

K820198 DERMA-AIDMar 5, 1982
39 days to decisionK820198 · Product code: **NAB** · General & Plastic SurgerySource: <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

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Device record: <https://www.510kdatabase.net/k820198/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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