

K925040 TELEPULSEAug 23, 1993
322 days to decisionK925040 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925040/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026