

Get your added value!

Pharma Business Development



Your Gateway to Swiss Technology

Within the framework of an API Project,
Dolder Pharma Business Development offers comprehensive services:

Regulatory Support

Evaluation and elaboration of Drug Master Files (DMF), Certificates of Suitability (CEP) and Quality Overall Summary (QOS)
Escort of registration procedures

Patent Investigations

Evaluation of patents in terms of potential infringements, supported by patent lawyers registered with the European Patent Office (EPO)

GMP Audit Reports

Reliable inspection by means of 3rd party audits conducted by auditors of the APIC audit team

Analytical Support

Structure elucidation of unknown impurities
Method development and transfer

Reference Substances

Supply of substances listed in leading pharmacopoeas, though not available through EDQM and USP

Polymorphy (in close cooperation with external partners)

Identification of polymorphic forms
Evaluation towards patented polymorphs

www.dolder.com/pbd



To maintain and continuously improve our high level of quality,
we are certified according to ISO 9001 and GDP.



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