



# DURECT Corporation Presenting at the Adams Harkness & Hill's 23rd Summer Seminar

CUPERTINO, Calif., Aug. 5 /PRNewswire-FirstCall/ — DURECT Corporation (Nasdaq: DRRX) announced today that it will present at the Adams, Harkness & Hill — 23rd Annual Summer Seminar. The conference is being held at the Boston Marriott Long Wharf Hotel. James E. Brown, DVM, Chief Executive Officer, will be presenting at the conference on Wednesday, August 6, 2003 at 2:30 p.m. EDT.

(Logo: <http://www.newscom.com/cgi-bin/prnh/20020717/DRRXLOGO> )

A live audio webcast of Dr. Brown's presentation will be available by accessing DURECT's homepage at <http://www.durect.com> and clicking "Investor Relations."

DURECT Corporation ([www.durect.com](http://www.durect.com)) is pioneering the development and commercialization of pharmaceutical systems for the treatment of chronic debilitating diseases and enabling biotechnology-based pharmaceutical products. DURECT's goal is to deliver the right drug to the right site in the right amount at the right time. In addition to its rights to the CHRONOGESIC(R) product, DURECT owns three proprietary drug delivery platform technologies, including the SABER(TM) Delivery System (a patented and versatile depot injectable useful for protein delivery), the MICRODUR(TM) Biodegradable Microparticulates (microspheres injectable system) and the DURIN(TM) Biodegradable Implant (drug-loaded implant system).

NOTE: CHRONOGESIC(R), SABER(TM), MICRODUR(TM) and DURIN(TM) are trademarks of DURECT Corporation.

The statements in this press release regarding DURECT's products in development and product development plans, are forward-looking statements involving risks and uncertainties that can cause actual results to differ materially from those in such forward-looking statements. Potential risks and uncertainties include, but are not limited to, DURECT's ability to complete the design, development, and manufacturing process development of its products, manufacture and commercialize its products, obtain product and manufacturing approvals from regulatory agencies, manage its growth and expenses, manage relationships with third parties, finance its activities and operations, as well as marketplace acceptance of DURECT's products. Further information regarding these and other risks is included in DURECT's Annual Report on Form 10-K for the fiscal year ended December 31, 2002 filed with the SEC on March 14, 2003, DURECT's Quarterly Report on Form 10Q and other periodic reports filed with the SEC under the heading "Factors that may affect future results."

CHRONOGESIC is under development by DURECT and has not been submitted or approved for commercialization by the US Food and Drug Administration or other health authorities.

SOURCE DURECT Corporation

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