

Supplementary Table 1: Previously characterized properties of MCL cell lines.

EBV, Epstein Barr Virus; ND, Not Determined; Del, Allele Deletion; Pm, polymorphism; Mut, Mutation; *, mutation in ATM kinase domain. References are given in parenthesis.

Cell Line	ATM status (Allele 1 / Allele 2)	p53 status (Allele 1 / Allele 2)	Ploidy	EBV	t(11;14)(q13;q32) translocation
Granta-519	Del (1) / Mut (R2832C)* (2)	WT / Del (3)	2N (4)	+ (5)	+ (6)
HBL-2	ND / ND	Del (7) / D281G (8)	2N (4)	- (9)	+ (9)
JVM-2	WT / WT (10)	WT / WT (10)	2N (4)	+ (7)	+ (11)
MAVER-1	Del / ND (12)	Del / Mut (D281E/H) (8, 12)	2N (4)	- (12)	+ (12)
UPN1	WT / Pm (S333F) (13)	Mut (E286K) / Del (8, 13)	2N (4)	- (13)	+ (13)
UPN2	Del / Mut (D2725V)* (13)	Del / Mut (R175H) (13)	3N (4)	- (13)	+ (13)
Z138	ND / ND	WT / WT (7)	2N (4)	- (14)	+ (14)

References for Supplementary Table 1

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Figure legends for Supplementary Figures**Supplementary Figure 1- ATM signaling in lymphoblastoid control cell lines**

Lymphoblastoid cell lines, A) C35ABR (BT) and B) L3 were exposed to 2 Gy IR and harvested following the indicated incubation times. Whole cell extracts (50 µg total protein) were analyzed by SDS PAGE and immunoblots were probed for autophosphorylation of ATM on Ser-1981, phosphorylation of SMC-1 on Ser-957 and Ser-966, and phosphorylation of KAP1 on Ser-824 as indicated. Immunoblots were also probed for total ATM, SMC-1 and KAP1. Actin and DNA-PKcs levels are shown as loading controls.

Supplementary Figure 2. ATM-dependent signaling in MCL cell lines

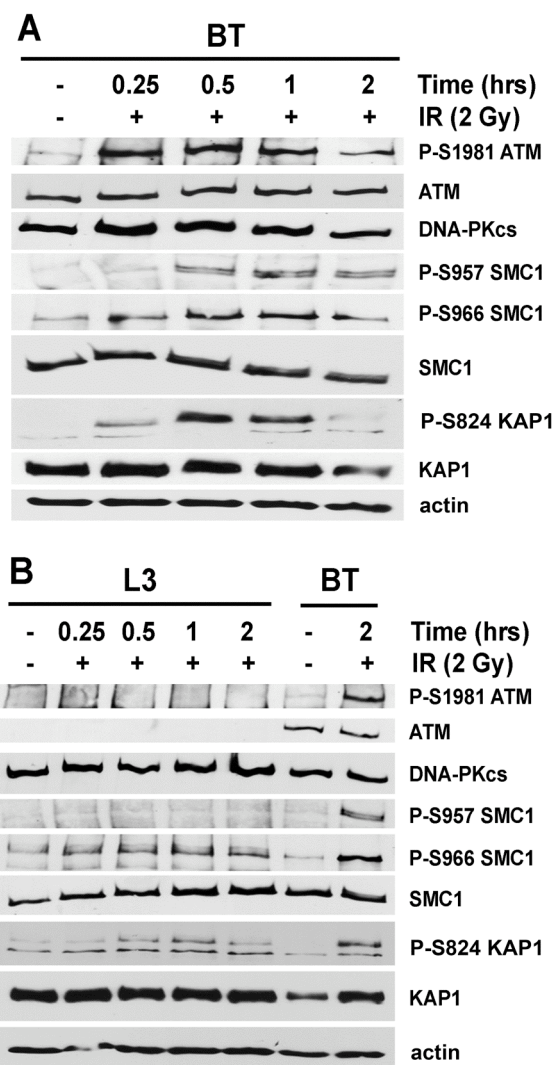
MCL cell lines, A) HBL-2, B) MAVER-1 and C) JVM-2 cells were exposed to 2 Gy of IR and analyzed as in Supplementary Figure 1.

Supplementary Figure 3. Quantitation of ATM knockdown in ZC-shATM

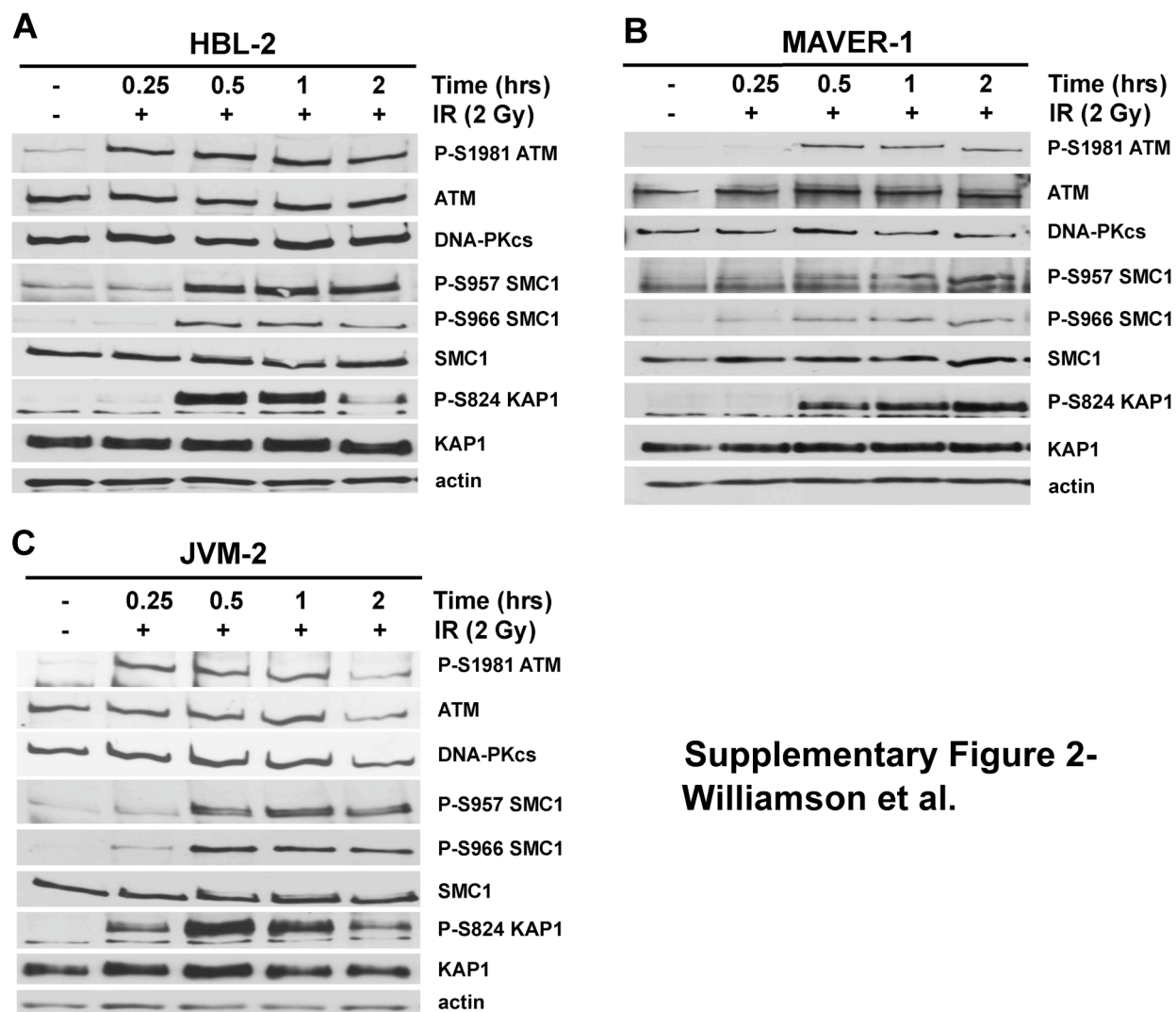
The ratio of ATM to DNA-PKcs protein expression for each cell line shown in Figure 4A was normalized to that in BT cells.

Supplementary Figure 4. Olaparib induces P-S1981 in UPN1 cells but not UPN2 cells

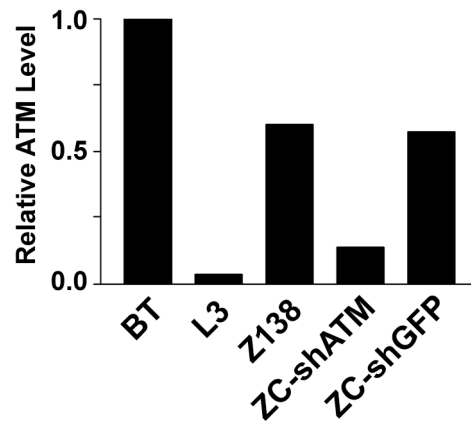
UPN1 (A) and UPN2 (B) cells were exposed to olaparib (2.5 µM) or vehicle (VEH, DMSO) for 24, 48, 72 or 96 hours as indicated. Whole cell extracts (50 µg total protein) were run on SDS PAGE, immunoblotted and probed for ATM autophosphorylation at Ser-1981 (P-S1981), total ATM and total SMC-1 as indicated. Quantitation of olaparib-induced P-S1981 ATM compared to total ATM for UPN1 is shown below the blot. As a positive control for P-Ser-1981, in the UPN2, BT cells were irradiated 2 Gy and harvested after 1 hour.



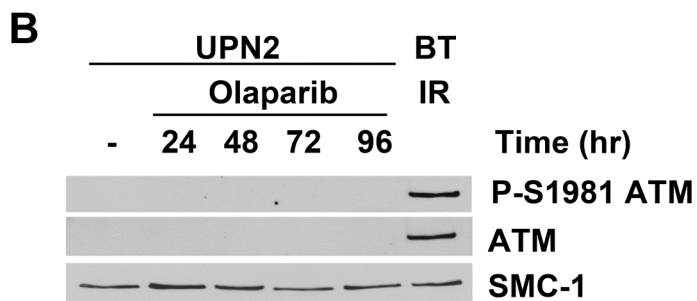
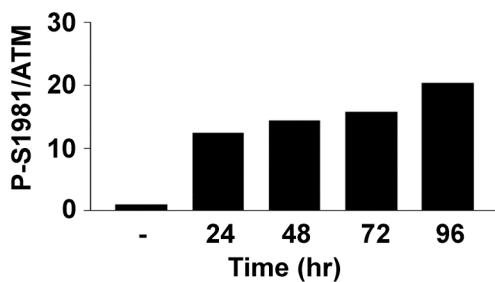
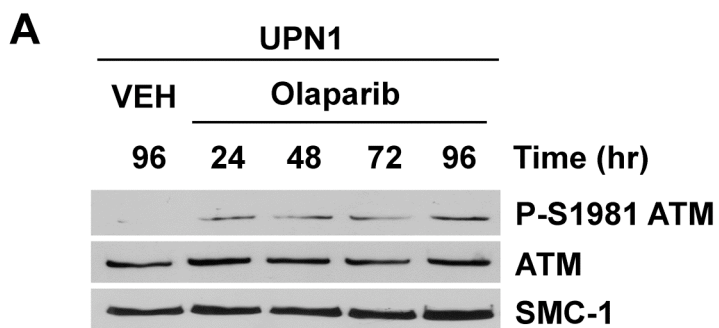
**Supplementary Figure 1-
Williamson et al.**



**Supplementary Figure 2-
Williamson et al.**



**Supplementary Figure 3-
Williamson et al.**



Supplementary Figure 4- Williamson et al.