READY CARE XL KIT - first aid kit
Dukal Corporation

Front Label

Physicians Care

FIRST AID

50 Person

ReadyCare XL Kit

Contains over 355 Pieces

Back Label

Physicians Care

CLEANSE

20 Antiseptic Wipes
20 Alcohol Pads

TREAT

6 Antibiotic Ointment Packets
6 Burn Ointment Packets
10 3" Cotton Tip Applicators

PROTECT

3 Knuckle Bandages
3 Fingertip Bandages
35 3/8” x 1 1/2” Bandages
16 1” x 3” Bandages
200 3/4” x 3” Bandages
1 1/2" Tape
1 Triangular Bandage
4 3" x 3" Sterile Gauze Pads
1 Abdominal Compress

Medicine

3 Acetaminophen Packets (packet of 2)
3 Aspirin Packets (packets of 2)
3 Antacid Packets (packets of 2)

Other

Gloves (2 Pair)
1 Cold Pack
1 First Aid Guide
10 Finger Splints
5 Disposable Thermometers
1 Tweezer

Acme United Corporation
60 Round Hill Road

Fairfield CT 06824

www.acmeunited.com

Designed in the USA | Made in China

CAUTION: This product may contain natural rubber latex which may cause allergic reactions. This kit contains products that have expiration dates. Please check before use.

BZK Towelette Labeling

Reorder 855

NDC 65517-0004-1

Dukal

BZK TOWELETTE

Contains Benzalkonium Chloride

For External Use Only

Helps Prevent Infection

1 / Pouch

DUKAL CORPORATION

(631) 656-3800

Ronkonkoma, NY 11779 www.dukal.com

Made in China

Drug Facts

Active Ingredients.....Benzalkonium Chloride 0.133% w/v

Purpose..... ***First Aid Antiseptic***

USE: Antiseptic Cleansing of face, hands and body without soap and water. Airs dries in seconds

DO NOT USE: in the eyes or apply over large areas of the body.

STOP USE: If irritation, redness or other symptoms develop. Consult a doctor if the conditions persists or gets worse.

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

DIRECTIONS: Tear open packet, unfold and use as a washcloth.

INACTIVE INGREDIENTS: Distilled Water

Prep Pad Labeling

Reorder 852

Dukal Corporation

NDC 65517-0001-1

NPN 80003156

ALCOHOL PREP PAD

Saturated with 70% Isopropyl Alcohol

For External Use Only

1 / Pouch

Dukal Corporation

Ronkonkoma, NY 11779

631-656-3800

www.dukal.com

Made in China

Drug Facts

Active Ingredients

Isopropyl Alcohol 70%

Purpose
Antiseptic Cleanser

Use
For Preparation of Skin prior to an injection

Warnings
- For External Use Only
- Flammable, Keep away from fire or flame

Do Not Use
- with electrocautery procedures
- In the Eyes. If contact occurs, flush eyes with water

Stop Use
If irritation and redness develop. If condition persists, consult your health care practitioner.

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

Directions
Wipe injection site vigorously and discard.

Other Information
Store at Room Temperature 15 - 30 C (59 - 86 F)

Inactive Ingredient
purified water

Triple Antibiotic Labeling

WaterJel

Triple Antibiotic

First Aid Ointment

To Help Prevent Infection

Each Gram contains

Bacitracin Zinc 400 units

Neomycin Sulphate 5 mg

(equivalent to 3.5 mg Neomycin base)

Polymyxin B Sulfate 5000 units

Water-Jel Technologies

Carlstadt, NJ 07072

Drug Facts

Uses to help prevent infection in

minor cuts, scrapes, burns

Warnings

For external use only

Do not use

in the eyes or apply over large areas of the body

If you are allergic to any of the ingredients

longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

the condition persists or gets worse

a rash or other allergic reaction develops

Keep out of reach of children

if ingested contact Poison Control Center right away

Directions

clean affected area apply a small amount of product

(an amount equal to the surface area of the tip of a finger)

on the area 1 to 3 times daily may be covered with a sterile bandage

First Aid Burn Cream Labeling

WaterJel

First Aid Burn Cream

Antiseptic Pain Relief with Aloe

Active Ingredients:

Benzalkonium Chloride 0.13%

Lidocaine HCL 0.5%

Water-Jel Technologies

Carlstadt, NJ 07072

Drug Facts

Purpose

First Aid Antiseptic, External analgesic

Uses

first aid to help prevent infection and for temporary

relief of pain an itching associated with minor cuts,

scrapes, burns

Warnings

For external use only

Do not use

in the eyes

in large quantities over raw or blistered areas or on

deep puncture wounds, animal bites, or serious burns

Keep out of reach of children

if ingested contact Poison Control Center right away

Directions

clean affected area apply a small amount not more

than 3 times daily may be covered with a sterile bandage

Other Information

Store at room temperature

APAP Labeling

Drug Facts
Active Ingredients (In each tablet).....Purposes
Acetaminophen 325 mg.....Pain Reliever/Fever Reducer
Uses:
For the temporary relief of minor aches and pains associated with:
• headache • muscular aches • minor arthritis pain
• common cold • toothache • menstrual cramps
For the reduction of fever.
Warnings:
Alcohol Warning: If you consume 3 or more alcoholic drinks every day, ask your doctor whether you should take acetaminophen or other pain relievers/fever reducers. Acetaminophen may cause liver damage.
Do not use:
• with any other product containing acetaminophen
• for more than 10 days for pain unless directed by a doctor
• for more than 3 days for fever unless directed by a doctor
S**top Using and Ask a Doctor If:**

• symptoms do not improve
• new symptoms occur
• pain or fever persists or gets worse
• redness or swelling is present
Do not exceed recommended dosage. Keep this and all drugs out of the reach of children.
In case of accidental overdose, contact a physician or Poison Control Center immediately.
Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.
I**f pregnant or breast feeding,** ask a health professional before use.
Directions:
Do not use more than directed
Adults and children: (12 years and older)
Take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours, or as directed by a doctor.
Children under 12 years:
Do not give to children under 12 years of age.
Other Information:
• Store at room temperature
• Tamper-Evident Sealed Packets
• Do Not Use any Opened or Torn Packets
Inactive Ingredients:
Corn starch*, croscarmellose sodium, crospovidone, hypromellose, microcrystalline cellulose, mineral oil, opadry clear, pharmaceutical glaze, polyethylene glycol, povidone*, pregelatinized starch*, sodium carboxymethylcellulose, sodium starch glycolate, stearic acid*, talc*, titanium dioxide.
*may contain
Questions or comments? 1-800-634-7680

Alcalak Labeling

Drug Facts
Active Ingredients (In each tablet).....Purposes
Calcium Carbonate 420 mg .....Antacid
Uses:
For the relief of the following symptoms associated with:
• acid indigestion • sour stomach • heartburn • upset stomach
Warnings:
Ask a doctor or health professional before use if you have:
• been taking a prescription drug. Antacids may interact with certain prescription drugs.
• kidney stones
• a calcium-restricted diet
Stop using this product and ask a doctor if:
• symptoms last more than 2 weeks
Do not exceed recommended dosage.
Keep this and all drugs out of the reach of children. If you are pregnant or breast feeding, ask a health professional before use.
Directions:
• Do not use more than directed
Adults and children: (12 years and older)
Take 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not take more than 19 tablets in a 24 hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician.
Children under 12 years:
Do not give to children under 12 years of age.

Other Information:
• Phenylketonurics: Contains Phenylalanine 1.5 mg per tablet
• Each tablet contains 168 mg of elemental calcium
• Store at room temperature in a dry place
• Tamper-Evident Sealed Packets
• Do not Use any Opened or Torn Packets
Inactive Ingredients:
aspartame*, carageenan*, croscarmellose sodium*, glycine*, magnesium stearate, mint flavor*, silica*, sorbitol, spearmint flavor*, stearic acid*, stevix*, xylitol*
*may contain
Questions or comments? 1-800-634-7680

Aspirin Labeling

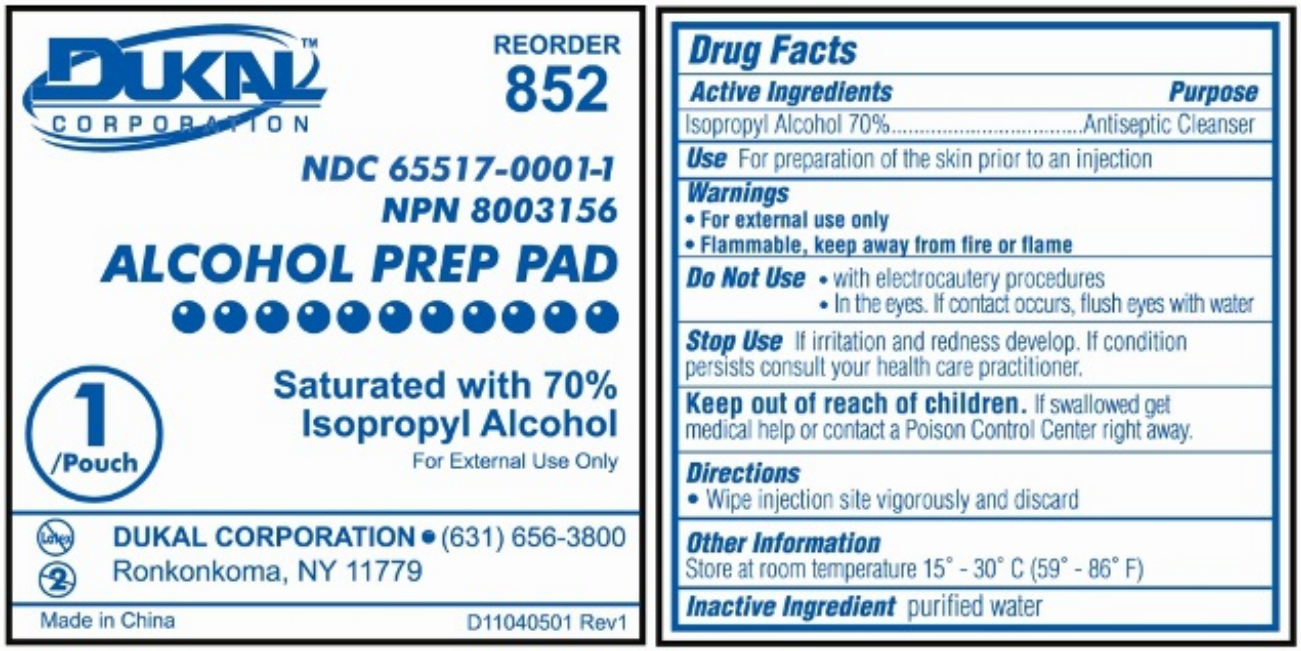
Drug Facts
Active Ingredients (In each tablet).....Purposes
Aspirin 325mg (NSAID*) .....Pain Reliever / Fever Reducer
*nonsteroidal anti-inflammatory drug
Uses:
Temporarily relieves minor aches and pains due:
• headache • muscular aches • minor arthritis pain • backache
• common cold • toothache • menstrual cramps
Temporarily reduces fever.
Warnings:
Reye's Syndrome: Children and teenagers should not use this medicine for chicken pox, or flu symptoms before a doctor is consulted about Reye's Syndrome, a rare but serious illness reported to be associated with aspirin.
Allergy Alert: Aspirin may cause a severe allergic reaction which may include:
• hives • skin reddening • facial swelling • rash • asthma (wheezing) • blisters • shock
If an allergic reaction occurs, stop use and seek medical help right away.
Stomach Bleeding Warning: This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause stomach bleeding.
The chance is higher if you:
• are age 60 or older • have had stomach ulcers or bleeding problems
• take a blood thinning (anticoagulant) or steroid drug
• take other drugs containing an NSAID (aspirin, ibuprofen, naproxen, or others)
• have 3 or more alcohol drinks every day while using this product
• take more or for a longer time than directed
Do not use:
• if you have ever had an allergic reaction to any other pain reliever/ fever reducer.
• right before or after heart surgery
Ask a Doctor before use if you have:
• problems or serious side effects from taking pain relievers or fever reducers
• stomach problems that last or come back, such as heartburn, upset stomach, or stomach pain • ulcers • bleeding problems • high blood pressure
• heart or kidney disease • taken a diuretic • reached age 60 or older
Ask a Doctor or Pharmacist before use if you are:
• taking any other drug containing and NSAID (prescription or nonprescription)
• taking a blood thinning (anticoagulant) or steroid drug
• under a doctor’s care for any serious condition • taking any other drug
When using this product:
• take with food or milk if stomach upset occurs
• long term continuous use may increase the risk of heart attack or stroke

Stop Using and Ask a Doctor If:
• you feel faint, vomit blood, or have bloody or black stools. These are signs of stomach bleeding. • pain gets worse or lasts more than 10 days
• fever gets worse or lasts more than 3 days • you have difficulty swallowing
• it feels like the pill is stuck in your throat • you develop heartburn
• stomach pain or upset gets worse or lasts
• redness or swelling is present in the painful area • any new symptoms appear
If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use aspirin during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.
Directions:
• Do not take more than directed.
• The smallest effective dose should be used.
• Do not take longer than 10 days, unless directed by a doctor (see Warnings)
• Drink a full glass of water with each dose.
Adults and children: (12 years and older)
Take 1 or 2 tablets with water every 4 hours as needed. Do not take more than 12 tablets in 24 hours, or as directed by a doctor.
Children under 12 years:
Do not give to children under 12 years of age.
Other Information:
• Read all product information before using
• Store at room temperature
• Avoid excessive heat and humidity
• Tamper evident sealed packets
• Do not use any opened or torn packets
Inactive Ingredients:
Corn starch, croscarmellose sodium*, hypromellose*, microcrystalline cellulose*, mineral oil*, polyethylene glycol, propylene glycol, silicon dioxide, starch, stearic acid, talc, titanium dioxide*.
*may contain
Questions or comments? 1-800-634-7680

Ready Care XL Kit Label

Ready Care XL Kit Back Label

Prep Pad Labeling



Antiseptic Towelette Label

DUKAL™ CORPORATION

REORDER 855

NDC 65517-0004-1

BZK TOWELETTE

1 /Pouch

Contains Benzalkonium Chloride
For External Use Only
Helps Prevent Infection

DUKAL CORPORATION ● (631) 656-3800
Ronkonkoma, NY 11779 ● www.dukal.com

Made in China

DRUG FACTS:

Active Ingredient	Purpose
Benzalkonium Chloride, 0.133% w/v	First Aid Antiseptic

USE: Antiseptic Cleansing of face, hands and body without soap and water. Air dries in seconds.

DO NOT USE: in the eyes or apply over large areas of the body.

STOP USE: if irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

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

DIRECTIONS: Tear open packet, unfold and use as a washcloth.

INACTIVE INGREDIENTS: Distilled Water

Tear Here

DUKAL CORPORATION D07110604 Rev1

Antacid Label

Acetaminophen Label

Aspirin Label

Triple Antibiotic Label

Burn Cream Label

READY CARE XL KIT

first aid kit kit

Product Information

Product Type	MEDICAL DEVICE	Item Code (Source)	NDC:65517-0021

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-0021-1	1 in 1 PACKAGE, COMBINATION		

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	20 POUCH	8.0 mL
Part 2	6 PACKET	5.4 g
Part 3	6 PACKET	5.4 g
Part 4	20 POUCH	28.0 mL
Part 5	3 PACKET	6
Part 6	3 PACKET	6
Part 7	3 PACKET	6

Part 1 of 7

DUKAL ALCOHOL PREP PAD

isopropyl alcohol swab

Product Information	
Item Code (Source)	NDC:65517-0001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.70 mL in 1.0 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65517-0001-1	0.4 mL in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2000	

Part 2 of 7

WATER-JEL 3-IN1 ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate and neomycin sulfate ointment

Product Information	
Item Code (Source)	NDC:59898-740
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B SULFATE	5000 [iU] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-740-01	0.9 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333	01/01/2010	

Part 3 of 7

FIRST AID BURN

lidocaine hydrochloride and benzalkonium chloride cream

Product Information

Item Code (Source)	NDC:59898-902
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59898-902-01	0.9 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333	01/01/2010	

Part 4 of 7

BZK TOWELETTE

benzalkonium chloride swab

Product Information

Item Code (Source)	NDC:65517-0004
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.00186 mL in 1.4 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	1.39814 mL in 1.4 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-0004-1	1.4 mL in 1 POUCH		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2006	

Part 5 of 7

MEDIQUE APAP

acetaminophen tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-145
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
HYPROMELLOSE (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47682-145-99	2 in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part343	12/30/2008	

Part 6 of 7

MEDIQUE ASPIRIN

aspirin tablet, film coated

Product Information	
Item Code (Source)	NDC:47682-116
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ASPIRIN (UNII: R16CO5Y76E) (ASPIRIN - UNII:R16CO5Y76E)	ASPIRIN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
HYPROMELLOSE (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47682-116-99	2 in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part343	12/30/2008	

Part 7 of 7

MEDIQUE ALCALAK

calcium carbonate tablet, chewable

Product Information	
Item Code (Source)	NDC:47682-088
Route of Administration	ORAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Calcium Carbonate (UNII: H0G9379FGK) (Calcium - UNII:SY7Q814VUP)	Calcium Carbonate	420 mg

Inactive Ingredients	
Ingredient Name	**Strength**
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	12mm
Flavor	MINT (MINT)	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47682-088-99	2 in 1 PACKET		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph final	part331	12/30/2008	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph final	part333	05/01/2010	

Labeler - Dukal Corporation (791014871)

Revised: 11/2010 Dukal Corporation