FDA 510(k) Premarket Notification

K813498 MEDIFIX

CLEARED

Jan 7, 1982
24 days to decision

K813498 · Product code: **KRK** · Cardiovascular
Source: https://www.510kdatabase.net/k813498/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manometer, Blood-pressure, Venous (KRK)
Date received	Dec 14, 1981
Decision date	Jan 7, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

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