

**K896019 MODIFIED DISP. CHIBA-TYPE NEEDLE DESIGNATED  
ASPI.**Jan 3, 1990  
79 days to decisionK896019 · Product code: **GAA** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k896019/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Disposable (GAA)
Date received	Oct 16, 1989
Decision date	Jan 3, 1990
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Dip, Inc.</b>
Location	Mchenry, IL, US
Contact	CHET VANHOF
510(k) history	56 submissions · 56 cleared · 1979-1997

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896019/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 2, 2026