FDA 510(k) Premarket Notification

K951901 MASTERSCREEN PFT BODY

CLEARED

Jul 31, 1995
98 days to decision

K951901 · Product code: **JEH** · Anesthesiology
Source: https://www.510kdatabase.net/k951901/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Volume (JEH)
Date received	Apr 24, 1995
Decision date	Jul 31, 1995
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Erich Jaeger GmbH & Co. KG**
Location	Wuerzburg, Germany, DE
Contact	DETLEF GROTHEER
510(k) history	4 submissions · 4 cleared · 1995-1998