FDA 510(k) Premarket Notification

K990288 FREE FLOW SAFETY DEVICE, MODEL MG 245052

CLEARED

Feb 24, 1999
26 days to decision

K990288 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k990288/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 29, 1999
Decision date	Feb 24, 1999
Days to decision	26 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Teva Medical, Ltd.**
Location	Kulpsville, PA, US
Contact	MARYANN MELUS
510(k) history	2 submissions · 2 cleared · 1997-1999