FDA 510(k) Premarket Notification

K801421 TUBE HOLDER

CLEARED

Jul 28, 1980
41 days to decision

K801421 · Product code: **LDQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801421/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jun 17, 1980
Decision date	Jul 28, 1980
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Devon Industries, Inc.**
Location	Mchenry, IL, US
Website	https://www.devonmedicalproducts.com
510(k) history	47 submissions · 46 cleared · 1976-1996