FDA 510(k) Premarket Notification

K801611 DEEP BALFOUR BLADE FOR WILKINSON RETRACT

CLEARED

Aug 4, 1980
20 days to decision

K801611 · Product code: **GAD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801611/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Jul 15, 1980
Decision date	Aug 4, 1980
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Edward Weck, Inc.**
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992