State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Mesa Biotech recalls Accula SARS-CoV-2 Test for Increased False Positives

Recall Date	Product Description	Recalling Firm	Recall Reason
4/06/2022	**Accula SARS-CoV-2 Test** REF: COV4100	**MESA BIOTECH INC** San Diego, California	SARS-CoV-2 Test has an increased potential for false positive results, which may lead to a delay in correct diagnosis and treatment.

Recall Class	Product Identification	Distribution	Affected Dates
I	UDI/DI: B540COV41000; P22006-026 - P22032-011	Nationwide including California	December 2021 – February 2022

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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