

K832063 ROBIN CORDLESS HANDPIECEAug 26, 1983
60 days to decisionK832063 · 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

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Device record: <https://www.510kdatabase.net/k832063/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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