

K841299 BIERMAN BIOPSY NEEDLEMay 2, 1984
33 days to decisionK841299 · Product code: **GDM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k841299/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Aspiration And Injection, Reusable (GDM)
Date received	Mar 30, 1984
Decision date	May 2, 1984
Days to decision	33 days
Third-party review	No

APPLICANT

Company	Popper & Sons, Inc.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1984-1991

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Device record: <https://www.510kdatabase.net/k841299/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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