FDA 510(k) Premarket Notification

K022681 AZOG, INC. HCG ONE-STEP URINE PREGNANCY TEST (DEVICE OR CASSETTE)

CLEARED

Nov 1, 2002
81 days to decision

K022681 · Product code: **DHA** · Chemistry
Source: https://www.510kdatabase.net/k022681/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Human Chorionic Gonadotropin (DHA)
Date received	Aug 12, 2002
Decision date	Nov 1, 2002
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Azog, Inc.**
Location	Phillipsburg, NJ, US
Contact	AZUBUIKE OGALA
510(k) history	3 submissions · 3 cleared · 2002-2004