

Prior Authorization Form

GEHA FEDERAL - STANDARD OPTION

Xenazine (FA-PA)

This fax machine is located in a secure location as required by HIPAA regulations.  
Complete/review information, sign and date. Fax signed forms to CVS/Caremark at **1-888-836-0730**.  
Please contact CVS/Caremark at **1-855-240-0536** with questions regarding the prior authorization process.  
When conditions are met, we will authorize the coverage of Xenazine (FA-PA).

Drug Name (select from list of drugs shown)

Xenazine (tetrabenazine)

Quantity

Frequency

Strength

Route of Administration

Expected Length of Therapy

Patient Information

Patient Name: \_\_\_\_\_

Patient ID: \_\_\_\_\_

Patient Group No.: \_\_\_\_\_

Patient DOB: \_\_\_\_\_

Patient Phone: \_\_\_\_\_

Prescribing Physician

Physician Name: \_\_\_\_\_

Physician Phone: \_\_\_\_\_

Physician Fax: \_\_\_\_\_

Physician Address: \_\_\_\_\_

City, State, Zip: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code: \_\_\_\_\_

Comments: \_\_\_\_\_

Please circle the appropriate answer for each question.

1. Has the patient failed treatment with the generic medication due to an intolerable adverse event (e.g., rash, nausea, vomiting)?

Y N

2. Was the intolerable adverse event an expected adverse event attributed to the ACTIVE ingredient as described in the prescribing information (i.e., known adverse reaction for both the brand and generic medication)?

Y N

3. Was this documented in the patient's chart? Documentation is required for approval. Provide SPECIFIC AND DETAILED chart documentation including description, date/time, and severity of the adverse event,

Y N

dosage and duration of generic medication treatment,  
required intervention (if any), and relevant tests or  
laboratory data (if any) OR MedWatch form of this trial and  
failure including the adverse reaction.

I affirm that the information given on this form is true and accurate as of this date.

**Prescriber (Or Authorized) Signature and Date**