FDA 510(k) Premarket Notification

K030456 VASC-ALERT

CLEARED

Jul 11, 2003
150 days to decision

K030456 · Product code: **KOC** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k030456/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Feb 11, 2003
Decision date	Jul 11, 2003
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vasc-Alert, LLC**
Location	Chicago, IL, US
Contact	JOHN KENNEDY
510(k) history	1 submissions · 1 cleared · 2003-2003