



All dear,

TAURET s.r.o. is a young, dynamic, independent and highly client oriented firm, created in 2003.

The company's mission is to provide professional and specialised advice and healthcare regulatory affairs consultancy and services to the domestic and international pharmaceutical industry.

Main goals are helping our customers in getting their products to the market economically and quickly, ensuring the highest level of performance and patient efficacy.

Consulting areas:

Ⓞ Drug registration services

- presubmission and other agency interaction and consultancy
- coordination and preparation of documents for submission to Regulatory Authorities
- product submission and registration process management
- maintenance up to the end of product life cycle /renewals and variations/
- PIL, SmPC and labelling in local language /Czech and Slovak/
- clinical trials submission and approval management
- expert report writing services /pharmaceutical, preclinical and clinical/

Ⓞ Regulatory Affairs consultancy

- market entry strategy development /recomendation of strategies to effect the earliest possible approval of regulatory applications/
- negotiation with Drug Regulatory Authorities
- impacts of new regulations and EU accesion /serving as informational resource, regulatory analysis and evaluation the impact of trends, relative to government regulatory activities
- current regulatory perspectives /ensure compliance with Drug Regulatory Authority requirements/
- pricing and reimbursement services

If you are interested in our services do not hesitate to contact us.

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