The Swiss GLP Monitoring Authorities

Swiss Federal Office of Public Health

Swiss Agency for the Environment, Forests and Landscape

Intercantonal Office for the Control of Medicines

Statement of GLP Compliance

It is hereby confirmed that

during the period of

January 25 and 26, and
May 31, 2001

the following Test Facilities of

Institute for Biopharmaceutical Research (IBR), Inc., Matzingen

were inspected by the Intercantonal Office for the Control of Medicines with respect to the compliance with the Swiss legislation on Good Laboratory Practice.

Test Facilities	**Areas of Expertise**
IBR	**ACC**

The inspection was performed in agreement with the OECD Guidelines for National GLP Inspections and Audits. It was found that the aforementioned test facilities were operating in compliance with the Swiss Ordinance relating to Good Laboratory Practice [RS 813.016.5] at the time they were inspected.

Intercantonal Office for the Control of Medicines
The Director

H. Stocker

Bern, June 2001

Dr. Hans Stocker