

End of Study report

I General information	
EudraCT number	2017-003860-11
NL number	NL60690.029.017
Title: <i>11C-erlotinib brain uptake in Diffuse Intrinsic Pontine Glioma imaged by PET</i>	
Sponsor	VU University Medical Center
Participating sites	<input type="checkbox"/> singlecenter:
	<input type="checkbox"/> multicenter, national
Principal Investigator	Dr. D.G. van Vuurden

II Study status	
<input type="checkbox"/> Terminated as planned	End date of study:
<input checked="" type="checkbox"/> Early Termination	End date early termination: 01-06-2018
Reason for early termination: <i>Moving of the pediatric oncology department from the Amsterdam UMC, location VUmc to the Princess Maxima Center in Utrecht due to centralization of pediatric oncology care in the Netherlands. ¹¹C-Erlotinib has a short half-life, making it logistically impossible to be administered at any other hospital than at the Amsterdam UMC, VUmc.</i>	
How many participants were included at the time it was decided to terminate the study early? What are the consequences for these participants? <i>No participants were included in the study, therefore there were no consequences for participants.</i>	
What are the consequences of the early termination for the study results? <i>This study was never formally open for inclusion. No data was collected, we won't be able to answer the primary and secondary research questions of the study.</i>	

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?	5-15
What is the cause for not reaching the intended number of participants? <i>This study was never formally open for inclusion.</i>	

IV Report drafted by	
Name and function:	Prof. Dr. G.J.L. Kaspers, Head of Pediatric Oncology at VUmc
Date report	19-11-2023
Signature	