

**K022681 AZOG, INC. HCG ONE-STEP URINE PREGNANCY TEST (DEVICE OR CASSETTE)**Nov 1, 2002  
81 days to decisionK022681 · Product code: **DHA** · Chemistry  
Source: <https://www.510kdatabase.net/k022681/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022681/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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