

DAQUMA

— MAKING IMPACTFUL CHANGES —

SARQA / DKG

2025

WHO IS DAQUMA

DAQUMA

=

DATA

+

QUALITY

+

MANAGEMENT

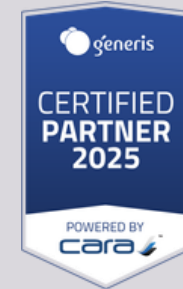
ABOUT US

we help Life Science organizations achieve operational excellence by combining AI-driven data and content innovation with industry expertise to streamline processes, ensure compliance, and accelerate drug-to-patient delivery

OUR CUSTOMER PROMISE

Our content and data automation delivers actual cost reduction and compliance improvement

TRUSTED BY



GxP
Consultants



Data &
IT Experts



Veeva and
Life Science Specialists



Copenhagen
Denmark



Frankfurt
Germany

TRANSFORMATIVE POWER OF GENAI

Quality paradigms and what we need to control



DEMOCRATIZE KNOWLEDGE
MAKING EVERYONE AN EXPERT

Key challenge:
Lack of transparency



HYPER AUTOMATION WITH MASSIVE
EFFICIENCY GAINS

Key challenge:
Dependency and power
concentration



ACCELERATED PRODUCT DISCOVERY
FROM SPEEDING UP RESEARCH

Key challenge:
Bias and discrimination
(age, sex, pregnancy)



PERSONALIZED BENEFITTING
SOCIETY AS A WHOLE

Key challenge:
Misinformation and
manipulation

Direct value to business

Save

TMF Document Intake with QC

Problem: Currently our team uploads and QC documents from internal and external teams in a combination of manual and automated tasks.

Solution: DAQUMA:AI intake automates meta-data population and document QC based on ALCOA+ standard checks to remove the need for human input



19 EUR
per doc

Continuous migration

Problem: Receives content in relation to local content preparation, M&A and product swap activities that require our system teams to do mass uploads

Solution: Setup meta-data automation and ingest documents automatically with AI supported workflows requiring less expertise and fewer man hours



70%
More efficient

Supplier Relations

Problem: Securing that our suppliers' procedures adhere to our contractual and QMS expectations require us to review and store supplier documentation

Solution: Automatically ingesting documents from shared inbox, automatically reviewing against internal procedures before, tagging and uploading to DMS



4.985 EUR
per review

Fill PQR and Safety Cases

Problem: We receive material from CMO's for PQR's and emails with safety cases that we manually interpret and refer and create in our data repositories

Solution: Automatically QC, populate meta-data for PQR and Safety Cases and upload backing documentation and generate PQR and Case record.



34.000 EUR
per month

Direct value to business

The screenshot displays the Veeva Clinical DAQUMA Document Inbox interface. The top navigation bar includes the Veeva Clinical logo and a search bar. The left sidebar shows the 'STUDY SELECTOR' with options for 'All Studies', 'VIEWS', and 'FILTERS'. The main content area is titled 'Document Inbox' and shows a list of documents. The table below represents the data shown in the interface.

Name	Document ID	Status	Created By	Created Date	Study	Study Country	Study Site	Type	Subtype	Classification
Doc11 (v0.1)	746	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc10 (v0.1)	745	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc09 (v0.1)	744	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc08 (v0.1)	743	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc06 (v0.1)	742	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc07 (v0.1)	741	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc04 (v0.1)	740	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc05 (v0.1)	739	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc03 (v0.1)	738	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc02 (v0.1)	737	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		
Doc01 (v0.1)	730	Incomplete	Peter Smedegaard Andersen	20 Oct 2025 8:35 PM EDT				Unclassified		

At the bottom left, there is a link to 'Access documents on Vault Mobile'.

AI content QC new risks?



RISK 1: Automation Control

Challenge: Making a multi-faceted and multi-outcome process driven by a machine rather than a human

Response: Human QC combined with non-AI tech verification to avoid autonomy in AI decisions



RISK 2: Transparency

Challenge: The outcome of AI enabled process makes traceability a high demand

Response: Required to trace the basis of all AI outcomes, with references to original literature/basis.



RISK 3: Bias

Challenge: Our Bias and quality testing happens at implementation, but models and data foundation evolve, and so may the AI agent bias

Response: We need to constantly track output quality and response drift

Track output quality over time as models or data change. Measure accuracy, turnaround time, and rejection rates. Feed QC outcomes back to model fine-tuning and refinement.



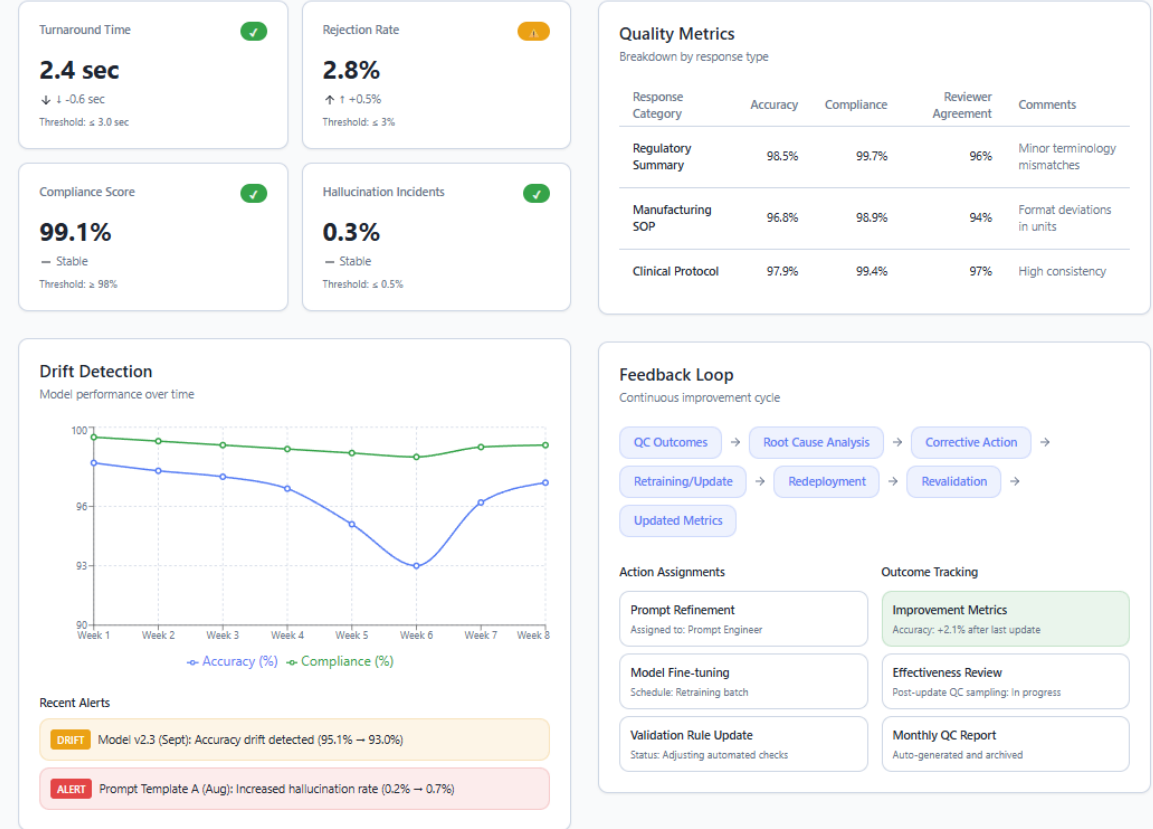
Drift Detection



Metrics Dashboard



Feedback Loop



Acknowledging reality: AI is a tool (not a solution) that augments, not replace quality expertise

Inspire urgency: Professionals and organizations who embrace AI will have competitive advantages (for a while)

Maintain positivity: Focus on positive impact on business and patient outcomes

NOTHING IS PERFECT