State of California—Health and Human Services Agency

California Department of Public Health

Food and Drug Branch – Device Recalls

CALIFORNIA DEVICE RECALL INFORMATION SHEET

Universal Meditech Recalls PrestiBio Ketone and Urinalysis Test Strips for Distributing without Marketing Authorization

Recall Date	Product Description	Recalling Firm	Recall Reason
11/1/2023	**PrestiBio KETONE TEST STRIPS** REF 900-1KET 100 STRIPS/+50 FREE	**UNIVERSAL MEDITECH INC.** Reedley, California	Universal Meditech Inc. was violatively distributing PrestiBio" Breastmilk Alcohol Test Strip, DiagnosUSÂ¿ SARS-CoV-2 Antibody (IgG/IgM) Test, HealthyWiser KetoFast" Ketone Test Strips and PrestiBio" Ketone Test Strips without marketing authorization. UMI is also recall their tests because they are going out of business and would not be able to continue fulfilling any post-market responsibilities of these distributed products. UMI claims to hold 510(k)s for the other devices, purchased the intellectual property in 2015 from the previous 510(k) holder, but FDA has been unable to verify this information. The devices may have unknown performance

			characteristics because the firm is unable to provide any documentation that the devices were manufactured in conformance with the Quality System regulation, including documentation (e.g., DHF) that the distributed devices had not been modified since original clearance in a way that could impact their safety and effectiveness, documentation of controlled storage temperature/humidity, and post-market surveillance documentation, such as complaints and adverse events.
11/1/2023	**Prestibio URINALASYS TEST STRIP** 10 PARAMETERS REF 900-10	**UNIVERSAL MEDITECH INC.** Reedley, California	Universal Meditech Inc. was violatively distributing PrestiBio" Breastmilk Alcohol Test Strip, DiagnosUSÂ¿ SARS-CoV-2 Antibody (IgG/IgM) Test, HealthyWiser KetoFast" Ketone Test Strips and PrestiBio" Ketone Test Strips without marketing authorization. UMI is also recall their tests because they are going out of business and would not be able to continue fulfilling any post-market responsibilities of these distributed products. UMI claims to hold 510(k)s for

			the other devices, purchased the intellectual property in 2015 from the previous 510(k) holder, but FDA has been unable to verify this information. The devices may have unknown performance characteristics because the firm is unable to provide any documentation that the devices were manufactured in conformance with the Quality System regulation, including documentation (e.g., DHF) that the distributed devices had not been modified since original clearance in a way that could impact their safety and effectiveness, documentation of controlled storage temperature/humidity, and post-market surveillance documentation, such as complaints and adverse events.

Recall Class	Product Identification	Distribution	Affected Dates
II	**PrestiBio KETONE TEST STRIPS** Model Number: 900-1KET UDI-DI Code: None Lot Numbers: All products manufactured after March 2021	N/A Units Nationwide including California	22 May 2023 and prior

II	**Prestibio Urinalasys Test Strip** Model Number: 900-10 UDI-DI Code: None Lot Numbers: All products manufactured after March 2021	N/A Units Nationwide including California	22 May 2023 and prior

FOR ADDITIONAL INFORMATION, PLEASE VISIT THE FDA WEBSITE

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