FDA 510(k) Premarket Notification

K832063 ROBIN CORDLESS HANDPIECE

CLEARED

Aug 26, 1983
60 days to decision

K832063 · Product code: **EKX** · Dental
Source: https://www.510kdatabase.net/k832063/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Handpiece, Direct Drive, Ac-powered (EKX)
Date received	Jun 27, 1983
Decision date	Aug 26, 1983
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Kaycor Intl., Ltd.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1997