

**COMPLIMENTS ADVANCED RELIEF EYE DROPS 15ML- dextran 70, povidone, polyethylene glycol 400, tetrahydrozoline hydrochloride solution/ drops
KC Pharmaceuticals, Inc.**

Compliments Advanced Relief Eye Drops 15mL (MFR)

Active Ingredients

Dextran 70 0.01%

Povidone 1%

Polyethylene glycol 400 1%

Tetrahydrozoline hydrochloride 0.05%

Purpose

Lubricant eye drop

Redness reliever

Warnings

For use in the eye only

Do not use

- if you use a monoamine oxidase inhibitor (MAOI)
- if solution changes colour or becomes cloudy.

Ask a doctor or pharmacist before use if you

- have narrow angle glaucoma
- are pregnant or breastfeeding.

When using this product

- pupils may become enlarged temporarily
- to avoid contamination, do not touch tip of the container to any surface
- replace cap after using
- overuse of this product may cause increased redness of the eye.

Stop use and ask a doctor if

- you have a headache, eye pain, changes in vision, redness or irritation of the eye
- symptoms worsen or last longer than 3 days.

Keep out of the reach of children.

If swallowed, call a poison control centre or get medical help right away.

Directions

Adults and children 6 years and over: Remove contact lenses before using. Squeeze 1 or 2 drops in the affected eye(s) up to 4 times a day or as directed by a doctor.

Store between 15°C and 30°C. Do not use if safety seal around cap is missing or broken.

Inactive ingredients

Benzalkonium Chloride, Boric acid, Disodium EDTA, Sodium Borate, Sodium Chloride, Water

Questions?

Call 1-866-672-0061

Compliments Advanced Relief Eye Drops 15mL



COMPLIMENTS ADVANCED RELIEF EYE DROPS 15ML

dextran 70, povidone, polyethylene glycol 400, tetrahydrozoline hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55651-989
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTRAN 70 (UNII: 7SA290YK68) (DEXTRAN 70 - UNII:7SA290YK68)	DEXTRAN 70	0.1 g in 100 mL

Drug Facts / Info-médicament	
Active ingredients (w/v)	Purposes
Dextran 70 0.1%, Povidone 1%	Lubricant eye drop
Polyethylene glycol 400 1%	Redness reliever
Tetrahydrozoline hydrochloride 0.05%	Utilités
Ingredients actifs (p/v)	Gouttes ophtalmiques lubrifiantes
Dextran 70 à 0.1 %, Povidone à 1 %	Gouttes ophtalmiques lubrifiantes
Polyéthylène glycol 400 à 1 %	Soulagement de la rougeur oculaire
Chlorhydrate de tétrahydrozoline à 0.05 %	

Warnings / Mises en garde
For use in the eye only / Pour usage ophtalmique seulement
Do not use • if you use a monoamine oxidase inhibitor (MAOI) • if solution changes colour or becomes cloudy.
Ne pas utiliser • si vous prenez un inhibiteur de la monoamine-oxydase (MAO) • si la solution change de couleur ou devient trouble.
Ask a doctor or pharmacist before use if you • have narrow angle glaucoma • are pregnant or breastfeeding.
Consultez un médecin ou un pharmacien avant l'utilisation si vous • avez un glaucome par fermeture de l'angle • êtes enceinte ou allaitez.
When using this product • pupils may become enlarged temporarily • to avoid contamination, do not touch tip of container to any surface • replace cap after using • overuse of this product may cause increased redness of the eye.
Lorsque vous utilisez ce produit • peut causer une dilatation temporaire des pupilles • pour éviter la contamination, il faut éviter que l'embout vienne en contact avec une surface • remettre le bouchon après l'usage • l'usage excessif de ce produit peut intensifier la rougeur oculaire.
Stop use and ask a doctor if • you have headache, eye pain, changes in vision, redness or irritation of the eye • symptoms worsen or last longer than 3 days.
Cessez d'utiliser et consultez un médecin si • vous avez des maux de tête ou une douleur aux yeux, si votre vision change ou que la rougeur ou l'irritation des yeux persistent • les symptômes s'aggravent ou persistent plus de 3 jours.
Keep out of the reach of children. If swallowed, call a poison control centre or get medical help right away.
Garder hors de la portée des enfants. En cas d'ingestion, appeler un centre antipoison ou obtenir une assistance médicale immédiate.

Directions / Mode d'emploi
Adults and children 6 years and over: Remove contact lenses before using. Squeeze 1 or 2 drops in the affected eye(s) up to 4 times a day or as directed by a doctor.
Adultes et enfants de 6 ans et plus : Retirez les lentilles de contact avant l'utilisation. Instillez 1 ou 2 gouttes dans l'œil ou les yeux touché(s), jusqu'à 4 fois par jour ou selon les directives d'un médecin.
Inactive ingredients / Ingrédients inactifs
 Benzalkonium Chloride, Boric acid, Disodium EDTA, Sodium Borate, Sodium Chloride, Water
 Acide borique, borate de sodium, chlorure de benzalkonium, chlorure de sodium, eau, EDTA disodique
Questions?
 Call 1-866-672-0061
 Composez le 1-866-672-0061

compliments

Triple Action Formula
ADVANCED RELIEF EYE DROPS
 Temporarily relieves • redness of the eye due to minor eye irritations • burning and irritation due to dryness of the eyes.

Formule à triple action
GOUTTES POUR LES YEUX SOULAGEMENT OPTIMAL
 Soulagement temporaire • de la rougeur de l'œil due à des irritations mineures • de la sensation de brûlure et de l'irritation causées par la sécheresse de l'œil.

Solution
15 mL STERILE
 STERILE

DIT 0244319

Store between 15°C and 30°C. Do not use if safety seal around cap is missing or broken.
 Conserver entre 15 °C et 30 °C. Ne pas utiliser si le sceau de sécurité autour du bouchon est brisé ou manquant.
 Manufactured for / Fabriqué pour: Sobey's, Mississauga, ON L4W 0C7 © 2020. Teva Canada Limited / Limitée Toronto, Ontario M1B 2K9



LOT:
EXP:

POVIDONE (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)	POVIDONE	1 g in 100 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: OYZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B6978945GQ) (POLYETHYLENE GLYCOL, UNSPECIFIED - UNII:3WJQ0SDW1A)	POLYETHYLENE GLYCOL 400	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-989-15	1 in 1 BOX	05/14/2022	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Export only		05/14/2022	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc		174450460	manufacture(55651-989) , pack(55651-989) , label(55651-989)

Revised: 3/2023

KC Pharmaceuticals, Inc.