

**K790093 MODULE PERSONAL EAR (SUPER)**Jan 26, 1979  
10 days to decisionK790093 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k790093/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 16, 1979
Decision date	Jan 26, 1979
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Qualitone</b>
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1996

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

Device record: <https://www.510kdatabase.net/k790093/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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