

**K782092 ARMBORD CVER, DISPOSABLE**Jan 10, 1979  
27 days to decisionK782092 · Product code: **FSE** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k782092/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Manual (FSE)
Date received	Dec 14, 1978
Decision date	Jan 10, 1979
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Anago, Inc.</b>
Location	Mchenry, IL, US
510(k) history	44 submissions · 42 cleared · 1979-1993

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

Device record: <https://www.510kdatabase.net/k782092/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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