FDA 510(k) Premarket Notification

K841275 FINE NEEDLE ASPIRATION DEVICE

CLEARED

Aug 17, 1984
143 days to decision

K841275 · Product code: **DRM** · Cardiovascular
Source: 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