

**K812868 FETAL PULSE DETECTORS, F50/F60**Nov 25, 1981  
42 days to decisionK812868 · Product code: **KNG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k812868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Oct 14, 1981
Decision date	Nov 25, 1981
Days to decision	42 days
Third-party review	No

**APPLICANT**

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Company	<b>South Carolina Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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

Device record: <https://www.510kdatabase.net/k812868/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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