

**K070265 DIAGNOSTIC HYBRIDS' D3 DFA HERPES
SIMPLEX VIRUS IDENTIFICATION AND TYPING KIT, MODEL
01-090000**

Oct 24, 2007
268 days to decision

K070265 · Product code: **GQN** · Microbiology
Source: <https://www.510kdatabase.net/k070265/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antigen, Cf (including Cf Control), Herpesvirus Hominis 1,2 (GQN)
Date received	Jan 29, 2007
Decision date	Oct 24, 2007
Days to decision	268 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Diagnostic Hybrids, Inc.
Location	Athens, OH, US
Contact	GAIL GOODRUM
510(k) history	37 submissions · 36 cleared · 1988-2014

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k070265/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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