APPENDIX

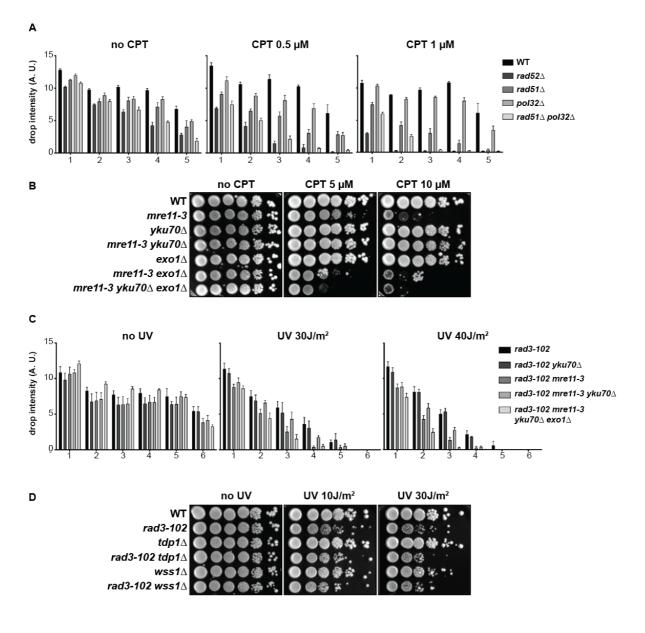
Homologous recombination and Mus81 promote replication completion in response to replication fork blockage

Benjamin Pardo^{1,2,*}, María Moriel-Carretero^{1,2,3}, Thibaud Vicat¹, Andrés Aguilera^{2,*} and Philippe Pasero¹.

Page 2: Appendix Figure S1

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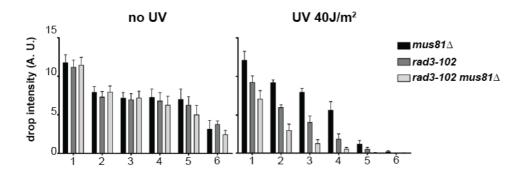
Page 4: Appendix Figure S3



Appendix Figure S1. DNA repair is similar in CPT-treated and *rad3-102* cells.

Related to Figure 1

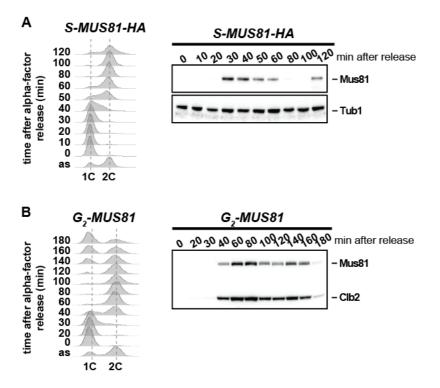
- (A) Quantification of four independent CPT sensitivity assays illustrated in Figure 1B. Mean and SD values are indicated for each yeast drop (1 to 5).
- **(B)** CPT sensitivity assayed by 10-fold serial dilutions of different mutant combinations on YPAD plates (n=3).
- (C) Quantification of four independent UV sensitivity assays illustrated in Figure 1D. Mean and SD values are indicated for each yeast drop (1 to 6).
- (**D**) UV sensitivity assayed by 10-fold serial dilutions of $tdp1\Delta$ and $wss1\Delta$ mutants in combination with rad3-102 allele on YPAD plates after exposure to the indicated UV-C doses (n=3).



Appendix Figure S2. The absence of Mus81 increases the UV sensitivity *rad3-102* cells.

Related to Figure 2

Quantification of four independent CPT sensitivity assays illustrated in Figure 2C. Mean and SD values are indicated for each yeast drop (1 to 5).



Appendix Figure S3. Cell-cycle restricted expression of *MUS81* using S and G₂ tags.

Related to Figure 5

- (A) Immunoblot analysis of S-Mus81 tagged with 3 HA epitopes at its C-terminus. *S-MUS81-HA* cells were synchronized in G_1 with α -factor and released into S phase (n=2). Cells were collected at the indicated time points and S-Mus81 was immunodetected with anti-HA antibodies. Tub1 immunodetection serves as a loading control. Flow cytometry profiles corresponding the experimental setup are shown.
- (B) Immunoblot analysis of G_2 -Mus81. G_2 -Mus81 cells were synchronized in G_1 with α -factor and released into S phase (n=2). Cells were collected at the indicated time points and G_2 -Mus81 and Clb2 were immunodetected with anti-Clb2 antibodies. Flow cytometry profiles corresponding the experimental setup are shown.