FDA 510(k) Premarket Notification

K925040 TELEPULSE

CLEARED

Aug 23, 1993
322 days to decision

K925040 · Product code: **JEH** · Anesthesiology
Source: https://www.510kdatabase.net/k925040/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Volume (JEH)
Date received	Oct 5, 1992
Decision date	Aug 23, 1993
Days to decision	322 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Biometrix , Ltd.**
Location	Greenwich, CT, US
Contact	MANFRED ASRICAN
510(k) history	4 submissions · 4 cleared · 1993-2015