S1 Table A. Serious Adverse Events

Study treatment	Adverse Event	MedDRA Preferred Term	NCI-CTCAE grade	Outcome	Action Taken
Ibuprofen[a]	Severe induration	Injection site induration	3	Resolved	None
Ibuprofen[a]	Respiratory infection	Respiratory tract infection	3	Resolved	None
Placebo[b]	Headache	Headache	3	Resolved	Permanently discontinued

Safety Population. All events were catalogued as unrelated to the study as regards to causality. [a] patient from abdominal surgery; [b] patient from orthopaedic surgery.