

**K052473 CX 250 C PLUS WITH GAMMAXP SOFTWARE, MODEL
0121 0000**Oct 28, 2005
49 days to decisionK052473 · Product code: **IYX** · Radiology
Source: <https://www.510kdatabase.net/k052473/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Sep 9, 2005
Decision date	Oct 28, 2005
Days to decision	49 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Inter Medical Medizintechnik GmbH
Location	Luebbecke, DE
Contact	HANS GUENTER OSIEK
510(k) history	3 submissions · 3 cleared · 2005-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052473/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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