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.

Regulatory

- 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

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

Safety

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

Clinical

- eTMF
- Medical Writing
- Clinical study management (CTMS) Q1 2021

Enterprise

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