

K812676 AMPLITONE-3Oct 13, 1981
21 days to decisionK812676 · Product code: **LDE** · CardiovascularSource: <https://www.510kdatabase.net/k812676/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Manual (LDE)
Date received	Sep 22, 1981
Decision date	Oct 13, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Progressive Technology, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-2015

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Device record: <https://www.510kdatabase.net/k812676/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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