FDA 510(k) Premarket Notification

K770383 STOPCOCK, ONE WAY

CLEARED

Mar 7, 1977
7 days to decision

K770383 · Product code: **KOC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k770383/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Feb 28, 1977
Decision date	Mar 7, 1977
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**Cobe Laboratories, Inc.**
Location	Mchenry, IL, US
Website	https://www.gambro.com
510(k) history	77 submissions · 77 cleared · 1976-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k770383/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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