

AnorMED Inc.
Form SC 14D9/A
October 23, 2006

**SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549**

**SCHEDULE 14D-9
SOLICITATION/RECOMMENDATION STATEMENT UNDER
SECTION 14(d)(4) OF THE SECURITIES EXCHANGE ACT OF 1934
(Amendment No. 18)**

AnorMED Inc.

(Name of Subject Company)

AnorMED Inc.

(Name of Persons Filing Statement)

Common Shares

(Title of Class of Securities)

035910108

(CUSIP Number of Class of Securities)

William J. Adams

Vice President, Finance, Chief Financial Officer,

Secretary and Treasurer

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(Name, address and telephone number of person authorized to receive notices
and communications on behalf of filing persons)

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☐ Check the box if the filing relates solely to preliminary communications made before the commencement of a tender offer.

This Amendment No. 18 amends and supplements the Solicitation/Recommendation Statement on Schedule 14D-9 (as amended, the Statement) initially filed by AnorMED Inc. with the U.S. Securities and Exchange Commission on September 5, 2006, relating to the tender offer (the Genzyme Offer) made by Dematal Corporation (Dematal), a Nova Scotia unlimited liability company and a wholly-owned subsidiary of Genzyme Corporation (Genzyme and together with Dematal, the Offerors), a Massachusetts corporation, for all of the common shares of AnorMED. The terms and conditions of the Genzyme Offer are set forth in the Offer to Purchase and Circular of the Offerors, dated September 1, 2006 (the Tender Offer Circular). The Tender Offer Circular has been filed by the Offerors with the U.S. Securities and Exchange Commission as part of a Tender Offer Statement on Schedule TO (as it may be amended or supplemented from time to time, the Schedule TO), which includes information required to be reported under Rule 14d-3 of the Securities Exchange Act of 1934, as amended. The Schedule TO was initially filed by the Offerors on September 1, 2006.

In connection with the Genzyme Offer, the Company's board of directors has prepared a directors' circular (the Directors' Circular), dated September 5, 2006, pursuant to applicable securities laws in Canada and the United States. The Company's board of directors has also prepared a Notice of Change to Directors' Circular (the Notice of Change), dated October 17, 2006, that relates to the Directors' Circular, pursuant to applicable securities laws in Canada and the United States. The Directors' Circular has been mailed to AnorMED shareholders and was filed as exhibit (a)(2)(A) to the initial filing of this Statement. The Notice of Change has been mailed to AnorMED shareholders and was filed as Exhibit (a)(2)(W) to Amendment No. 16 to this Statement. The Directors' Circular, as amended and supplemented by the Notice of Change, is incorporated by reference into this Statement in its entirety.

Capitalized terms used herein and not defined herein have the respective meanings assigned to such terms in the Directors' Circular and the Notice of Change.

Item 8. ADDITIONAL INFORMATION

Item 8 is hereby amended and supplemented as follows:

On October 23, 2006, AnorMED Inc. announced that it has completed enrollment in the second pivotal Phase III clinical trial evaluating its proprietary product MOZOBIL in cancer patients, and that AnorMED is on track to meet its schedule of releasing top-line data from both Phase III trials in the first half of 2007. If the Phase III trials are successful, it would be possible to file for marketing approval with the United States Food and Drug Administration (FDA) in the second half of 2007 and with regulators in Canada and Europe in 2008.

The Phase III trials are evaluating MOZOBIL's capacity to improve stem cell transplantation treatment options for multiple myeloma (MM) and non-Hodgkin's lymphoma (NHL) patients whose immune systems have a reduced capacity to generate stem cells as a result of extensive chemotherapy treatments.

The second pivotal Phase III trial has enrolled 300 NHL patients and the first pivotal Phase III trial met its enrollment target of 300 MM patients on July 10, 2006.

The most recently enrolled patients will undergo their transplants over the next four to six weeks and each patient will be followed over a period of 100 days. The results of the study will be unblinded for analysis after all MM and NHL patients have completed their 100-day follow-up. The trials design is in accordance with a Special Protocol Assessment from the FDA.

The two Phase III trials are being conducted at up to 45 major centres in the U.S., Canada and Europe. Both trials are randomized, double-blind, placebo-controlled, comparative trials of MOZOBIL plus G-CSF versus placebo plus G-CSF, the current standard drug used to stimulate additional stem cells within bone marrow.

A copy of AnorMED's press release, dated October 23, 2006, with respect to the matters described above is attached hereto as Exhibit (a)(2)(Y). The section of the press release entitled FORWARD LOOKING STATEMENTS is incorporated herein by reference.

Item 9. EXHIBITS

Item 9 is hereby amended and supplemented to include the following exhibits:

Exhibit	Description
(a)(2)(Y)	News release, dated October 23, 2006

SIGNATURES

After due inquiry and to the best of my knowledge and belief, I certify that the information set forth in this Statement is true, complete and correct.

Dated: October 23, 2006

ANORMED INC.

By: /s/ William J. Adams

Name: William J. Adams

Title: Vice President, Finance, Chief

Financial

Officer, Secretary and Treasurer

EXHIBIT INDEX

Exhibit	Description
(a)(2)(A)*	Directors Circular, dated September 5, 2006
(a)(2)(B)*	Press release of AnorMED Inc., dated September 5, 2006
(a)(2)(C)*	News release, dated September 7, 2006
(a)(2)(D)*	Material Change Report, dated September 11, 2006
(a)(2)(E)*	Material Change Report, dated September 11, 2006
(a)(2)(F)*	Limited Duration Shareholder Rights Plan Agreement
(a)(2)(G)*	News release, dated September 15, 2006
(a)(2)(H)*	News release, dated September 18, 2006
(a)(2)(I)*	News release, dated September 19, 2006
(a)(2)(J)*	PowerPoint slides from Annual General Meeting presentation
(a)(2)(K)*	Transcript of portions of Annual General Meeting
(a)(2)(L)*	News release, dated September 20, 2006
(a)(2)(M)*	News release, dated September 25, 2006
(a)(2)(N)*	News release, dated September 26, 2006
(a)(2)(O)*	Material Change Report, dated September 28, 2006, in connection with an amended license agreement between AnorMED Inc. and Poniard Pharmaceuticals, Inc.
(a)(2)(P)*	Amendment No. 1 to License Agreement, effective as of September 18, 2006, between AnorMED Inc. and Poniard Pharmaceuticals, Inc.
(a)(2)(Q)*	Material Change Report, dated September 28, 2006, in connection with the Support Agreement, dated as of September 26, 2006, between AnorMED Inc. and Millennium Pharmaceuticals, Inc.
(a)(2)(R)*	News release, dated October 4, 2006
(a)(2)(S)*	Directors Circular, dated October 5, 2006
(a)(2)(T)*	News release, dated October 5, 2006
(a)(2)(U)*	News release, dated October 10, 2006

- (a)(2)(V)* News release, dated October 11, 2006
 - (a)(2)(W)* Notice of Change, dated October 17, 2006
 - (a)(2)(X)* News release, dated October 17, 2006
 - (a)(2)(Y) News release, dated October 23, 2006
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Exhibit	Description
(e)(1)*	Confidentiality Agreement, dated October 4, 2006, among Genzyme Corporation, Dematal Corp. and AnorMED Inc.
(e)(2)*	Form of Support Agreement, dated as of October 17, 2006, among AnorMED Inc., Genzyme Corporation and Dematal Corp.
(e)(3)*	Form of Shareholder Support Agreement, dated as of October 17, 2006, among specified shareholders, Genzyme Corporation and Dematal Corp.
(e)(4)*	Form of Shareholder Support Agreement, dated as of October 17, 2006, among Kenneth Galbraith, Genzyme Corporation and Dematal Corp.
(g)(1)*	Information Agent Script for Incoming Calls
(g)(2)*	Information Agent Script for Outgoing Calls
(g)(3)*	Information Agent Script for Outgoing Calls
(g)(4)*	Information Agent Script for Incoming Calls
(g)(5)*	Information Agent Script for Incoming Calls
(g)(6)*	Information Agent Script for Outgoing Calls

* Previously filed.