

**K780921 MM-CE DM (BEHIND THE EAR)**Jun 14, 1978  
9 days to decisionK780921 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k780921/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jun 5, 1978
Decision date	Jun 14, 1978
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Argosy Intl. (Usa), Inc.</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1978-1982

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

Device record: <https://www.510kdatabase.net/k780921/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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