FDA 510(k) Premarket Notification

K903913 PARAMAX URINE/CSF CALIBRATOR LEVEL I AND II

CLEARED

Sep 27, 1990
34 days to decision

K903913 · Product code: **JIX** · Chemistry
Source: https://www.510kdatabase.net/k903913/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrator, Multi-analyte Mixture (JIX)
Date received	Aug 24, 1990
Decision date	Sep 27, 1990
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	DANA JONES
510(k) history	505 submissions · 496 cleared · 1977-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k903913/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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