LBEL EFFET PARFIT- octinoxate lotion
VENTURA INTERNATIONAL LTD.,

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

L'BEL Effet Parfait fond de teint

Drug Facts

Active Ingredients

Octinoxate 7.0 %

Purpose

sunscreen

Uses

- Helps prevent sunburn
- Higher SPF gives more sunburn protection
- Provides minimal protection against sunburn

Warnings

For external use only.

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply liberally before sun exposure and as needed.

Other information

- Sun alert: Limiting sun exposure, wearing protective clothing, and using sunscreen may reduce the risk of skin aging, skin cancer, and other harmful effects of the sun.

Inactive ingredients

aqua (water), cyclomethicone, cyclopentasiloxane, cyclohexasiloxane, sodium chloride, peg/ppg-18/18 dimethicone, dimethicone, nylon-12 fluorescent brightener 230 salt, dimethiconol, diazolidinyl urea, methicone, cetearyl dimethicone crosspolymer, methylparaben, parfum (fragrance), propylparaben, polyvinylalcohol crosspolymer, sodium hyaluronate. **May contain:** ci77891 (titanium dioxide), ci77492 (iron oxides), ci77491 (iron oxides), ci77499 (iron oxides).

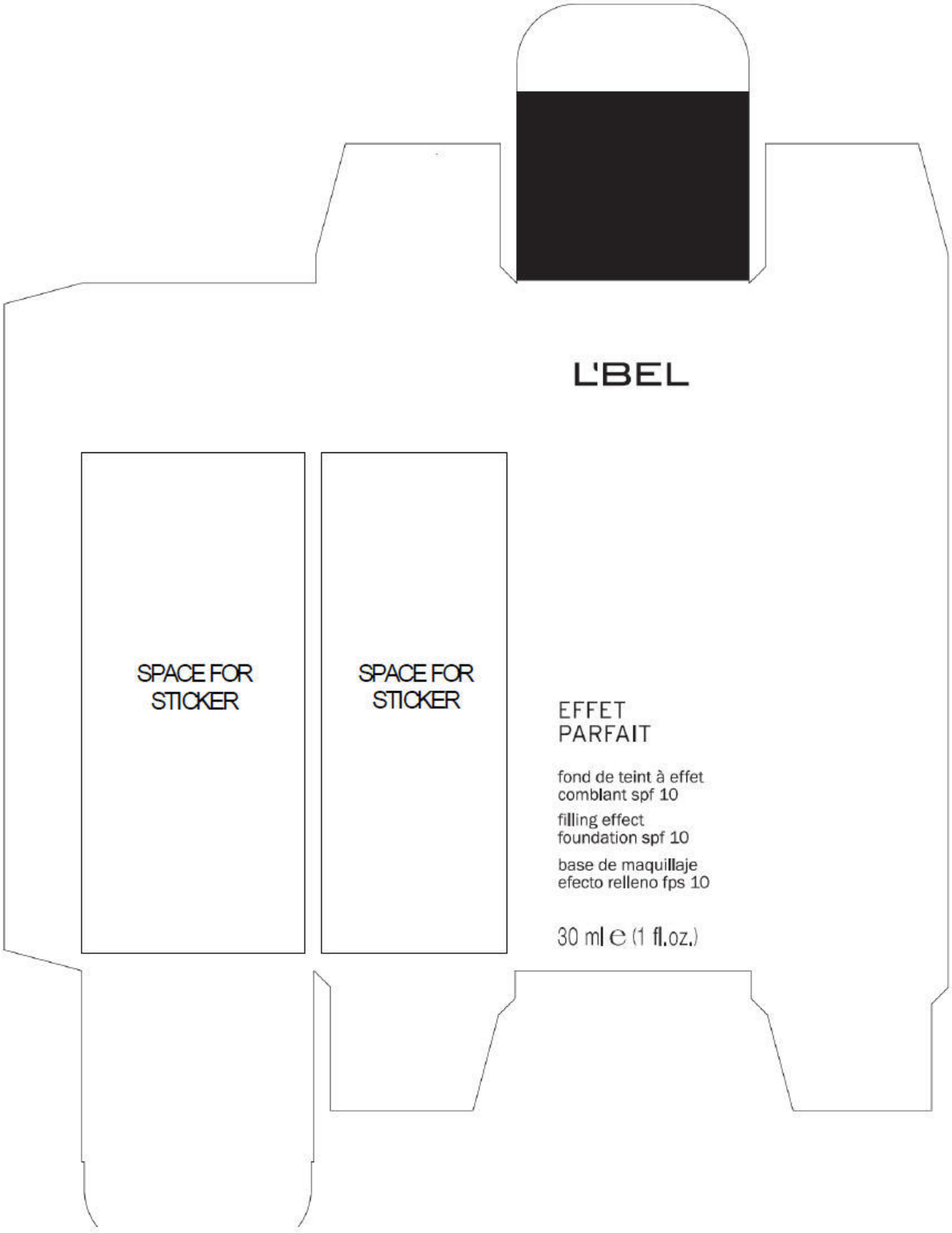
PRINCIPAL DISPLAY PANEL - 30 ml Bottle Carton

L'BEL

EFFET
PARFAIT

filling effect
foundation spf 10

30 ml e (1 fl.oz.)



LBEL EFFET PARFIT

octinoxate lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13537-230
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	2.1 g in 30 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
cyclomethicone (UNII: NMQ347994Z)	
cyclomethicone 5 (UNII: 0THT5PCI0R)	
cyclomethicone 6 (UNII: XHK3U310BA)	
sodium chloride (UNII: 451W47IQ8X)	
dimethicone (UNII: 92RU3N3Y1O)	
diazolidinyl urea (UNII: H5RIZ3MPW4)	
methylparaben (UNII: A2I8C7HI9T)	
propylparaben (UNII: Z8IX2SC1OH)	
hyaluronate sodium (UNII: YSE9PPT4TH)	
titanium dioxide (UNII: 15FIX9V2JP)	
ferric oxide red (UNII: 1K09F3G675)	
ferric oxide yellow (UNII: EX438O2MRT)	
ferrosoferric oxide (UNII: XM0M87F357)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:13537-230-62	1 in 1 BOX		
1	NDC:13537-230-61	30 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	08/15/2010	

Labeler - VENTURA INTERNATIONAL LTD., (602751344)

Revised: 8/2010 VENTURA INTERNATIONAL LTD.,