

**Table S1.** Gene List.

<b>Number</b>	<b>Gene</b>
1	ATM
2	BRIP1
3	MLH1
4	NBN
5	RAD50
6	TP53
7	BARD1
8	CDH1
9	MRE11A
10	PALB2
11	RAD51C
12	XRCC2
13	BRCA1
14	CHEK2
15	MSH2
16	PMS2
17	RAD51D
18	BRCA2
19	EPCAM
20	MSH6
21	PTEN
22	RINT1

**Table S2.** Distribution of P/LP-Vs in multiple P/LP-Vs carriers.

Carriers ID	Genes	HGVsg	Carriers ID	Genes	HGVsg
brca14_rscm_17	ATM	11:g.108282707delinsCATACAACACTAAAAAATG	brca79_rscm_17	BRCA1	17:g.43091349delinsTTTAAAGTGCAGCTTTTC
	BRCA2	13:g.32333103delinsTA		PMS2	7:g.5987583_5987584delinsC
	MSH6	2:g.47806453delinsCTTAGAT		STK11	19:g.1219400_1219456delinsT
brca25_rscm_17	BRCA2	13:g.32379885delinsCA	brca80_rscm_17	ATM	11:g.108345760delinsTCAGTAGCTCAAGGG
	STK11	19:g.1219400_1219456delinsT		BRCA2	13:g.32316515_32316516delinsT
brca28_rscm_17	BRCA2	13:g.32379885delinsCA		STK11	19:g.1219400_1219456delinsT
	PMS2	7:g.5987525_5987526delinsC	brca82_rscm_17	BRCA2	13:g.32379885delinsCA
brca37_rscm_17	BRCA2	13:g.32332277delinsGCATACAT		PMS2	7:g.5987525_5987526delinsC
	CDKN2	9:g.21974732_21974737delinsC	brca85_rscm_17	BRCA2	13:g.32379885delinsCA
	RAD50	5:g.132595759_132595760delinsT		PMS2	7:g.5987525_5987526delinsC
	TP53	17:g.7674917delinsTC		STK11	19:g.1219400_1219456delinsT
brca48_rscm_17	PMS2	7:g.5987525_5987526delinsC	brca86_rscm_17	ATM	11:g.108326058delinsCCTTCTTCCAACAGAAACGATTGT
	PTEN	10:g.87965294delinsTCTTATCA		BRCA2	13:g.32379885delinsCA
brca58_rscm_17	PALB2	16:g.23635659_23635660delinsA	brca89_rscm_17	BRCA2	13:g.32379885delinsCA
	PMS2	7:g.5987525delinsCT		PMS2	7:g.5987525delinsCT
brca59_rscm_17	BRCA1	17:g.43093581_43093582delinsT		STK11	19:g.1219400_1219456delinsT
	MSH6	2:g.47803500delinsAC	brca90_rscm_17	BRCA2	13:g.32379885delinsCA
	RAD50	5:g.132595759_132595760delinsT		PMS2	7:g.5987525_5987526delinsC
	RAD51	17:g.58734130delinsAATCCAGGAAATGCAGAAGAG	brca91_rscm_17	PALB2	16:g.23635659_23635660delinsA
brca67_rscm_17	MSH6	2:g.47803500delinsAC		STK11	19:g.1219400_1219456delinsT
	PALB2	16:g.23635659_23635660delinsA	brca93_rscm_17	PMS2	7:g.5987525delinsCT
	RAD50	5:g.132595759_132595760delinsT		STK11	19:g.1219400_1219456delinsT
brca72_rscm_17	BRCA1	17:g.43093821_43093822delinsT	brca98_rscm_17	BRCA2	13:g.32379885delinsCA
	STK11	19:g.1219400_1219456delinsT		BRIP1	17:g.61683605_61683606delinsA
brca73_rscm_17	ATM	11:g.108245025_108245026delinsA		MSH6	2:g.47803657_47803658delinsT
	BRIP1	17:g.61683605_61683606delinsA		RAD50	5:g.132595759_132595760delinsT
	STK11	19:g.1219400_1219456delinsT		BRCA2	13:g.32379885delinsCA
brca74_rscm_17	BRCA2	13:g.32338277delinsGACTTTGACAGAAA	brca100_rscm_17	PMS2	7:g.5987525_5987526delinsC
	PALB2	16:g.23635659_23635660delinsA		RAD50	5:g.132595759_132595760delinsT
	TP53	17:g.7674917delinsTC			

Carriers ID with bold text have familial history (mother/aunt/grandmother) in breast cancer

**Table S3.** Potentially pathogenic VUS detected in breast cancer susceptibility genes.

Gene	HGVScg	Type of variant	dbSNP / ClinVar ID	Number of carriers
BRCA1	17:g.43093594delinsCAAAA	Frameshift	-	1
BRCA2	13:g.32354874C>T	Missense	rs41293505	1
STK11	19:g.1221967C>T	Missense	-	1
	19:g.1226589G>A	Missense	rs775978755	1
	19:g.1222992C>T	Missense	rs750366043	1
	19:g.1219346G>A	Missense	rs567769257	1
ATM	11:g.108247122C>T	Missense	950587	1
	11:g.108271274G>A	Missense	rs749471737	1
	11:g.108335101C>T	Missense	rs1591192429	1
PALB2	16:g.23624080C>G	Missense	rs372931676	1
BRIP1	17:g.61808594C>T	Missense	rs758360637	1
	17:g.61799232C>T	Missense	rs786202780	1
	17:g.61799193C>T	Missense	rs772570870	1
	17:g.61780973T>C	Missense	rs1567813893	1
CHEK2	22:g.28725253C>T	Missense	rs587781667	1
BARD1	2:g.214781051A>T	Missense	rs776157713	1
	2:g.214730494G>T	Missense	rs1553612535	1
	2:g.214728819G>A	Missense	rs76744638	3
MLH1	3:g.36993576G>T	Missense	rs777971423	1
	3:g.37028831C>T	Missense	rs532873141	1
MSH2	2:g.47414338C>G	Missense	953109	3
	2:g.47414418G>T	Missense	rs587779197	2
MSH6	2:g.47798716A>T	Missense	rs762168786	1
	2:g.47799065G>A	Missense	rs63750440	1
	2:g.47799639T>A	Missense	rs745937181	2
	2:g.47800636A>G	Missense	rs587782593	1
	2:g.47806650C>T	Missense	rs773763465	3
PMS2	7:g.5999217C>T	Missense	rs876658387	1
RAD51C	17:g.58703259G>A	Missense	rs200857129	2
RAD51	17:g.58703259G>A	Missense	rs200857129	1
MUTYH	1:g.45333414C>T	Missense	rs587782683	1
	1:g.45331466C>T	Missense	rs748700385	1