**The statutory declaration accompanying an application for an approval of the veterinary non-medicinal product and its entry into the register of approved veterinary non-medicinal products**

*………………………………. (name of the product)* **manufactured or marketed in some EU Member State**

**or any of the Contracting Parties of the Agreement on the EEA**

The applicant for an approval of the product *…………. (Applicant’s identification data)* represented by

Authorised person*………… (*Authorised person *identification data)* declares hereby:

* that this product is marketed in EU Member States,
* that the manufacturer of the product complies with the principles of quality assurance
of manufacturing by following the principles set out in the relevant production standards
and guidelines (ISO, GMP) and has processed in writting, implemented and adhered to system procedures of Hazard Analysis and Critical Control Points (HACCP); that this product complies with the requirements of the EU regulations on general safety and TSE/BSE safety.

In………..……, Date…….………. ..……….…………………………………………………………

 Signature of the applicant or authorised person