FDA 510(k) Premarket Notification

K791660 PREPACELL-BONE MARROW SAMPLER

CLEARED

Sep 12, 1979
22 days to decision

K791660 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k791660/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 21, 1979
Decision date	Sep 12, 1979
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Applied Medical Devices, Inc.**
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1979-1984