FDA 510(k) Premarket Notification

K072050 PROLONG-EX, MODELS: PLEX-50CS, PLEX-100CS, PLEX-150CS

CLEARED

Oct 18, 2007
84 days to decision

K072050 · Product code: **BSO** · Anesthesiology
Source: https://www.510kdatabase.net/k072050/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Conduction, Anesthetic (BSO)
Date received	Jul 26, 2007
Decision date	Oct 18, 2007
Days to decision	84 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