FDA 510(k) Premarket Notification

K833819 M305

CLEARED

May 9, 1984
189 days to decision

K833819 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k833819/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Nov 2, 1983
Decision date	May 9, 1984
Days to decision	189 days
Third-party review	No

APPLICANT

Company	**Burkick Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1984-1984

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k833819/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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