

K801605 BIMECO FILTERED EXTENSION SETAug 12, 1980
28 days to decisionK801605 · Product code: **FPB** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k801605/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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