ATENOLOL SCOPOLAMINE- atenolol scopolamine tablet
TPS

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Take one tablet sublingually or orally as directed by your physician as needed for symptoms of panic or anxiety.

Do not exceed two tablets per day.

TPS LLC

3524 DECATUR HWY FULTONDALE, AL 35068

877-608-4995 1-877-608-4995 BT9752747

Caution: Federal law prohibits transfer of this drug to any other person than patient for whom prescribed

Rx 263095 Jack Doe/Dr. Jane Doe MD

JOHN DOE

123 MAIN ST AUBURN, AL 12345

ATENOLOL SCOPOLAMINE TABLET TRITURATE

25 MG/0.25 MG

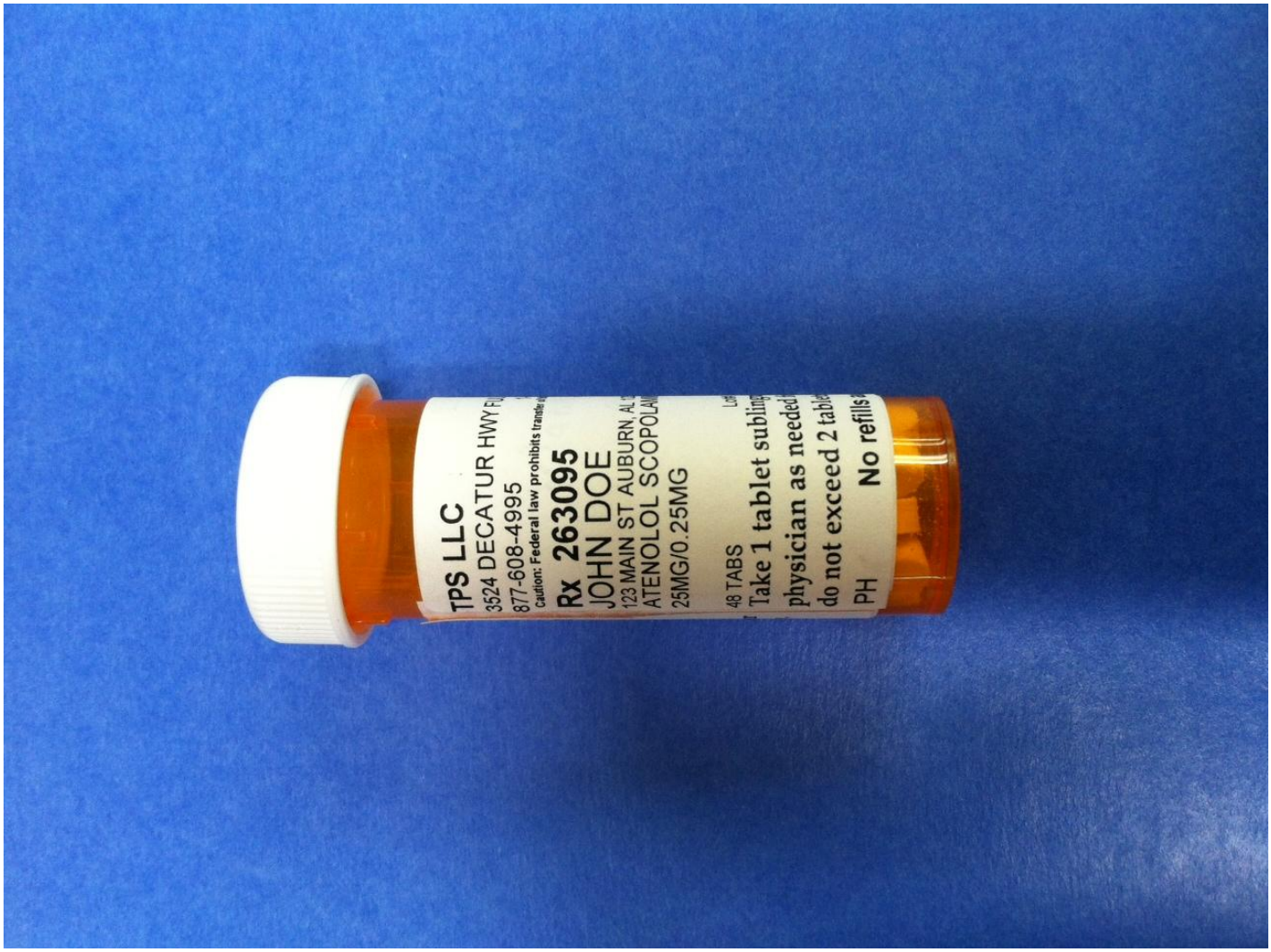
48 TABS Lot# Exp

Take 1 tablet sublingually or orally as directed by your physician as needed for symptoms of panic or anxiety.

Do not exceed 2 tablets per day

PH No refills authorized 10/8/2014

Pill bottle low res.jpg



ATENOLOL SCOPOLAMINE

atenolol scopolamine tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:69267-101
Route of Administration	SUBLINGUAL, BUCCAL, TRANSMUCOSAL, ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ATENOLOL (UNII: 50VV3VW0TI) (ATENOLOL - UNII:50VV3VW0TI)	ATENOLOL	25 mg in 25.25 mg
SCOPOLAMINE HYDROBROMIDE (UNII: 451IFR0GXB) (SCOPOLAMINE - UNII:DL48G20X8X)	SCOPOLAMINE HYDROBROMIDE	.25 mg in 25.25 mg

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND (The diameter depends on dies)	**Size**	5mm

Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69267-101-24	606 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
2	NDC:69267-101-06	151.5 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
3	NDC:69267-101-12	303 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
4	NDC:69267-101-48	1212 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2014	

Labeler - TPS (044805267)

Establishment

Name	Address	ID/FEI	Business Operations
TPS		044805267	manufacture(69267-101)

Revised: 10/2014 TPS