

**K911724 ACA(R) URINE DRUGS OF ABUSE CONTROL, MODIFICATION**

May 22, 1991  
35 days to decision

K911724 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k911724/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Apr 17, 1991
Decision date	May 22, 1991
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>E.I. Dupont DE Nemours &amp; Co., Inc.</b>
Location	Mchenry, IL, US
Contact	CHRISTOPHER BENTSEN
510(k) history	253 submissions · 252 cleared · 1976-1996

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

Device record: <https://www.510kdatabase.net/k911724/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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