

**K812604 BIOVATION URINALYSIS REPORTER**Oct 2, 1981  
17 days to decisionK812604 · Product code: **JQP** · Chemistry  
Source: <https://www.510kdatabase.net/k812604/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calculator/data Processing Module, For Clinical Use (JQP)
Date received	Sep 15, 1981
Decision date	Oct 2, 1981
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Biovation, Inc.</b>
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1978-1993

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

Device record: <https://www.510kdatabase.net/k812604/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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