FDA 510(k) Premarket Notification

K800015 AXIOM VESSEL LOOPS

CLEARED

Mar 25, 1980
81 days to decision

K800015 · Product code: **KDC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k800015/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Jan 4, 1980
Decision date	Mar 25, 1980
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**Axiom Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-2000