FDA 510(k) Premarket Notification

K840424 HEMOCATH REPAIR KIT

CLEARED

Mar 5, 1984
33 days to decision

K840424 · Product code: **KNZ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k840424/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, A-v Shunt (KNZ)
Date received	Feb 1, 1984
Decision date	Mar 5, 1984
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003