

**K790878 SCREENING AUDIOMETER, HIGH FREQUENCY**Aug 3, 1979  
88 days to decisionK790878 · Product code: **EWO** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k790878/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Audiometer (EWO)
Date received	May 7, 1979
Decision date	Aug 3, 1979
Days to decision	88 days
Third-party review	No

**APPLICANT**

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Company	<b>Vicon Instrument Co.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-1981

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

Device record: <https://www.510kdatabase.net/k790878/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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