State of California—Health and Human Services Agency
California Department of Public Health

Food and Drug Branch – Device Recalls

California Device Recall Information Sheet

Stryker Updates Use Instructions For Patient-Fitted Tmj Implants

Recall Date	Product Description	Recalling Firm	Recall Reason
4/15/2026	**Tmj Bilateral Implant** REF: CHG010-32 Sterile EO, Rx Only	**STRYKER/TMJ SOLUTIONS INC** Ventura, California	Patient-fitted temporomandibular joint implant contained discrepancy in screw hole position

Recall Class	Product Identification	Distribution	Affected Dates
I	**Tmj Bilateral Implant** UDI: 07613327626551,07613327626568,07613327626575,07613327626605,07613327627749 07613327627848/ Lot #s: 2410301062-2501021011	63 Units Nationwide Including California	October 2025 and prior

For additional information, please visit the FDA website