



## **RECALL STOCK RESPONSE FORM**

**Product RECALL February 3<sup>rd</sup>, 2017  
(Travoprost Ophthalmic Solution USP 0.004%)**

### **VOLUNTARY RECALL TO THE RETAIL LEVEL**

**Please fill out this form completely.** By doing so, this will acknowledge that you have read and understand the recall instructions and have taken the appropriate action.

Returned by \_\_\_\_\_ DEA # \_\_\_\_\_  
*\*DEA Registration # is required, if not provided the processing of your form will be delayed.*

Address \_\_\_\_\_

City \_\_\_\_\_ State \_\_\_\_\_ Zip \_\_\_\_\_

Contact Name (please print) \_\_\_\_\_ Telephone # \_\_\_\_\_

Contact Signature \_\_\_\_\_ Date \_\_\_\_\_

**I have checked my stock and:**

\_\_\_\_\_ Do not have any stock of the recalled **items**.

**OR**

Have quarantined and listed in the box below the qty of recalled units I will be returning to Inmar, as soon as possible. Upon receipt of this Response Form, Inmar, will issue return authorization label(s) Please indicate the # of needed box labels \_\_\_\_\_.

| Item Description  | NDC          | Lot     | Exp.    | Qty returning |
|---|--------------|---------|---------|---------------|
| Travoprost Ophthalmic Solution USP 0.004%<br>2.5 mL Bottles | 49884-044-48 | G40814  | 09/2016 |               |
|   |              | GA50089 | 12/2016 |               |
|   |              | GA50259 | 01/2017 |               |
|   |              | GA50657 | 03/2017 |               |
|   |              | GA51073 | 06/2017 |               |
|   |              | GA51651 | 09/2017 |               |
|   |              | GA51652 | 09/2017 |               |
|   |              | GA51722 | 09/2017 |               |
|   |              | GA51723 | 09/2017 |               |
|   |              | GA51760 | 10/2017 |               |
|   |              | GA51761 | 10/2017 |               |

| Item Description  | NDC          | Lot     | Exp.    | Qty returning |
|---|--------------|---------|---------|---------------|
| Travoprost Ophthalmic Solution USP 0.004%<br>5 mL Bottles | 49884-044-63 | GA45033 | 10/2016 |               |
|   |              | GA50258 | 01/2017 |               |
|   |              | GA50944 | 05/2017 |               |
|   |              | GA50174 | 06/2017 |               |
|   |              | GA51340 | 07/2017 |               |
|   |              | GA51479 | 08/2017 |               |
|   |              | GA51762 | 10/2017 |               |

**In addition, please check the appropriate response below:**

\_\_\_\_\_ We **HAVE** received complaints of adverse events associated with use of the product.

\_\_\_\_\_ We **HAVE NOT** received complaints of adverse events associated with use of the product.

**If you did not purchase the product directly from the Manufacturer please complete the below section.**

Purchased From: Wholesaler Name \_\_\_\_\_ DEA # \_\_\_\_\_

City \_\_\_\_\_ State \_\_\_\_\_

If you have any questions regarding this form or product return please contact Inmar at 1-800-967-5952, prompt # 1 for recall. Office hours: 9am to 5pm EST Mon thru Fri.

**Please fax this form to: 1-817-868-5362 or E-mail: [rxrecalls@inmar.com](mailto:rxrecalls@inmar.com)**