

U.S. ENVIRONMENTAL PROTECTION AGENCY  
OFFICE OF PESTICIDES PROGRAMS  
REGISTRATION DIVISION (WH-567)  
WASHINGTON, D.C. 20460

EPA REGISTRATION NO.

43351-8

DATE OF ISSUANCE

February 21, 1985

TERM OF ISSUANCE

NOTICE OF PESTICIDE:  REGISTRATION  
 REREGISTRATION

(Under the Federal Insecticide, Fungicide,  
and Rodenticide Act, as amended)

NAME OF PESTICIDE PRODUCT

Magna CP-1

NAME AND ADDRESS OF REGISTRANT (Include ZIP code)

Abram Pharmaceutical, Inc.  
P. O. Box 13588  
Research Triangle Park  
N. C. 27709

BEST AVAILABLE COPY

NOTE: Changes in labeling formula differing in substance from that accepted in connection with this registration must be submitted to and accepted by the Registration Division prior to use of the label in commerce. In any correspondence on this product always refer to the above U.S. EPA registration number.

On the basis of information furnished by the registrant, the above named pesticide is hereby Registered/Reregistered under the Federal Insecticide, Fungicide, and Rodenticide Act.

A copy of the labeling accepted in connection with this Registration/Reregistration is returned herewith.

Registration is in no way to be construed as an indorsement or approval of this product by this Agency. In order to protect health and the environment, the Administrator, on his motion, may at any time suspend or cancel the registration of a pesticide in accordance with the Act. The acceptance of any name in connection with the registration of a product under this Act is not to be construed as giving the registrant a right to exclusive use of the name or to its use if it has been covered by other pesticide registration notices. This registration notice is being issued in accordance with Section 3(c)(7)(A) of the FIFRA subject to the following conditions:

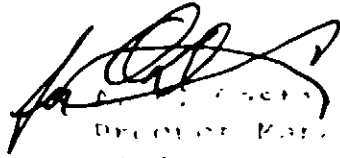
(1) It is understood that you will submit and/or cite all data required for registration or re-registration of your product under Section 3(c)(5) of the FIFRA when the Agency requires all registrants of similar products to submit such data.

(2) You will submit five copies of your finished printed labels complying with the requirements or revisions specified in the attachment to this registration notice. These labels must be submitted to the Agency before the product is released for shipment as required by PE Notice 82-2 of June 19, 1982.

A stamped copy of the draft label is enclosed for your records. Your release of the product for shipment constitutes acceptance of these conditions. If the conditions are not complied with, the registration will be subject to cancellation in accordance with Section 6(c) of the FIFRA.

ATTACHMENT IS APPLICABLE

SIGNATURE OF APPROVING OFFICIAL

  
Director, Registration Division  
U.S. Environmental Protection Agency  
Washington, D.C. 20460

DATE  
FEB 21 1985

ATTACHMENT TO EPA REGISTRATION NOTICE

EPA Registration No. 43351-8

Date Issued February 21, 1985

Company Name Abram Pharmaceutical, Inc.

BEST AVAILABLE COPY

Product Name Magna CP-1

The following requirements or revisions apply to the label which was accepted with this registration notice:

1. An edited copy of the proposed label is enclosed indicating the revisions which must be made to your label to satisfy our requirements.
2. The registration number which has been assigned to this product must appear on the label as EPA Reg. No. 43351-8.
3. Delete the words "General Classification" which appear under the use directions. Classification of this product has been deferred indefinitely.
4. The labels which will be used for this product in channels of trade must reflect these revisions.
5. Five copies of the finished printed labels, as defined in the A-79 enclosure, must be submitted to the Agency before the product is released for shipment to comply with PR Notice 82-2 of June 18, 1982.
6. The finished printed labels must comply with the general labeling requirements of Section 162.10 of Title 40 of the Code of Federal Regulations. If the label does not satisfy these requirements it will be rejected.
7. Any collateral literature proposed for this product must be submitted for approval before it is circulated in channels of trade.

