FDA 510(k) Premarket Notification

K790953 BORDISPO BRAND HYPODERMIC NEEDLE

CLEARED

Jun 27, 1979
36 days to decision

K790953 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k790953/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	May 22, 1979
Decision date	Jun 27, 1979
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Borda Products, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979