

# **LESSON 1.8: DOCUMENTATION STANDARDS | WHAT MUST BE WRITTEN DOWN**

# LESSON 1.8: SUPPLEMENTAL READING

Documentation is often viewed as administrative overhead, something created to satisfy compliance checklists, completed after decisions have already been made, or treated as a burdensome formality that slows operational efficiency. This perception fundamentally misunderstands the regulatory purpose of documentation.

From a supervisory perspective, documentation is a **control mechanism**. It's the evidence that governance occurred, that supervision was applied, that risks were considered, and that accountability was assigned. Without documentation, regulators can't verify that any of these things happened, regardless of how thoughtful or rigorous the firm's internal processes may have been.

The regulatory principle is straightforward: undocumented decisions are treated as if they never occurred. Firms may exercise sound judgment, conduct thorough reviews, and apply appropriate oversight, but if those actions are not documented, they're invisible to examiners. In the absence of documentation, regulators must assume that controls didn't exist, that supervision wasn't applied, and that decisions were made informally or haphazardly.

This lesson explains what regulators expect firms to document, why informal or assumed records are rarely sufficient, and how documentation standards must be established before analytics, automation, or AI amplify the consequences of inadequate record-keeping.

## What Regulators Expect to See Documented

Regulators don't expect exhaustive records of every data interaction, every conversation, or every routine decision. That would be impractical and would generate volumes of documentation with little supervisory value. Instead, they expect documentation that reflects **material decisions**, interpretive choices, and governance action. These are the moments where judgment was exercised, assumptions were made, or accountability was assigned.

Effective documentation captures:

**Why a dataset was selected for a particular purpose:** What business need, analytical objective, or regulatory requirement justified collecting or using this data? What alternatives were considered? Why was this dataset deemed appropriate while others were not?

**How the data was interpreted or transformed:** What cleaning, aggregation, derivation, or enrichment steps were applied? What logic governed these transformations? Who decided how categories should be defined or how thresholds should be set?

**What assumptions or limitations were recognized:** What constraints, biases, sampling limitations, or contextual factors were identified? How did these limitations affect what conclusions could be drawn? Were there known gaps in coverage, accuracy, or representativeness?

**Who reviewed or approved its use:** Who was responsible for validating that the data was fit for purpose? What review process was followed? Who had the authority to approve this use case, and did they exercise that authority?

**What controls governed reuse or escalation:** If the data was later repurposed, what approval process applied? If exceptions or anomalies were detected, what escalation path was followed? How were edge cases or unexpected outcomes handled?

The level of documentation required increases as data moves closer to influencing decisions, communications, or client outcomes. For example, exploratory analysis conducted for internal understanding may require lighter documentation than data used to inform client recommendations, credit decisions, or regulatory filings. But even exploratory uses should leave some record of purpose, scope, and limitations.

Documentation should be **proportional to risk and impact**, not uniform across all contexts. High-stakes uses demand rigorous documentation. Low-stakes uses require less. But the absence of documentation altogether signals weak governance.

## Documentation Must Follow the Data

A common and costly documentation failure is documenting only the final output or communication while leaving upstream data decisions undocumented. Firms create polished reports, client presentations, or regulatory filings with extensive supporting materials, but they don't document the earlier decisions about which data to use, how to transform it, or what assumptions to apply.

Regulators evaluate documentation **holistically**. They expect firms to be able to trace decisions back through the entire data lifecycle, from original collection or acquisition through transformation, analysis, and eventual application. If documentation exists only at the endpoint (the final model output, the completed report, the published communication), then it becomes impossible to determine whether earlier interpretive choices were appropriate, supervised, or even deliberate.

This gap often leads regulators to question **the reliability of the entire workflow**. If a firm can't explain how data was selected, validated, or transformed, the fact that the final output is well-documented provides little assurance. The foundation remains opaque, and opaque foundations cannot support defensible conclusions.

Documentation should therefore be **aligned with data movement**, not just endpoints:

- When data is extracted from a source system, there should be a record of what was extracted, when, by whom, and for what purpose
- When data is cleaned or transformed, there should be documentation of what changed, why those changes were necessary, and who validated them
- When datasets are combined or enriched, there should be a record of what sources were used, what logic governed the combination, and what assumptions were introduced
- When **derived outputs** are created, there should be documentation of the derivation logic, the interpretive choices embedded in it, and who approved its use

This doesn't require creating extensive documentation at every step. It requires creating sufficient documentation at key decision points so that the narrative of data use can be reconstructed when needed.

### **Informal Documentation Is Rarely Sufficient**

In practice, many data decisions are discussed in emails, chat messages, verbal conversations, or undocumented meetings. These informal exchanges may reflect real decision-making, genuine oversight, and thoughtful judgment. But they don't reliably meet regulatory expectations for documentation.

Informal records are:

**Difficult to retrieve:** Emails and chat messages are often scattered across systems, accounts, and timeframes. Reconstructing a decision narrative from fragmented communications is labor-intensive and often incomplete.

**Inconsistent in content:** Informal communications vary widely in what they capture. One email chain may document rationale in detail; another may simply approve a decision without context. There's no standardization, making it difficult to ensure that key elements are consistently recorded.

**Rarely structured for audit replay:** Informal documentation is created for immediate operational needs, not for future reconstruction. It may omit critical context, assume shared knowledge, or refer to decisions made elsewhere without explanation.

**Subject to retention gaps:** Email retention policies, chat message expiration, and turnover of personnel can result in informal documentation disappearing over time, even when formal retention obligations apply.

## Why This Matters Before Analytics or AI

Analytics platforms, machine learning models, and AI-driven systems can generate outputs at scale and speed, but they often **obscure the human decisions embedded in their design and use.**

Models make predictions, rankings, or classifications based on data inputs, feature engineering choices, training methodologies, and optimization objectives, which often reflect interpretive judgment.

Without documentation of these upstream decisions, firms struggle to explain how automated outputs were produced:

- What data was used to train the model, and why?
- What **transformations** were applied before training?
- What assumptions were embedded in feature selection or weighting?
- Who validated that the data was appropriate for this purpose?
- How were edge cases, exceptions, or anomalies handled?

Regulators don't accept automation as a substitute for documentation. In fact, they often expect more documentation, not less, when advanced tools are involved. The opacity of automated systems heightens the need for clear records of the human judgments that shaped them.

Common regulatory questions about AI and analytics include:

- How was the training data selected, and was it representative?
- What interpretive choices were made during data preprocessing?
- Who approved the use of this data for this purpose?
- How do you know the model is performing as intended?
- What happens when the model encounters data it was not trained on?

Firms that haven't documented data selection, transformation, and oversight decisions before deploying AI systems find themselves unable to answer these questions. The automation created efficiency, but it also created accountability gaps that can't be closed retroactively.

Establishing documentation standards **before introducing analytics or AI** ensures that accountability remains visible even as processes scale. It creates the audit trail that allows firms to demonstrate governance, not just outcomes.

## **Conclusion**

Documentation isn't administrative overhead. It's evidence that governance, supervision, and accountability existed. Without documentation, regulators can't verify that decisions were deliberate, that risks were considered, or that oversight was applied.

Effective documentation captures intent, rationale, and responsibility at the time decisions are made. It follows data through its lifecycle, not just to its endpoint. It's deliberate, accessible, and retained according to policy.

**Before introducing analytics, automation, or AI**, firms must establish documentation standards that ensure accountability remains visible even as systems scale. This isn't a constraint on innovation. It's the foundation that allows advanced tools to be deployed responsibly, with the audit trail that regulators expect and that firms need to defend their practices.