

K924348 AD-TECH'S BRAIN BIOPSY NEEDLE FOR STEREOTAXIC

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k924348/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026