

K801328 THE EQUALIZERJun 9, 1980
6 days to decisionK801328 · Product code: **MDM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k801328/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Surgical, General Use (MDM)
Date received	Jun 3, 1980
Decision date	Jun 9, 1980
Days to decision	6 days
Third-party review	No

APPLICANT

Company	The Anspach Effort, Inc.
Location	Mchenry, IL, US
510(k) history	60 submissions · 60 cleared · 1980-2022

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Device record: <https://www.510kdatabase.net/k801328/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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