FDA 510(k) Premarket Notification

K950386 MODEL 3200P/3200PR HYPERBARIC CHAMBER

CLEARED

Jun 13, 1995
132 days to decision

K950386 · Product code: **CBF** · Anesthesiology
Source: https://www.510kdatabase.net/k950386/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Hyperbaric (CBF)
Date received	Feb 1, 1995
Decision date	Jun 13, 1995
Days to decision	132 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sechrist Industries, Inc.**
Location	Mchenry, IL, US
Contact	KAORU LEE ADAIR
510(k) history	23 submissions · 22 cleared · 1978-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k950386/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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