Strategic consultancy

- regulatory strategies for early projects: "from transcription to prescription"
- development planning: pharmacology, toxicology, clinical development,
- scientific advise with EMEA, FDA and national bodies
- orphan drug applications with EMEA and FDA, SME applications
- regulatory intelligence in / for your company
- CE-certification of medicinal products according to EU Medicinal Products Acts
- contacts to / meetings with European, national and local / state authorities
- evaluation of R&D portfolios with focus on scientific value, developmental planning, regulatory compliance

Do not hesitate to ask us for special solutions and further assistance.