

K812166 PERCUSSORAug 18, 1981
19 days to decisionK812166 · Product code: **BYI** · Anesthesiology
Source: <https://www.510kdatabase.net/k812166/>**SUBMISSION DETAILS**

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

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

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Device record: <https://www.510kdatabase.net/k812166/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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