



# STATEMENT OF GLP COMPLIANCE

Registration number: 2/2010/GLP

## Assessment of conformity with GLP according to the Directive 2004/9/EC of the European Parliament and of the Council

On the basis of the inspection which was held on 25<sup>th</sup> and 26<sup>th</sup> November 2009, and on 26<sup>th</sup> January 2010 in accordance with criteria specified in the Order of the Inspector for Chemical Substances and Preparations of 24<sup>th</sup> November 2003 concerning the rules on control and verification of compliance with principles of Good Laboratory Practice, as well as in accordance with Directive 2004/9/EC and the relevant OECD regulations, the Inspector for Chemical Substances and Preparations hereby confirms that

### PHARMACOLOGY DEPARTMENT PHARMACEUTICAL RESEARCH INSTITUTE

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Poland

complies with the OECD and the EU principles of Good Laboratory Practice in the fields of:

**analytical phase of pharmacokinetic studies including bioavailability and bioequivalence studies**

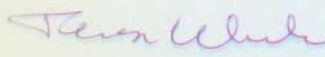
The Certificate is valid from: 12<sup>th</sup> January 2010

to: 12<sup>th</sup> January 2012

Łódź, date: 5<sup>th</sup> February 2010

Good Laboratory Practice  
Inspectors:

  
Mirosław Muszyński

  
Tomasz Wasiela

Inspector for Chemical Substances  
and Preparations:



  
Jerzy Majka