On the basis of obligations arising from § 23 par. 1 let. d) and § 77 par. 1 let. f) of the Act on Medicinal Products No. 378/2007 Coll., as amended, USKVBL in this section more detailed information and documents concerning the reporting of the volume of distribution (reporting of consumption of VMP). Reports are sent by VMP distributors, manufacturers of medicinal products who, pursuant to Section 75 (4) of Act No. 378/2007 Coll. on Medicinal Products authorized to act as a distributor in the case of medicinal products manufactured or imported from third countries, as well as manufacturers and distributors of medicated feedingstuffs. In addition to the Basic Information, there is a Manual, which describes in detail two possible ways of reporting VMP consumption, a form for registration in the VMP consumption reporting system and sample MS Excel reports, used to send VMP consumption reports in the form of an Excel workbook.

- Distribution volume reporting - basic information
- Manual
- Registration form
- MS Excel report (without medicinal products)
- MS Excel report (contains all registered VMP packages)
- MS Excel report (contains all approved exceptions)