FDA 510(k) Premarket Notification

K810530 PERI-PATCH GLUE MOLD

CLEARED

Mar 24, 1981
26 days to decision

K810530 · Product code: **KGZ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810530/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter (KGZ)
Date received	Feb 26, 1981
Decision date	Mar 24, 1981
Days to decision	26 days
Third-party review	No

APPLICANT

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