FDA 510(k) Premarket Notification

K022774 IQ 200 SYSTEM

CLEARED

Oct 21, 2002
61 days to decision

K022774 · Product code: **LKM** · Pathology
Source: https://www.510kdatabase.net/k022774/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter, Urine Particle (LKM)
Date received	Aug 21, 2002
Decision date	Oct 21, 2002
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Intl. Remote Imaging Systems**
Location	Mchenry, IL, US
Contact	HARVEY L KASDAN
510(k) history	14 submissions · 14 cleared · 1982-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k022774/ 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 14, 2026