

Supplemental Online Content

Sanfilippo R, Hayward RL, Musoro J, et al. Activity of cabazitaxel in metastatic or inoperable locally advanced dedifferentiated liposarcoma: a phase 2 study of the EORTC Soft Tissue and Bone Sarcoma Group (STBSG). *JAMA Oncol*. Published online August 18, 2022. doi:10.1001/jamaoncol.2022.3218

eFigure A. Patient disposition for the Phase II trial of cabazitaxel

eFigure 1. Time to progression

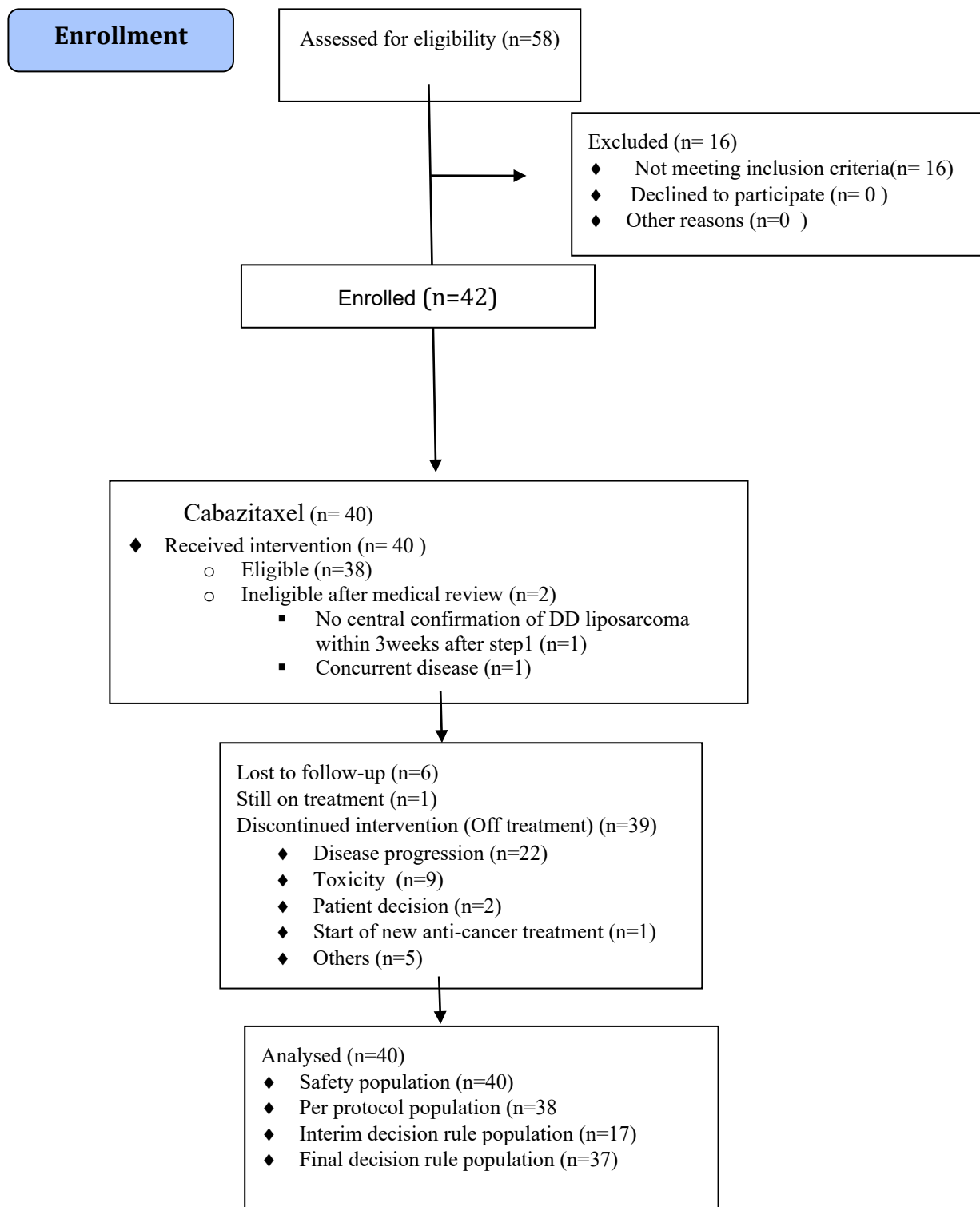
eFigure 2. Overall survival

eTable 1. Summary of main clinical efficacy endpoints

eTable 2. Cabazitaxel-related clinical AEs (all grades) occurring in $\geq 10\%$ of patients in the safety population

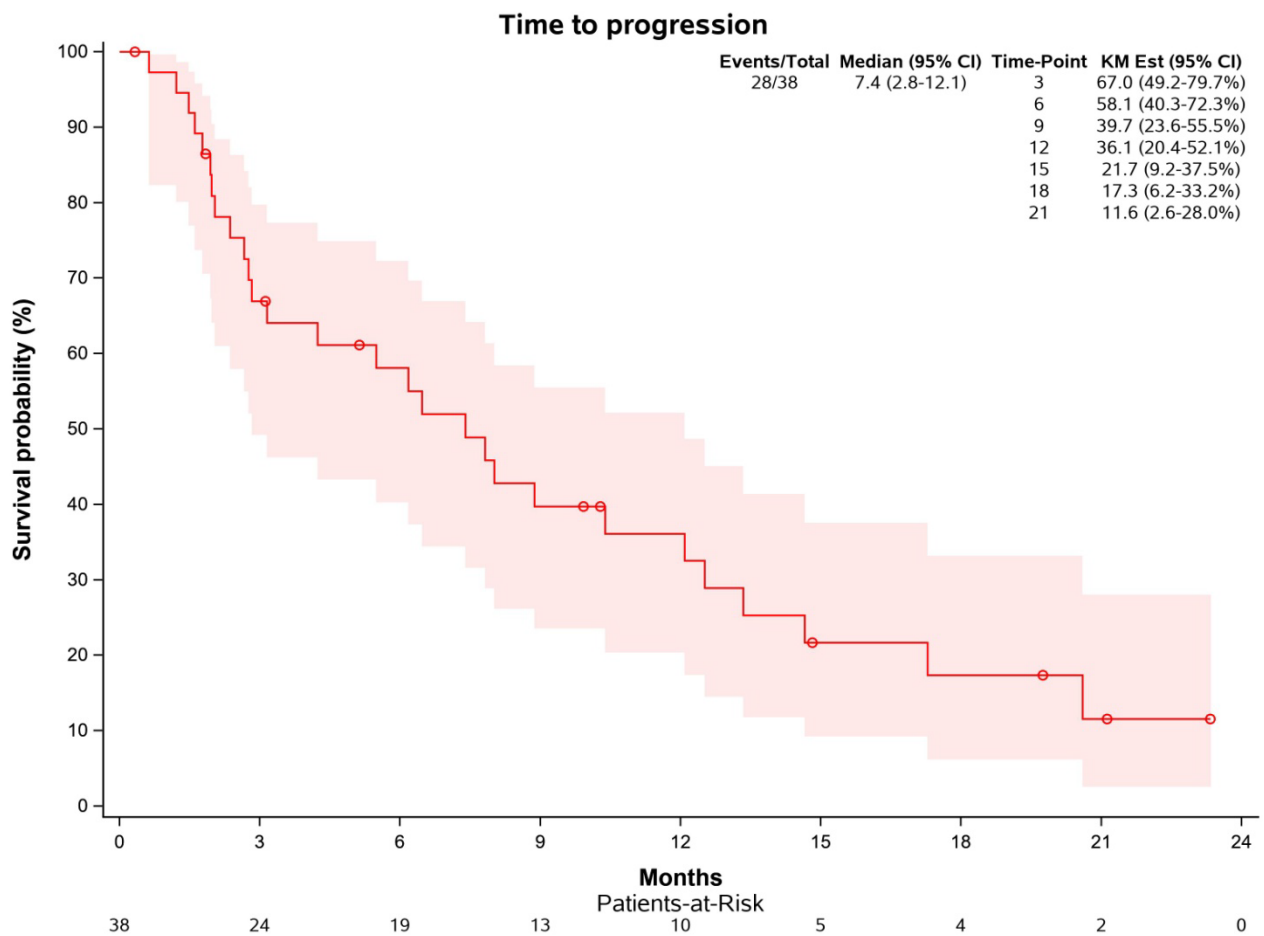
This supplemental material has been provided by the authors to give readers additional information about their work.

eFigure A.

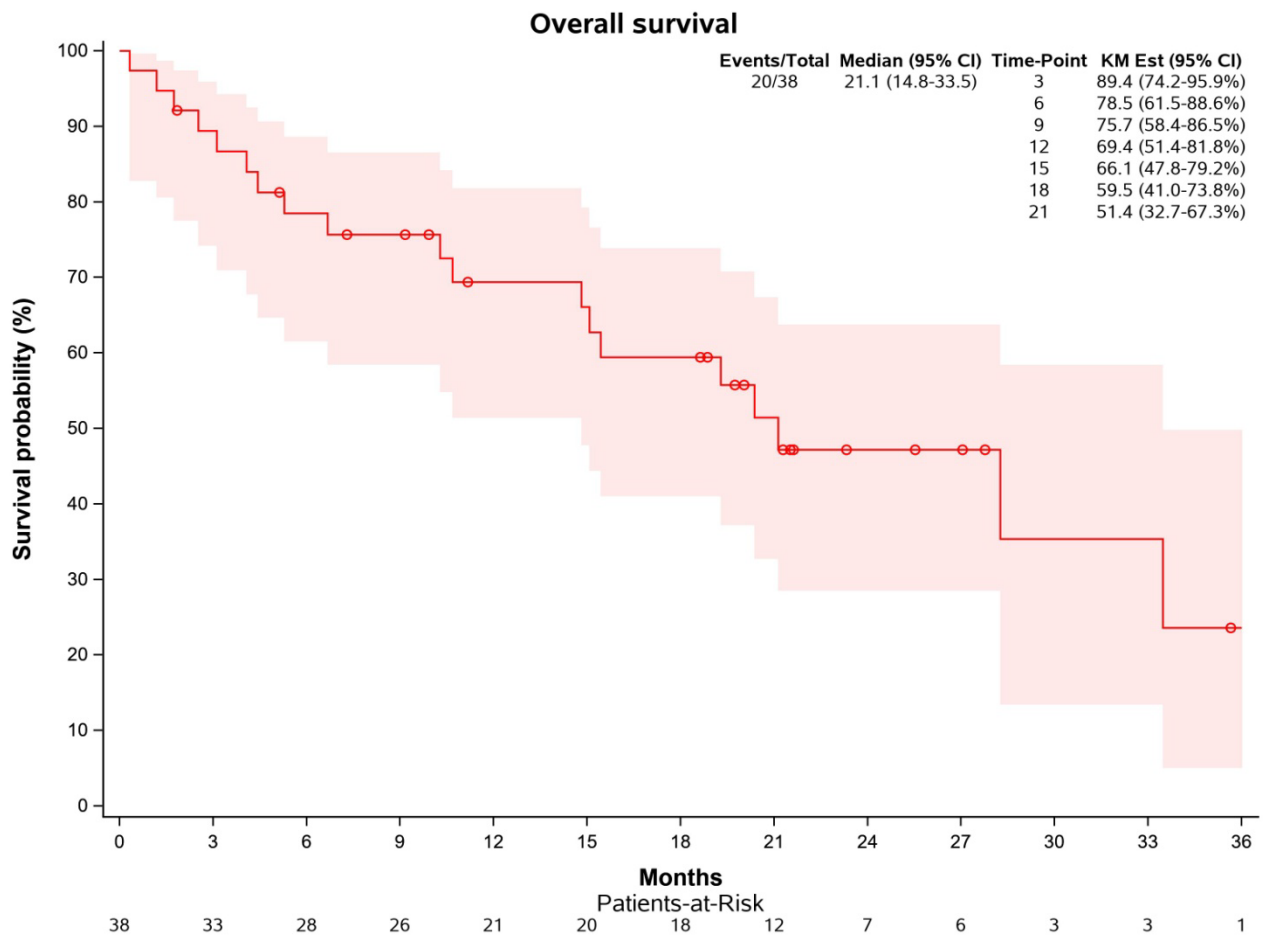


eFigure A: Patient disposition for the Phase II trial of cabazitaxel in metastatic or inoperable locally advanced dedifferentiated liposarcoma.

eFigure 1



eFigure 2



SUPPLEMENTARY MATERIAL			
eTable 1. Summary of main clinical efficacy endpoints.			
	Per protocol population (n=38) n(%)	Final decision rule population (n=37) n(%)	Interim decision rule population (n=17) n(%)
<u>Progression free survival at 12 weeks (binary)</u>			
Progression free at 12 weeks (binary)			
Success (%) [conditional 1-sided 95 % CI]	21 (55.3) [0.408 - 1]	21 (56.8)	11 (64.7)
Response at 12 weeks			
Stable	21 (55.3)	21 (56.8)	11 (64.7)
Progression	12 (31.6)	11 (29.7)	3 (17.6)
Early death	1 (2.6)	1 (2.7)	1 (5.9)
Not evaluable	4 (10.5)	4 (10.8)	2 (11.8)
<u>Objective tumor response rate</u>			
Best overall response			
Complete response	1 (2.6)		
Partial response	2 (5.3)		
Stable disease	22 (57.9)		
Progressive disease	11 (28.9)		
Early death	1 (2.6)		
Not evaluable	1 (2.6)		
Objective response (CR+PR)			
CR+PR (%) [Exact binomial 95 % CI]	(7.9) [1.7% to 21.4%]		
Median time to onset of response	8 mo (range: 5.1 - 8.3 mo)		
<u>Median survival estimate</u>			
Progression free survival (95% CI)	6.5 mo (2.8-10.3 mo)		
Time to progression (95% CI)	7.4 mo (2.8-12.1 mo)		
Overall survival (95% CI)	21.1 mo (14.8-33.5 mo)		
Abbreviations: ; CI, confidence interval; CR, complete response; PR, partial response; Mo, months			

SUPPLEMENTARY MATERIAL**eTable 2:** Cabazitaxel-related clinical AEs (all grades) occurring in $\geq 10\%$ of patients in the safety population.

	Grade 1 N (%)	Grade 2 N (%)	Grade 3 N (%)	Grade 4 N (%)	Grade 5 N (%)	Grade ≥ 1 N (%)
All AEs	4 (10.0)	3 (7.5)	13 (32.5)	17 (42.5)	1 (2.5)	38 (95.0)
Anemia		8 (20)	4 (10)			12 (30)
Febrile Neutropenia			9 (22.5)	1 (2.5)		10 (25)
Abdominal Pain		5 (12.5)	1 (2.5)			6 (15)
Constipation	3 (7.5)	1 (2.5)				4 (10)
Diarrhea	11 (27.5)	3 (7.5)	3 (7.5)			17 (42.5)
Dyspepsia	4 (10)	2 (5)				6 (15)
Nausea	10 (25)	6 (15)	1 (2.5)			17 (42.5)
Vomiting	3 (7.5)	2 (5)	1 (2.5)			6 (15)
Fatigue	6 (15)	5 (12.5)	5 (12.5)			16 (40)
Fever	4 (10)	1 (2.5)		1 (2.5)		6 (15)
Pain	3 (7.5)		1 (2.5)			4 (10)
Alanine Aminotransferase Increased	3 (7.5)	1 (2.5)				4 (10)
Lymphocyte Count Decreased	1 (2.5)	3 (7.5)				4 (10)
Neutrophil Count Decreased		1 (2.5)	4 (10)	16 (40)		21 (52.5)
Weight Loss	7 (17.5)	4 (10)				11 (27.5)
White Blood Cell Decreased		1 (2.5)		3 (7.5)		4 (10)
Anorexia	8 (20)	4 (10)	1 (2.5)			13 (32.5)
Arthralgia	1 (2.5)	3 (7.5)				4 (10)
Dysgeusia	5 (12.5)		1 (2.5)			6 (15)