FDA 510(k) Premarket Notification

# K772376 NEIVERT OSTEOTOME

**CLEARED**

Mar 9, 1978
72 days to decision

K772376 · Product code: **KDG** · General & Plastic Surgery
Source: 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 |