FDA 510(k) Premarket Notification

# K914621 IONTOPHORESIS DEVICE, MODIFICATION

**CLEARED**

Dec 20, 1991
100 days to decision

K914621 · Product code: **EGJ** · Physical Medicine
Source: https://www.510kdatabase.net/k914621/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent - KD |
| Submission type | Traditional |
| Device classification | Device, Iontophoresis, Other Uses (EGJ) |
| Date received | Sep 11, 1991 |
| Decision date | Dec 20, 1991 |
| Days to decision | 100 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

| | |
|---|---|
| Company | **Iomed, Inc.** |
| Location | Salt Lake City, UT, US |
| Contact | ANNE T CARTER |
| 510(k) history | 17 submissions · 12 cleared · 1990-2007 |