FDA 510(k) Premarket Notification

K882405 HYPODERMIC NEEDLE SHEATH HOLDER

CLEARED

Jun 29, 1988
19 days to decision

K882405 · Product code: **FMJ** · General Hospital
Source: https://www.510kdatabase.net/k882405/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manometer, Spinal-fluid (FMJ)
Date received	Jun 10, 1988
Decision date	Jun 29, 1988
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Ansr Ent., Inc.**
Location	Franktown, CO, US
Contact	WAYNE R TURNER
510(k) history	1 submissions · 1 cleared · 1988-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k882405/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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