FDA 510(k) Premarket Notification

K820454 THE RED CAP

CLEARED

Mar 8, 1982
18 days to decision

K820454 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k820454/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 18, 1982
Decision date	Mar 8, 1982
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Burron Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820454/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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