

K960295 LONG LIFE RX-12Mar 22, 1996
60 days to decisionK960295 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k960295/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Hearing Aid, Air-conduction, Prescription (ESD) |
| Date received | Jan 22, 1996 |
| Decision date | Mar 22, 1996 |
| Days to decision | 60 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Rexton, Inc. |
| Location | Mchenry, IL, US |
| Contact | DANIEL I ANDERSON |
| 510(k) history | 21 submissions · 21 cleared · 1983-1997 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960295/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 5, 2026