

K813422 SCALPELJan 5, 1982
29 days to decisionK813422 · Product code: **GDX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k813422/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scalpel, One-piece (GDX)
Date received	Dec 7, 1981
Decision date	Jan 5, 1982
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Abco Dealers, Inc.
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991

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Device record: <https://www.510kdatabase.net/k813422/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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