FDA 510(k) Premarket Notification

K842075 CRITIKON I.V. FILTER

CLEARED

Jun 28, 1984
36 days to decision

K842075 · Product code: **FPB** · General Hospital
Source: https://www.510kdatabase.net/k842075/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Infusion Line (FPB)
Date received	May 23, 1984
Decision date	Jun 28, 1984
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Critikon Company, LLC**
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000