



ICON8



Diagnosis of Stage IC-IV EOC/PPC/FTC
 After immediate primary surgery or planned to receive NACT plus delayed primary surgery

N=1485

Randomise 1:1:1

Arm A1
6 cycles

Arm A2
6 cycles

Arm A3
6 cycles

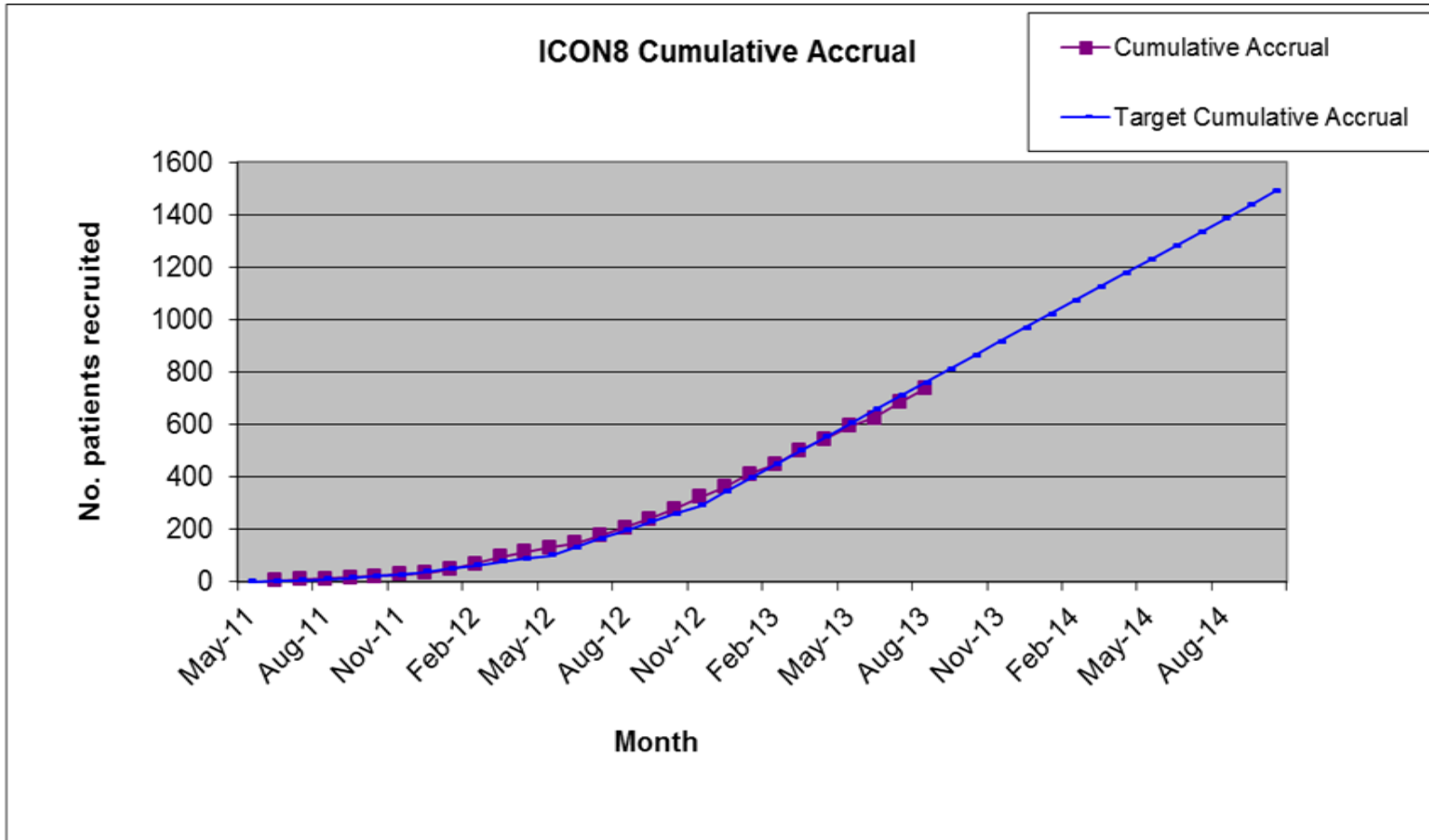
Arm 1 vs. Arm 2

Arm 1 Carboplatin AUC 5 q3w
Paclitaxel 175mg/m² q3w

Arm 2 Carboplatin AUC 5 q3w
Paclitaxel 80mg/m² q1w

Arm 1 vs. Arm 3

Arm 3 Carboplatin AUC 2 q1w
Paclitaxel 80mg/m² q1w



@24Sept2013: n=778 (UK=742, KGOG=18, GICOM= 10, ICORG=6)



ICON8



ICON8 trials programme, revised design

N=1485

ICON8A

Diagnosis of Stage IC-IV EOC/PPC/FTC

Randomise 1:1:1

Arm A1
6 cycles

Arm A2
6 cycles

Arm A3
6 cycles

Arm A1 Carboplatin AUC 5 q3w
 Paclitaxel 175mg/m² q3w

Arm A2 Carboplatin AUC 5 q3w
 Paclitaxel 80mg/m² q1w

Arm A3 Carboplatin AUC 2 q1w
 Paclitaxel 80mg/m² q1w

N=1170

ICON8B

Diagnosis of Stage III-IV EOC/PPC/FTC with residual disease after surgery or planned for neoadjuvant chemotherapy

Randomise 1:1:1

Arm B1
6 cycles

Arm B2
6 cycles

Arm B3
6 cycles

16 cycles maintenance Bevacizumab

Arm B1 Carboplatin AUC 5 q3w
 Paclitaxel 175mg/m² q3w
 Bevacizumab 15mg/kg q3w

Arm B2 Carboplatin AUC 5 q3w
 Paclitaxel 80mg/m² q1w

Arm B3 Carboplatin AUC 5 q3w
 Paclitaxel 80mg/m² q1w
 Bevacizumab 15mg/kg q3w

NB. Patients with Stage III & residual disease after surgery or who are planned to receive neoadjuvant chemotherapy OR any patients with stage IV disease are still eligible for ICON8A as well as B so that they may still enter the trial if:

- they have contra-indications to or decline bevacizumab
- their site does not have access to bev, e.g. in Australia