FDA 510(k) Premarket Notification

K924348 AD-TECH'S BRAIN BIOPSY NEEDLE FOR STEREOTAXIC

CLEARED

Nov 23, 1992
88 days to decision

K924348 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k924348/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Aug 27, 1992
Decision date	Nov 23, 1992
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ad-Tech Medical Instrument Corp**
Location	Racine, WI, US
Contact	ANTHONY PUTZ
510(k) history	18 submissions · 17 cleared · 1985-2012