



## Statement of GLP Compliance

According to Article 14 paragraph 3 Ordinance on Good Laboratory Practice [OGLP, SR 813.112.1]

The notification authority for chemicals confirms that the following test facility was inspected with respect to the compliance with the Swiss Ordinance on Good Laboratory Practice, adopted on 18th May 2005 [OGLP, SR 813.112.1]. This Ordinance is based on the OECD Principles of Good Laboratory Practice, as revised in 1997 and adopted on 26th November 1997 by decision of the OECD Council [C(97)186/Final].

Unequivocal name and address  
of the test facility:

Areas of expertise according to  
article 3 paragraph 1 letter d OGLP:

Institute for Biopharmaceutical  
Research (IBR) Inc.  
Lauchefeld 31

2. mutagenicity studies,  
8. analytic and clinical chemistry testing.

9548 Matzingen, Switzerland

Inspection authority: Swiss Agency for Therapeutic Products (Swissmedic)

Date of inspection: 1<sup>st</sup> and 2<sup>nd</sup> September 2008

Date of decision: 7<sup>th</sup> November 2008

Based on the above mentioned decision it can be confirmed that the above mentioned test facility is able to conduct studies according to the aforementioned areas of expertise in compliance with the principles of GLP. The above mentioned test facility is listed in the register and GLP list according to the Article 14 OGLP and is inspected on a regular basis according to Article 6 paragraph 2 OGLP.

Swiss Federal Office of Public Health  
Consumer protection directorate  
Notification authority for chemicals  
CH-3003 Bern



Bern, 12<sup>th</sup> December 2008, The Head, Dr. Dag Kappes.