FDA 510(k) Premarket Notification

K982018 LIGHT SABER ASPIRATION NEEDLE

CLEARED

Aug 7, 1998
60 days to decision

K982018 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k982018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 8, 1998
Decision date	Aug 7, 1998
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Minrad, Inc.**
Location	Washington, DC, US
Contact	JOHN MCNEIRNEY
510(k) history	14 submissions · 14 cleared · 1997-2007