FDA 510(k) Premarket Notification

K822784 HEMO-DE

CLEARED

Nov 5, 1982
52 days to decision

K822784 · Product code: **KEM** · Pathology
Source: https://www.510kdatabase.net/k822784/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Clearing (KEM)
Date received	Sep 14, 1982
Decision date	Nov 5, 1982
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Pmp Industries**
Location	Anaheim, CA, US
510(k) history	1 submissions · 1 cleared · 1982-1982

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k822784/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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