

**K812817 NEUTRAL DUFFERED FORMA-SCENT**Nov 10, 1981  
34 days to decisionK812817 · Product code: **IFP** · Pathology  
Source: <https://www.510kdatabase.net/k812817/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Formalin, Neutral Buffered (IFP)
Date received	Oct 7, 1981
Decision date	Nov 10, 1981
Days to decision	34 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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k812817/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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