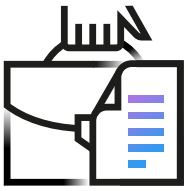
Navigate global guidelines and efficiently drive strategic decisions

Clarivate™

Be compliant, increase approval rates and expand to new markets

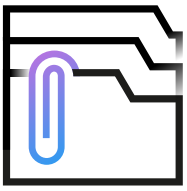
Staying current with the ever-changing regulatory landscape is challenging, often introducing uncertainty around maintaining compliance. Cortellis Regulatory Intelligence™ is a timely and comprehensive database that spans all regulatory functions across the R&D lifecycle, providing a single point of access to:

Daily updates to maintain network of historic and current source documents without language barriers.



Helpful context from proprietary regulatory intelligence reports to support strategy.

Detailed summaries (in English) of local regulatory practices and reference documents.



Full coverage of FDA AdComm meetings since 2001 and FDA workshops since 2003.

Leverage high quality, comprehensive data supported by extensive domain knowledge and expertise

Robust data:

297K+

Regulatory documents for drugs, biologics, medical devices and in vitro diagnostics (IVDs)

81

Modules for country- and region-specific regulations

15K+

Human English translations

39K+

English machine translations

6K+

Inspection documents

Expert team:

Geographically diverse with global breadth and local depth.

Deep knowledge on key issues and trends impacting your submissions.

Translations by regulatory professionals who are also native English speakers.

Manual report curation ensures attention to detail and high quality.

"Cortellis Regulatory Intelligence supports strategic decision making and submission preparation, and avoids the need to review multiple data sources and handle translations internally."

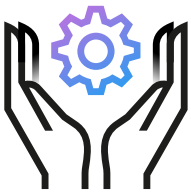
Regulatory Professional, mid-size pharma

Unlock the hidden insights in data

Increase the chances of approval and rapidly expand to new markets using our comprehensive, timely, accurate, central source of regulatory information including FDA483s, approval documents, inspection reports and submission forms.

Reports and analytics

Prepare for inspections, understand approval trends, discern submission needs and more.



Global comparisons

Efficiently compare regulations across countries — all from a single point of database access.

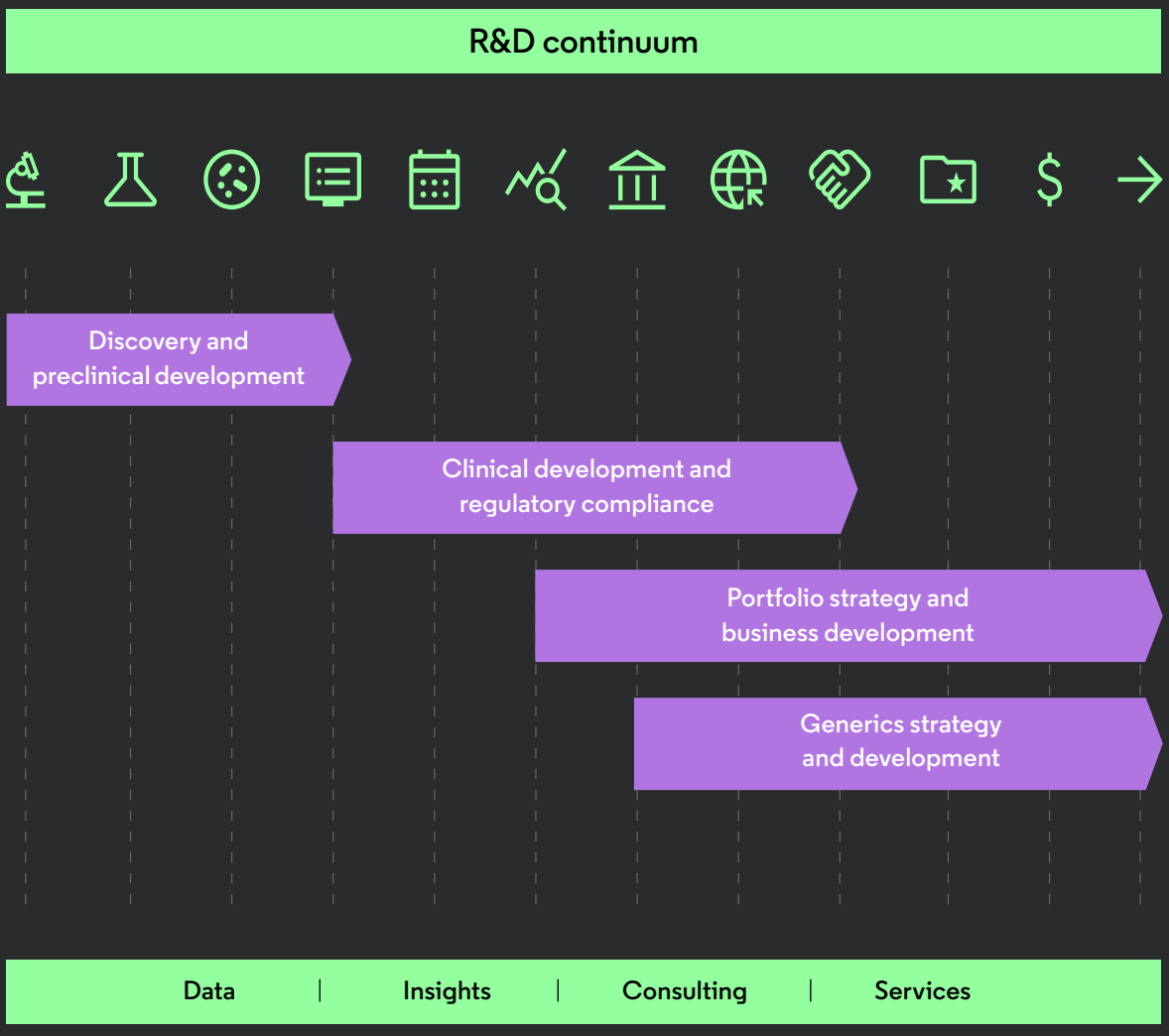
Regulatory summaries

Quickly understand regulatory requirements across all aspects of therapy development.

About Cortellis

Make data-driven decisions with speed and certainty across the drug and device development lifecycle with the Cortellis™ suite of life science intelligence solutions — including customized consulting and services delivered to your specific requirements. Only Cortellis delivers the insights needed to accelerate innovation with confidence.

Figure 1: R&D continuum



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About Clarivate

Clarivate™ is a leading global provider of transformative intelligence. We offer enriched data, insights & analytics, workflow solutions and expert services in the areas of Academia & Government, Intellectual Property and Life Sciences & Healthcare. For more information, please visit **clarivate.com**.

Make better decisions and accelerate innovation

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

clarivate.com/CortellisRegulatoryIntelligence

Contact our experts today:

clarivate.com

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