FDA 510(k) Premarket Notification

K911202 ATRAUMATIC STANDARD SPROTTE NEEDLE

CLEARED

May 4, 1992
412 days to decision

K911202 · Product code: **BSP** · Anesthesiology
Source: https://www.510kdatabase.net/k911202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Conduction, Anesthetic (w/wo Introducer) (BSP)
Date received	Mar 19, 1991
Decision date	May 4, 1992
Days to decision	412 days
Third-party review	No

APPLICANT

Company	**Pajunk GmbH**
Location	Washington, DC, US
Contact	BRUCE MACKLER
510(k) history	19 submissions · 19 cleared · 1991-2012