

K810115 DOPPLEX-PLB & PXBJul 16, 1981
181 days to decisionK810115 · Product code: **KNG** · Radiology
Source: <https://www.510kdatabase.net/k810115/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Jan 16, 1981
Decision date	Jul 16, 1981
Days to decision	181 days
Third-party review	No

APPLICANT

Company	Huntleigh Technology, Inc.
Location	Walker, MI, US
510(k) history	23 submissions · 23 cleared · 1981-1999

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Device record: <https://www.510kdatabase.net/k810115/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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