



## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:55259-0002
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>CALCIUM</b> (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)	CALCIUM	0.99

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>MAGNESIUM</b> (UNII: I38ZP9992A)	
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z)	
<b>SELENIUM</b> (UNII: H6241UJ22B)	
<b>GERMANIUM</b> (UNII: 00072J7XWS)	
<b>IRON</b> (UNII: E1UOL152H7)	
<b>POTASSIUM</b> (UNII: RWP5GA015D)	
<b>MANGANESE</b> (UNII: 42Z2K6ZL8P)	

## Product Characteristics

<b>Color</b>	white (white)	<b>Score</b>	no score
<b>Shape</b>	OVAL (tablet)	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	3;hp;x
<b>Contains</b>			

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:55259-0002-1	120 in 1 PACKAGE; Type 0: Not a Combination Product	03/18/2021	

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved drug other		03/18/2021	

**Labeler** - Apexel Co., Ltd (687287979)

**Registrant** - Apexel Co., Ltd (687287979)

## Establishment

Name	Address	ID/FEI	Business Operations
Apexel Co., Ltd		687287979	manufacture(55259-0002)

Revised: 3/2021

Apexel Co., Ltd