



Submission Archive



Publish back to the CARA Submission Archive

From your submission publishing tool (e.g. from Extedo, Lorenz or Parexel) publish the dossier to the CARA Submission Archive, or put the published output on a "watched" folder for automatic ingestion



Submission viewing / Current Approved View

Use the built-in preview tool for content, with the optional addition of submission viewing tools from Lorenz, Extedo, Qdossier and others. See the Current Approved View of your products across markets



Linking to source components and RIM data

As part of the CARA Life Science platform, the Submission archive is seamlessly interconnected to the source submission documents and RIM data, allowing easy reporting, and viewing of related information



Cloud or On Premise / unlimited storage

CARA is available as a SaaS / cloud solution or for installation on premise. Our cloud is GxP, offered as multi-tenant or VIP single tenant based on your requirements. Storage is unlimited

The CARA Life Science Platform

What is it?

A True Platform

Data and content can flow between any solution on the platform **without** integration, enabling seamless collaboration, discovery, and traceability across the business.

Everything is connected in a Common Interface

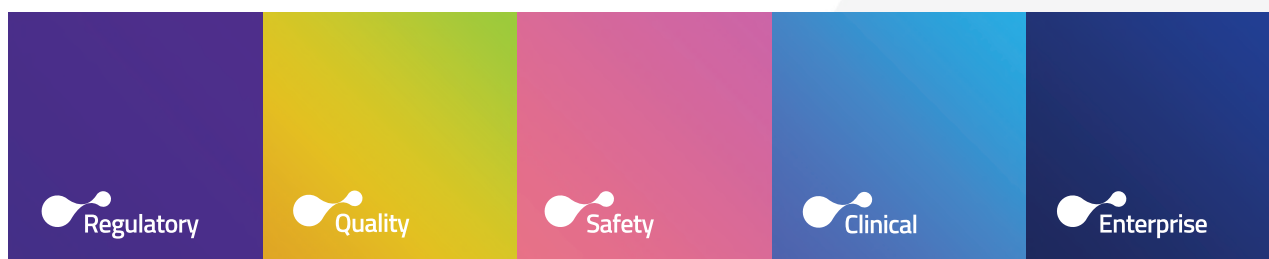
The UI is common between all business processes, reducing training and improving adoption. All data and documents are seamless connected without being in separate vaults / repositories.

Powerful Process Automation

The platform supports different, powerful functionality for each use case, e.g. security, versioning, workflows, lifecycles, tools, automations, and integrations.

One User, One License

Users who work across multiple parts of the business can have access to multiple use cases on the same platform.



RIM
Registrations and Applications
Submission Planning and Tracking
Events and Activities
Product Management including IDMP and SPOR
Health Authority Correspondence and Commitments
Submission documents (Pharma, Devices, Consumer, Veterinary)
Labelling
Promotional Materials
Regulatory Intelligence (Cortellis)

Quality documents
QMS
Deviations management
Complaints Management
Learning Management (LMS)
Audits Management
Laboratory Management
APQR

Pharmacovigilance Case Management
Adverse Event Management
Medical Inquiries
PSURs & Educational Materials
Safety Data Exchange Agreements
PSMF

eTMF
Medical Writing
Clinical study management (CTMS) Q1 2021

Content Management
Records Management
Case Management
Internal & External Collaboration
Financial Documents
Legal / Contracts
Structured Content Authoring
Human Resources