FDA 510(k) Premarket Notification

K880342 UROSTHESIA

CLEARED

Feb 25, 1988
30 days to decision

K880342 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k880342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 26, 1988
Decision date	Feb 25, 1988
Days to decision	30 days
Third-party review	No

APPLICANT

Company	**Medical Research Innovators, Inc.**
Location	El Cajon, CA, US
Contact	SEAN ROWLAND
510(k) history	1 submissions · 1 cleared · 1988-1988