

**K881185 ANNULAR ARRAY PROBES FOR ARTIS 3200 DIAG.
ULTRA.**Jun 1, 1988
72 days to decisionK881185 · Product code: **HEM** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k881185/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Imager, Ultrasonic Obstetric-gynecologic (HEM)
Date received	Mar 21, 1988
Decision date	Jun 1, 1988
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Philips Medical Systems (Cleveland), Inc.
Location	Mchenry, IL, US
Contact	WILLIAM J GERBER
510(k) history	190 submissions · 190 cleared · 1977-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881185/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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