SUPREMACIE NX JOUR- ensulizole, homosalate, octinoxate, and oxybenzone cream
Ventura Corporation, LTDA

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.

Suprémacie Nx Tour Replenishing Treatment Daytime Face Cream SPF 15 Normal to Dry Skin

ACTIVE INGREDIENTS: ENSULIZOLE 2.0%, HOMOSALATE 2.0%, OCTINOXATE 7.0%, OXYBENZONE 4.0%.

WARNINGS: For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. When using this product keep out of eyes. Rinse with water to remove. Stop use and ask a doctor if irritation develops and lasts.

USES: Sun Protection Factor 15. Helps prevent sunburn.

DIRECTIONS: Apply evenly before sun exposure and as needed.

PRINCIPAL DISPLAY PANEL - 50 g jar

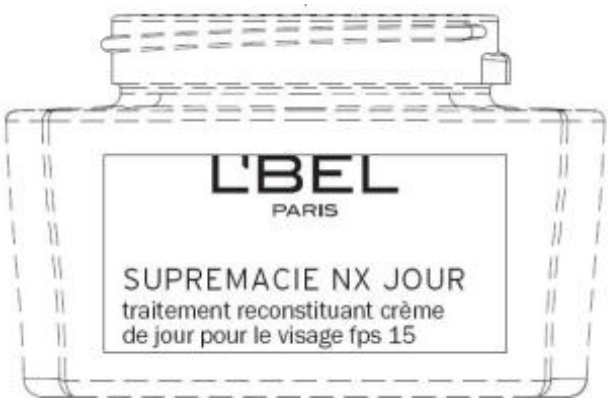
L'BEL
PARIS

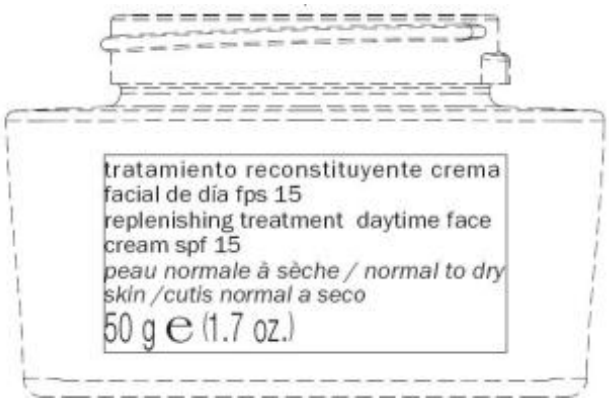
SUPREMACIE NX JOUR

traitement reconstituant crème
de jour pour le visage fps 15
tratamiento reconstituyente crema
facial de día fps 15
replenishing treatment daytime face
cream spf 15

peau normale à sèche / normal to dry
skin / cutis normal a seco

50 g e (1.7 oz.)





SUPREMACIE NX JOUR

ensulizole, homosalate, octinoxate, and oxybenzone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13537-261
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Ensulizole (UNII: 9YQ9DI1W42) (Ensulizole - UNII:9YQ9DI1W42)	Ensulizole	0.001 g in 5 g
Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	0.001 g in 5 g
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	0.00375 g in 5 g
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	0.002 g in 5 g

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:13537-261-64	1 in 1 BOX		
1	NDC:13537-261-61	5 g in 1 TUBE		
2	NDC:13537-261-65	1 in 1 JAR		
2	NDC:13537-261-62	50 g in 1 JAR		
3	NDC:13537-261-63	1 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	02/19/2010	

Labeler - Ventura Corporation, LTDA (602751344)

Revised: 2/2010 Ventura Corporation, LTDA