FDA 510(k) Premarket Notification

K050499 PROBLOC, HN SERIES AND PROLONG, PL SERIES

CLEARED

May 17, 2005
78 days to decision

K050499 · Product code: **BSP** · Anesthesiology
Source: https://www.510kdatabase.net/k050499/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Conduction, Anesthetic (w/wo Introducer) (BSP)
Date received	Feb 28, 2005
Decision date	May 17, 2005
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Life-Tech, Inc.**
Location	Stafford, TX, US
Contact	JEFF KASOFF
510(k) history	14 submissions · 14 cleared · 2005-2012