

**K912007 BAXTER SINGLE PATIENT USE BLOOD PRESSURE CUFF**Jul 22, 1991  
77 days to decisionK912007 · Product code: **DXQ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k912007/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	May 6, 1991
Decision date	Jul 22, 1991
Days to decision	77 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Tecnadyne Scientific, Inc.</b>
Location	Vero Beach, FL, US
Contact	JOHN GARRETT
510(k) history	5 submissions · 5 cleared · 1990-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912007/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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