FDA 510(k) Premarket Notification

K992610 GENDEX 765DC, MODEL 110-0154

CLEARED

Sep 7, 1999
34 days to decision

K992610 · Product code: **EHD** · Radiology
Source: https://www.510kdatabase.net/k992610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, X-ray, Extraoral With Timer (EHD)
Date received	Aug 4, 1999
Decision date	Sep 7, 1999
Days to decision	34 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dentsply Intl.**
Location	Walker, MI, US
Contact	DANIEL P MURPHY
510(k) history	279 submissions · 279 cleared · 1976-2013