

S3 Table. Patients with DNA damage repair gene alteration detected at baseline (n=10)

Study ID	Best response to mFOLFIRINOX by RECIST v1.1	Gene	Mutation setting	Variant type	Chromosome	Mutated amino acid
FOL005	SD	<i>BRCA2</i>	Germline	SNV	13	M784V
FOL006	SD	<i>ATM</i>	Germline	SNV	11	N1650S
FOL012	PR	<i>ATM</i>	Somatic	SNV	11	Q1627R
FOL015	SD	<i>ATM</i>	Germline	SNV	11	K92T
FOL016	PR	<i>MLH1</i>	Somatic	SNV	3	T451A
FOL020	SD	<i>ATM</i>	Somatic	Indel	11	F1036fs
FOL023	PR	<i>ATM</i>	Somatic	SNV	11	A2986S
FOL024	PR	<i>ATM</i>	Germline	SNV	11	Y2036Y
		<i>BRCA1</i>	Germline	Indel	17	K653fs
FOL034	SD	<i>BRCA2</i>	Germline	SNV	13	Q1886*
FOL037	SD	<i>ATM</i>	Germline	SNV	11	I2861V
		<i>ATM</i>	Germline	SNV	11	N1650S
		<i>ATM</i>	Somatic	SNV	11	S2162R
		<i>ATM</i>	Somatic	SNV	11	P2699A
		<i>ATM</i>	Somatic	SNV	11	G2891D
		<i>ATM</i>	Somatic	SNV	11	G3030E
		<i>ATM</i>	Somatic	Indel	11	T915fs
		<i>ATM</i>	Somatic	SNV	11	L439P
		<i>ATM</i>	Somatic	Indel	11	D588fs

Indel, insertions and deletions; mFOLFIRINOX, modified fluorouracil, leucovorin, oxaliplatin, and irinotecan; PR, partial response; RECIST, Response Evaluation Criteria in Solid Tumor; SD, stable disease; SNV, single nucleotide variation.