FDA 510(k) Premarket Notification

K791756 VDC

CLEARED

Nov 27, 1979
83 days to decision

K791756 · Product code: **DXG** · Cardiovascular
Source: https://www.510kdatabase.net/k791756/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Sep 5, 1979
Decision date	Nov 27, 1979
Days to decision	83 days
Third-party review	No

APPLICANT

Company	**Life Sciences Manufacturing, Inc.**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1979-1991

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k791756/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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