

**K061979 MODIFICATION TO MYALLERGY TEST**Sep 22, 2006  
71 days to decisionK061979 · Product code: **JKA** · Immunology  
Source: <https://www.510kdatabase.net/k061979/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Jul 13, 2006
Decision date	Sep 22, 2006
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Immunetech Corporation</b>
Location	Rancho Santa Fe, CA, US
Contact	NANCY BENSON
510(k) history	4 submissions · 4 cleared · 2002-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061979/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026