

**K792428 COE-SIL**Jan 23, 1980  
57 days to decisionK792428 · Product code: **LDG** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k792428/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Earmold, Impression (LDG)
Date received	Nov 27, 1979
Decision date	Jan 23, 1980
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Coe Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1976-1990

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

Device record: <https://www.510kdatabase.net/k792428/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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