FDA 510(k) Premarket Notification

K802248 DISARP ORIGINAL

CLEARED

Sep 26, 1980
10 days to decision

K802248 · Product code: **GAO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k802248/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable (GAO)
Date received	Sep 16, 1980
Decision date	Sep 26, 1980
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Unoplast A/S**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1990