FDA 510(k) Premarket Notification

K932178 AIA-PACK PA CALIBRATION VERIFICATION/LINEARITY TES

CLEARED

Sep 21, 1993
139 days to decision

K932178 · Product code: **JJX** · Chemistry
Source: https://www.510kdatabase.net/k932178/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single (specified) Analyte Controls (assayed And Unassayed) (JJX)
Date received	May 5, 1993
Decision date	Sep 21, 1993
Days to decision	139 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tosoh Medics, Inc.**
Location	Washington, DC, US
Contact	JUDITH E LOEBEL
510(k) history	41 submissions · 41 cleared · 1990-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k932178/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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