Prematurely ended-statement

EudraCT number	2008-007031-41
Full title of the study	Beta-Blocker in Acute Ischemic Stroke – a prospective, randomized, double-blinded, placebo-controlled safety and efficacy trial of early treatment
Sponsor ID	BIAS
Study Contact	Prof. Dr. Wilhelm Haverkamp PD Dr. Benjamin Hotter
Sponsor	Center for Stroke Research Berlin, Charité - Universitätsmedizin Berlin
Contact email address	benjamin.hotter@charite.de
Product	Propranolol
Date of the early termination of the trial	31 – AUG - 2012
Statement on discontinuation of the study	Study prematurely ended due to lack of recruitment. Only 20 of 250 planned patients were included in the trial. A formal statistical evaluation could not be conducted due to the very low recruitment.
Publication	Hotter B., Jegzentis K., Steinbrink J., Schmidt W.U., Endres M., Meisel A., Haverkamp W., Jungehulsing G.J. Impact of Selection Criteria on Recruitment in an Interventional Stroke Trial. Cerebrovasc Dis 2013, 36:344-350

Signature: Prof. Dr. Andreas Meisel

Job role: Director of Sponsor Center for Stroke Research Berlin