

ALIV Sample Documentation Packet

Illustrative partner review copy

Status: Redacted sample for workflow and document-layout review

Important: This packet is illustrative only. It is not a Certificate of Analysis, release record, sterility certificate, or evidence that a specific product has been manufactured or tested. Product-specific documentation is issued only through the applicable approved program and qualified partner.

1. Sample Certificate of Analysis

Field	Illustrative entry
Product	ALIV Partner Program — Illustrative Peptide A
Product code	[REDACTED]
Lot / batch number	[REDACTED]
Manufacturing partner	[REDACTED — qualified partner]
Manufacturing date	[REDACTED]
Report date	[REDACTED]
Specification	Approved program specification on file
Testing laboratory	[REDACTED — qualified laboratory partner]
Final disposition	Illustrative layout only — not a release decision

Illustrative analytical summary

Test	Method	Specification	Result	Status
Identity	Method defined by approved program	Conforms to approved specification	[REDACTED]	Illustrative
Purity	HPLC / UPLC where applicable	Program-specific	[REDACTED]	Illustrative
Mass confirmation	Mass spectrometry where applicable	Program-specific	[REDACTED]	Illustrative
Sterility	Where applicable to product and channel	Program-specific	[REDACTED]	Illustrative
Endotoxin	Where applicable to product and channel	Program-specific	[REDACTED]	Illustrative

Authorized approval: [REDACTED]

Signature / date: [REDACTED]

Actual test methods, specifications, results, signatures, and release decisions vary by product, manufacturing partner, intended market, and approved program.

2. Sample lot traceability sheet

Traceability field	Illustrative entry
Product code	[REDACTED]
Product presentation	[REDACTED]
Lot / batch number	[REDACTED]
Source / manufacturing partner	[REDACTED]
Packaging lot	[REDACTED]
Label version	[REDACTED]
Quantity received	[REDACTED]
Date received	[REDACTED]
Storage instruction	Per documented product requirement
Documentation reference	[REDACTED]
Shipment reference	[REDACTED]

Chain-of-custody checkpoints

1. **Program setup:** Product, presentation, label, and documentation requirements confirmed.
 2. **Receipt:** Incoming product counted and matched to the configured product and lot records.
 3. **Packaging:** Approved packaging and label version recorded against the applicable lot.
 4. **Order fulfillment:** Shipment linked to the applicable product and lot information where required.
 5. **Record retention:** Supporting documentation retained according to the approved program workflow.
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3. Sample release summary

Program review

The applicable product, manufacturing pathway, testing plan, packaging requirements, documentation set, and fulfillment channel are reviewed before a program begins.

Release package may include

- Batch-specific Certificate of Analysis
- Applicable identity, purity, assay, or related-substance results
- Sterility or endotoxin documentation where applicable
- Lot and packaging traceability
- Manufacturing and release records supplied through the applicable qualified partner
- Storage and stability information where available for the approved program
- Fulfillment and shipment records associated with the order

Partner review notes

- The exact documentation set is product- and program-specific.
 - ALIV coordinates documentation through qualified manufacturing and laboratory partners.
 - Product availability, eligibility, permitted use, and regulatory obligations vary by product, customer type, jurisdiction, and applicable channel.
 - The ALIV corporate address is not represented as a manufacturing facility.
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Request the applicable documentation set

Approved partners can review program-specific product documentation, inventory, ordering, and fulfillment tools through secure account access. For program questions, submit an inquiry through the ALIV website contact form.

ALIV

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