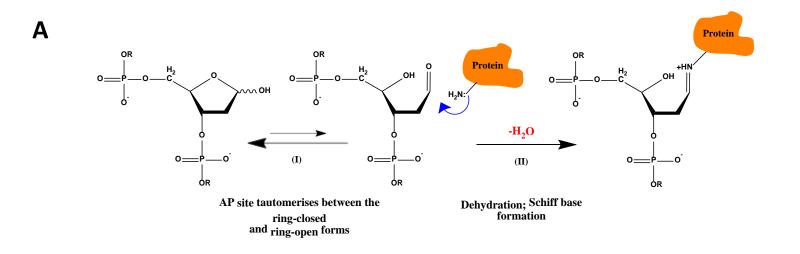
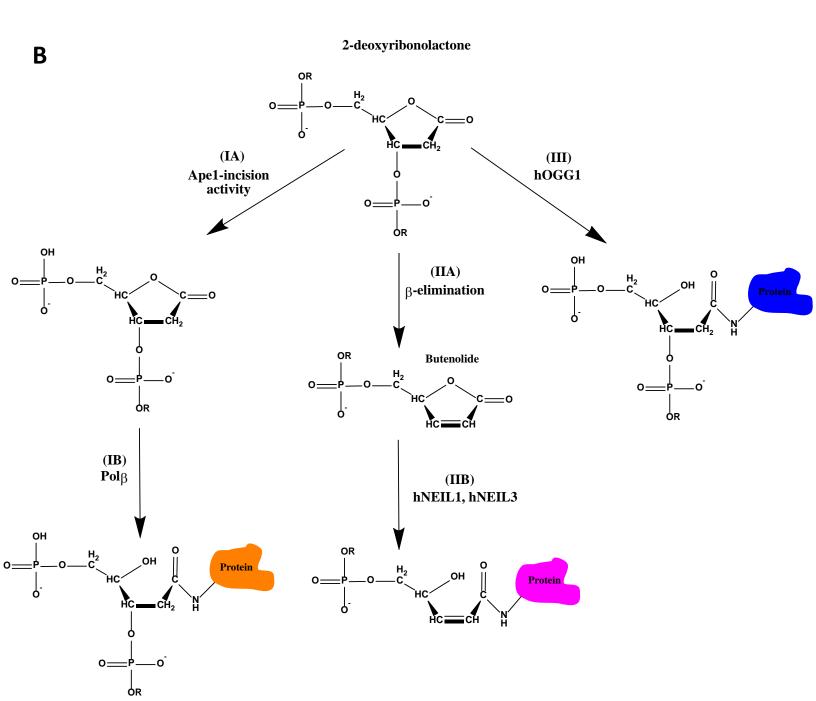
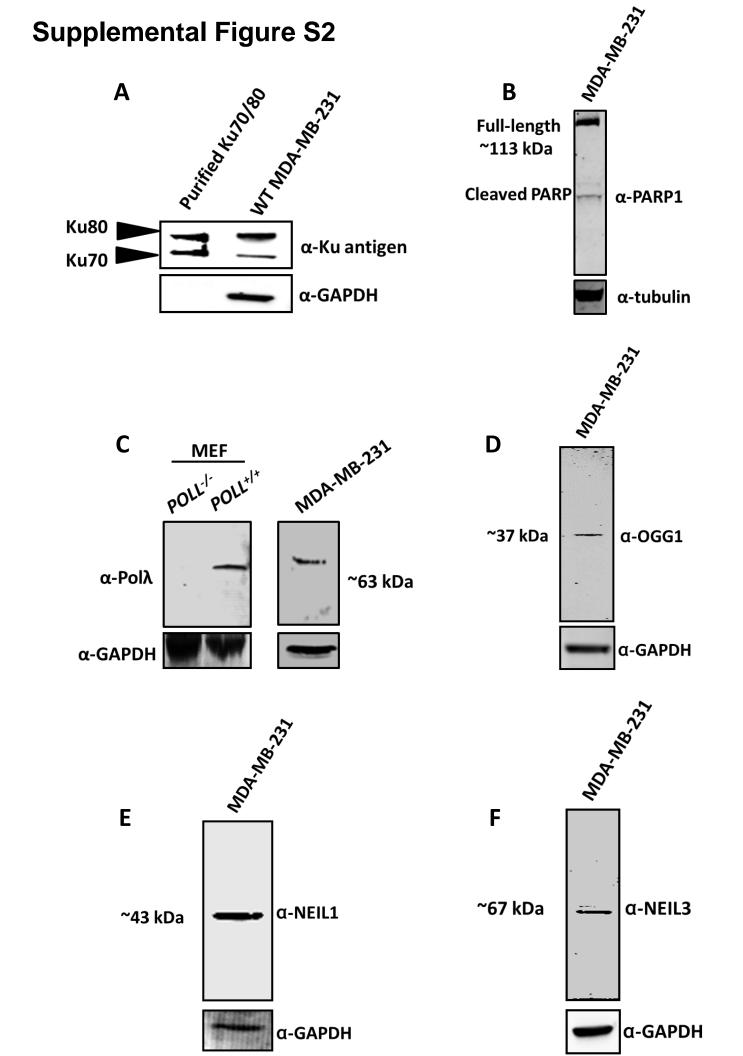
Supplemental Figure S1







Supplemental Figure S3

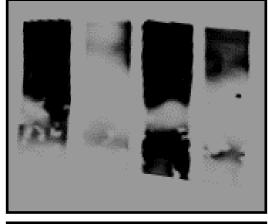
MDA-MB-231

Veg. Ctrl.

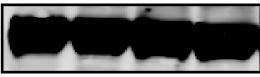
1 μM Olaparib

10 µM cuop

10 μΜ CuOP 1 μΜ Olaparib



 α -PAR



 α -Tubulin



Figure S1. Trapping of BER AP Lyases in DPC. A. Schematic showing reaction mechanism for trapping 5'-dRp/AP lyases in DPC with sodium borohydride (NaBH₄). **B.** Trapping of 5'-dRp/AP lyases by dL residues.

Figure S2. Confirmation of AP Lyase Expression. Whole cell extracts were prepared from MDA-MB-231 cells (A., B., D.-F.) or MEF cells (C.) as described in the material and methods section, then 50 μg of total protein were resolved by SDS-PAGE, and electroblotted onto PVDF membranes. Membranes were then blocked in 3% BSA, then probed using specific anti-sera raised against eother A. the Ku 70/80 heterodimer, B. PARP1, C. Polλ, D. OGG1, E. NEIL1, or F. NEIL3.

Figure S3. PARylation in MDA-MB-231 Cells Following Olaparib Treatment with or without CuOP. Cells were treated with 1 μ M olaparib in the presence or absence 10 μ M CuOP for 2 h, then whole cell extracts were prepared as described in the materials and methods section. For immunoblot analysis of PARylation, a total of 100 μ g of protein were resolved by SDA-PAGE, electroblotted onto PVDF membranes, then probed with anti-serum against PAR (1:500).