

**K792681 BLASTOMYCES DERMATITIDIS POS/CONTROL**Jan 17, 1980  
21 days to decisionK792681 · Product code: **KFH** · Microbiology  
Source: <https://www.510kdatabase.net/k792681/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antiserum, Positive Control, Blastomyces Dermatitidis (KFH)
Date received	Dec 27, 1979
Decision date	Jan 17, 1980
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>Meridian Diagnostics, Inc.</b>
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999

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Device record: <https://www.510kdatabase.net/k792681/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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