

Quality & documentation

A review framework for ingredient identity, analytical evidence and project-specific documentation.

SCOPE - This document describes questions to ask during qualification. It is not a COA, test report, certificate or declaration that Nerivanta holds any accreditation or regulatory file.

Review the evidence as a set

Assess what a file covers, who issued it and whether it applies to the intended material. A product name or certificate label is not a substitute for product, site, method, date and batch details.

Review area	Evidence to examine	Review question
Identity	Product name, source, composition and molecular form.	Does the material match the agreed brief?
Oxidation	PV, AV, TOTOX and the associated analytical methods.	Are units, methods and limits explicit?
Contaminants	An application-specific panel with methods and limits.	Does the test scope address the target specification?
Credentials	Issuer, holder, site, product scope and validity.	Is the claimed coverage supported by the actual file?
Traceability	Lot identifiers and links between production and shipment records.	Can the reviewed sample be related to the supplied lot?
Packaging	Container, closure, headspace and handling instructions.	Are the conditions defined for the proposed shipment?

Set acceptance criteria before sample assessment and record any missing evidence. This framework does not replace product-specific technical or regulatory review.

A product-specific file set

Organize the requested evidence around the intended product, manufacturing site and destination market.

Project type	Files to request	Additional checks
Nutrition ingredient	Specification, COA, TDS, SDS and relevant test reports.	Sensory requirements, storage and packaging.
API / regulated	Approved technical and regulatory files applicable to the project.	Holder, site, product coverage and market applicability.
Softgel project	Oil and finished-product specifications, formula and stability information.	Capsule format, packaging and release criteria.
Animal nutrition	Composition, analytical evidence and handling instructions.	Intended species/application and shipping conditions.

Certificate names are not certificates

Terms such as GMP, CEP and DMF may appear in a project brief. They are review requirements, not confirmed Nerivanta credentials. No certification logo, certificate number or accreditation claim has been added to this draft.

Approval record

Before external publication	Required confirmation
Company identity	The registered entity and approved contact details.
Product information	The current specification, its owner and release status.
Credentials	The actual evidence and authorization to make any claim.
Document control	An approver, revision identifier and publication date.

REVIEW COPY ONLY - Product and company details remain unapproved. Keep this version out of the public download area until those details and claims have been signed off.