

K831648 ANCHOR BRAND SURGICAL NEEDLESAug 16, 1983
85 days to decisionK831648 · Product code: **GAB** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k831648/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Suturing, Disposable (GAB)
Date received	May 23, 1983
Decision date	Aug 16, 1983
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Anchor Products Co.
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1979-2009

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k831648/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 22, 2026