End of Study report

I General information	
EudraCT number	2014-003303-30
NL number	NL50440.078.14
Title: *Thromboprophylaxis in Children treated for Actule Lymphoblastic Leukemia with Low molecular-weight heparin: a randomized controlled trial (TropicALL)*	
Sponsor	Stichting Kinderoncologie Nederland (SKION)
Participating sites	☐ singlecenter:
	☒ multicenter, national
Principal Investigator	Dr. C.H. van Ommen / Dr. M.D. van de Wetering

II Study status	
☐ Terminated as planned	End date of study:
☒ Early Termination	End date early termination: 18-DEC-2018
Reason for early termination: *After 3.5 years inclusion no subjects included. There are competitive trials in the same population and trials with the same primary endpoint. We do not expect improvement of inclusion rate the next few year and will not reach the inclusion targets.*	
How many participants were included at the time it was decided to terminate the study early? What are the consequences for these participants? *No participants were included in the study, no consequences for participants.*	
What are the consequences of the early termination for the study results? *No statistical analysis on these results is feasible. We won't be able to answer the primary and secondary research questions of the study.*	

III Status inclusion	
How many participants are included in the study?	0
What is the total number of participants that needed to be included according to protocol?	354
What is the cause for not reaching the intended number of participants? *There are competitive trials in the same population and trials with the same primary endpoint.*	

IV Report drafted by	
Name and function:	J. Mur, Trialmanager
Date report	04-MAR-2021