

K801611 DEEP BALFOUR BLADE FOR WILKINSON RETRACTAug 4, 1980
20 days to decisionK801611 · Product code: **GAD** · General & Plastic SurgerySource: <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

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Device record: <https://www.510kdatabase.net/k801611/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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