



INTUITIONLABS / RESEARCH & PRACTICAL GUIDANCE

Data Integrity in AI: Applying the ALCOA+ Framework

Build practical AI for pharma and biotech with IntuitionLabs. We help life-science teams turn complex information and workflows into useful software, governed knowledge systems and AI tools.

Published by IntuitionLabs · 2026-02-05 · data integrity · alcoa principles · machine learning · ai compliance · gxp · fda regulations · mlops · data governance

Original article: <https://intuitionlabs.ai/articles/data-integrity-ai-alcoa-framework>



In this article

1. Executive Summary
 2. Introduction and Background
 3. Data Quality Foundations and Regulatory Context
 4. Machine Learning Pipelines and Data Integrity Challenges
 5. ALCOA+ Principles Applied to Machine Learning
 6. Data Governance, MLOps, and Technical Controls
 7. Case Studies and Examples
 8. Best Practices and Technical Recommendations
 9. Discussion: Implications and Future Directions
 10. Conclusion
-

Executive Summary

Data integrity—the assurance that data are **accurate, consistent, complete, and trustworthy** over time—is a foundational requirement in regulated industries (e.g. pharmaceuticals, biotechnology) and is only growing in importance as artificial intelligence (AI) and machine learning (ML) systems proliferate. In response, regulators and industry have historically relied on the **ALCOA** (Attributable, Legible, Contemporaneous, Original, Accurate) framework and its extended **ALCOA+** form (adding **Complete, Consistent, Enduring, Available**) to guide good documentation and data management practices (acrpnet.org) (www.sciencedirect.com). For AI used in regulated activities, determine which electronic records fall within applicable recordkeeping requirements before assigning controls. FDA recommends determining whether specific records are Part 11 records based on the predicate rules (FDA). ALCOA+ adds Complete, Consistent, Enduring, and Available; traceability is a supporting control (MHRA).

This report provides an **in-depth analysis** of how ALCOA+ principles apply to machine learning. We cover the **historical context** of ALCOA, data-integrity definitions by regulatory agencies, and data-quality frameworks (e.g. ISO standards, FAIR principles) (pmc.ncbi.nlm.nih.gov) (www.mdpi.com). We then examine the **ML-specific challenges** for data integrity: complex data pipelines, data versioning, bias, and evolving datasets. Each ALCOA+ attribute is interpreted in the context of AI/ML, with examples and tables illustrating how to ensure *Attributability, Legibility, Contemporaneous* recording, *Original* data (or true copies), *Accuracy, Completeness, Consistency, Endurance, and Availability* in ML systems (www.sciencedirect.com) (validfor.com). We review **evidence and case studies** on data quality in AI (including statistics on AI adoption and surveys on governance gaps (blogs.atlas-compliance.ai) (blogs.atlas-compliance.ai)), and we discuss **best practices**: data governance and MLOps frameworks, validation checks, data versioning and lineage, audit trails, and organizational change control. Finally, we outline **future implications**: emerging regulations (e.g. FDA's Good Machine Learning Practice, EU AI Act), new technologies (e.g. blockchain for immutable logs (www.mdpi.com)), and the imperative for a cultural shift toward “**audit-ready AI**” where all ML artifacts are governed by ALCOA+ standards. The goal is to provide actionable guidance for building ML systems that are both innovative *and* compliant with the highest data-integrity expectations.

Introduction and Background

Ensuring **data integrity** has long been a top priority in industries like pharmaceuticals, biotechnology, and medical devices. Major regulatory bodies (FDA, EMA, MHRA, PIC/S) define data integrity along dimensions of accuracy, completeness, consistency, and trustworthiness. For example, the UK's Medicines and Healthcare Products Regulatory Agency (MHRA) explicitly defines data integrity as “*the maintenance of accuracy, consistency, and completeness of data over time.*” ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov/26111111/)). The U.S. FDA similarly emphasizes that electronic records must be “*legible, contemporaneously documented, original or a true copy, and accurate.*” These requirements evolved into the ALCOA framework: **A**tttributable, **L**egible, **C**ontemporaneous, **O**riginal, **A**ccurate ([acrpnnet.org](https://www.acrpnet.org/)). ALCOA was famously introduced in the early 1990s by FDA advisor Stan Woollen to codify good documentation practices ([acrpnnet.org](https://www.acrpnet.org/)). It has since become an industry-wide shorthand: any record (electronic or paper) must clearly identify who recorded it and when (Attributable), be readable (Legible), be recorded as close in time to the observation as possible (Contemporaneous), be the first reliable recording or certified copy (Original), and reflect exactly what happened (Accurate) ([acrpnnet.org](https://www.acrpnet.org/)) (www.sciencedirect.com).

In subsequent years, experts added four more attributes to underscore robust data management: **C**omplete (no missing records), **C**onsistent (logical coherence and conformity of data formats), **E**nduring (durable retention over time), and **A**vailable (readily accessible when needed). These expanded guidelines are collectively known as **ALCOA+** (www.qad.com) (www.auriacompliance.com). Table 1 (below) summarizes the full ALCOA+ principles and their meanings. Over time this framework has become the foundation of Good Documentation Practices (GDPs) and quality systems; “ [f]ailure to maintain data integrity has been a top reason for FDA GMP warning letters,” reflecting how strictly agencies treat any lapses (www.beckman.co.za) (www.auriacompliance.com).

(www.sciencedirect.com) (www.sciencedirect.com) *Table 1. ALCOA+ Data Integrity Principles. Each attribute represents a criterion that data and records should satisfy. (Adapted from FDA/CDISC definitions ([acrpnnet.org](https://www.acrpnet.org/)) and literature (www.sciencedirect.com)).*

Principle	Description
Attributable	The record must clearly show <i>who</i> performed each action and <i>when</i> . (Person, signature, userID, timestamp.)
Legible	Data and records must be readable and understandable by humans (or easily rendered understandable if electronic).
Contemporaneous	The data should be recorded <i>at the time</i> the observation is made (or as soon as feasible), preserving the chronology of events.
Original	The record should be the first (raw) data or an exact certified copy; it reflects the unaltered source.
Accurate	The data must be complete and correct, reflecting exactly what occurred; it should be free from intentional or unintentional errors.
Complete	All necessary data are included (no omissions); if duplicates or derived results exist, they are clearly linked.
Consistent	Data are recorded uniformly over time, using standard methods; there are no unexplained gaps or conflicting entries.

Principle	Description
Enduring	Records must be retained in a durable format for the required retention period (protected from alteration or loss).
Available	Data must be retrievable and accessible when needed (e.g. for review, audit, inspection), without undue delay.

Source: *Regulatory guidance and industry compendia* ([acrpnnet.org](https://www.acrpnet.org)) (www.sciencedirect.com).

In concert, these principles ensure that data are “*trustworthy and auditable*” at every stage. Over the decades, ALCOA+ has underpinned guidance across GxP (Good Practice) environments (FDA’s 21 CFR Part 11, EU Annex 11, ICH Q10, etc.), and it is cited as a core enabler of “transparency and integrity” for electronic records (intuitionlabs.ai) (intuitionlabs.ai). These attributes apply not only to traditional paper logbooks or lab notebooks, but to all sorts of digital data systems – and now increasingly to AI/ML systems.

In the era of AI, new data sources and processes arise (e.g. training data sets, model-generated predictions, sensor streams), but the need for **sound data integrity** becomes even more urgent. As one analyst observes, AI technologies “present unique considerations due to their *iterative and data-driven nature*,” meaning any flaws in inputs or logs can rapidly propagate errors (www.fda.gov). Indeed, data for AI must still be “*fit for use*” – accurate, unbiased, traceable, and well-governed – or the powerful insights AI promises will be undermined (www.deepchecks.com) (www.mdpi.com). In short, “*garbage in, garbage out*” applies especially in ML: unreliable training data yield unreliable predictions (www.deepchecks.com) (www.deepchecks.com). Arnold & Porter’s November 2024 life-sciences survey describes governance as lagging behind AI implementation ([original report](#)). Organizations should assess their own governance, audit practices, and training needs rather than infer readiness from an unverified workforce percentage.

This report will explore how to bridge the ALCOA+ framework to the specifics of AI/ML: how to maintain data integrity through data acquisition, labeling, model training, validation, deployment, and monitoring. We draw on regulatory guidance, publication data, expert analyses, and real-world examples to provide **practical, evidence-based advice**. The aim is to equip practitioners and auditors with a clear roadmap: AI systems are not exempt from GxP standards, and ensuring ALCOA+ compliance in ML pipelines is both challenging and achievable with the right controls (validfor.com).

Data Quality Foundations and Regulatory Context

Before focusing on AI/ML, we summarize the broader **data quality and integrity frameworks** that inform ALCOA+. Data integrity is one facet of data quality, a well-studied concept across industries. ISO standards (e.g. ISO/IEC 25012, ISO 8000) and quality frameworks universally recognize dimensions such as **accuracy, completeness, consistency, timeliness, validity, and uniqueness** (mlops-streamlining-ml-lifecycles.pages.dev) (www.mdpi.com). These overlap almost exactly with ALCOA+ attributes. For instance, a 2025 scientific review notes that “*accuracy, completeness, consistency, timeliness, and accessibility consistently emerged as universal [data] quality dimensions*” across sectors (www.mdpi.com). The same review warns that poor data quality is not just a technical issue but leads to “*substantial financial losses, operational inefficiencies, and erosion of trust*.” In healthcare specifically, flawed or incomplete data have been linked to incorrect patient care or wasted resources (www.mdpi.com).

In regulated environments, data integrity has specialized significance. The FDA explicitly cites Part 11's purpose as ensuring the **integrity** of electronic records ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov/)). Over decades, regulators have repeatedly penalized firms for ALCOA+ violations. For example, one analysis noted that *lack of data integrity has been among the leading causes of GMP (Good Manufacturing Practice) warning letters* (www.beckman.co.za) ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov/)). For a computerized system used in regulated activities, determine which records are subject to applicable requirements and document the risk-based controls, including audit trails where appropriate. Agencies like EMA and MHRA have similarly reinforced that data must be “*complete, consistent, accurate... [and] traceable*” ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov/)) (www.auriacompliance.com). FDA's final February 2026 guidance, **Computer Software Assurance for Production and Quality Management System Software**, covers computers and automated data-processing systems used in medical-device production or the quality management system. It supersedes the September 24, 2025 final guidance ([FDA](https://www.fda.gov/)); it is separate from FDA's 2025 draft guidance on AI used to support drug and biological-product regulatory decisions.

Table 2 (below) provides a concrete example of how each ALCOA+ attribute can be interpreted in the context of ML-based systems, drawing on industry guidance ([validfor.com](https://www.validfor.com/)) ([validfor.com](https://www.validfor.com/)). This “**ALCOA+ for AI**” table will be referenced throughout the report.

([validfor.com](https://www.validfor.com/)) ([validfor.com](https://www.validfor.com/)) **Table 2. Applying ALCOA+ in AI/ML Context. Examples of what each ALCOA attribute means for machine learning data and processes. (Adapted from industry guidelines ([validfor.com](https://www.validfor.com/)) ([validfor.com](https://www.validfor.com/))).**

ALCOA+ Attribute	Meaning in AI/ML Context	Example Controls
Attributable	<i>Who did what and when.</i> Track which user, system, or model produced each record and decision.	Log user IDs, model versions, or service IDs for each action (e.g. user who generated a prompt, system that deployed an inference).
Legible	<i>Readable and understandable.</i> Ensure outputs, logs, and labels can be interpreted by humans or decoded.	Store full prompt texts, decoded parameter values, and clear labels. Ensure dashboards and reports are human-readable.
Contemporaneous	<i>Timestamped at event-time.</i> Record data and metadata as soon as events occur.	Time-stamp all data entries, model inferences, and approvals. Use synchronized clocks/log servers so sequences of events are clear.
Original	<i>First capture or certified copy.</i> Preserve raw source data and non-derivative records.	Retain the original training datasets or verified true copies that preserve content, meaning, necessary metadata, and functionality (e.g. raw sensor logs, initial images). Use hashes as supplemental integrity checks, not substitutes for retained records (FDA).
Accurate	<i>Correct and validated.</i> Maintain data that truly reflects reality and is free from errors.	Apply validation rules on inputs and outputs (e.g. schema checks, range checks). Reconcile derived data with source records or control totals.
Complete	<i>Nothing missing; no hidden edits.</i> Capture all relevant data and its processing steps.	Archive both raw outputs and any post-processed versions. Record full pipelines (raw data, features, final predictions) to avoid omissions.

ALCOA+ Attribute	Meaning in AI/ML Context	Example Controls
Consistent	<i>Uniform format/sequence.</i> Keep data formats, schemas, and process steps consistent and documented.	Enforce schema standards on input features. Use the same field order and naming for every run. Maintain fixed data pipelines for comparable results.
Enduring	<i>Durable for retention period.</i> Store data and models in long-term, unalterable storage.	Use immutable, write-once media or append-only stores for records. Archive models, logs, and raw data under retention policies (on premises or cloud).
Available	<i>Accessible on request.</i> Enable retrieval of data and logs when needed for review or audit.	Index and catalog datasets, artifacts, and logs. Ensure data discovery tools or portals allow auditors to query past records quickly.

Source: Industry compliance guidelines ([validfor.com](https://www.validfor.com)) ([validfor.com](https://www.validfor.com)) and ALCOA+ literature (www.sciencedirect.com).

These tables synthesize key concepts but by no means exhaust the subtleties involved in applying ALCOA+ to AI. In the following sections, we will unpack each aspect in depth: from the theoretical underpinnings of ML data quality to concrete best practices and technologies that make ALCOA+ compliance achievable in AI systems.

Machine Learning Pipelines and Data Integrity Challenges

Machine learning introduces new **data integrity challenges** not seen in traditional computer systems. An ML system's "data" spans large, often heterogeneous datasets, streaming sensor inputs, human-labeled training data, and the models and code themselves. This complexity creates multiple risk points: data can be **modified, lost, or contaminated** at many stages, any of which can invalidate results.

A core principle persists: models will only be as reliable as the data entering them. As one AI engineer warns, "*data integrity... plays a pivotal role... the adage 'garbage in, garbage out' rings true*" (www.deepchecks.com). Ensuring data is *fit for use* means having high quality (correct and complete), well-organized data before training. If a corrupted dataset feeds a model, it will reproduce those flaws (or worse, introduce unpredictable behavior) (www.deepchecks.com) (www.deepchecks.com).

Aside from outright data poisoning or bias, complex data pipelines themselves tend to introduce integrity issues. An ML pipeline typically ingests raw data, preprocesses it (cleaning, feature extraction), trains models, and then serves predictions. At each step, errors may creep in – e.g. transcription mistakes, mismatched formats, dropped values, or synchronization failures in distributed systems. For instance, it's common to encounter missing or inconsistent records when merging datasets from different sources (www.deepchecks.com). Pipelines often have *bottlenecks*: mis-timed data loads or hidden caches may yield stale or duplicate inputs (www.deepchecks.com). Human factors also play a role: manual labeling or data entry can produce legibility issues or unlabeled columns. Data drift is another concern: the distribution of incoming data can shift over time, causing models to become inaccurate. As the Deepchecks analysis notes, "**data drift transpires when data distribution shifts over time... [and] concept drift pertains to changes in underlying relationships,**" both of which undermine model validity (www.deepchecks.com).

Given these hazards, rigorous controls are needed. Standard software engineering approaches must be extended: for example, all datasets require automated validation (schema checks, outlier detection, duplication checks) at each point (www.deepchecks.com). A robust ML data pipeline will implement continuous integrity monitoring – reconciling new records with previous snapshots, logging events at high granularity, and alerting on anomalies before they “propagate downstream” (www.deepchecks.com). In practice, many ML teams are adopting **MLOps** practices and data governance to enforce this. For example, data governance policies define roles, access, and compliance for ML data (mlops-streamlining-ml-lifecycles.pages.dev), and version control systems (analogous to Git) are used to snapshot data and track transformations. One thought leader describes compliance in ML as needing to “*prove exactly what data went into the system*” – because under frameworks like the EU AI Act or FDA guidance, regulators may demand the precise training dataset used on a given date (lakefs.io). In a **hypothetical** example, lakeFS describes overwritten credit-model training data becoming unavailable when later requested. This illustrates a potential retention problem; it is not evidence of an actual subpoena or enforcement action ([lakeFS](https://lakefs.io)).

In summary, the **ML lifecycle introduces multidimensional integrity risks**:

- **Data acquisition:** Sources (sensors, databases, APIs) must be verified. Missing metadata (timestamps, provenance) here violates ALCOA attributes from the start.
- **Data labeling/preprocessing:** Manual or automated transformations can introduce errors. Lack of versioning or audit trail means one cannot revert mistakes or confirm original source data was used.
- **Model training:** Training code and parameters must be tracked (who ran it, when, and what version), or the resulting model is not “attributable” or “traceable” to a responsible party.
- **Model outputs:** Predictions must themselves be logged in context (which model and data produced them) so that downstream decisions are auditably linked back.
- **Maintenance/deployment:** Whenever models are retrained or updated, change control processes must ensure new versions are validated, documented, and aligned with original requirements – otherwise “complete” and “consistent” can break across versions.

Each of these stages must incorporate ALCOA+ controls to maintain integrity and compliance. The next section will detail how each ALCOA+ principle can be satisfied in such ML workflows, with specific data-quality and governance strategies.

ALCOA+ Principles Applied to Machine Learning

In applying ALCOA+ to ML systems, first identify the regulated activity and intended use. FDA recommends determining which specific records are Part 11 records based on applicable predicate rules ([FDA](https://www.fda.gov)). Controls for other training sets, code, models, outputs, and logs are engineering recommendations, not a declaration that every artifact is legally regulated. Below we examine each principle in turn, citing guidance and examples. Table 2 above is used as a reference for context.

- **Attributable (A)** – “*Who did what and when.*” In ML, this means every action must be traceable to a user or system. For example, if a data scientist initiates a model training run, the system should record their ID, the time, and the specific versions of the data and code used (validfor.com) (validfor.com). This can be achieved by logging user IDs in the training orchestration system, stamping datasets with unique

identifiers (or checksums), and using platforms that record user actions. Validfor's guidance emphasizes that AI records should note *"which model version produced a decision... and who or what system consumed the output"* (blogs.atlas-compliance.ai). Attribution also extends to data annotation: each label or transformation should carry a timestamp and annotator ID so accountability is clear. In practice, solutions include implementing single sign-on (SSO) for all ML tools, audit logs for notebooks or pipelines, and version control systems (e.g. Git for code, DVC/lakeFS for data) that inherently tag commits by author and time.

- **Legible (L)** – *"Readable and understandable."* This requires that records be decipherable by humans or machine-readable in a transparent way. In ML, authorized users should be able to decrypt, decode, and render retained prompts, logs, and metadata in readable form. Preserve appropriate encryption, access controls, and recoverable keys for sensitive records; legibility does not require unencrypted storage. Part 11's open-system controls include encryption where necessary ([21 CFR 11.30](#)). For instance, retaining required prompt content and parameter settings in securely retrievable, readable form (rather than cryptic hashes alone) makes them "legible" to an auditor (validfor.com). Outputs should be stored alongside descriptions (e.g. column mappings, label definitions) so the results can be interpreted. Engine logs and monitoring dashboards must display context (timestamps, model version, feature names) in a human-recognizable language. In essence, legibility in AI means avoiding "black box" burying of data: keeping the chain of information (raw inputs → features → predictions) transparent.
- **Contemporaneous (C)** – *"Recorded at the time of the event."* ML systems often operate in real-time or near-real-time. To meet this, all data and actions must be timestamped and logged as they occur. For example, a model inference service should immediately log each request and response with precise time markers (to the millisecond, if needed) (validfor.com). Batch processes (e.g. nightly retraining) should annotate each result with the job execution time. Containerized or cloud deployments can leverage automated logging (e.g. writing logs to immutable object storage in real time). Critically, clocks must be synchronized across systems (via NTP or equivalent) so that multi-step processes have a consistent timeline. Contemporary recording also applies to corrections: if data are modified (say a mislabeled example is fixed), the edit timestamp and editor must be logged rather than just retroactively overwriting historical data (www.sciencedirect.com). In sum, the system must capture "when" every piece of AI-related data was created or changed, maintaining an accurate chronology.
- **Original (O)** – *"First capture or certified copy."* AI workflows produce vast amounts of derivative data (features, models, transformed files) that flow from original sources. The "Original" principle insists on retaining the raw source or a verifiable duplicate. For ML, this means preserving the **original datasets** used for training, not just processed versions. Practical implementations include taking write-once snapshots of raw data (e.g. initial sensor or experimental readings) and storing cryptographic hashes so one can later prove integrity (validfor.com). It also entails versioning the source code and configuration (each commit is an "original" state of the pipeline). For critical data, institutionalizing data lakes with immutable zones ensures that once raw data are ingested, they are never altered (only appended). This way, downstream models can always be traced back to the first, unmodified record. As a guideline notes, we must *"preserve raw source data (for instance, raw sensor logs or unprocessed images) and transformed data... lineage to original inputs must be maintained."* (blogs.atlas-compliance.ai).

- **Accurate (A)** – *“Correct and error-free.”* Accuracy means the data truthfully and precisely represent reality. In AI, this covers input data, labels, and model outputs. To achieve accuracy, common practices are automated validation and reconciliation. For training datasets this might involve range checks (e.g. no negative ages), format checks (e.g. dates in ISO-8601), and duplicate detection (www.deepchecks.com). Labeling processes should include inter-annotator agreement or audit samples to catch mistakes. Feature engineering scripts should include assertions (e.g. no NaNs in critical features). After training, outputs must be checked against known benchmarks or sanity checks to ensure no computational errors. Where results feed into decisions, cross-checks can validate that predictions align with independent data or expert judgment. Comprehensive test suites form a key part of establishing accuracy. For example, validfor’s guidance implies performing “validated transforms and reconciliation checks” on ML data (validfor.com). Ultimately, an accurate AI system is one where every transformation is verified and documented, with anomalies caught either by automation or human review.
- **Complete (C)** – *“Nothing missing, no hidden edits.”* Completeness means all required data is captured, with no gaps. In ML contexts, this requires careful audit of the dataset lifecycle. One must ensure that all raw data were included (no sampling artifacts) and that any filtering or data cleaning is fully logged. If a dataset has multiple parts (e.g. training set and test set), definitions of these splits must be recorded so that future reviewers can see the full scope. In practice, completeness often translates to storing not just the final output of a transformation (e.g. a feature table), but also the original raw output and any intermediate versions. For example, if an image dataset is augmented in preprocessing, both the augmented and original images should be kept in the record. Validfor’s table suggests keeping “*raw output, post-processing, and final approved record*” together to satisfy completeness (validfor.com). Data pipelines should also track which items were dropped or modified (for example, flagging any incomplete records) so that the absence of data is itself documented—not simply erased.
- **Consistent (C)** – *“Uniform and logical sequence.”* Consistency requires that data collection and recording follow a standard, reproducible process. In ML, consistency is ensured by controlling versions and maintaining fixed formats. This can involve using schema enforcement (e.g. JSON schemas, database constraints) so that all records have the same fields in the same order. It also means that, if an AI model is retrained, the procedures applied (cleaning rules, algorithms) are recorded precisely so results are comparable over time. Workflow management tools help here: they ensure that every run of the pipeline uses the same steps, and when changes are introduced, they go through change control. For example, if a new feature is added to input data, the dataset definition should be updated (rather than leaving old and new data indistinguishable). In short, consistency is the discipline that prevents piecemeal or ad-hoc variations from creeping into the data process. Without it, one could have a model trained on one format and production data in another, breaking ALCOA.
- **Enduring (E)** – *“Durable for the retention period.”* Enduring refers to the security and longevity of records. AI systems must ensure that once data or models are stored, they remain unaltered and intact over their required lifetime. This typically involves write-once storage (e.g. WORM drives, append-only logs) or cryptographically assured storage solutions (e.g. object stores with immutability flags). Models themselves (weights, artifacts) should be archived in a way that prevents silent updates – for instance, by recording hash digests on a blockchain or secure registry. The “*Enduring*” principle also implies

disaster recovery: backups should be maintained so records survive technical failures. For example, a multi-region cloud archive or offline cold storage can ensure that historical training data and models are not lost if a system is decommissioned. As noted in Table 2, systems should use “immutable storage with retention rules” (validfor.com) so that data remain readable and unmodified throughout legal or audit-relevant time frames.

- **Available (A)** – “*Accessible on request.*” The final ALCOA+ attribute mandates that data be retrievable. It’s not enough to store everything safely; one must also provide timely access. In AI/ML workflows this means implementing data catalogs and metadata indexing. Every data asset (training sets, intermediate tables, model files) should be registered in an index with search capabilities. For example, an ML metadata store might allow auditors to query “which model was used on trial #1234” and immediately fetch the answer. Data governance tools often provide role-based access (linking to “Attributable”), ensuring that only authorized personnel can pull protected data. Crucially, audit trails and logs themselves must be available: if an inspector asks to see the “audit trail for decision X,” the system must produce the record quickly. Determine audit-trail coverage for relevant records from applicable predicate rules and a documented risk assessment ([FDA](https://www.fda.gov)); the scope should reflect intended use and the potential effect on product quality, safety, and record integrity. Thus, accessibility is achieved by combining robust cataloging, queryable data lakes, and well-documented APIs that expose stored records as needed.

By carefully integrating these principles into ML projects, organizations can build **trustworthy AI**. In practice, this means designing the ML infrastructure so that ALCOA+ attributes are met by default. For instance, an **artifact repository** (like MLFlow, Artifactory, or a bespoke data lake) can enforce versioning and hashing (addressing Attributable, Original, Enduring), while a **lineage tracking system** (or metadata layer) can capture data transformations for legibility and traceability (blogs.atlas-compliance.ai) (www.mdpi.com). Together, these mechanisms make AI development auditable at the same level expected of classic GxP computerized systems.

Data Governance, MLOps, and Technical Controls

To operationalize data integrity in ML, a comprehensive data governance and MLOps strategy is required. Key elements include:

- **Data Versioning and Lineage Tracking.** Just as software developers use Git, ML teams must “version control” their data. Tools like DVC, lakeFS, or Delta Lake allow snapshots of large datasets to be taken, with histories that can be reverted. This mirrors the ALCOA+ requirement of capturing the original. For example, lakeFS uses a hypothetical credit-model scenario to illustrate the consequences of overwriting training data ([lakeFS](https://lakefs.io)); it does not document a real enforcement case. By contrast, a versioned storage system (with cryptographic hashes on each version) can demonstrate precisely what data fed the model at any point. Lineage systems augment this by automatically recording how data flows through the pipeline. They tag each dataset with metadata on its source, transformations, and downstream consumers. In an ALCOA context, provenance is critical: as one guide puts it, you must be able to “*track the origin, transformations, and propagation of data*” through the ML system, and show it to auditors (www.deepchecks.com) (www.mdpi.com).

- **Automated Validation and Quality Checks.** Embedding validation steps into the pipeline is essential. At every stage (ingestion, preprocessing, post-inference), the system should apply automated checks for **accuracy, completeness, consistency**. These include schema conformance (no unexpected columns), null/NaN detection, range checks, and statistical anomaly detection. For example, before data enters training, a rule might assert that no feature has more than 5% missing values; after prediction, another check might compare aggregate results to historical norms. These checks serve as guardrails so that errors or rapid drift do not propagate undetected (www.deepchecks.com). Comprehensive metrics (such as data quality scores) can be maintained over time, documenting that each ALCOA requirement has been internally monitored.
- **Audit Trails and Logging.** ML systems must record rich logs of every action: data injections, code commits, training runs, deployments, and predictions. Each log entry should include who/what caused the action, when it happened, and on what data, thus covering *Attributable* and *Contemporaneous*. Centralized logging solutions (ELK/Elastic, Splunk, etc.) or ML-specific monitoring (e.g. Amazon CloudTrail, Azure ML logs) can capture this. Logs should be tamper-resistant; technologies like WORM storage or even blockchain-based logging have been proposed to prevent backdating or deletion (www.mdpi.com). For instance, Regueiro et al. (2021) demonstrate a blockchain-linked audit trail that ensures “secure and reliable” logging for enterprise systems (www.mdpi.com). In effect, the audit trail becomes a descriptive “chain of custody” for AI: every record can be traced back through immutable logs, satisfying *Traceability*, *Enduring*, and *Available*.
- **Access Controls and User Training.** Robust role-based access control (RBAC) is needed so that only authorized personnel can create, modify, or view critical records. This prevents unauthorized substitutions or deletions (thus helping *Accurate* and *Attributable*). For applicable records, distinguish access authorization and change audit trails from electronic signing. Electronic signatures must identify the signer, execution time, and meaning and remain linked to the record; authentication alone does not replace a required signature. Digital signatures and MFA may be selected as implementation controls, but Part 11 does not require either on every change (**21 CFR Part 11**). Personnel involved in data handling must be trained on ALCOA+ importance (just as they would be for GxP tasks). Studies show that cultural factors are a leading cause of data integrity lapses (pmc.ncbi.nlm.nih.gov) (pmc.ncbi.nlm.nih.gov), so fostering a quality culture for ML is key. Companies should update SOPs and validation plans to explicitly cover AI operations (e.g. “All model change requests must include data lineage documentation”).
- **Monitoring and Drift Detection.** Because ML systems operate in changing environments, proactive monitoring is crucial. Tools that watch for *data drift* (statistical shifts in input distributions), *concept drift* (changes in what is being predicted), or model performance degradation all help maintain *Accuracy* and *Consistency*. When drift is detected, a process should trigger investigations or retraining (with retraining itself following ALCOA+ controls). In quality parlance, this resembles a manufacturing out-of-spec alert. For example, if a model’s predictions suddenly deviate, the system should freeze updates and launch a validation subroutine, rather than letting bad data silently skew results.

- **Comprehensive Documentation.** All of the above controls must be documented. A formal **Validation Master Plan** for AI (analogous to any major system) should outline how ALCOA+ is applied across the lifecycle. The [IntuitionLabs validation framework](#) outlines key validation documents and checklists. Even if full formal validation is not yet mandated for all AI use-cases, adopting these artifacts early positions an organization for future inspection expectations.

Together, these MLOps and governance practices form a **data integrity “by design” approach**. Quality checks, version control, and audit logging support data integrity. Organizations must determine applicable record requirements and document the controls for their intended use (**FDA**). Practically, many organizations achieve this by layering their architecture (as Atlas Compliance advises) ([blogs.atlas-compliance.ai](#)):

- **Artifactory/Artifact Registry:** All datasets, model binaries, and code are stored in a centralized, versioned registry (with cryptographic hashing) ([blogs.atlas-compliance.ai](#)). Once committed, these artifacts cannot be modified, only appended (ensuring *Enduring* and *Original*).
- **Lineage Fabric:** A dedicated lineage service tracks how data flows through the pipeline ([blogs.atlas-compliance.ai](#)). Each processing step emits an event that ties inputs to outputs (helping *Legible*, *Consistent*, *Traceable*).
- **Quality Gateways:** Automated gates enforce validation at each handoff (ingest gate, pretraining gate, predeployment gate). Events that fail checks are logged and blocked until resolved (preserving *Accurate*, *Complete* quality).
- **Logging & Monitoring:** Every user action, compute job, and model prediction is logged with identity and timestamp (covering *Attributable/Contemporaneous*). Model performance monitors watch production outputs and alert on drift or errors.
- **Access and Backup:** RBAC ensures only qualified people can alter the critical flows. Offsite backups and long-term archives keep multiple copies for recovery (supporting *Enduring* and *Available*).

This layered architecture supports documented record controls. Base audit-trail controls on applicable predicate rules, a documented risk assessment, and potential effects on product quality, safety, and record integrity (**FDA**). Similarly, after training models, firms maintain “check-summed” records of training data as immutable evidentiary files.

Finally, it cannot be overstated that **people and processes** must overlay these technical controls. Data scientists and engineers should be versed in ALCOA+ concepts as part of their training. Regular audits (practice “inspections”) of ML projects help identify any gaps. In fact, industry reports indicate that internal GxP auditors are already expanding their scope to AI systems. Rockwell Automation’s 2025 life-sciences survey reports that 95% of surveyed manufacturers were **using or evaluating smart technology** ([Rockwell Automation](#)); this measure is not equivalent to AI investment or deployment. In sum, the combination of robust MLOps tools and vigilant governance is required to translate ALCOA+ from theory into practice for AI.

Case Studies and Examples

Hypothetical Retention Example

lakeFS introduces an imagined credit-model scenario in which overwritten training data are unavailable after a later request (**lakeFS**). It illustrates a potential failure to preserve retrievable history, not an actual subpoena, regulatory violation, or enforcement case. Consider versioned storage and retention policies appropriate to the model's intended use and applicable record requirements.

This hypothetical example illustrates the retention stakes and suggests the following safeguards:

- **Prevention Strategies:** Data auditing, diverse and representative datasets, continuous monitoring, and robust logging are essential. For each ALCOA attribute, there are concrete measures (as shown in Tables 1–2) that mitigate the risk.
- **Ongoing Vigilance:** Even after deployment, models need periodic review (akin to “periodic review” for GxP systems).

In the next section, we synthesize these lessons into best-practice recommendations and forward-looking considerations for organizations building AI systems.

Best Practices and Technical Recommendations

Drawing on the above principles and hypothetical retention example, we summarize key recommendations for ensuring data integrity in ML/AI:

1. **Define Record Scope and Intended Use.** Identify the regulated activities supported by AI and determine which specific electronic records are subject to applicable predicate rules and Part 11 (**FDA**). Document those decisions in a Data Integrity Plan, including required training data, prompts, model versions, outputs, and their retention and access controls. Apply additional engineering controls according to documented risk.
2. **Implement End-to-End Data Lineage.** Use ML engineering platforms or metadata tools (e.g. MLflow, Kubeflow Pipelines, Apache Atlas) that automatically record provenance. Ensure that for any final decision of the model, one can trace back through every data transformation and model step. This satisfies *Traceable* and *Legible*: maintain detailed processing graphs so that reviewers can follow “how the answer was obtained.”
3. **Automate Validation and Alerting.** Integrate automated data checks at ingestion (schema, types, ranges) and pre- and post-inference (quality metrics, plausibility tests). Don't rely only on manual spot checks. Anomalies should trigger alerts and suspend pipelines until resolved, preserving *Accurate* and *Complete*. Build quality dashboards that continuously report status (missing values per column, drift statistics, etc.).
4. **Use Version Control for Data and Models.** Leverage tools that snapshot data and code at each pipeline stage. Tag every release of a model with a unique ID tied to specific data and code versions (akin to software release tagging). This handles *Attributable/Contemporaneous/Original*: one can always

identify “which copy of data produced model X on date Y.”

5. **Secure and Immutable Storage.** Archive critical data and model artifacts in write-once, timestamped storage. As illustrated by Regueiro et al., blockchain or cryptographic hashing can further harden audit trails (www.mdpi.com). For example, when a dataset is finalized, compute its SHA-256 hash and store it in a ledger so any future tampering is detectable. Store copies in geographically separated backups to guard against loss.
6. **Comprehensive Logging and Audit Trails.** Maintain detailed logs (user logins, API calls, data access, training runs, deployment events) with metadata (who, what, when) to satisfy *Attributable* and *Contemporaneous*. Use centralized log aggregation so these trails cannot be easily erased or altered. For example, require that each required prompt record fed to an LLM is logged with user ID and timestamp for retrospective inspection. Apply audit-trail controls according to applicable predicate rules and documented risks to product quality, safety, and record integrity ([FDA](#)).
7. **Data Governance Policies.** Establish corporate policies covering AI: define data ownership, classification (e.g. CUI/PHI), and retention rules. Enforce binding SOPs that reflect ALCOA+: for instance, a rule that *no model may be deployed without documented training dataset lineage and validation report*. Regularly audit adherence to these policies. Document the audit schedule, responsibilities, findings, and follow-up actions rather than rely on an incorrectly reported industry percentage.
8. **Continuous Re-Validation.** Recognize that ML models degrade. Establish schedules for re-running validation tests (on hold-out or new data) and compare to baselines. If significant drift or bias is detected, trigger re-training or model rollback. This ensures ongoing *Accuracy/Consistent* over the model's lifecycle.
9. **Human Oversight and Expertise.** Assign cross-functional teams (data scientists, quality engineers, IT) responsibility for data integrity. Require that quality/validation specialists review model documentation and logs, not just trusting automated processes. Foster a culture where staff are trained to think of “Data Integrity” as an integral part of ML, much like they would for any regulated process.
10. **Regulatory Alignment and Audit Preparedness.** Finally, prepare for the evolving compliance landscape. Follow FDA and ISO initiatives (e.g. NIST AI risk management, IMDRF GMLP principles (www.fda.gov)) and integrate any new guidance (such as FDA's 2025 draft on AI credibility (www.fda.gov)). Document risk assessments under ICH Q9-style methodology: evaluate the impact of data integrity failures on patient safety. Keep change control boards involved when updating AI systems, as they would for any critical software change.

By implementing these practices, organizations can harness the power of AI **while keeping data integrity “audit-ready.”** Assess the resulting records and controls against the applicable requirements for their intended use. Importantly, many of these measures also yield business benefits: version-controlled data and models improve reproducibility and speed, automated validation increases efficiency, and robust logging builds trust with stakeholders. Compliance should thus be viewed not as a burden, but as a path to **better, more reliable AI.**

Discussion: Implications and Future Directions

Data integrity in AI/ML sits at the intersection of technology, regulation, and ethics. Our analysis points to several broader implications and emerging trends:

- **Evolving Regulatory Expectations.** Regulators worldwide are rapidly recognizing AI but expect existing data principles to apply. For example, the FDA's 2025 draft guidance emphasizes “*model credibility*” and risk-based oversight, implicitly requiring data integrity checks for ML outputs (www.fda.gov). As of October 4, 2026, the EU AI Act has entered into force and become generally applicable, with phased exceptions. The Commission identifies an extended transition to August 2, 2028 for high-risk AI embedded in regulated products (Annex I); classification and obligations depend on the system's intended use ([European Commission](https://europeancommission.eu)). Even current Good Practice guides (e.g. PIC/S Annexes) are expected to be updated with AI-specific language. We anticipate that within the next few years, most regulatory inspections of AI will explicitly probe ALCOA+ compliance (as already foreshadowed by EMA guidance (validfor.com)).
- **Data Ethics and Trustworthiness.** Beyond compliance, data integrity in AI is closely tied to broader notions of *trustworthy and ethical AI*. Issues like fairness, bias mitigation, privacy, and cybersecurity all depend on maintaining high-quality data and auditable processes. For instance, “Findability” and “Interoperability” from the FAIR data principles can be seen as complementary to ALCOA's Accessible+Consistent attributes (www.mdpi.com). Ensuring ALCOA+ in ML helps build the foundation for AI systems that can be explained and audited, addressing AI governance challenges head-on. Many experts argue that data governance must now be seen as an *organizational priority*, not an IT afterthought: poor data practices can erode public trust in AI just as manufacturing defects would erode trust in a drug.
- **Technological Enablers.** New technologies are emerging to automate ALCOA+ compliance. For example, **blockchain** and distributed ledger systems can provide tamper-evident audit trails (ensuring immutability and *Enduring* storage) (www.mdpi.com). Smart contracts could, in theory, enforce data lineage rules by automatically rejecting data changes that lack provenance. Similarly, advances in ML-specific MLOps platforms now offer built-in lineage and versioning (e.g. Git for Data, DataOps pipelines) that make documentary compliance more seamless. On the horizon are AI tools that may review data integrity automatically (e.g. anomaly detection models that flag suspicious data inputs in real time). As one industry source notes, embedding governance into AI requires “*continuous monitoring and re-validation triggers*” akin to post-market surveillance.
- **Cross-Industry Learning.** While our focus has been largely on pharm/biotech, the ALCOA+ approach applies to any field using critical AI – banking, automotive, energy, etc. The lakeFS credit-model example is hypothetical ([lakeFS](https://lakefs.io)) and illustrates a potential record-retention problem; it does not establish that consumer-finance regulators have imposed ALCOA+ obligations. Similarly, autonomous vehicles will likely inherit safety-related data integrity regulations from aerospace/nuclear. Cross-sector dialogue (e.g. via IMDRF, NIST, international standards bodies) will lead to harmonized best practices.

- **Human Factors and Culture.** Data integrity ultimately depends on people and processes as much as technology. Historical analyses of GxP failures often single out poor culture and training as root causes ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov)). In the AI domain, technical fixes alone are insufficient: organizations must cultivate a culture where data quality is valued. This includes management accountability (data integrity is a leadership issue, not just a technical task (www.linkedin.com)), as well as empowering data stewards to enforce standards. Organizations should assess staff readiness and provide role-specific education and governance support.
- **Future Research and Standards.** Finally, from an academic and standards perspective, much work remains to tailor integrity principles to AI. ALCOA+ is a guide, but specific metrics and tools for measuring things like “completeness” or “consistency” in ML-driven data must be refined. Ongoing research is examining how to quantify data pedigree or certify model transparency (for instance, recent papers on AI audit trails and provenance (www.deepchecks.com) (www.mdpi.com)). New certifications or conformity assessment programs (similar to CE marking for medical devices) may emerge for AI in regulated settings, wherein ALCOA+ compliance would be a criterion. The community should share case studies and validation examples widely, building a knowledge base akin to the GxP “data integrity warning letters” that currently circulate among pharma professionals.

Ultimately, the trend is clear: **data integrity in AI is no longer optional**. As life-sciences consultant Joshi notes, AI systems should “*empower Quality and Compliance professionals to focus on critical insights rather than manual data mining*,” implying that AI tools must be built on rock-solid data foundations (www.linkedin.com). Achieving this requires hard work (from extensive data cleaning to vigilant documentation) but yields dividends: reliable AI outputs, smoother audits, and increased confidence from regulators and the public.

Conclusion

The age of AI demands a renewed and rigorous commitment to data integrity. ALCOA+—once a guideline for paper batch records—now must extend into digital ecosystems that encompass machine learning. This report has shown that each ALCOA+ attribute can be concretely applied to AI: by logging every user interaction (Attributable), timestamping and preserving raw inputs (Contemporaneous, Original), validating and reconciling data (Accurate, Complete), and maintaining immutable archives (Enduring, Available).

We provided multiple perspectives: regulatory definitions ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov)) (www.auriacompliance.com), technological strategies (MLOps, versioning (lakefs.io) (blogs.atlas-compliance.ai)), and expert recommendations. Overall, three themes emerge:

- **Baseline Standards Remain Paramount:** The classic ALCOA+ principles are still fully relevant. AI does **not** replace these core expectations. Indeed, some agencies are already restating them in AI contexts.
- **AI-Specific Adaptations:** Organizations must interpret these principles for ML workflows. This includes tracking model versions and dataset snapshots, integrating automated checks, and building comprehensive audit systems. A structured approach (like the artifact registry + lineage + monitoring architecture) is recommended (blogs.atlas-compliance.ai).

- **Integration of Effort:** Technical measures must be matched with process controls and culture. Training, documentation, and continuous review are as crucial as any software tool. Companies should document applicable record requirements and risk-based controls as part of their AI lifecycle management (FDA).

In practical terms, implementing ALCOA+ in ML yields concrete benefits: reproducibility of experiments, defensible model audits, and (ultimately) better product outcomes. As regulators and the public demand trustworthy AI, organizations with mature data-integrity practices will have a competitive advantage. Conversely, those who neglect it risk not only compliance violations, but also damage to their AI investments and reputations.

Looking ahead, the landscape will only grow more demanding. New laws (EU AI Act, AI in medical devices) and guidance (FDA's evolving AI frameworks) all emphasize “explainable, audited, verifiable” AI – in other words, AI that meets the spirit of ALCOA+ even if it doesn't say so by name. The industry must therefore keep pushing the envelope: developing better data observability tools, refining standards for AI data governance, and perhaps sharing anonymized integrity data as part of regulatory submissions.

In conclusion, **data integrity and AI are inseparable** in modern practice. Applying ALCOA+ principles supports data integrity; compliance must be assessed against the applicable requirements for the system's intended use. The evidence and expert guidance reviewed throughout this report make one thing clear: the age of AI is also the age of data accountability, and ALCOA+ provides a vital foundation for that accountability (validfor.com).

References: Credible sources cited in the text, including regulatory guidance and peer-reviewed publications, back up all claims ([pmc.ncbi.nlm.nih.gov](https://pubmed.ncbi.nlm.nih.gov)) (www.sciencedirect.com) (validfor.com) (www.auriacompliance.com) (www.mdpi.com) (www.deepchecks.com) (www.deepchecks.com) (blogs.atlas-compliance.ai) (www.mdpi.com). (See inline citations for direct links to each source.)

Source links

Links cited in this article, in order of first appearance. Full addresses are provided for readers using extracted text.

1. <https://acrpnet.org/glossary/attributable-legible-contemporaneous-original-accurate-alcoa>
2. <https://www.sciencedirect.com/science/article/pii/S2949866X24001060>
3. <https://www.fda.gov/regulatory-information/search-fda-guidance-documents/part-11-electronic-records-electronic-signatures-scope-and-application>
4. https://assets.publishing.service.gov.uk/media/5aa2b9ede5274a3e391e37f3/MHRA_GxP_data_integrity_guide_March_edited_Final.pdf
5. <https://pubmed.ncbi.nlm.nih.gov/articles/PMC10997167/>
6. <https://www.mdpi.com/2306-5729/10/12/201>
7. <https://validfor.com/ai-in-the-age-of-regulated-work-with-alcoa-principles/>
8. <https://blogs.atlas-compliance.ai/how-is-alcoa-evolving-with-ai-and-digital-systems-in-2025/>
9. https://www.mdpi.com/1999-4893/14/12/341?type=check_update&version=1
10. <https://www.qad.com/blog/2024/09/using-alcoa-to-ensure-data-integrity-in-the-age-of-ai>

11. <https://www.auriacompliance.com/gmp-blog/data-integrity-and-ai-integration-key-considerations-for-compliance-in-gmp-pharmaceutical-manufacturing>
12. <https://www.beckman.co.za/resources/industry-standards/alcoa>
13. <https://intuitionlabs.ai/articles/generative-ai-gxp-validation-part-11>
14. <https://www.fda.gov/medical-devices/software-medical-device-samd/good-machine-learning-practice-medical-device-development-guiding-principles>
15. <https://www.deepchecks.com/why-data-integrity-is-crucial-for-effective-ml-monitoring/>
16. https://www.arnoldporter.com/-/media/files/perspectives/publications/2024/11/report_the-convergence-of-life-sciences-and-artificial-intelligence_november-2024.pdf
17. <https://mlops-streamlining-ml-lifecycles.pages.dev/data-governance-mlops.html>
18. <https://www.fda.gov/regulatory-information/search-fda-guidance-documents/computer-software-assurance-production-and-quality-management-system-software>
19. <https://lakefs.io/blog/building-compliant-ml-pipelines/>
20. <https://www.ecfr.gov/current/title-21/chapter-I/subchapter-A/part-11>
21. <https://www.rockwellautomation.com/en-pr/company/news/press-releases/state-of-smart-manufacturing-life-sciences-2025.html>
22. <https://www.fda.gov/news-events/press-announcements/fda-proposes-framework-advance-credibility-ai-models-used-drug-and-biological-product-submissions>
23. <https://digital-strategy.ec.europa.eu/en/policies/regulatory-framework-ai>
24. https://www.linkedin.com/posts/dr-dushyant-joshi-msc-phd-mba_pharma-dataintegrity-alcoa-activity-7393633629014233088-6ME9

THE PUBLISHER

About IntuitionLabs

IntuitionLabs is an AI consulting, custom software development and data engineering firm serving pharmaceutical, biotechnology, medical-device and other life-science organizations. We work with clinical, regulatory, medical-affairs, commercial, quality and IT teams to connect technology decisions with the work people need to accomplish.

AI consulting and adoption

Our AI enablement services cover readiness assessments, use-case selection, governance and policies, team workshops, adoption measurement and ongoing advisory support. We help organizations structure the information layer behind AI: source material, context, permissions and maintained knowledge that make generated answers useful and reviewable. Private LLM inference and hosted AI options support teams evaluating how to operate AI with appropriate control over their data and infrastructure.

Software, data and life-science workflows

IntuitionLabs develops custom software for pharma and biotech, integrates enterprise systems, and builds data engineering and business intelligence solutions. Areas of focus include AI agents, regulatory research, medical writing, medical affairs, CMC information, competitive intelligence and clinical-document workflows. Our eTMF intelligence work includes cross-system reconciliation and inspection-readiness support.

Enterprise platforms and regulated delivery

We provide Veeva services, application support, managed services, integrations and custom applications, alongside enterprise content work involving platforms such as Egnyte. For regulated workflows, our services include GxP enablement, computer-system validation and software development addressing 21 CFR Part 11 requirements. The applicable controls, validation responsibilities and acceptance criteria are defined for each engagement.

Work with IntuitionLabs

Explore [AI enablement](#), [pharma and biotech software development](#), [data engineering and BI](#), and [Veeva services](#). [Contact IntuitionLabs](#) to discuss your workflow, information sources and implementation needs.

IntuitionLabs publishes educational research to help life-science teams make informed technology decisions. Coverage of a product or organization does not imply a client relationship, endorsement or partnership.

Website: <https://intuitionlabs.ai>

This article is educational. Verify consequential medical, legal, financial, regulatory and technical decisions with appropriate professionals and the cited primary sources. Company services are described separately from the article's research findings.