

SupplierCheckFDA - Supplier Evidence Report

Ferrara Candy Company | FEI 3010631837
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SAMPLE REPORT - a real firm, built from FDA's public records, scored by the published rubric at suppliercheckfda.com/methodology/

This report summarizes FDA enforcement history for **Ferrara Candy Company** based on publicly available data from the FDA Data Dashboard. It is designed to support the supplier evaluation component of your Foreign Supplier Verification Program (FSVP) under 21 CFR §1.505(a)(1)(iii).

No Adverse Findings

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Next review recommended by: **August 27, 2029** (per 21 CFR §1.505(c) reevaluation requirements)

Domain Analysis - per 21 CFR §1.505(a)(1)(iii)

Domain	Level	Finding
Inspections	LOW	NAI-only inspections
Compliance Actions	LOW	No compliance actions on record
Import Refusals	LOW	No import refusals on record
Citations	LOW	No citations on record

FDA Data Sources

Firm Profile: <https://datadashboard.fda.gov/oii/firmprofile.htm?FEI=3010631837>
Inspections: <https://datadashboard.fda.gov/ora/cd/inspections.htm>
Compliance Actions: <https://datadashboard.fda.gov/ora/cd/complianceactions.htm>
Import Refusals: <https://datadashboard.fda.gov/ora/cd/imprefusals.htm>
Citations: <https://datadashboard.fda.gov/ora/cd/citations.htm>
Scoring methodology (published rubric): <https://suppliercheckfda.com/methodology/>

Appendix - FDA Records Detail

Inspections (2)

Date	Classification	Product Type	Location
2026-08-03	Voluntary Action Indicated (VAI)	Food/Cosmetics	Fairfield, California
2023-02-28	Voluntary Action Indicated (VAI)	Food/Cosmetics	Fairfield, California

Disclaimer: This report is generated from publicly available FDA Data Dashboard records and is provided for informational purposes only. References to 21 CFR Part 1, Subpart L (FSVP) and specific regulatory sections are included for informational context and do not indicate that this report constitutes a compliance determination, legal advice, or a substitute for professional FSVP evaluation. The importer of record is solely responsible for supplier verification decisions under FSVP. FDA records may be incomplete, delayed, or subject to correction. Always verify critical findings directly with the FDA and the supplier.