

K992610 GENDEX 765DC, MODEL 110-0154Sep 7, 1999
34 days to decisionK992610 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992610/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026