FDA 510(k) Premarket Notification

K962010 EPI-SCOPE OPTION FOR Q SERIES CT SYSTEMS

CLEARED

Mar 21, 1997
302 days to decision

K962010 · 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	**Philips Medical Systems (Cleveland), Inc.**
Location	Mchenry, IL, US
Contact	ROBERT L TUROCY
510(k) history	190 submissions · 190 cleared · 1977-2017