

K791649 TETRAHYDROCANNALBINOL DIRECT BLOOD KITOct 11, 1979
48 days to decisionK791649 · Product code: **LAT** · Toxicology
Source: <https://www.510kdatabase.net/k791649/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radioimmunoassay, Cannabinoid(s) (LAT)
Date received	Aug 24, 1979
Decision date	Oct 11, 1979
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Immunalysis Corporation
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1978-2025

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k791649/> 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 June 28, 2026