

**K983372 BIOPORT CLOSED BLOOD SAMPLING SYSTEM,
MODEL BT XXXXXX- BP**

Feb 5, 1999
134 days to decision

K983372 · Product code: **DRS** · Cardiovascular
Source: <https://www.510kdatabase.net/k983372/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 24, 1998
Decision date	Feb 5, 1999
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sunscope Intl., Inc.
Location	Westlake Village, CA, US
Contact	PHIL TRIOLO
510(k) history	7 submissions · 7 cleared · 1991-2000

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k983372/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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