

K841275 FINE NEEDLE ASPIRATION DEVICEAug 17, 1984
143 days to decisionK841275 · Product code: **DRM** · CardiovascularSource: <https://www.510kdatabase.net/k841275/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Mar 27, 1984
Decision date	Aug 17, 1984
Days to decision	143 days
Third-party review	No

APPLICANT

Company	Gyneco, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1984-1986

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Device record: <https://www.510kdatabase.net/k841275/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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