FDA 510(k) Premarket Notification

# K770305 CUTTER, P607 BONE

**CLEARED**

Mar 9, 1977
21 days to decision

K770305 · Product code: **FZT** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k770305/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Cutter, Surgical (FZT) |
| Date received | Feb 16, 1977 |
| Decision date | Mar 9, 1977 |
| Days to decision | 21 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Dentronix, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 9 submissions · 9 cleared · 1977-1991 |