

DIGITAL TRANSFORMATION FOR REGULATED INDUSTRIES CREATING THE CONNECTED CONTENT NETWORK FROM AUTHORING TO GLOBAL SUBMISSIONS DOSSIER COMPILATIONS

In regulated industries, information is the source of innovation, and many content and information sources feed the Research, Development and Submissions pipeline. Digital Transformation, from content creation to compiling traceable, global submissions dossiers requires solutions with business agility in mind. Our platform and solutions deliver consistent, contextual, traceable, and connected content through this entire lifecycle to accelerate time-to-market.

ABOUT US

InteliNotion is a state-of-the-art, Cloud native, highly secure and scalable Software-as-a-Service (SaaS) solution, built using the most advanced Web technologies on the leading trusted Enterprise Cloud platform. We deliver a new generation of groundbreaking solutions for Structured Content Authoring, Component Content Management and Enterprise Information Management (EIM) for highly regulated industries.

BUSINESS CHALLENGES

During the product research and development lifecycle, a large variety of documents that describe the product design and definition, analysis plans, results, and safety and summary reports are developed and submitted to regulatory health agencies for approval to market. Today, the information created, shared, reviewed, and approved is captured in unstructured documents developed throughout the product research and development phases, which can span many years. A large percentage of these documents are made up of reusable sub-components that are generated by experts, scientists, and systems, increasing both the complexity, volume, and consistency of information. The ability to trace the information through the entire product development cycle is also difficult. To ensure quality, consistency, and improved efficiencies, structuring the information within documents into meaningful components and reusing information within and across documents, as well as the compiled global submissions dossiers with full traceability is essential to achieving transformational improvements in capacity and efficiency.

THE *INTELINOTION* PLATFORM

InteliNotion is a feature rich, cloud-based compliance platform used to govern and manage the end-to-end content-oriented, lifecycle driven business processes of regulated products and services. The platform delivers an innovative, configurable, model-driven set of services for component-based collaborative content authoring, review/approvals, anonymization and redaction, conditional publishing for multiple audiences, and global variant management with full traceability of the content through the lifecycle, from creation to global submissions.

APPLICATION MODULES

In partnership with ArborSys, *InteliNotion* delivers a suite of functional modules with a preconfigured baseline set of models, templates, content policies, lifecycles, workflows and content libraries. Our baseline configurations allow clients to rapidly adopt, adjust and deploy capabilities across the enterprise at their own pace. Current Biopharma Industry-specific modules delivered include: Clinical Documentation, Labeling, CMC, and Dossier Management.