FDA 510(k) Premarket Notification

# K812166 PERCUSSOR

**CLEARED**

Aug 18, 1981
19 days to decision

K812166 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k812166/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Percussor, Powered-electric (BYI) |
| Date received | Jul 30, 1981 |
| Decision date | Aug 18, 1981 |
| Days to decision | 19 days |
| Third-party review | No |

## APPLICANT

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|---|---|
| Company | **Dhd Medical Products Div. Diemolding Corp.** |
| Location | Mchenry, IL, US |
| 510(k) history | 43 submissions · 43 cleared · 1979-1988 |