

Resilient Knowledge Management in Life Sciences

Knowledge Graphs, Semantic Integration, and AI-Driven Master Data



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Industry Challenge

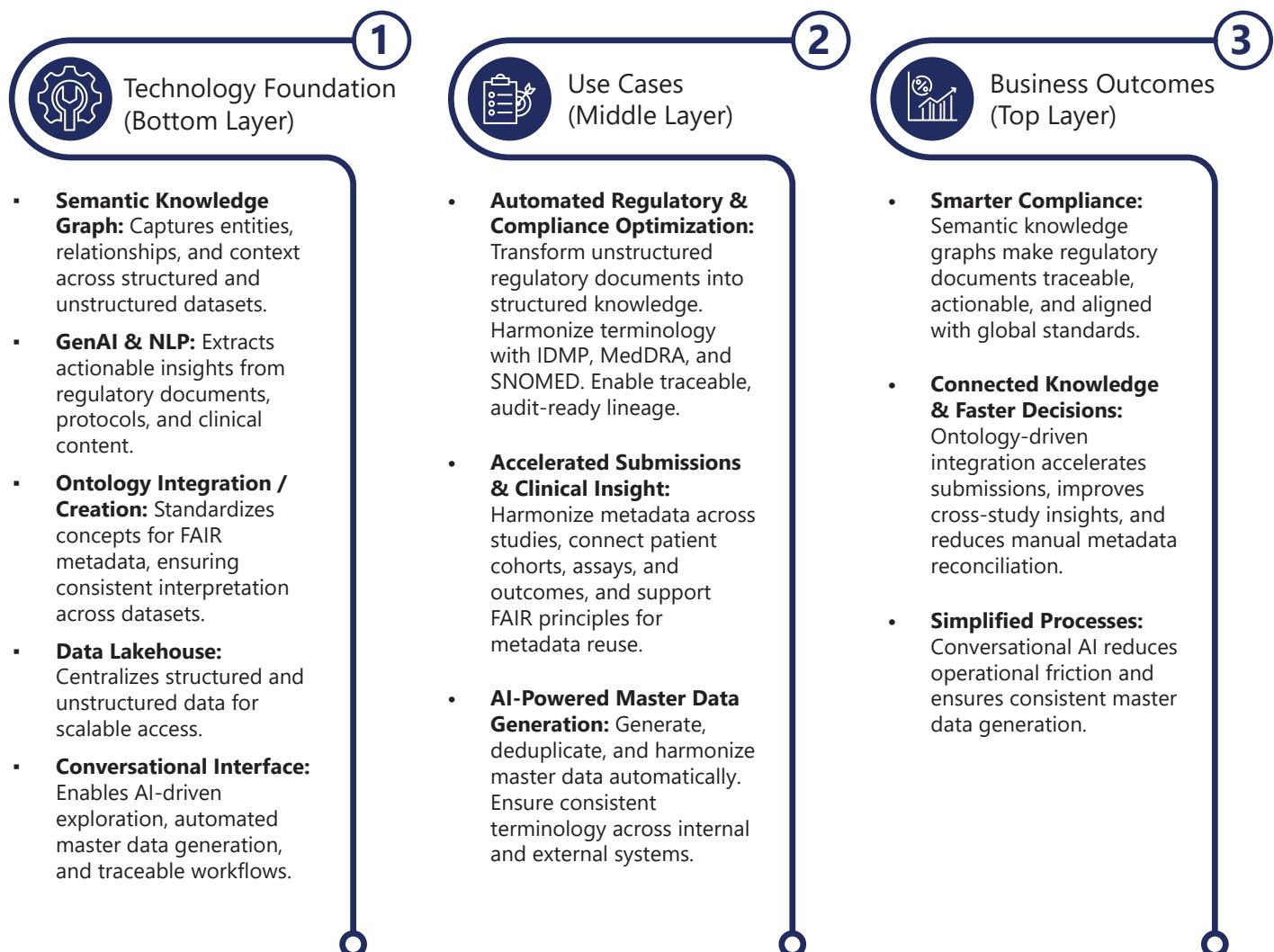
Life sciences organizations generate vast volumes of regulatory, clinical, and operational data. Yet, this data is often fragmented across multiple systems—LIMS, CTMS, EDC, ERP, SOP repositories, and document stores. Variability in metadata, inconsistent identifiers, and siloed documentation creates operational bottlenecks:

- Regulatory teams manually extract product attributes, indications, and references from SmPCs, protocols, and validation reports, increasing risk of errors and delays in submissions.
- Clinical and R&D teams struggle to reconcile assay results, patient cohorts, and biomarker data across trials, slowing evidence evaluation and decision-making.
- Manual reconciliation and lack of semantic integration prevent efficient reuse of historical knowledge and limit AI-driven insights.

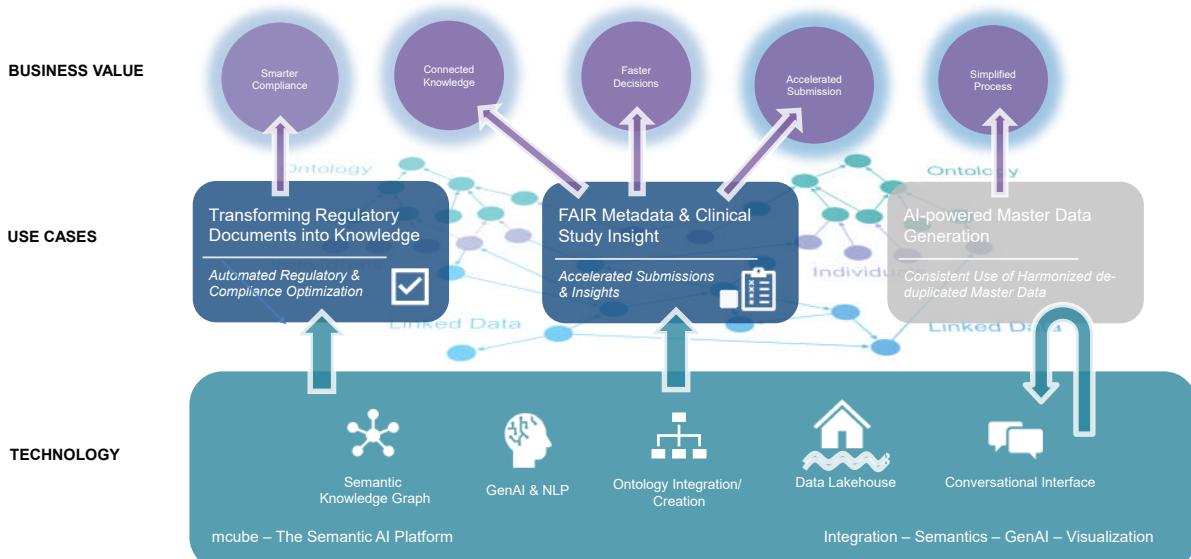
These challenges result in slower regulatory submissions, higher compliance risk, operational inefficiencies, and limited ability to leverage AI for actionable insights.

Solution

mcube™ addresses these challenges by unifying fragmented life sciences data into a connected, AI-ready semantic fabric. The platform combines semantic knowledge graphs, ontology integration, GenAI, NLP, and centralized data architecture to provide a single source of truth:



One Platform – Infinite Possibilities (Ecosystem of Semantic AI Technologies)



Use Cases

1

Structured Regulatory Intelligence

Challenge

Regulatory teams face fragmented repositories of SmPCs, protocols, and validation reports. Extracting product attributes and ensuring traceability is manual and error-prone. Maintaining audit-ready documentation and terminology alignment is difficult.

Solution

- mcube™ ingests unstructured documents and extracts structured information using LLM & GenAI.
- Ontology-based mapping ensures alignment with regulatory standards.
- Alerts flag missing fields or inconsistencies.
- Dashboards enable cross-product, cross-region analysis with full lineage.

Outcome

- Automated extraction and harmonization of regulatory data.
- Audit-ready, traceable documentation.

Use case1: Transforming SmPCs & Regulatory Documents Into Structured Knowledge

SmPCs trapped in PDFs; manual extraction	Inconsistent terminology across products & regions	No search or filtering across large SmPC datasets	Difficult to view complete product profile	Siloed regulatory knowledge	Time-consuming compliance checks & variations
Upload doc → LLM Extraction	Semantic Enrichment (ontology mapping, synonyms, multilingual labels)	Graph Linking (product → indication → class → substance)	Product Detail Page	Knowledge Graph Construction	Semantic Search & Comparison
Automated extraction of product, indication, dosage, safety fields	Standardized disease, vaccine, chemical descriptors	Browsable product catalog with filters: vaccine type, indication, dosage form, year	Unified view: key fields, ontology mapping, GenAI abstract, related documents	Connected regulatory knowledge base for small molecules, antibodies, vaccine etc	Faster impact analysis across SmPC versions and products

2

Cross-Study Clinical Metadata Integration

Challenge

Clinical and R&D teams struggle to integrate data across multiple trials, assays, patient cohorts, and biomarkers. Manual reconciliation delays insight generation and decision-making.

Solution

- mcube™ consolidates clinical metadata and protocols into a unified semantic knowledge graph.
- Ontologies standardize endpoints, sample types, and assay definitions.
- Automated alerts surface missing or inconsistent data.

Outcome

- Instant access to cross-study insights.
- FAIR-compliant, consistent metadata usable across teams.

Use case 2: FAIR Clinical Metadata & Study Intelligence

Ingest→ Extract → Normalize	Knowledge Graph Linking (drug-arm-patient → biomarker sample)	Study Search (filter by condition sample type, assy)	Graph-Powered Comparison	Info Access Control	Advanced Analytics (heat maps, dashboards, study reports)
Clinical data scattered across CTMS, EDC, UMS PDFs	No visibility into relationships between study components	Slow, manual study discovery	Hard to compare trials or reuse design elements	Restricted patient data access	Complex biomarker/clinical data hard to interpret

Conclusion

Life sciences organizations face high volumes of disconnected data, impeding compliance, operational efficiency, and AI adoption.

By transforming fragmented data into a connected intelligence fabric, mcube™ enables organizations to accelerate submissions, streamline operations, reduce manual effort, and make faster, smarter decisions.

01

02

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04

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A unified, FAIR semantic foundation connecting regulatory, clinical, and operational domains.

Consistent, reusable master data across multiple systems.

Automated regulatory alignment with global standards.

End-to-end traceability for audit and compliance.

AI-powered insights grounded in explainable, semantic context.



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mcube™ is TCG Digital's flagship Data, AI, and Analytics platform—engineered at the intersection of deep industry knowledge and digital innovation. Built on a domain-driven design, it seamlessly navigates the most complex and disparate data landscapes. With AI 2.0 at its core, mcube™ fuses advanced models with real-world context to solve high-impact business challenges. By integrating mcube.data, mcube.ai, and mcube.agents, it transforms enterprise data into intelligent ecosystems. mcube™ drives semantic discovery through ontologies, and empowers agentic applications for autonomous decision-making and continuous learning. From harmonizing data to deploying intelligent agents, mcube™ accelerates 'velocity to value'—delivering clarity, intelligence, and outcomes at scale.

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