

STAQUIS- crisaborole ointment
Pharmacia & Upjohn Company LLC

Staquis® 2%

PRINCIPAL DISPLAY PANEL - Shipping Label - Label 0009-2211-30

Tubo con 30 g

Staquis®

Crisaborol Ungüento 2%

Vía de administración: Cutánea.

Lote:

Caducidad:

The image shows a shipping label for a tube of Staquis Crisaborol Ungüento 2% ointment. The label is rectangular with a white background and a black border. At the top right, it says "Tubo con 30 g". The product name "Staquis®" is written in a large, purple, sans-serif font. Below it, "Crisaborol Ungüento 2%" is written in a smaller, black, sans-serif font. Underneath that, "Vía de administración: Cutánea." is written in a small, black, sans-serif font. A decorative horizontal line of 20 dots, alternating between purple and light blue, separates the header from the body text. The body text is arranged in columns. On the left, it lists: "Fórmula: Cada 100 g contienen: Crisaborol 2.0 g", "Excipiente csp 100 g", "Dosis: La que el médico señale.", "Vía de administración: Cutánea.", "Consérvese el tubo bien cerrado a no más de 25°C.", "Su venta requiere receta médica.", "No se deje al alcance de los niños.", "No se use durante el embarazo o lactancia.", "●Marca Registrada", "Reg. No.", "Reporte las sospechas de reacción adversa al correo: farmacovigilancia@cofepris.gob.mx, MEX.AEReporting@pfizer.com y a la línea Pfizer 800 401 2002". In the center, it says "Hecho en Estados Unidos por: Pharmacia & Upjohn Company LLC, 7000 Portage Road Kalamazoo, MI 49001, Estados Unidos". On the right, it says "Distribuido por: Pfizer, S.A. de C.V. Km. 63 Carretera México Toluca, Zona Industrial, C.P. 50140, Toluca, México, México". At the bottom left, there is a small white box with the number "14674000" and a black square to its right. On the right side of the label, there are two vertical labels: "Lote:" and "Caducidad:", both followed by blank space for handwritten information.

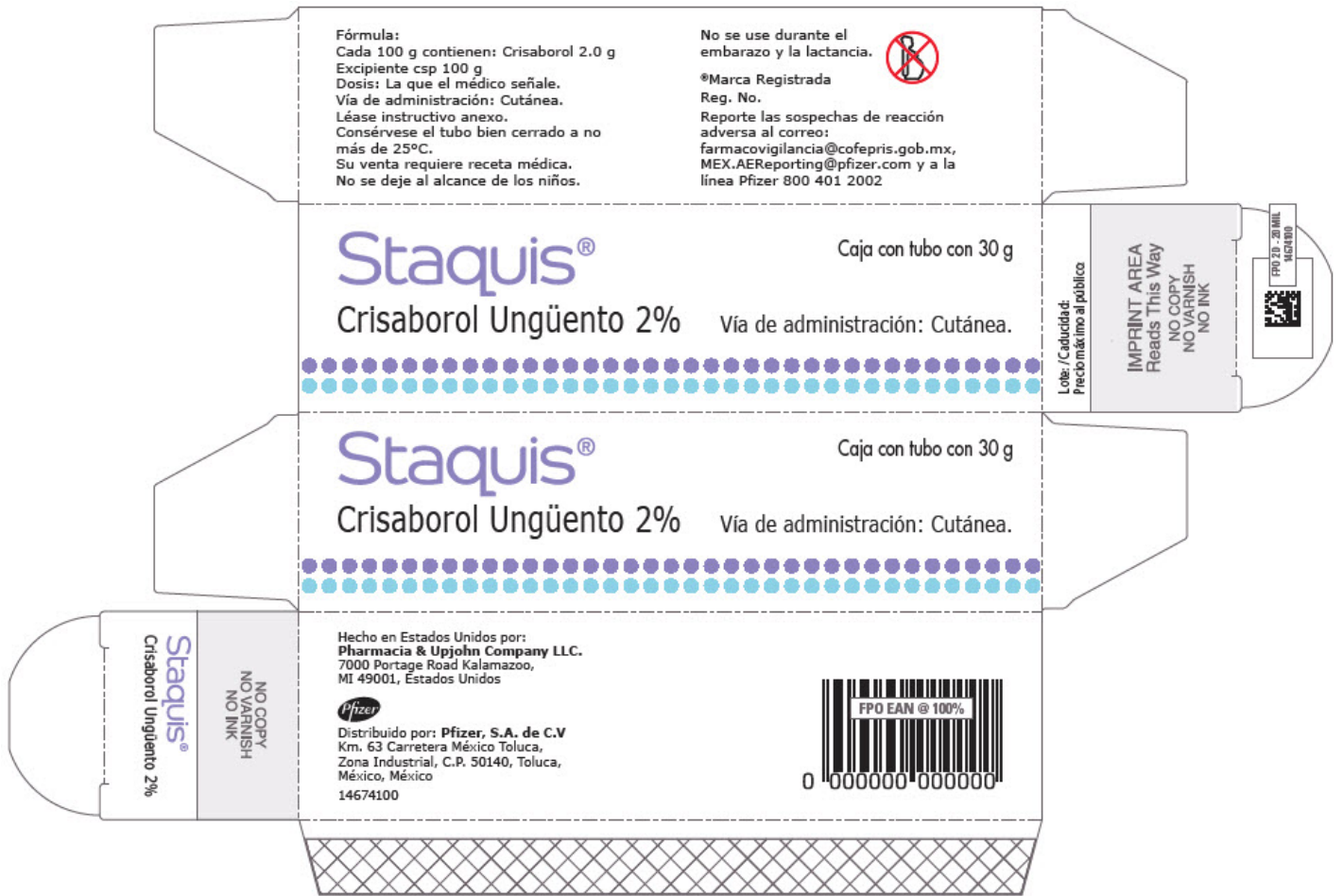
PRINCIPAL DISPLAY PANEL - Shipping Label - Carton 0009-2211-30

Staquis®

Crisaborol Ungüento 2%

Caja con tubo con 30 g

Vía de administración: Cutánea.



PRINCIPAL DISPLAY PANEL - Shipping Label - Label 0009-2211-12

Staquis®

30 g

Rx only

(crisaborole) ointment 2%

For Topical Use Only.

Not for ophthalmic, oral, or intravaginal use.

Lot

EXP

Staquis®

(crisaborole) ointment 2%

30 g
Rx only

For Topical Use Only.

Not for ophthalmic, oral, or intravaginal use.



DOSAGE AND USE: Apply twice daily to the affected areas. See accompanying prescribing information.

Each gram contains: 20 mg of crisaborole in an ointment containing white petrolatum USP, propylene glycol USP, mono- and di-glycerides NF, paraffin NF, butylated hydroxytoluene NF, and edetate calcium disodium USP.

Store below 30°C.
[see USP Controlled Room Temperature].
Keep out of reach of children.
Keep tube tightly closed.



Manufacturer:
Pharmacia & Upjohn
Company LLC,
subsidiary of Pfizer Inc,
7000 Portage Road,
Kalamazoo, Michigan (MI)
49001, USA

PAA202203

Lot

EXP

MFD



STAQUIS

crisaborole ointment

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0009-2211
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CRISABOROLE (UNII: Q2R47HGR7P) (CRISABOROLE - UNII:Q2R47HGR7P)	CRISABOROLE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PARAFFIN (UNII: I9O0E3H2ZE)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
EDETATE CALCIUM DISODIUM (UNII: 25IH6R4SGF)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0009-2211-30	1 in 1 CARTON	11/01/2020	
1		30 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0009-2211-12	1 in 1 CARTON	06/21/2023	
2		30 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Export only		11/01/2020	

Labeler - Pharmacia & Upjohn Company LLC (618054084)

Registrant - Pfizer Inc (113480771)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmacia & Upjohn Company LLC		618054084	ANALYSIS(0009-2211) , MANUFACTURE(0009-2211) , API MANUFACTURE(0009-2211) , PACK(0009-2211) , LABEL(0009-2211)

Revised: 6/2023

Pharmacia & Upjohn Company LLC