical characterization of the enzymes. YIP did genetic experiments with yeast. YIP, SM, HM and TK participated in the design of the study. TK coordinated the study. All authors read and approved the final manuscript.

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