

K230603 Arrow Non-Stimulating SnapLock Adapter (K-05520-005C)

Aug 30, 2023
180 days to decision

K230603 · Product code: **BSO** · Anesthesiology
Source: <https://www.510kdatabase.net/k230603/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Conduction, Anesthetic (BSO)
Date received	Mar 3, 2023
Decision date	Aug 30, 2023
Days to decision	180 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Arrow Non-Stimulating Next Gen SnapLock Adapter (Luer Connection) (CA-000010-19); Arrow Non-Stimulating Next Gen SnapLock Adapter (Neuraxial Connection) (CA-000014-19); Arrow Stimulating SnapLock Adapter (with cable) (TZ-02060-001); Arrow Stimulating SnapLock Adapter (with tab) (TZ-05000-002)

APPLICANT

Company	Teleflex Medical
Location	Fall River, MA, US
Contact	Kristen Bisanz
510(k) history	39 submissions · 39 cleared · 2003-2025

Teleflex Medical is an American medical device company headquartered in Wayne, Pennsylvania, with operations in Fall River, US. The company is a major provider of specialty medical devices for critical care and surgical procedures. Teleflex Medical has received FDA 510(k) clearances from total submissions since 2003. The company maintains active regulatory engagement, with the latest clearance in 2025. Its cleared devices span multiple specialties including anesthesiology, general and plastic surgery, cardiovascular, and vascular access systems. The company's product port...