

**K073516 DISCYPHOR DIRECT CATHETER SYSTEM,
DISCYPHOR DIRECT NEEDLES, LONG, DISCYPHOR DIRECT
NEEDLES, REGULAR**

Feb 21, 2008
69 days to decision

K073516 · Product code: **BSP** · Anesthesiology
Source: <https://www.510kdatabase.net/k073516/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Conduction, Anesthetic (w/wo Introducer) (BSP)
Date received	Dec 14, 2007
Decision date	Feb 21, 2008
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kyphon, Inc.
Location	Sunnyvale, CA, US
Contact	PAMELA SEGALE
510(k) history	9 submissions · 9 cleared · 1998-2008

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k073516/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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