FDA 510(k) Premarket Notification

K822631 ARGYLE DUO-TUBE

CLEARED

Sep 9, 1982
9 days to decision

K822631 · Product code: **KNT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k822631/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Aug 31, 1982
Decision date	Sep 9, 1982
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Sherwood Medical Co.**
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998