FDA 510(k) Premarket Notification

K913098 APIRATION BIOPSY SYRINGE GUN, CAT. CODE #54501

CLEARED

Nov 5, 1991
116 days to decision

K913098 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k913098/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Jul 12, 1991
Decision date	Nov 5, 1991
Days to decision	116 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dlp, Inc.**
Location	Mchenry, IL, US
Contact	ROGER W BRINK
510(k) history	56 submissions · 56 cleared · 1979-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k913098/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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