FDA 510(k) Premarket Notification

CLEARED

K072854 DIACHEX BASIC BLOOD GLUCOSE MONITORING SYSTEM, DIACHEX SUPERB BLOOD GLUCOSE MONITORING SYSTEM

Jun 30, 2008
269 days to decision

K072854 · Product code: **NBW** · Chemistry
Source: https://www.510kdatabase.net/k072854/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Blood Glucose, Over The Counter (NBW)
Date received	Oct 5, 2007
Decision date	Jun 30, 2008
Days to decision	269 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tyson Bioresearch, Inc.**
Location	Washington, DC, US
Contact	WEN-HAI TSAI
510(k) history	13 submissions · 13 cleared · 2006-2018

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k072854/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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