FDA 510(k) Premarket Notification

K823693 MYELOGRAPHY NEEDLE

CLEARED

Feb 7, 1983
61 days to decision

K823693 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k823693/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Dec 8, 1982
Decision date	Feb 7, 1983
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Ocean Medical Products, Ltd.**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1981-1988