FDA 510(k) Premarket Notification

K822318 ARGYLE MEDIASTINAL DRAIN

CLEARED

Aug 27, 1982
24 days to decision

K822318 · Product code: **GBS** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k822318/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Ventricular, General & Plastic Surgery (GBS)
Date received	Aug 3, 1982
Decision date	Aug 27, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

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