FDA 510(k) Premarket Notification

K831648 ANCHOR BRAND SURGICAL NEEDLES

CLEARED

Aug 16, 1983
85 days to decision

K831648 · Product code: **GAB** · General & Plastic Surgery
Source: 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