The following areas are covered by documents provided in this section:

- manufacture of veterinary medicinal products including autogenous vaccines
- manufacture of active substances
- activities of control laboratories for medicinal products.

This sections is divided into three parts:

- GMP Guidance documents (EU GMP Guide) and other guidance documents published by the ÚSKVBL according Act No. 378/2007 Coll., on pharmaceuticals
- Forms and templates
- Registers and databases (actual registers of manufacturers, control laboratories including scope of activities, register of API manufacturers in the Czech Republic)