

K801653 MAXI CAMERA 37Aug 20, 1980
33 days to decisionK801653 · Product code: **IYX** · Radiology
Source: <https://www.510kdatabase.net/k801653/>**SUBMISSION DETAILS**

| | |
|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Camera, Scintillation (gamma) (IYX) |
| Date received | Jul 18, 1980 |
| Decision date | Aug 20, 1980 |
| Days to decision | 33 days |
| Third-party review | No |

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801653/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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