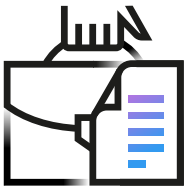
Navigate global guidelines and efficiently drive strategic decisions

Clarivate

Be compliant, increase approval rates and expand to new markets

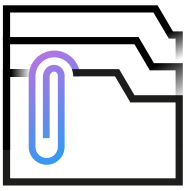
Staying current with the ever-changing regulatory landscape is challenging, often introducing uncertainty around maintaining compliance. Cortellis Regulatory Intelligence is a timely and comprehensive database that spans all regulatory functions across the R&D lifecycle, providing a single point of access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

310K+

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

81

Modules for country- and region-specific regulationss

15K+

Human English translations

40K+

English machine translations

16K+

Compliance and inspection documents

2K+

Proprietary regulatory summaries

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 has provided our organization with the necessary insights to confidently comply with regulatory guidelines. It's used every day to keep us up to date and stay compliant."

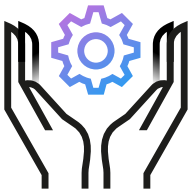
Caroline Alba,
Regulatory Professional, Paion.

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.

Coming soon

AI-powered Regulatory Assistant

The assistant will boost your efficiency by guiding you to the right actions and save you time performing day-to-day tasks.

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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