BATCH TEST LABORATORIES LIMITED

Independent Analytical Testing & Quality Verification

SPECIMEN — ILLUSTRATIVE TEMPLATE. NOT A REAL BATCH RESULT.
This document is a worked example showing how Batch Test Labs handles third-party results that fall below the agreed specification. All product names, batch numbers, laboratory data, and outcomes on this document are fictional. It exists so that our supplier can see, in advance and in our standard format, exactly what happens when agreed numbers are not met.

BATCH REJECTED

Specification Not Met | Illustrative Outcome Document

BATCH TEST VERIFICATION REPORT — SPECIMEN

Report Ref: BTL-SPEC-SHORTFALL-0000
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Page: 1 of 2

PURPOSE OF THIS SPECIMEN

This specimen is issued to our supplier as part of our quality agreement. It shows, using fictional data, the standard Batch Test Labs process that follows when an independent third-party laboratory returns results below the numbers agreed between us. It applies to every batch, without exception, and is shared in the spirit of the agreement itself: **agreed numbers are the standard, results are verified independently, and both sides know exactly what happens when they are not met.** Nothing on this document refers to a real batch.

1. PRODUCT INFORMATION (ILLUSTRATIVE)

Product Name	Illustrative Peptide (fictional)	**Batch Number**	SPEC-0000-00 (fictional)
Catalog / SKU	BTL-SPEC (fictional)	**Quantity**	10mg × 10vials (fictional)
Intended Use	Research Use Only (RUO) – Not for human or veterinary consumption	**Sample ID**	BTL-SMP-SPEC-0000 (fictional)

2. BATCH TEST IN-HOUSE VERIFICATION (ILLUSTRATIVE)

In this worked example the batch passes all seven physical checks. This is realistic — and it is the point: physical checks describe the container, not the contents. Only independent HPLC/MS testing confirms what is inside.

Check Item	Result	Status
Vial Integrity	PASS	PASS
Seal / Cap Security	PASS	PASS
Vacuum Seal (Suction)	PASS	PASS
Powder Appearance	PASS	PASS
Reconstitution Clarity	PASS	PASS
Mixing Behaviour	PASS	PASS
Packaging Integrity	PASS	PASS

3. AGREED SPECIFICATION

The acceptance criteria below are the numbers agreed between Batch Test Labs and our supplier and recorded in correspondence. Every batch is held to these numbers, verified by an independent third-party laboratory.

Parameter	Agreed Standard	Verified By
Identity	Named peptide confirmed (HPLC/MS)	Independent third-party laboratory
Purity	>= 99.0% (HPLC)	Independent third-party laboratory
Net Content per Vial	At stated mass or above; substantially under is not acceptable	Independent third-party laboratory

4. INDEPENDENT THIRD-PARTY VERIFICATION — ILLUSTRATIVE BELOW-SPEC RESULTS

Illustrative example: the laboratory confirms the peptide is genuine, but purity and net content fall below the agreed numbers. The verify link below is fictional; real reports carry a live public verify link.

Test Item	Agreed	Illustrative Result	Status
Identity (HPLC/MS)	Named peptide detected	Named peptide detected	PASS
Purity (HPLC)	>= 99.0%	98.20% (fictional)	FAIL
Net Content per Vial	At stated mass or above	9.31mg vs 10mg stated — 93.1% (fictional)	FAIL
Lab Report Reference	Publicly verifiable	uzorak.com/verify — report ID SPEC-000 (fictional)	VERIFY

5. WHAT HAPPENS NEXT — SUPPLIER ACCOUNTABILITY PROCESS

1. Immediate hold. The batch is placed on hold the day results arrive. If listed, its sale listing is withdrawn the same day.

2. No Certificate of Analysis. A batch below the agreed specification never carries the green COA badge. No exceptions.

3. Retired designation. The batch designation is retired. The stock is not sold, in whole or in part, and sealed units are retained as evidence.

4. Supplier investigation. We share the full laboratory report and its public verify link with our supplier and expect a root-cause account: manufacturing, handling, or logistics.

5. Remedy. Replacement stock or credit, at the supplier's cost, within an agreed window — our customer's orders must still be met.

6. Full re-testing of any replacement. Replacement stock enters as a NEW batch: full 7-point in-house checks plus a third-party baseline before any sale.

7. Publication. Results are published — pass or fail — with the verify link, because our commitment is to test everything and publish everything. Our supplier's identity is never disclosed; the data speaks for itself.

8. Escalation. A second consecutive below-specification batch triggers a formal review of the supply arrangement.

QUALITY STATEMENT

Batch Test Laboratories Limited tests every batch it receives and publishes the results of independent third-party verification, whether they pass or fail. This specimen shows the standard that applies when agreed numbers are not met. It is shared with our supplier so that expectations are clear in advance — the accountability process is not a threat, it is the agreement working as designed.

Verified By

_____

Signature & Date

Approved By (QA)

_____

Signature & Date

REGULATORY DISCLAIMER

Caution: This product has not been approved for human or veterinary consumption. It is intended for research and laboratory use only. The purchaser assumes all liability for the use of this product in compliance with applicable laws and regulations. Batch Test Laboratories Limited assumes no liability for misuse or misrepresentation of tested materials.

This specimen is a process template. It is not a Certificate of Analysis, does not certify any product, and must not be presented as a test result for any real batch.

HOW TO VERIFY ANY COA — A QUICK GUIDE

1. **Check who performed the test** — Was it the seller themselves, or an independent accredited laboratory?
2. **Look for a verifiable reference** — Can you cross-check the lab report number directly with the testing facility?
3. **Ask about accreditation** — Does the testing laboratory hold ISO 17025 or equivalent certification?
4. **Be cautious of perfection** — Claims of 99.99% purity with no supporting data or lab contact should prompt questions.

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