

TERCICA INC  
Form 8-K  
December 14, 2005

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**UNITED STATES**  
**SECURITIES AND EXCHANGE COMMISSION**  
**WASHINGTON, D.C. 20549**

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**FORM 8-K**

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**CURRENT REPORT**

**Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934**

**Date of Report (Date of earliest event reported): December 12, 2005**

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**TERCICA, INC.**

(Exact name of registrant as specified in its charter)

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**Delaware**

(State or other jurisdiction of incorporation)

**000-50461**  
(Commission File Number)

**26-0042539**  
(IRS Employer Identification No.)

**2000 Sierra Point Parkway, Suite 400**

**Brisbane, CA 94005**

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(Address of principal executive offices, including Zip Code)

**Registrant's telephone number, including area code: (650) 624-4900**

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Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
  - Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
  - Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
  - Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
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**Item 8.01. Other Events.**

On December 12, 2005, Inmed Incorporated announced that the U.S. Food and Drug Administration (FDA) approved Inmed's product, IPLEX(TM) (mecasermin rinfabate (rDNA origin) injection), for the treatment of growth failure in children with severe primary IGF-1 deficiency (Primary IGFD) or with growth hormone (GH) gene deletion who have developed neutralizing antibodies to GH. Inmed also announced that the FDA designated Inmed's product as an orphan drug for the treatment of Primary IGFD.

