

**K972312 SIEMENS ORTHOPHOS DS/DS CEPH**Sep 15, 1997  
87 days to decisionK972312 · Product code: **EHD** · Radiology  
Source: <https://www.510kdatabase.net/k972312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, X-ray, Extraoral With Timer (EHD)
Date received	Jun 20, 1997
Decision date	Sep 15, 1997
Days to decision	87 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Siemens AG</b>
Location	Iselin, NJ, US
Contact	STEVEN CLARKE
Website	<a href="http://www.siemens.it/">http://www.siemens.it/</a>
510(k) history	15 submissions · 15 cleared · 1995-2015

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972312/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 16, 2026