

K810305 DATASON DB ULTRASOUND IMAGING SYSTEMMar 13, 1981
36 days to decisionK810305 · Product code: **HEM** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026