

**K800095 MGD-3**Feb 11, 1980  
28 days to decisionK800095 · Product code: **KNG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k800095/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Jan 14, 1980
Decision date	Feb 11, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Kt Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1980

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800095/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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