FDA 510(k) Premarket Notification

K810305 DATASON DB ULTRASOUND IMAGING SYSTEM

CLEARED

Mar 13, 1981
36 days to decision

K810305 · Product code: **HEM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k810305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Imager, Ultrasonic Obstetric-gynecologic (HEM)
Date received	Feb 5, 1981
Decision date	Mar 13, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

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