

**K791354 CODE QAS 103 ANTI-DNA CONTROLS**Sep 17, 1979  
61 days to decisionK791354 · Product code: **KTL** · Immunology  
Source: <https://www.510kdatabase.net/k791354/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anti-dna Indirect Immunofluorescent Solid Phase (KTL)
Date received	Jul 18, 1979
Decision date	Sep 17, 1979
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>Amersham Corp.</b>
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1976-1992

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Device record: <https://www.510kdatabase.net/k791354/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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