

**K790122 RUBANON**Feb 15, 1979  
23 days to decisionK790122 · Product code: **GOK** · Microbiology  
Source: <https://www.510kdatabase.net/k790122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Hai (including Hai Control), Rubella (GOK)
Date received	Jan 23, 1979
Decision date	Feb 15, 1979
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Organon, Inc.</b>
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1977-1995

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

Device record: <https://www.510kdatabase.net/k790122/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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