FDA 510(k) Premarket Notification

K781814 SULFATE, FIBRIQUIK PROTAMINE

CLEARED

Nov 8, 1978
13 days to decision

K781814 · Product code: **JBN** · Hematology
Source: https://www.510kdatabase.net/k781814/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fibrin Monomer Paracoagulation (JBN)
Date received	Oct 26, 1978
Decision date	Nov 8, 1978
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**General Diagnostics**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988