FDA 510(k) Premarket Notification

K820198 DERMA-AID

CLEARED

Mar 5, 1982
39 days to decision

K820198 · Product code: **NAB** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k820198/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze / Sponge,nonresorbable For External Use (NAB)
Date received	Jan 25, 1982
Decision date	Mar 5, 1982
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**Winfield Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 2 cleared · 1982-1997