

TRAMANT: phase II study of maintenance therapy with trabectedin vs trabectedin + PLD after trabectedin + PLD in relapsed ovarian cancer recurring 6-12 months after platinum based chemotherapy

OC/fallopian tube/primary peritoneum cancer relapsed 6-12 months after the last platinum based treatment

6 cycles of PLD 30 mg/m² +
Trabectedin 1.1 mg/m² q3w

RANDOMIZATION
1:1

Trabectedin 1,1 mg/m²
q3w up to toxicity or
progression

PLD 30 mg/m² day 1 +
Trabectedin 1,1 mg/m²
q3w up to toxicity or
progression

Main inclusion/exclusion criteria:

- patients must be in SD/PR/CR after 6-8 cycles of PLD+trabectedin
- patients must have received no more than 3 previous lines of chemotherapy
- patients must have measurable or evaluable disease
- patients must be able to receive dexamethasone or its equivalent
- patients with prior resistance to anthracyclines or PLD will be excluded
- patients with prior exposure to trabectedin will be excluded

CONTATTI

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MITO 30: phase II study of trabectedin and olaparib for platinum-resistant ovarian cancer

High grade serous or endometrioid platinum resistant/refractory ovarian cancer



Trabectedin 1.1 mg/mq i.v. (24h infusion) q3w + olaparib 150 mg BID until toxicity or progression

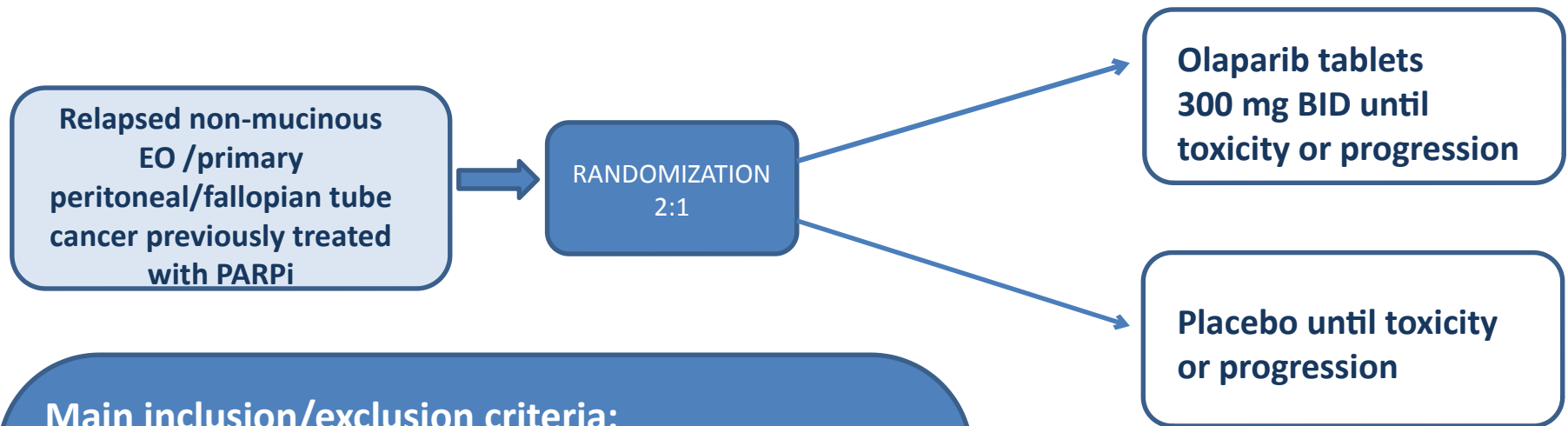
Main inclusion/exclusion criteria:

- at least one previous platinum-based regimen and no more than two previous lines of chemotherapy
- availability of tumor samples for central revision and translational analysis
- measurable disease (RECIST 1.1)
- patients must be able to receive dexamethasone or its equivalent
- patients with symptomatic metastatic brain or meningeal tumors will be excluded

CONTATTI

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OReO: phase IIIb of olaparib maintenance retreatment in patients with EOC previously treated with a PARPi



Main inclusion/exclusion criteria:

- documented *BRCA1/2* status (**only BRCA-negative cohort open**)
- patients must have received one prior PARPi therapy
- patients must be in response after a platinum-based chemotherapy regimen and have received at least 4 cycles of treatment
- patients must **not** have received bevacizumab during this course of treatment
- patients must have available tumor samples
- patients with sintomatic metastatic brain or meningeal tumors will be excluded

CONTATTI

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