

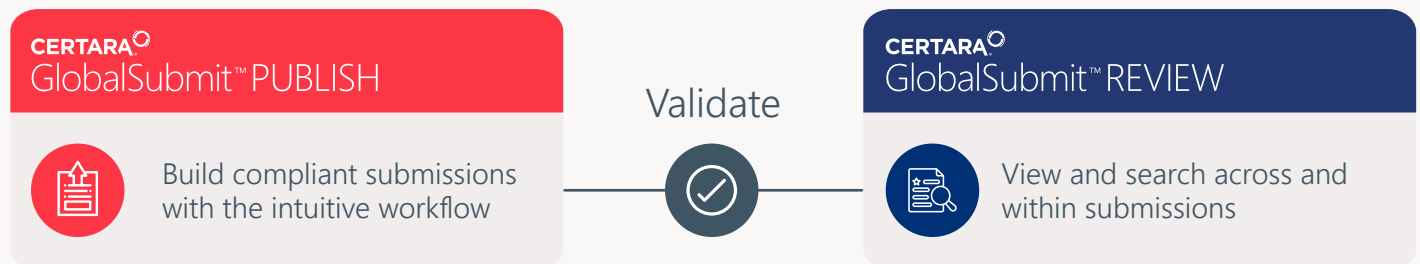
GlobalSubmit™ eCTD Platform

Simple. Efficient. Transparent.

Certara's GlobalSubmit eCTD Platform is a submission software that provides regulatory teams the tools they need to efficiently publish, validate, and review eCTD submissions. This web-based, cloud native and 21 CFR Part 11 compliant software allows you to seamlessly manage electronic submissions.

Simplify complex regulatory processes

GlobalSubmit enables teams to manage the regulatory process across a global footprint of regulatory health authorities, so they can accelerate submission timelines while reducing risk with this easy-to-use publishing tool and global reviewer.

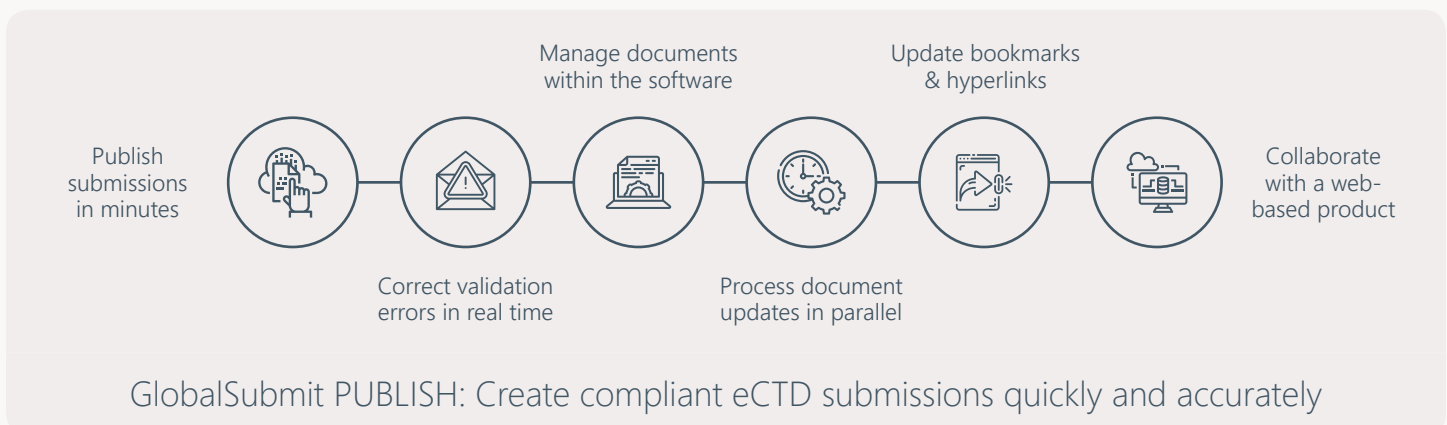


CERTARA[®] GlobalSubmit™ PUBLISH

PUBLISH is a desktop software that enables you to create submissions that align with a health authority's required structure, XML, and validation report.

Eliminate
≥4
publishing tasks
per document

Hyperlink and
bookmark QC
8x
faster compared with
manual methods

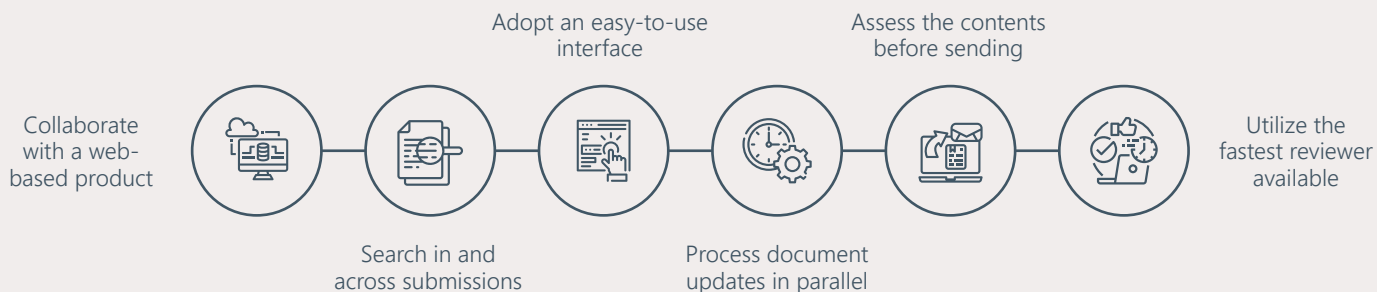


CERTARA[®] GlobalSubmit™ REVIEW

REVIEW is a web-based software that enables you to facilitate timely reviews of eCTD submissions across stakeholders prior to submitting to a regulatory health authority.

Utilize
200+
error conditions including
40+ PDF checks

Load Submissions
50%+
faster reducing
wait time



GlobalSubmit REVIEW: Simplified eCTD submission review

United States (FDA)
European Union (EMA)
China (NMPA)
Japan (PMDA)
Canada (HC)
Brazil (ANVISA)
Australia (TGA)
South Africa (MCC)
Switzerland (CH)
Gulf Cooperation Council (GCC)



Simplify your regulatory process with GlobalSubmit™
Learn more and request a demo certara.com/globalsubmit



About Certara

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