**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)*

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

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

* that the manufacturer of the product complies with the principles of quality assurance
of manufacturing by following the principles set out in good manufacturing practice as described in the relevant production standards and guidelines (ISO, GMP) and has processed in writing, implemented and adhered to system procedures of Hazard Analysis and Critical Control Points (HACCP); that the raw materials used for manufacturing of the product are not of ruminant origin; (or if so, an evidence that the animals were free from BSE / TSE should be provided),
* that the product has been assessed in term of its declared efficacy and is efficient if used according to the instruction in the package leaflet,
* that the product, including the packaging material, has been assessed for safety and is safe for animal health and for the health and safety of persons handling it, for property and the environment,
* that, provided that the prescribed storage conditions and handling of the product are observed, the product is stable and retains its declared composition and characteristics for its entire intended shelf life.

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

 Signature of the applicant or authorised person