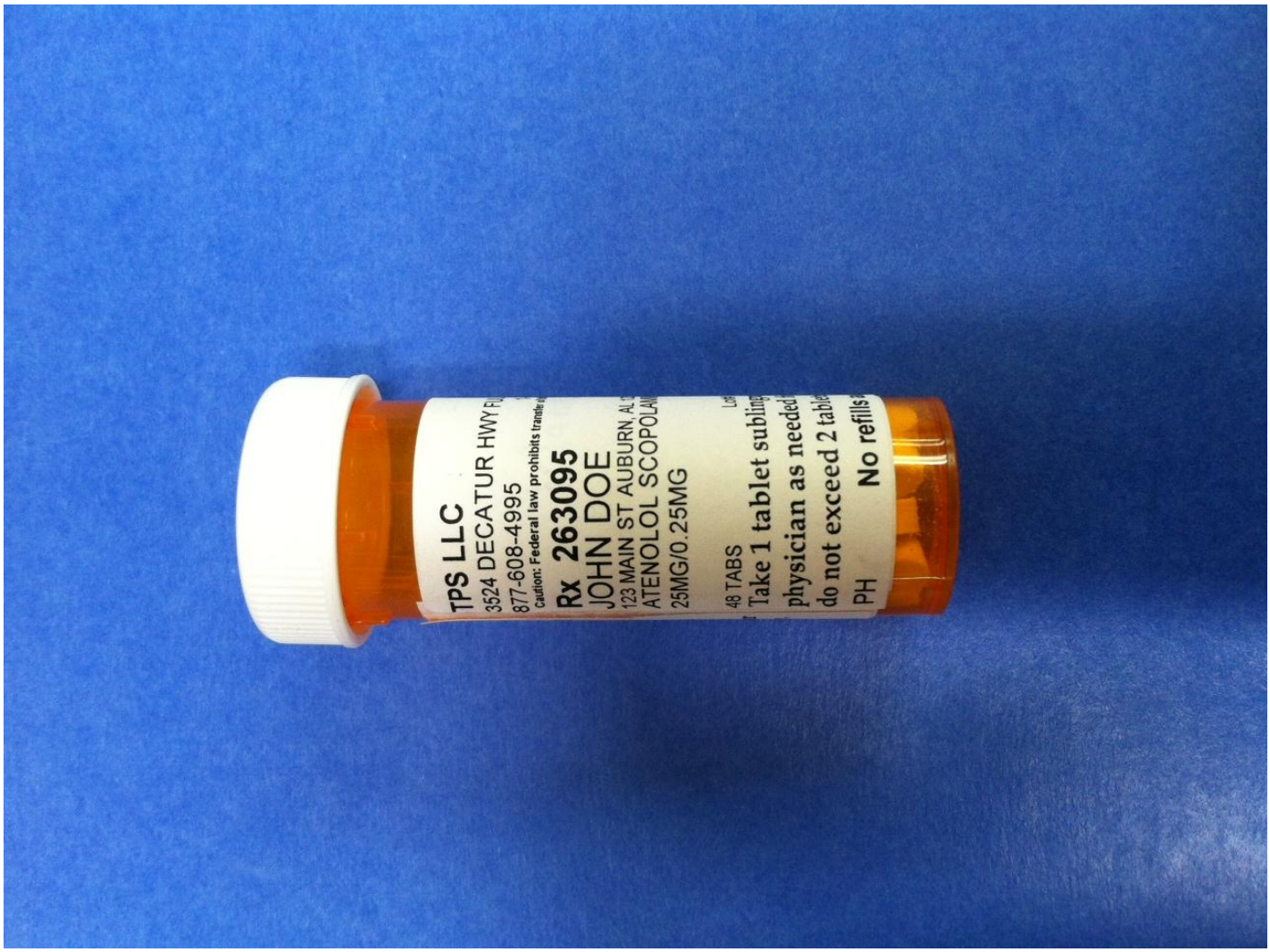
PROPRANOLOL SCOPOLAMINE- propranolol scopolamine tablet
TPS

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 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

PROPRANOLOL SCOPOLAMINE TABLET TRITURATE

20 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

PROPRANOLOL SCOPOLAMINE

propranolol scopolamine tablet

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:69267-201
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SCOPOLAMINE HYDROBROMIDE (UNII: 451IFR0GXB) (SCOPOLAMINE - UNII:DL48G20X8X)	SCOPOLAMINE HYDROBROMIDE	.25 mg in 20.25 mg
PROPRANOLOL HYDROCHLORIDE (UNII: F8A3652H1V) (PROPRANOLOL - UNII:9Y8NXQ24VQ)	PROPRANOLOL HYDROCHLORIDE	20 mg in 20.25 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:69267-201-06	121.5 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
2	NDC:69267-201-12	243 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
3	NDC:69267-201-24	486 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2014	
4	NDC:69267-201-48	972 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(69267-201)

Revised: 10/2014 TPS