

# FSVP VERIFICATION ACTIVITY — SAMPLING AND TESTING OF THE FOOD

21 CFR 1.506(e)(1)(ii) — documentation of a supplier verification activity

**Importer**

SAMPLE — Example Foods Co.

**Foreign supplier**

Example Nut Processing Ltd. (fictional)

**Food**

SAMPLE-001 — Roasted cashews, 25 kg cartons (fictional)

**THE ACTIVITY****Date(s) the test(s) were conducted (21 CFR 1.506(e)(1)(ii))**

2026-09-12

**Conducted by ( § 1.506(e)(2))**

Another entity, relied on under 21 CFR 1.506(e)(2)(i)

Example Testing Laboratory (fictional)

**Documentation that the activity was conducted by a qualified individual ( § 1.503(a))**

Accredited testing laboratory (fictional accreditation certificate on file).

**Food tested, including lot number ( § 1.506(e)(1)(ii))**

SAMPLE-001 — Roasted cashews, 25 kg cartons (fictional)

Lot EX-2026-0912 (fictional)

**Number of samples tested**

5

**Test(s) conducted, including analytical method(s)**

Salmonella by the laboratory's validated method; aflatoxin by HPLC (fictional report).

**Date of the report of the testing**

2026-09-14

**Laboratory conducting the testing**

Example Testing Laboratory, 100 Example Way, Sampletown (fictional)

**RESULTS****Results of the testing (21 CFR 1.506(e)(1)(ii))**

Salmonella not detected in the samples tested; aflatoxin below the importer's specification (fictional results).

**Corrective actions taken in response (21 CFR 1.506(e)(1)(ii))**

None recorded

**REVIEW AND ASSESSMENT OF THE RESULTS (21 CFR 1.506(e)(3))****Date reviewed**

2026-09-14

**Assessment ( § 1.506(e)(3))**

Results reviewed by the importer's qualified individual; the lot was released (fictional).

**SIGNATURE (21 CFR 1.510(a)(2))**

I reviewed this record. It is accurate and complete to the best of my knowledge, and I sign and date it as required by 21 CFR 1.510(a)(2).

**Signed by**

Alex Sample, FSVP qualified individual (fictional)

**Date**

2026-09-14

Sample — not signed or filed. The signer named above is fictional; no signature, filing record or hash exists for this document.

Prepared in Aleph from the importer's own answers; Aleph did not assess them. FDA prescribes no form for this record (21 CFR 1.510(a)(1)).

FOR REFERENCE ONLY