

**MODEL 6240-35-001CS- first aid kit with drug**  
**Aerospace Accessory Service, Inc**

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**Model 6240-35-001CS**

**Aerospace Accessory Service**

Aerospace Accessory Service

P/N:

S/N:

EXP:

Prep. By:

# AEROSPACE **AA** ACCESSORY



F.A.A. No XM4R653M

**SERVICE**

E.A.S.A. No EASA 145.5194

**P/N:** \_\_\_\_\_ **EXP:** \_\_\_\_/\_\_\_\_/\_\_\_\_

**S/N:** \_\_\_\_\_ **PREP. BY:** \_\_\_\_\_

IF SEAL IS BROKEN OR EXPIRATION DATE ARRIVES,  
REMOVE IMMEDIATELY FOR RE-CERTIFICATION

**Aerospace Accessory Service, Inc.**

**Ph: 305-594-1955 Fax: 305-470-2026**

**[www.aerospaceaccessory.com](http://www.aerospaceaccessory.com)**

# MODEL 6240-35-001CS

first aid kit with drug kit

## Product Information

Product Type	MEDICAL DEVICE	Item Code (Source)	NHRIC:27860-012
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## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:27860-012-09	1 in 1 PACKAGE; Type 1: Convenience Kit of Co-Package		

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1		1
Part 2		2
Part 3		3
Part 4		7
Part 5		5
Part 6		6
Part 7		7
Part 8		8
Part 9		9

## Part 1 of 9

### MOORE MEDICAL NON ASPIRIN

acetaminophen tablet, film coated

## Product Information

Item Code (Source)	NDC:55670-467
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Route of Administration	ORAL
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## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

## Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

**Product Characteristics**

<b>Color</b>	white (white)	<b>Score</b>	no score
<b>Shape</b>	ROUND (ROUND)	<b>Size</b>	10 mm
<b>Flavor</b>		<b>Imprint Code</b>	AZ;234
<b>Contains</b>			

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part343	12/30/2008	

**Part 2 of 9**

**NASAL DECONGESTANT**  
oxymetazoline hcl spray

**Product Information**

<b>Item Code (Source)</b>	NDC:0904-6761
<b>Route of Administration</b>	NASAL

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>OXYMETAZOLINE HYDROCHLORIDE</b> (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM</b> (UNII: K679OBS311)	
<b>SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM</b> (UNII: 3980JIH2SW)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	

## Product Characteristics

Color	white (to off white, viscous)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	10/12/2018	

## Part 3 of 9

### MOORE MEDICAL ANTACID

calcium carbonate tablet, chewable

## Product Information

Item Code (Source)	NDC:55670-142
Route of Administration	ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	420 mg

## Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SUCROSE (UNII: C151H8M554)	

## Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (ROUND)	Size	12mm
Flavor	MINT (MINT)	Imprint Code	FR;8
Contains			

## Marketing Information

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

## Part 4 of 9

### MOTION SICKNESS

meclizine hcl tablet

#### Product Information

Item Code (Source)	NDC:70677-0026
Route of Administration	ORAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MECLIZINE HYDROCHLORIDE (UNII: HDP7W44CIO) (MECLIZINE - UNII:3L5TQ84570)	MECLIZINE HYDROCHLORIDE	25 mg

#### Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

#### Product Characteristics

Color	yellow	Score	no score
Shape	ROUND	Size	9mm
Flavor		Imprint Code	44;403
Contains			

#### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part336	06/24/2002	

## Part 5 of 9

### AMMONIA INHALANTS

ammonia inhalants inhalant

**Product Information**

Item Code (Source)	NDC:46414-3333
Route of Administration	RESPIRATORY (INHALATION)

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
AMMONIA (UNII: 5138Q19F1X) (AMMONIA - UNII:5138Q19F1X)	AMMONIA	0.045 g in 0.3 mL

**Inactive Ingredients**

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
WATER (UNII: 059QF0KO0R)	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/14/1976	

**Part 6 of 9****EASY CARE FIRST AID DIPHENHYDRAMINE**

diphenhydramine hydrochloride tablet, film coated

**Product Information**

Item Code (Source)	NDC:44224-0017
Route of Administration	ORAL

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

**Inactive Ingredients**

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	

<b>LACTOSE</b> (UNII: J2B2A4N98G)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	pink (pink)	<b>Score</b>	no score
<b>Shape</b>	OVAL (OVAL)	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	048;D
<b>Contains</b>			

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/01/2012	

### Part 7 of 9

#### BZK PADS

benzalkonium chloride swab

### Product Information

<b>Item Code (Source)</b>	NDC:67777-245
<b>Route of Administration</b>	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/05/2011	



## Part 8 of 9

### POVIDONE-IODINE

povidone-iodine solution

#### Product Information

Item Code (Source) NDC:46414-7777

Route of Administration TOPICAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
NONOXYNOL-9 (UNII: 48Q180SH9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0K00R)	

#### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/14/1976	

## Part 9 of 9

### MOORE MEDICAL BISMUTH

bismuth subsalicylate tablet, chewable

#### Product Information

Item Code (Source) NDC:55670-474

Route of Administration ORAL

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ, BISMUTH CATION - UNII:ZS9CD118YE)	BISMUTH SUBSALICYLATE	262 mg

## Inactive Ingredients

Ingredient Name	Strength
ACACIA (UNII: 5C5403N26O)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ASPARTAME (UNII: Z0H242BBR1)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
DEXTRATES (UNII: G263M44RU)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

## Product Characteristics

Color	pink	Score	no score
Shape	ROUND	Size	16mm
Flavor		Imprint Code	RH;046
Contains			

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part335	04/01/2014	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
exempt device	ABC	01/01/2015	

**Labeler** - Aerospace Accessory Service, Inc (859100547)

**Registrant** - Aerospace Accessory Service, Inc (859100547)

## Establishment

Name	Address	ID/FEI	Business Operations
Aerospace Accessory Service, Inc		859100547	manufacture, repack