FDA 510(k) Premarket Notification

K863286 ERS SYSTEM

CLEARED

Oct 27, 1986
63 days to decision

K863286 · Product code: **LMB** · Radiology
Source: https://www.510kdatabase.net/k863286/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Digital Image Storage, Radiological (LMB)
Date received	Aug 25, 1986
Decision date	Oct 27, 1986
Days to decision	63 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k863286/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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