

K791660 PREPACELL-BONE MARROW SAMPLERSep 12, 1979
22 days to decisionK791660 · Product code: **FMF** · General HospitalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k791660/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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