FDA 510(k) Premarket Notification

CLEARED

K873121 GE INTEGRATED DIAGNOSTIC SYSTEM (DRS.PRS)

Aug 23, 1988
378 days to decision

K873121 · Product code: **LMD** · Radiology
Source: https://www.510kdatabase.net/k873121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Aug 11, 1987
Decision date	Aug 23, 1988
Days to decision	378 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	KROGER, PHD
510(k) history	254 submissions · 254 cleared · 1976-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k873121/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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