

**K201370 Multiflex Tinnitus Technology**Jun 19, 2020  
28 days to decisionK201370 · Product code: **KLW** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k201370/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Masker, Tinnitus (KLW)
Date received	May 22, 2020
Decision date	Jun 19, 2020
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Starkey Laboratories, Inc.</b>
Location	Mchenry, IL, US
Contact	Ka Zoua Xiong
510(k) history	60 submissions · 60 cleared · 1977-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201370/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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