

K011393 RANDOX PHENYTOINDec 4, 2001
211 days to decisionK011393 · Product code: **DIP** · Toxicology
Source: <https://www.510kdatabase.net/k011393/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Diphenylhydantoin (DIP)
Date received	May 7, 2001
Decision date	Dec 4, 2001
Days to decision	211 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Radox Laboratories, Ltd.
Location	Antrim, N. Ireland, IR
Contact	P. ARMSTRONG
Website	http://www.radox.com/
510(k) history	116 submissions · 115 cleared · 1992-2025

Radox Laboratories, Ltd. is a global diagnostic company specializing in chemistry devices and laboratory solutions. The company operates with a manufacturing facility in Antrim, Northern Ireland, and has over 40 years of expertise in diagnostic innovation. Radox has received FDA 510(k) clearances from total submissions since its first clearance in 1992. The company's regulatory portfolio is dominated by chemistry devices, including clinical chemistry analyzers, reagents, controls, and electrodes. The latest clearance on record dates to 2023. Recent FDA 510(k) cleared de...
