

**K802423 PHADEBACT HAEMOPHILUS TEST**Nov 12, 1980  
40 days to decisionK802423 · Product code: **GRO** · Microbiology  
Source: <https://www.510kdatabase.net/k802423/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                      |
| Submission type       | Traditional   |
| Device classification | Antisera, Fluorescent, All Types, Hemophilus Spp. (GRO) |
| Date received         | Oct 3, 1980   |
| Decision date         | Nov 12, 1980  |
| Days to decision      | 40 days   |
| Third-party review    | No  |

**APPLICANT**

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| Company        | <b>Pharmacia, Inc.</b>                    |
| Location       | Mchenry, IL, US                           |
| 510(k) history | 129 submissions · 126 cleared · 1976-1998 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802423/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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