Boehringer Ingelheim

Clinical Study Synopsis for Public Disclosure

This clinical study synopsis is provided in line with **Boehringer Ingelheim's *Policy on Transparency and Publication of Clinical Study Data***.

The synopsis - which is part of the clinical study report - had been prepared in accordance with best practice and applicable legal and regulatory requirements at the time of study completion.

The synopsis may include approved and non-approved uses, doses, formulations, treatment regimens and/or age groups; it has not necessarily been submitted to regulatory authorities.

A synopsis is not intended to provide a comprehensive analysis of all data currently available regarding a particular drug. More current information regarding a drug is available in the approved labeling information which may vary from country to country..

Additional information on this study and the drug concerned may be provided upon request based on **Boehringer Ingelheim's Policy on Transparency and Publication of Clinical Study Data**.

Name of Company: Boehringer Ingelheim		**Statement on discontinuation of the study**	
BI Proprietary Name: NA		**EudraCT No.:** 2015-004625-14	
BI Investigational Product: Volasertib (BI 6727)		**Page:** 1	
Report Date: NA	**Trial No. / Doc. No.:** 1230.28	**Dates of Trial:** NA	**Date of Revision:** NA

Title of Trial:	Open-label, dose-escalating trial to evaluate the tolerability, toxicity, safety, pharmacokinetics, pharmacodynamics and activity of volasertib added to the standard intensive salvage chemotherapy regimen with liposomal daunorubicine, fludarabine and cytarabine (DNX-FLA) followed by fludarabine and cytarabine (FLA) in children from 3 months to less than 18 years of age with acute myeloid leukaemia after failure of the front-line therapy
Trial Sites:	NA
Publications:	NA
Clinical Phase:	I
Statement on discontinuation of the study:	Discontinued by Boehringer Ingelheim during preparation of the trial. No patient entered the study, therefore no results / data are available.