

K772376 NEIVERT OSTEOTOMEMar 9, 1978
72 days to decisionK772376 · Product code: **KDG** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k772376/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chisel (osteotome) (KDG)
Date received	Dec 27, 1977
Decision date	Mar 9, 1978
Days to decision	72 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/k772376/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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