

Dexamethasone added to induction and post-remission therapy in older patients with newly diagnosed acute myeloid leukemia: a multicenter, phase II trial (DEXAML-02)

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Supplementary Information

Dexamethasone added to induction and post-remission therapy in older patients with newly diagnosed acute myeloid leukemia: a multicenter, phase II trial (DEXAML-02). *Sarah Bertoli et al.*

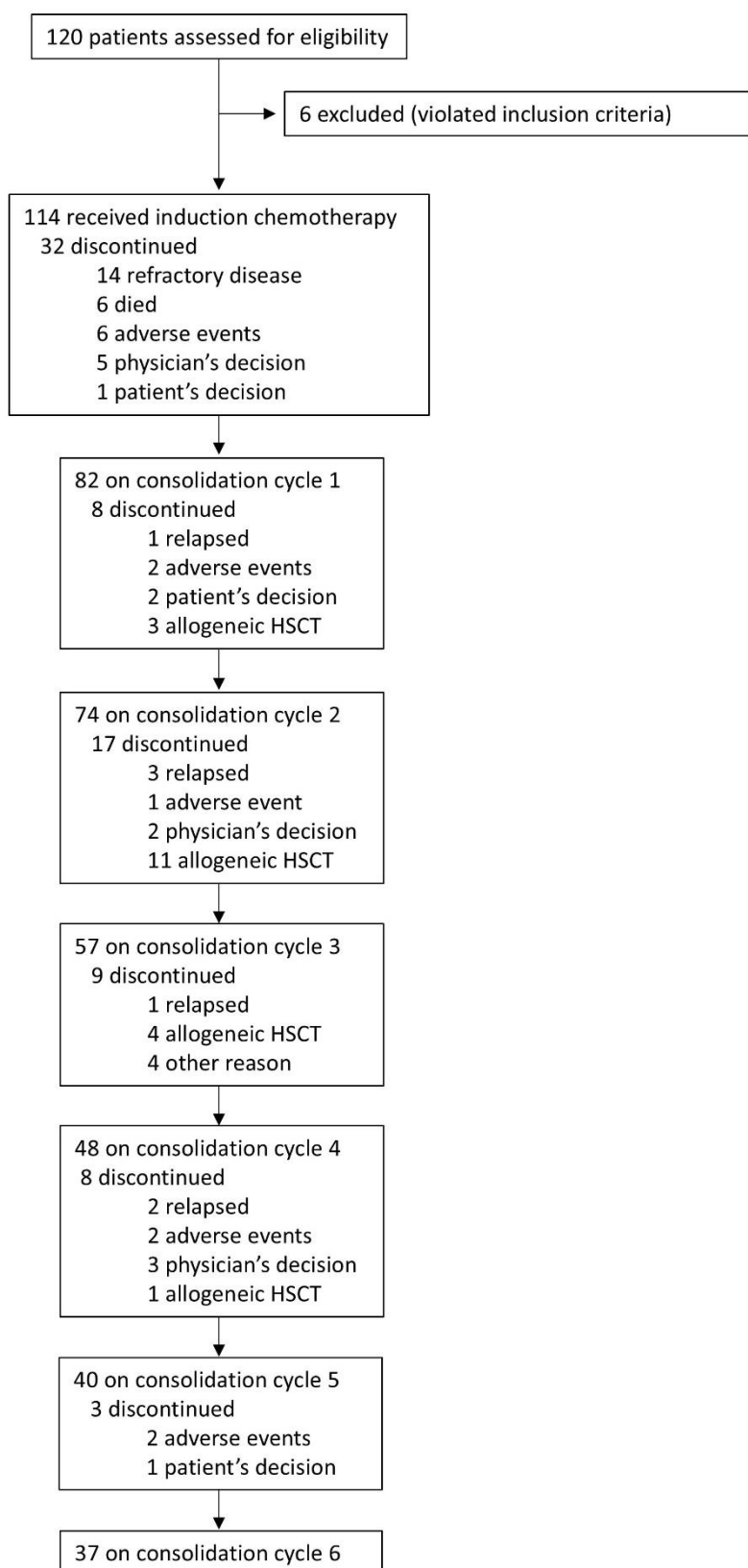
Supplementary table 1: Hematologic toxicity and adverse events during induction chemotherapy*

	DEXAML-02
	N=114
Number of days with platelets < 10 x 10⁹/L	
Median	10·5
IQR	1·50; 19·00
Number of days between day 1 of induction and day when ANC ≥ 0·5 x 10⁹/L	
Median	26
IQR	24; 30
Number of RBC transfusions during induction	
Median	9
IQR	6; 13
Number of platelets transfusions during induction	
Median	9·5
IQR	7; 15
Number of days with fever ≥ 38 degrees C	
Median	6
IQR	3; 11
Number of days with antibiotics	
Median	23
IQR	18; 29
Infections, n (%)	
Documented infection	65 (57%)
Grade 3-4	57 (50%)
Septic shock	3 (3%)
Grade 3-4 cutaneous, n (%)	6 (5%)
Grade 3-4 mucositis, n (%)	20 (18%)
Grade 3-4 haemorrhage, n (%)	5 (4%)
Grade 3-4 nausea or diarrhea or vomiting, n (%)	10 (9%)
Grade 3-4 pain, n (%)	4 (4%)
Grade 3-4 constipation , n (%)	2 (2%)
Grade 3-4 pulmonary events, n (%)	8 (7%)
Grade 3-4 arrhythmia, n (%)	2 (2%)
Grade 3-4 other cardiac toxicity, n (%)	7 (6%)
Grade 3-4 central nervous system, n (%)	3 (3%)
Grade 3-4 peripheral nervous system, n (%)	1 (1%)
Grade 3-4 alkaline phosphatases increased, n (%)	4 (4%)
Grade 3-4 ASAT increased, n (%)	4 (4%)
Grade 3-4 ALAT increased, n (%)	5 (4%)
Grade 3-4 bilirubin increased, n (%)	7 (6%)
Grade 3-4 creatinine increased, n (%)	1 (1%)
Grade 3-4 hyperglycemia, n (%)	12 (11%)
Grade 3-4 lipase increased, n (%)	1 (1%)
Grade 3-4 amylase increased, n (%)	1 (1%)

N=number. ANC=absolute neutrophil count. IQR=inter-quartile range. RBC=red blood cell. C=Celsius. NA=not applicable.

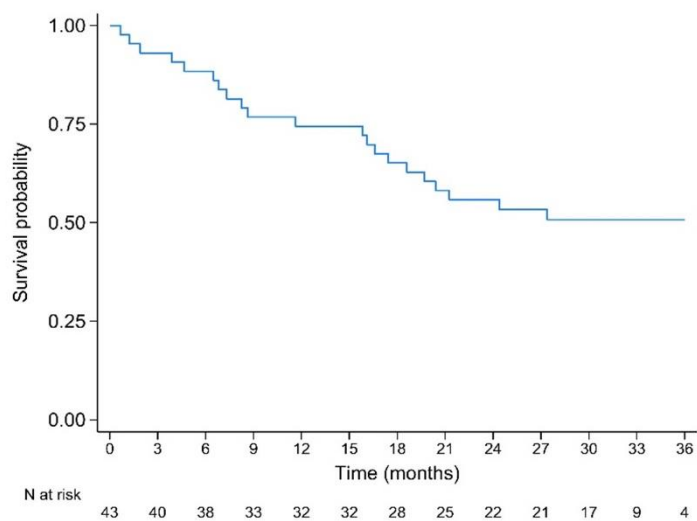
*Investigators were requested to monitor for and report adverse events as defined by the National Cancer Institute Common Terminology Criteria for Adverse Events (version 4.0). Safety reports were monitored by the sponsor.

Supplementary figure 1: Consort diagram

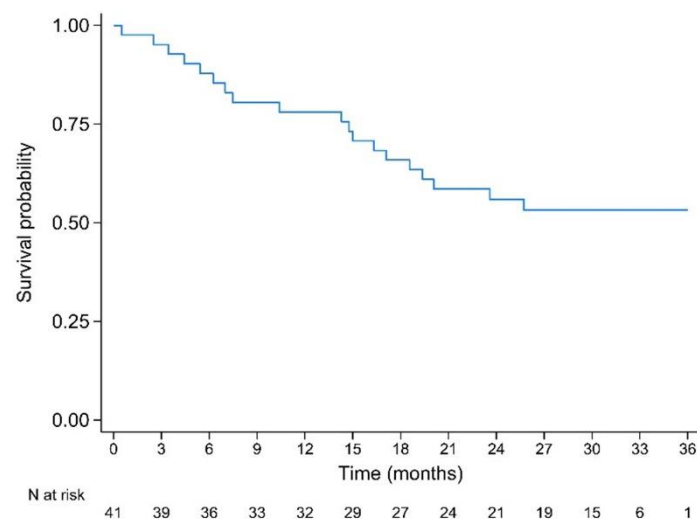


Supplementary figure 2: Event-free survival, relapse-free survival and overall survival in patients with *NPM1* mutations. A. Event-free survival. B. Relapse-free survival. C. Overall survival.

A



B



C

