FDA 510(k) Premarket Notification

K930217 BAXTER DISPOSABLE INTRAOSSEOUS INFUSION NEEDLE

CLEARED

Feb 28, 1994
409 days to decision

K930217 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k930217/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jan 15, 1993
Decision date	Feb 28, 1994
Days to decision	409 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Baxter Healthcare Corp**
Location	Mchenry, IL, US
Contact	JAN ROBERTSON
510(k) history	505 submissions · 496 cleared · 1977-2019