

K811218 ACMI ASPIRATOR KIT #8808Jul 1, 1981
58 days to decisionK811218 · Product code: **EOQ** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k811218/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	May 4, 1981
Decision date	Jul 1, 1981
Days to decision	58 days
Third-party review	No

APPLICANT

Company	American Cystoscope Makers, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1976-1983

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Device record: <https://www.510kdatabase.net/k811218/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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