

**TIME SECRET MINERAL PRESSED POWDER MEDIUM OCHRE- octyl methoxycinnamate, titanium dioxide, zinc oxide powder  
MSH 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.*

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**TIME SECRET Mineral Pressed Powder Medium Ochre**

**Active Ingredients**

Octyl Methoxycinnamate 5.63%

Titanium Dioxide 13.73%

Zinc Oxide 9.00%

**Purpose**

Sunscreen

**Warnings**

For external use only

**Do not use**

Do not use on broken or damaged skin

**When using**

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

**Stop use and ask a doctor**

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

**Keep out of reach of children**

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
- Children under 6 months of age: ask a doctor

**Inactive Ingredients**

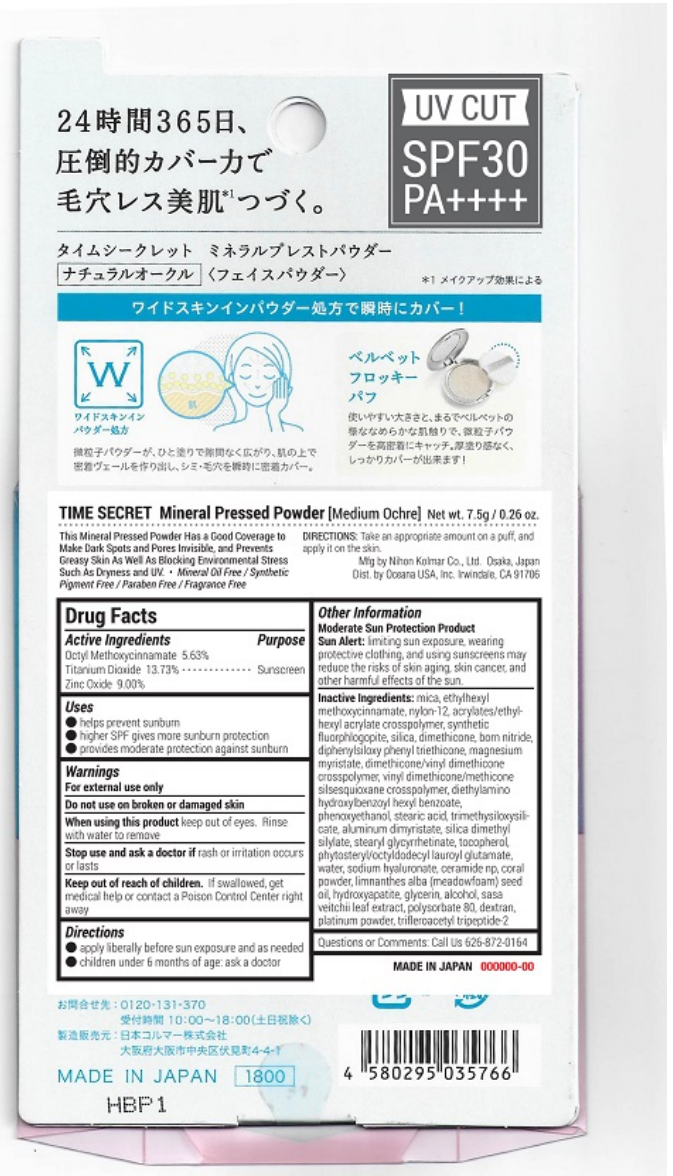
mica, nylon-12, acrylates/ethylhexyl acrylate crosspolymer, synthetic fluorphlogopite, silica, dimethicone, boron nitride, diphenylsiloxy phenyl trimethicone, magnesium myristate, dimethicone/vinyl dimethicone cross polymer, vinyl dimethicone/methicone silsesquioxane crosspolymer, diethylamino hydroxybenzoyl hexyl benzoate, phenoxyethanol, stearic acid, trimethyl

siloxysilicate, aluminum dimyristate, stearyl glycyrrhetinate, tocopherol, phytosteryl/octyldodecyl lauroyl glutamate, water, sodium hyaluronate, ceramide np, coral powder, limnanthes alba (meadow roam) seed oil, hydroxyapatite, glycerin, alcohol, sasa veitchii leaf extract, polysorbate 80, dextran, platinum powder, trifluoro acetyl tripeptide-2

## Questions

Questions or Comments: Call Us 626-872-0164

TimeSecret Mineral Pressed Powder Medium Ochre.jpg



## TIME SECRET MINERAL PRESSED POWDER MEDIUM OCHRE

octyl methoxycinnamate, titanium dioxide, zinc oxide powder

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72372-203
Route of Administration	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	422.25 mg in 7500 mg
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	1029.75 mg in 7500 mg
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	675 mg in 7500 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>NYLON-12</b> (UNII: 446U8J075B)	
<b>MICA</b> (UNII: V8A1AW0880)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:72372-203-01	7500 mg in 1 CASE; Type 0: Not a Combination Product	06/19/2018	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part352	06/19/2018	

**Labeler** - MSH Inc. (691891271)**Registrant** - MSH Inc. (691891271)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
MSH Inc.		691891271	manufacture(72372-203)

Revised: 6/2018

MSH Inc.