California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

DSAART LLC – AART Calf Implant Recall

Recall Date	Product Description	Recalling Firm	Recall Reason
12/18/2025	AART Calf Implant, silicone elastomer implants intended for cosmetic calf augmentation, available in multiple styles and sizes (models 502-103 through 502-304).	DSAART LLC Carson City, NV, USA	Products deviated from established specifications and were distributed without required FDA premarket approval or clearance.

Recall Class	Product Identification	Distribution	Affected Dates
II	**Model No.:** 502-103, 502-104, 502-105, 502-106, 502-201, 502-303, 502-304 **UDI-DI**: **502-103**: B490502103 **502-104**: B490502104 **502-105**: B490502105 **502-106**: B490502106 **502-201**: B490502201 **502-303**: B490502303 **502-304**: B490502304 **Lot Codes**: **502-103**: 23-11-014, 24-07-016, 25-04-018 **502-104**: 24-02-011, 24-07-017, 25-04-003 **502-105**: 24-05-019 **502-106**: 24-09-006, 24-12-011 **502-201**: 24-01-001 **502-303**: 23-06-019 **502-304**: 24-12-013, 25-03-005 **S/N range:** NA	35 Units; Nationwide including California	September 2025 and prior.

For additional information, please visit the FDA Website.