FDA 510(k) Premarket Notification

K820453 THE INT-STOPPER

CLEARED

Mar 8, 1982
18 days to decision

K820453 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k820453/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 18, 1982
Decision date	Mar 8, 1982
Days to decision	18 days
Third-party review	No

APPLICANT

Company	**Burron Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987