FDA 510(k) Premarket Notification

# K812676 AMPLITONE-3

**CLEARED**

Oct 13, 1981
21 days to decision

K812676 · Product code: **LDE** · Cardiovascular
Source: https://www.510kdatabase.net/k812676/

## SUBMISSION DETAILS

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|---|---|
| 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

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