

## STATEMENT OF GLP COMPLIANCE

Assessment of conformity with Good Laboratory Practice (GLP) according to Directive 2004/9/EC of the European Parliament and the Council of 11 February 2004.

Name of company: **Visionar Preclinical AB**

Site address: Rapskatan 25  
Uppsala

Postal address: P.O. Box 6373  
SE-751 36 Uppsala  
Sweden

According to the criteria specified in Articles 5 to 10 of the Swedish Provision LVFS 1996:10 Visionar Preclinical AB was found to operate in compliance with the principles of Good Laboratory Practice as established by the OECD and the European Community at the time of the inspection.

Date of latest inspection: 26 March, 2009

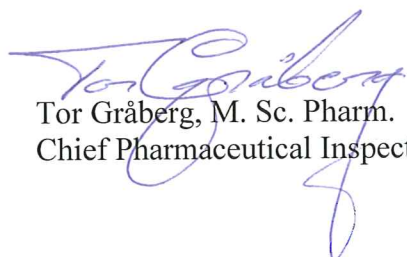
Area of expertise: Toxicity studies (2)


Type of inspection: Facility inspection and study audit

Inspection interval: Every 2 years

Visionar Preclinical AB has to inform the Medical Products Agency if major personnel or organisational changes will take place.

On behalf of the Medical Products Agency

  
Tor Gråberg, M. Sc. Pharm.  
Chief Pharmaceutical Inspector

  
Bengt Berglund, M. Sc. Pharm.  
Pharmaceutical Inspector