

**K812882 RHEUMATEX**Oct 26, 1981  
12 days to decisionK812882 · Product code: **DHR** · Microbiology  
Source: <https://www.510kdatabase.net/k812882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Rheumatoid Factor (DHR)
Date received	Oct 14, 1981
Decision date	Oct 26, 1981
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Armkel, LLC</b>
Location	Mchenry, IL, US
510(k) history	68 submissions · 68 cleared · 1979-2004

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Device record: <https://www.510kdatabase.net/k812882/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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