



Clinical Study Synopsis for Public Disclosure

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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**.

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Name of company: Boehringer Ingelheim		Statement	
Name of finished product:			
Name of active ingredient: BI 691751		Page: 1	
Report date:	Trial-Number: 1334.2	Study period (dates):	
Proprietary confidential information			
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Title of study:	Safety, tolerability, pharmacokinetics and pharmacodynamics of multiple rising oral doses of BI 691751 in healthy male subjects
Investigators	[REDACTED]
Study center(s):	[REDACTED] Germany
Publication (reference):	N/A
Clinical phase:	Phase I
Statement on discontinuation of the study:	The disease indication terminated by Boehringer Ingelheim International GmbH for strategic reasons