Clinician-reported event	Baseline (N=20)	3 months (N=20)	6 months (N=20)	12 months (N=10)	p
Abdominal pain	-	20%	25%	30%	0.05
Alopecia	-	5%	15%	10%	0.40
Arthralgia	40%	65%	60%	30%	0.20
Asthenia/Fatigue	50%	90%	80%	80%	0.04
Constipation	25%	35%	45%	20%	0.48
Decreased appetite	15%	50%	50%	30%	0.06
Diarrhoea	-	5%	20%	50%	<0.01
Dysphonia	60%	55%	40%	40%	0.59
Dysgeusia	5%	30%	30%	20%	0.14
Hand-foot syndrome	-	30%	55%	40%	<0.01
Headache	30%	15%	20%	10%	0.63
Myalgia	35%	55%	60%	40%	0.42
Nausea	5%	15%	25%	10%	0.39
Peripheral edema	5%	5%	15%	10%	0.78
Rash	-	10%	-	10%	0.22
Stomatitis	5%	30%	25%	20%	0.18
Xerostomia	10%	55%	25%	60%	<0.01
Vomiting	-	5%	15%	10%	0.40

Supplementary Table 1. Clinician's assessment of symptomatic events (all grades) at baseline and at 3, 6 and 12 months after the start of treatment. Only AEs included in the PRO-CTCAE questionnaire were reported.

Abbreviations: AEs, adverse events; PRO-CTCAE, Patient-Reported Outcomes Version of the Common Terminology Criteria for Adverse Events