Clarivate™

Cortellis CMC Intelligence

Increase submission rates and avoid costly delays

Developing a drug is difficult. Going to market shouldn't be.

Your trusted partner in planning CMC (chemistry, manufacturing, and controls).

Cortellis CMC Intelligence™ is a comprehensive database that organizes global CMC regulations and highlights any variances in local practices to provide:

- Content coverage for both biologics and small molecules dosage forms
- Curated submission procedures with estimated and official timelines
- The ability to automate the process of CMC regulatory monitoring through user configured alerts
- Detailed summaries (in English) from experts in local regulatory practices paired with reference source documents to provide a complete picture
- Capabilities to efficiently compare regulations across countries, territories, and organizations — including the ability to export vital comparison tables

Trust a CMC solution, compiled by specialized industry experts, to prepare your dossiers accurately and swiftly.

Robust data:

5K+

Source documents

64

Territories, countries, and regions for biologics content

900+

links to Cortellis Regulatory Intelligence, providing access to expanded detail

137

Countries, territories, and organization specific regulations for small molecule content

Small molecule content covers:

- Cream
- Drops
- Inhaler
- Liquid injectable
- Liquid oral
- Powder
- Solid oral
- Solid, modified release
- Spray

Biologics content covers:

- Biosimilars
- Blood derivatives
- Cell therapy
- Drug – devices combination products
- Gene therapy
- Monoclonal antibodies
- Recombinant hormones and proteins
- Vaccines
- Tissue Therapy

XBU942982079 / 07

Make better decisions and accelerate innovation

Contact a representative to learn how Cortellis CMC Intelligence can accelerate innovation for your organization, or visit:

clarivate.com/cortelliscmc