OHUI WHITE EXTREME ILLUMINATING PACT NO.10 - titanium dioxide, octinoxate, arbutin, atractylodes japonica root oil powder
LG Household and Healthcare, Inc.

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.

Drug Fact

TITANIUM DIOXIDE	9.84%
OCTINOXATE	4.5%
ARBUTIN	2%
ATRACTYLODES JAPONICA ROOT OIL	0.1%

For external use only

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

Keep out of eyes. Rinse with water to remove

Stop use if a rash or irritation develops and lasts.

OHUI White Extreme Illuminating Pact #20

OHUI WHITE EXTREME ILLUMINATING PACT NO.10

titanium dioxide, octinoxate, arbutin, atractylodes japonica root oil powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53208-534
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	9.84 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	4.5 g in 100 g
ARBUTIN (UNII: C5INA23HXF) (ARBUTIN - UNII:C5INA23HXF)	ARBUTIN	2 g in 100 g
ATRACTYLODES JAPONICA ROOT OIL (UNII: EC228KGY00) (ATRACTYLODES JAPONICA ROOT OIL - UNII:EC228KGY00)	ATRACTYLODES JAPONICA ROOT OIL	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
TALC (UNII: 7SEV7J4R1U)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
BORON NITRIDE (UNII: 2U4T60A6YD)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
MICA (UNII: V8A1AW0880)	
TALC (UNII: 7SEV7J4R1U)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM MYRISTATE (UNII: Z1917F0578)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAMOND (UNII: 6GRV67N0U2)	
COPPER (UNII: 789U1901C5)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PALMITOYL OLIGOPEPTIDE (UNII: HO4ZT5S86C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53208-534-02	1 in 1 BOX		
1	NDC:53208-534-01	11 g in 1 CONTAINER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	05/25/2011	

Labeler - LG Household and Healthcare, Inc. (688276187)

Registrant - LG Household and Healthcare, Inc. (688276187)

Establishment

Name	Address	ID/FEI	Business Operations
LG Household and Healthcare, Inc.		688276187	manufacture

Revised: 5/2011 LG Household and Healthcare, Inc.