SUPPLIER EVIDENCE PILOT

Resolve the supplier before relying on the record.

A bounded public-data pilot for multilingual supplier identity, FDA record matching and review-ready evidence requests.

PREPARED FOR

Weee
Food Safety & Compliance
Global+ Vendor Operations

The operational problem

A marketplace listing may identify a seller, consumer brand and country of origin without identifying the exact legal manufacturer, physical facility, US owner/consignee or FSVP importer. A brand-name search can then attach an FDA record to the wrong entity. The result is either false confidence or repeated manual research.

PILOT PRINCIPLE

Resolve identity before attaching a record.

Unproven matches remain visible, are explained, and become a minimum supplier evidence request.

01 What Origami Lab will deliver

Identity map	Separate seller, brand owner, legal manufacturer, physical facility, exporter, US owner/consignee and FSVP importer.
FDA evidence screen	Search relevant public records using confirmed names, aliases, addresses, countries and product scope.
False-match exclusions	Preserve plausible candidates that were rejected, with the exact reason they were not attached to the SKU.
Bilingual evidence request	Turn every unresolved field into a precise English/Japanese question and the smallest supporting-document request.
Review-ready memo	Provide source URLs, retrieval dates, decision boundaries and an audit trail for qualified human review.

02 Why this sample is useful

The attached Raoh sample shows two problems at once: inconsistent sell-unit imagery on the listing, and an unrelated Nissin-named entity appearing in a public FDA record. The memo refuses to collapse that record into the Japan-origin SKU and asks for the evidence needed to continue.

03 Bounded pilot scope

Included

01 20 Weee-listed Japanese shelf-stable food SKUs

02 Up to 20 supplier or manufacturer identity groups

03 Public English and Japanese sources

04 One memo per SKU plus one portfolio review queue

05 One review meeting and one correction round

Not included

01 Legal advice or an FSVP compliance determination

02 Supplier approval, shipment release or FDA clearance

03 Private system access or Seller Center integration

04 Authentication of supplier-provided documents

05 Continuous monitoring beyond the pilot window

04 Ten-business-day workflow

DAYS 01–02

Confirm cohort

Agree on the 20 public listing URLs and the reviewer who will receive unresolved items.

DAYS 03–08

Build evidence

Resolve identities, screen FDA records, document exclusions and generate supplier questions.

DAYS 09–10

Review and correct

Deliver the portfolio queue, review exceptions and apply one factual correction round.

05 Commercial test

Completed sample

1 SKU attached

The Raoh memo demonstrates the method before purchase. No confidential Weee data was used.

Introductory evaluation

$750 fixed fee

20 SKUs, portfolio review queue, bilingual evidence requests, review meeting and one correction round.

To start: written approval, the 20 listing URLs and one designated reviewer. Invoiced on delivery, Net 15. Origami Lab can complete US sole-proprietor vendor setup and provide a W-9. Offer valid for 30 days from submission.

06 Success measures

Review time	Median manual research minutes saved per SKU.
Identity quality	Useful confirmed links, rejected false matches and reviewer overrides.
Actionability	Unresolved items converted into document requests a vendor-operations team can send.

Responsibility boundary. Origami Lab assembles and explains evidence. Public agencies remain the source authorities. Weee and its qualified reviewers retain all supplier, product, import and compliance decisions.