FDA 510(k) Premarket Notification

# K811103 GE 900

**CLEARED**

Jun 9, 1981
47 days to decision

K811103 · Product code: **EHD** · Radiology
Source: https://www.510kdatabase.net/k811103/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Unit, X-ray, Extraoral With Timer (EHD) |
| Date received | Apr 23, 1981 |
| Decision date | Jun 9, 1981 |
| Days to decision | 47 days |
| Third-party review | No |

## APPLICANT

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