

K813498 MEDIFIXJan 7, 1982
24 days to decisionK813498 · Product code: **KRK** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k813498/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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