

**Table S1. Proportion of classified variant for 128 HGSC patients**

<b>Clinvar Classification</b>	<b>Proportion</b>
Benign	40 (3%)
Benign / Likely Benign	67 (5%)
Likely Benign	25 (2%)
VUS	991 (73%)
Likely Pathogenic	26 (2%)
Likely pathogenic / Pathogenic	19 (1%)
Pathogenic	181 (13%)

*\*HGSC – High grade serous ovariancancer; VUS – variants of uncertain significance.*

**Table S2. Pathogenic or likely pathogenic mutations per patient (N = 128)**

	<b>Number of patients (%)</b>	<b>Number of patients with druggable targets (%)</b>
None	10 (8%)	0
1 mutation	64 (50%)	1 (1%)
2 mutations	35 (27%)	18 (14%)
3 mutations	12 (9%)	4 (3%)
4 mutations	3 (3%)	1 (1%)
≥ 5 mutations	3 (3%)	3 (2%)

**Table S3. All pathogenic and likely pathogenic mutations**

**Table S4. Clinical trials evaluating PARP inhibitors in platinum-resistant patients**

PARP inhibitor	Trial	Analysis population	No. of patients	Results
Olaparib	Fong et al., 2010 (24)	<i>BRCA1/2</i> mutated ovarian cancer	50 - 13 platinum-sensitive - 24 platinum-resistant - 13 platinum-refractory	RECIST response - 6 (46.2%) - 8 (33.5%) - 0 (0%)
Olaparib	Kaufmann et al., 2015 (25)	PROC with germline and somatic <i>BRCA</i> mutation	193 platinum-resistant	Tumor response rate 31,1 (95% CI, 24,6-38,1) Median PFS 7 months Median OS 16,6 months
Niraparib	QUADRA, 2018 (26)	Relapsed ovarian cancer with PARP-naïve germline or somatic <i>BRCA</i> mutation	55 - 18 platinum-sensitive - 21 platinum-resistant - 16 platinum-refractory	ORR 39% ORR 33% ORR 19%
Olaparib	BAROCCO, 2019 (27)	High-grade PROC of whom 86% (109/123) are <i>BRCAwt</i>	123 platinum-resistant	Median PFS - Paclitaxel arm 3,1 months - Olaparib-cediranib arm 5,7 months - Cediranib arm 3,8 months HR for PFS in the paclitaxel vs. olaparib-cediranib arm was 0,76 (90% CI: 0,49-1,17).  Median PFS in the <i>gBRCAwt</i> patients (n=109) - Paclitaxel arm 2,1 months - Olaparib-cediranib arm 5,8 months - Cediranib arm 3,8 months HR for PFS in the paclitaxel vs. olaparib-cediranib arm was 0,63 (95% CI: 0,36-1,10).
Rucaparib	ARIEL4, 2021 (28)	<i>BRCA1/2</i> mutated high grade recurrent EOC.	345 - 176 platinum-resistant	Median PFS - Rucaparib arm 7,4 months (95% CI, 7,3-9,1) - Chemotherapy arm 5,7 months (95% CI, 5,5-7,3) HR 0,64 (95% CI, 0,49-0,84)
Olaparib	NRG-GY005 (NCT02502266)	Recurrent platinum-resistant or -refractory HGSC or germline <i>BRCA</i> mutant ovarian cancer	680	Estimated study completion June 2023.
Olaparib	OCTOVA (NCT03117933)	<i>BRCA1/2</i> mutated PROC	132	Estimated study completion Nov 2021. Results pending.

Figure S1. Kaplan-Meier curve for overall survival in patient groups with *BRCA1/2* mutation (upper curve) compared with the *BRCAwt* patients (lower curve).

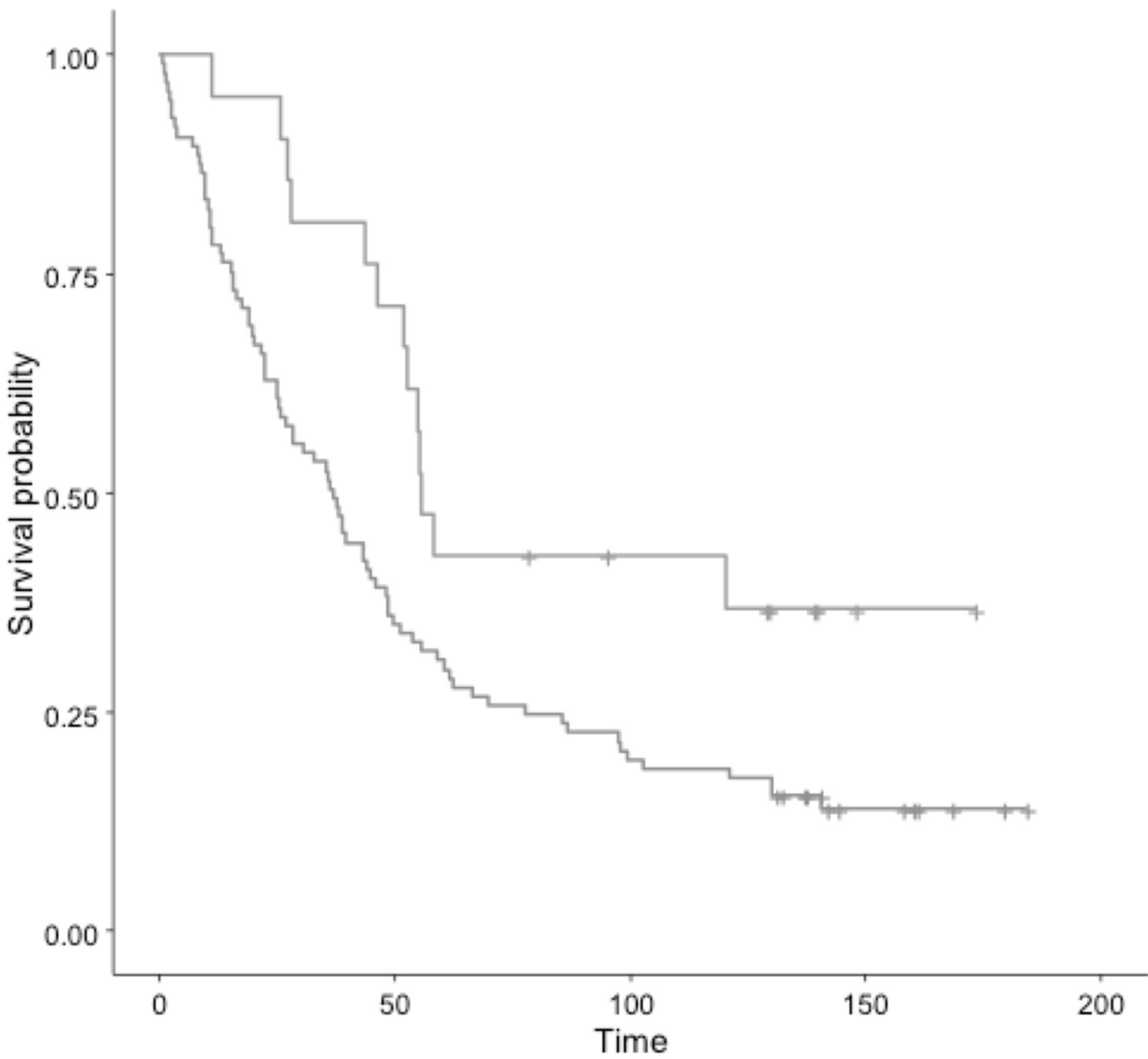


Figure S2. Mutations at the gene level in the groups with druggable targets

