FDA 510(k) Premarket Notification

K822420 REPLAM HYDROXYAPATATITE

CLEARED

Sep 28, 1982
47 days to decision

K822420 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k822420/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 12, 1982
Decision date	Sep 28, 1982
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Replam Corp.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983