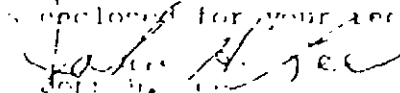


**BEST AVAILABLE COPY**

|  |   |   |
|--|---|---|
| <b>U.S. ENVIRONMENTAL PROTECTION AGENCY</b><br><b>OFFICE OF PESTICIDES PROGRAMS</b><br><b>REGISTRATION DIVISION (WH-567)</b><br><b>WASHINGTON, D.C. 20460</b><br><br><b>NOTICE OF PESTICIDE:</b> <input checked="" type="checkbox"/> REGISTRATION<br><input type="checkbox"/> REREGISTRATION<br><i>(Under the Federal Insecticide, Fungicide, and Rodenticide Act, as amended)</i>   | <b>EPA REGISTRATION NO.</b><br>5185-366               | <b>DATE OF ISSUANCE</b><br><b>19 APR 1985</b> |
|  | <b>TERM OF ISSUANCE</b>                               |   |
|  | <b>NAME OF PESTICIDE PRODUCT</b><br>Bio Guard HTD-256 |   |
| <b>NAME AND ADDRESS OF REGISTRANT (Include ZIP code)</b><br><br><div style="border: 1px solid black; padding: 10px; margin: 10px;">         Bio-Lab Inc.<br/>         P.O. Box 1489<br/>         Decatur, GA 30031       </div>  |   |   |
| <b>NOTE:</b> 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.  |   |   |
| <p>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.</p> <p>A copy of the labeling accepted in connection with this Registration/Reregistration is returned herewith.</p> <p>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 others.</p> <p>This product is conditionally registered in accordance with FIFR sec. 3(c)(7)(A) provided that you:</p> <ol style="list-style-type: none"> <li>1. Submit and/or cite all data required for registration/reregistration of your product under FIFPA sec. 3(c)(5) when the Agency requires all registrants of similar products to submit such data.</li> <li>2. Make the labeling changes listed below before you release the product for shipment:         <ol style="list-style-type: none"> <li>a. Add the phrase "EPA Registration No. 5185-366."</li> <li>b. Revise the statement:<br/>                 Perpes Simplex Type II<br/>                 to read:<br/>                 Perpes Simplex Type 2</li> <li>c. The referral statement should be placed directly below the Note to Physician section.</li> </ol> </li> <li>3. Submit five (5) copies of your final printed labeling before you release the product for shipment. Refer to the A-79 Enclosure for a further description of final printed labeling.</li> </ol> <p>If these conditions are not complied with, the registration will be subject to cancellation in accordance with FIFPA sec. 6(e). Your release for shipment of the product constitutes acceptance of these conditions.</p> <p>A carbon copy of the label is enclosed for your records.</p> <div style="text-align: right; margin-top: 20px;"> <br/>       John H. Lee<br/>       Product Manager (31)<br/>       Manufacturers Branch<br/>       Registration Division (TS-767C)     </div> |   |   |
| <div style="display: flex; justify-content: space-between;"> <div> <input type="checkbox"/> ATTACHMENT IS APPLICABLE<br/> <b>SIGNATURE OF APPROVING OFFICIAL</b> </div> <div> <b>DATE</b> </div> </div>  |   |   |