FDA 510(k) Premarket Notification

K840490 FLUID REMOVAL ASSEMBLY

CLEARED

Apr 17, 1984
71 days to decision

K840490 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k840490/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 6, 1984
Decision date	Apr 17, 1984
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**Ackrad Laboratories**
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1979-2002