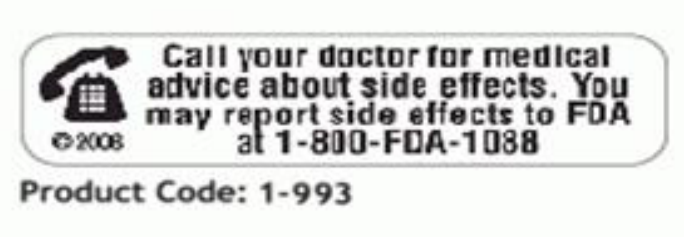
Effective July 1, 2009, as mandated by the FDA Amendments Act of 2007, pharmacies must provide the patient with the 1-800 phone number for reporting adverse events each time in a new prescription and REFILL is dispensed.

The statement should include the following: **"Call your doctor for medical advice about side effects. You may report side effects to the FDA at 1-800-FDA-1088."**

The statement can be on the label, the vial, a separate piece of paper, or the monograph.

PHARMEX HAS THE LABEL TO HELP YOU MEET THE REQUIREMENT.



H D Smith 224-5256
Kinray 325-332
McKesson 321-3048