

**OXAZEPAM- oxazepam capsule**  
**TruPharma, LLC**

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**Oxazepam Capsules, USP**

**Principal Display Panel**

**Oxazepam Capsules, USP**

NDC 10mg 52817-290-10

CIV Oxazepam Capsules, USP

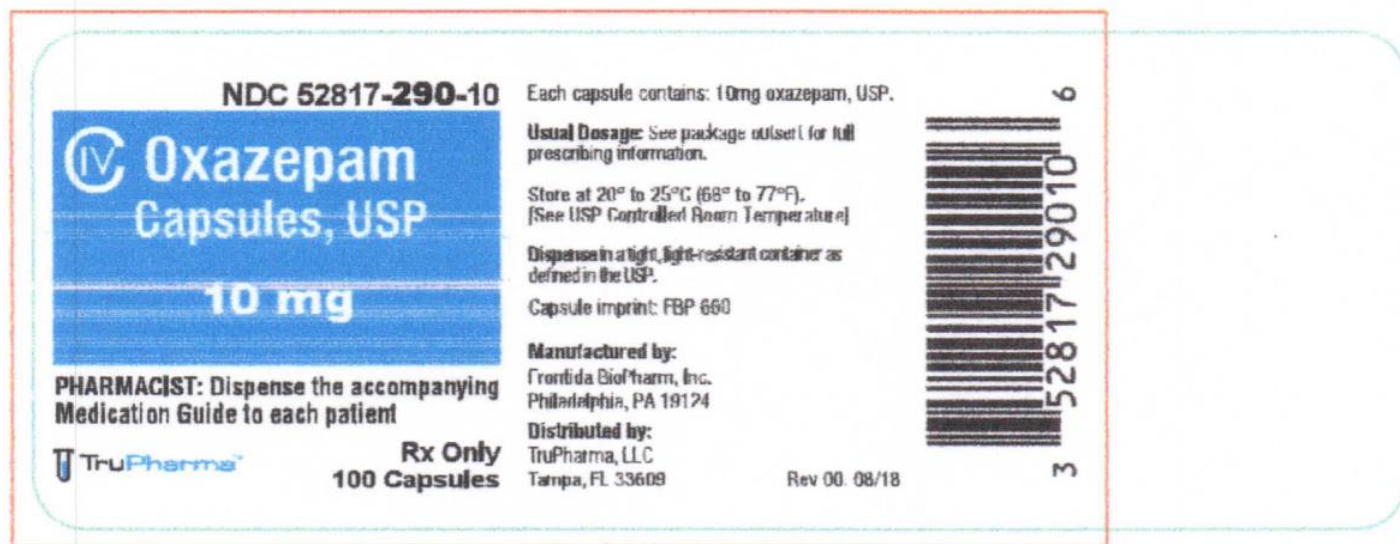
10mg

**PHARMACIST: Dispense the accompanying Medication Guide to each patient**

Rx only

100 Capsules

TruPharma



**Principal Display Panel**

NDC 15 mg 52817-291-10

CIV Oxazepam Capsules, USP

15mg

**PHARMACIST: Dispense the accompanying Medication Guide to each patient**

Rx only

100 Capsules

TruPharma

**NDC 52817-291-10** Each capsule contains: 15mg oxazepam, USP.

**Oxazepam Capsules, USP**  
**15 mg**

**PHARMACIST: Dispense the accompanying Medication Guide to each patient**

**Rx Only**  
**100 Capsules**

Usual Dosage: See package insert for full prescribing information.

Store at 20° to 25°C (68° to 77°F).  
[See USP Controlled Room Temperature]

Dispense in a tight, light-resistant container as defined in the USP.

Capsule imprint: FBP 661

Manufactured by:  
Frontida BioPharm, Inc.  
Philadelphia, PA 19124

Distributed by:  
TruPharma, LLC  
Tampa, FL 33609

Rev 00: 08/18

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### Principal Display Panel

NDC 30 mg 52817-292-10

CIV Oxazepam Capsules, USP

30mg

**PHARMACIST: Dispense the accompanying Medication Guide to each patient**

Rx only

100 Capsules

TruPharma

**NDC 52817-292-10** Each capsule contains: 30mg oxazepam, USP.

**Oxazepam Capsules, USP**  
**30 mg**

**PHARMACIST: Dispense the accompanying Medication Guide to each patient**

**Rx Only**  
**100 Capsules**

Usual Dosage: See package insert for full prescribing information.

Store at 20° to 25°C (68° to 77°F).  
[See USP Controlled Room Temperature]

Dispense in a tight, light-resistant container as defined in the USP.

Capsule imprint: FBP 662

Manufactured by:  
Frontida BioPharm, Inc.  
Philadelphia, PA 19124

Distributed by:  
TruPharma, LLC  
Tampa, FL 33609

Rev 00: 08/18

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## OXAZEPAM

oxazepam capsule

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:52817-290
<b>Route of Administration</b>	ORAL	<b>DEA Schedule</b>	CIV

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
OXAZEPAM (UNII: 6GOW6DWN2A) (OXAZEPAM - UNII:6GOW6DWN2A)	OXAZEPAM	10 mg

**Inactive Ingredients**

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
SHELLAC (UNII: 46N107B71O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

**Product Characteristics**

Color	WHITE	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	FBP660
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52817-290-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA071026	05/15/2020	

**OXAZEPAM**

oxazepam capsule

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:528 17-29 1
<b>Route of Administration</b>	ORAL	<b>DEA Schedule</b>	CIV

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
O XAZEPAM (UNII: 6GOW6DWN2A) (O XAZEPAM - UNII:6GOW6DWN2A)	O XAZEPAM	15 mg

### Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
SHELLAC (UNII: 46N107B71O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

### Product Characteristics

<b>Color</b>	RED	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	14mm
<b>Flavor</b>		<b>Imprint Code</b>	FBP661
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:528 17-29 1-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA071026	05/15/2020	

# OXAZEPAM

oxazepam capsule

## Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:528 17-292
Route of Administration	ORAL	DEA Schedule	CIV

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXAZEPAM (UNII: 6GOW6DWN2A) (OXAZEPAM - UNII:6GOW6DWN2A)	OXAZEPAM	30 mg

## Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
GELATIN (UNII: 2G86QN327L)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
SHELLAC (UNII: 46N107B71O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	

## Product Characteristics

Color	RED	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	FBP662
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52817-292-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/19/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA071026	03/19/2020	

**Labeler** - TruPharma, LLC (078533947)

Revised: 5/2020

TruPharma, LLC