FDA 510(k) Premarket Notification

K801605 BIMECO FILTERED EXTENSION SET

CLEARED

Aug 12, 1980
28 days to decision

K801605 · Product code: **FPB** · General Hospital
Source: https://www.510kdatabase.net/k801605/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	Jul 15, 1980
Decision date	Aug 12, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

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