FDA 510(k) Premarket Notification

K001522 TRANSQ3

CLEARED

Oct 18, 2000
155 days to decision

K001522 · Product code: **KTB** · Physical Medicine
Source: https://www.510kdatabase.net/k001522/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Iontophoresis, Specific Uses (KTB)
Date received	May 16, 2000
Decision date	Oct 18, 2000
Days to decision	155 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Iomed, Inc.**
Location	Salt Lake City, UT, US
Contact	W. TIM MILLER
510(k) history	17 submissions · 12 cleared · 1990-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k001522/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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