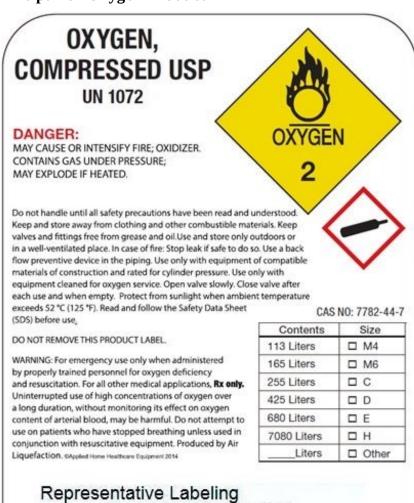
OXYGEN- oxygen gas Stone's Drugs Inc.

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.

Oxygen, USP

Principal for Oxygen Product



OXYGEN

COMPRESSED

USP

UN 1072

PRODUCED BY AIR LIQUIFICATION

are printed in this box

pursuant to 21 CFR 207.3 (10)

Company name and contact information

WARNING: HIGH PRESSURE OXIDIZING GAS, VIGOROUSLY ACCELERATES COMBUSTION. NO SMOKING IN THE PRESENCE OF OXYGEN OR A FIRE MAY RESULT.

WARNING: For emergency use only when administered by properly trained personnel for oxygen deficiency and resuscitation. For all other medical applications, Rx only.

WARNING: Federal law prohibits dispensing without prescription. Uninterrupted use of high concentrations of oxygen over a long duration, without monitoring its effect on oxygen content of arterial blood, may be harmful. Use only with pressure reducing equipment and apparatus designed for oxygen. Do not attempt to use on patients who have stopped breathing, unless used in conjuction with resuscitative equipment.

WARNING: Keep oil and grease away. Use only with equipment cleaned for oxygen service and rated for cylinder pressure. Open valve slowly. Close valve after each use and when empty.

USE IN ACCORDANCE WITH MATERIAL SAFETY DATA SHEET.

DO NOT REMOVE THIS PRODUCT LABEL.



Representative Labeling pursuant to 21 CFR 207.3 (10)

OXYGEN

oxygen gas

Product Information	
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Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:61752-001

Route of Administration RESPIRATORY (INHALATION)

Active Ingredient/Active Moiety

1	Ingredient Name	Basis of Strength	Strength
ı	OXYGEN (UNII: S88TT14065) (OXYGEN - UNII:S88TT14065)	OXYGEN	99 L in 100 L

Pac	kag	ing
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П	P	Packaging				
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
ı	1	NDC:61752-001-03	680 L in 1 CYLINDER; Type 0: Not a Combination Product	05/09/2017		
ı	2	NDC:61752-001-01	164 L in 1 CYLINDER; Type 0: Not a Combination Product	05/09/2017		

3 NDC:61752-001-02	255 L in 1 CYLINDER; Type 0: Not a Combination Produc	t 05/09/2017		
Marketing Information				
Marketing Init	ormation			
Marketing Categor		Marketing Start Date	Marketing End Date	
	y Application Number or Monograph Citation	Marketing Start Date 05/09/2017	Marketing End Date	

Labeler - Stone's Drugs Inc. (024105652)

Establishment			
Name	Address	ID/FEI	Business Operations
Stone's Drugs Inc.		024105652	manufacture (61752-001)

Revised: 10/2020 Stone's Drugs Inc.