FDA 510(k) Premarket Notification

K982668 PHORESOR II,MODEL PM900

CLEARED

Mar 2, 1999
214 days to decision

K982668 · Product code: **EGJ** · Physical Medicine
Source: https://www.510kdatabase.net/k982668/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Iontophoresis, Other Uses (EGJ)
Date received	Jul 31, 1998
Decision date	Mar 2, 1999
Days to decision	214 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