## **SUPPLEMENTARY DATA**

**Supplementary Table S1.** Dose-response and exposure-response parameters.

Talazoparib parameter	<b>A</b> <sub>0</sub>	I <sub>max</sub>	ID <sub>50</sub> /IAUC <sub>50</sub>	h
Dose (mg)	300	0.950	0.0331	0.865
AUC <sub>0-24</sub> (pg·hr/mL)	242	0.978	19,000	1.23

Abbreviations:  $A_0$ , estimated baseline activity;  $AUC_{0.24}$ , area under the plasma concentration-time curve from 0 to 24 h;  $ID_{50}$ , inhibitory dose 50%;  $IAUC_{50}$ , area under the curve at 50% of inhibition;  $I_{max}$ , maximum inhibitory effect; PARP, poly(ADP-ribose) polymerase; PBMC, peripheral blood mononuclear cell.

As illustrated in the dose-response curve, a fair degree of variability was observed across patients at doses of 0.4 mg/day and below, with near complete inhibition being achieved in one patient at 0.1 mg/day and all three patients at 0.2 mg/day dose levels, while two of the three patients at the 0.4 mg/day dose level showed little inhibition. PBMC PARP inhibition was consistently inhibited across all patients at doses of 0.6 mg/day and above with an  $IC_{50}$  of 0.0331 mg/day.