

**K962010 EPI-SCOPE OPTION FOR Q SERIES CT SYSTEMS**Mar 21, 1997  
302 days to decisionK962010 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k962010/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 23, 1996
Decision date	Mar 21, 1997
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Philips Medical Systems (Cleveland), Inc.</b>
Location	Mchenry, IL, US
Contact	ROBERT L TUROCY
510(k) history	190 submissions · 190 cleared · 1977-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962010/> 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