Public Summary

Summary for ARTG Entry: 15590 SUISSE LIQUID CHLOROPHYLL 20mg/g Solution bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

Products

1. LIQUID CHLOROPHYLL 20mg/g solution

Product Type Single Medicine Product **Effective date** 28/09/2001

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
500mL

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Oral Liquid, solution
Route of Administration	Oral
Visual Identification	Dark green aqueous liquid

Active Ingredients

Chlorophyll	**20 mg/g**

Public Summary

Summary for ARTG Entry: 15598 SUISSE ZESTABS Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

The label of this product's container and primary pack (if any) is to carry the following statement ' Warning - this product contains royal jelly and is not recommended for asthma and allergy sufferers as it can cause severe allergic reactions'.

Products

1. Zestabs Tablet

Product Type Single Medicine Product **Effective date** 27/03/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
100 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, round, convex faces, diameter 11.17 mm, brown speckled

Active Ingredients

Centella asiatica	**25 mg**
Eleutherococcus senticosus	**10 mg**
Pollen	**150 mg**
Royal jelly	**16.7 mg**
Smilax species	**37.5 mg**

Public Summary

Summary for ARTG Entry: 15597 SWISSE WOMEN'S MULTI-VITAMIN & MINERAL FORMULA Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

The label of this product's container and primary pack (if any) is to carry the following statement ' Warning - this product contains royal jelly and is not recommended for asthma and allergy sufferers as it can cause severe allergic reactions'.

Products

1. WOMEN'S MULTI-VITAMINS & MINERAL FORMULA Tablet

Product Type Single Medicine Product **Effective date** 1/04/1997

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size

60 tablets

Poison Schedule

Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Tablet, film-coated, oval with convex faces, approx. 24mm long x 11mm wide, approx. 1500mg, opaque apricot colour.

Active Ingredients

Aminobenzoic acid	**25 mg**
Berberis vulgaris	**15 mg**
Betacarotene	**10 mg**
Bioflavonoids	**25 mg**
Biotin	**50 microgram**
Calcium ascorbate	**200 mg**
Calcium citrate	**200 mg**
Calcium pantothenate	**75 mg**
Carica papaya	**5 mg**
Centella asiatica	**2.5 mg**
Chlorophyllin-copper complex	**5 mg**
Cholecalciferol	**201.4 IU**
Choline bitartrate	**25 mg**

Chromium amino acid chelate	10 microgram
Copper amino acid chelate	417 microgram
Cyanocobalamin	75 microgram
d-alpha-Tocopheryl acid succinate	83 mg
Equisetum arvense	6 mg
Evening Primrose Oil	10 mg
Ferrous fumarate	15.2 mg
Foeniculum vulgare	15 mg
Folic acid	180 microgram
Ginger	15 mg
Ginkgo biloba	625 microgram
Inositol	25 mg
Magnesium oxide	85 mg
Manganese amino acid chelate	8 mg
Matricaria recutita	15 mg
Mentha X cardiaca	1.5 mg
Nicotinamide	50 mg
Oryzanol	1 mg
Panax ginseng	8.3 mg
Petroselinum crispum	5 mg
Potassium amino acid chelate	10 mg
Potassium iodide	66 microgram
Pyridoxine hydrochloride	50 mg
Raspberry	15 mg
Riboflavine	25 mg
Royal jelly	7 mg
Silica - colloidal anhydrous	50 mg
Thiamine hydrochloride	50 mg
Ubidecarenone	1 mg
Zinc amino acid chelate	25 mg

Public Summary

Summary for ARTG Entry: 15595 SUISSE ULTI C Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

Products

1. Ulti C tablet

Product Type Single Medicine Product **Effective date** 27/03/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size	Poison Schedule
50 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Tablet, capsule shape 22.8 mm long x 9.4 mm wide, convex faces, film coated, cream colour, mottled.

Active Ingredients

Bioflavonoids	**100 mg**
Calcium ascorbate	**1000 mg**
Calcium carbonate	**41.3 mg**
Hesperidin	**25 mg**
Rutin	**25 mg**

Public Summary

Summary for ARTG Entry: 15593 SUISSE ULTI B COMPLEX Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Ulti B Complex Tablet

Product Type Single Medicine Product **Effective date** 27/03/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
30 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. SUISSE ULTI B COMPLEX Tablet

Dosage Form	Tablet
Route of Administration	Oral
Visual Identification	Tablet, round diameter 10.9 mm, convex faces, coated brown (chocolate)

Active Ingredients

Aminobenzoic acid	**50 mg**
Ascorbic acid	**100 mg**
Biotin	**50 microgram**
Calcium pantothenate	**50 mg**
Choline bitartrate	**50 mg**
Cyanocobalamin	**50 microgram**
Folic acid	**400 microgram**
Inositol	**100 mg**
Methionine	**50 mg**
Nicotinamide	**50 mg**
Pyridoxine hydrochloride	**50 mg**
Riboflavine	**50 mg**
Thiamine hydrochloride	**50 mg**

Public Summary

Public Summary

Summary for ARTG Entry: 15592 SWISSE ODOURLESS GARLIC & HORSERADISH Capsule bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. SWISSE HORSERADISH & ODOURLESS GARLIC Capsule

Product Type Single Medicine Product **Effective date** 25/11/1999

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size	Poison Schedule
75 capsules	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	Hard capsule size 0 clear containing an off white powder

Active Ingredients

Allium sativum	**250 mg**
Allium sativum	**83.4 mg**
Althaea officinalis	**20 mg**
Althaea officinalis	**5 mg**
Armoracia rusticana	**50 mg**
Armoracia rusticana	**250 mg**
Betacarotene	**1.5 mg**
Calcium ascorbate	**241 mg**
Histidine	**50 mg**
Thymus vulgaris	**20 mg**
Trigonella foenum-graecum	**5 mg**
Trigonella foenum-graecum	**20 mg**

Public Summary

Summary for ARTG Entry: 15596 SUISSE ULTIMINERAL Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. SUISSE Ultimineral tablet bottle

Product Type Single Medicine Product **Effective date** 16/10/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
100 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, capsule shaped, faces convex, grey, speckled, dimensions 17.3mm long x 6.9mm wide

Active Ingredients

Calcium amino acid chelate	**350 mg**
Chromium amino acid chelate	**50 microgram**
Copper amino acid chelate	**2 mg**
Ferrous fumarate	**1.65 mg**
Fucus vesiculosus	**10 mg**
Magnesium amino acid chelate	**175 mg**
Manganese amino acid chelate	**5 mg**
Potassium amino acid chelate	**30 mg**
Zinc amino acid chelate	**10 mg**

Public Summary

Summary for ARTG Entry: 15585 SWISSE HALIBUT LIVER OIL 145mg A PLUS D Capsule bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. SWISSE Halibut Liver Oil 145mg A Plus D capsule

Product Type Single Medicine Product **Effective date** 21/07/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
100 capsules

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, soft
Route of Administration	Oral
Visual Identification	Capsule SEG (soft elastic gelatin), round seamed 4minim, colour clear white

Active Ingredients

Halibut-liver oil	**145 mg**

Public Summary

Summary for ARTG Entry: 15589 SUISSE LIVER EXTRACT 400mg Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Liver Extract 400mg tablet

Product Type Single Medicine Product **Effective date** 27/03/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size

70 tablets

Poison Schedule

Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, oval, dimensions 23.4 mm long x 10.6 mm wide. Brown (light) with brown (dark) speckle.

Active Ingredients

Liver extract	**400 mg**

Public Summary

Summary for ARTG Entry: 15586 SUISSE HERBAL DIURETIC Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

Products

1. SUISSE Herbal Diuretic tablet

Product Type Single Medicine Product **Effective date** 28/09/1998

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size
60 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, round, convex faces, diameter 12mm, green speckled.

Active Ingredients

Arctostaphylos uva-ursi	**50 mg**
Equisetum arvense	**77.5 mg**
Juniperus communis	**50 mg**
Petroselinum crispum	**300 mg**
Potassium gluconate	**25 mg**

Public Summary

Summary for ARTG Entry: 15581 SUISSE CO ENZYME Q 10 Capsule bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989".

Products

1. CO ENZYME Q 10 Capsule

Product Type Single Medicine Product **Effective date** 25/11/1999

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size
75 capsules

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	Two piece capsule (hard) clear size o filled with a cream powder

Active Ingredients

Calcium hydrogen phosphate	**200 mg**
dl-alpha tocopheryl acetate	**10 IU**
Magnesium phosphate	**100 mg**
Potassium chloride	**50 mg**
Ubidecarenone	**12 mg**

Public Summary

Summary for ARTG Entry: 15583 SUISSE GINKGO BILOBA 200mg Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Ginkgo Biloba Tablet

Product Type Single Medicine Product **Effective date** 27/03/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size
90 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, round, diameter 11.26 mm, convex faces, grey with brown mottles

Active Ingredients

Ginkgo biloba	**200 mg**

Public Summary

Summary for ARTG Entry: 15587 SUISSE INOSINE Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. INOSINE tablet

Product Type Single Medicine Product **Effective date** 16/10/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size
50 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, capsule shape convex faces, 22.6 mm x 9.3 9.3mm, white

Active Ingredients

Inosine	**500 mg**
Magnesium aspartate	**30 mg**
Potassium aspartate	**10 mg**

Public Summary

Summary for ARTG Entry: 15588 SUISSE LIPOTROPIC FAT EMULSIFIERS Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Lipotropic Fat Emulsifiers Tablet

Product Type	Single Medicine Product	**Effective date**	16/10/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
69 tablets,	Not scheduled. Not considered by committee,
40 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Tablet, capsule shape, dimensions 22.7 mm long x 9.3 mm wide. Convex faces, film coated. Light brown speckled.

Active Ingredients

Betaine hydrochloride	**100 mg**
Choline bitartrate	**500 mg**
Fucus vesiculosus	**10 mg**
Inositol	**250 mg**
Lecithin	**20 mg**
Liver extract	**5 mg**
Medicago sativa	**10 mg**
Methionine	**100 mg**
Pancreatin	**20 mg**
Pyridoxine hydrochloride	**25 mg**
Stellaria media	**5 mg**

Public Summary

Public Summary

Summary for ARTG Entry: 15582 SUISSE GARLIC & HORSERADISH COMPLEX Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Garlic & Horseradish Complex Tablet

Product Type Single Medicine Product **Effective date** 16/10/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
60 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, uncoated
Route of Administration	Oral
Visual Identification	Tablet, round, convex faces, diameter 12.08 mm, grey speckled

Active Ingredients

Allium sativum	**167 mg**
Althaea officinalis	**12.5 mg**
Armoracia rusticana	**100 mg**
Petroselinum crispum	**50 mg**
Trigonella foenum-graecum	**50 mg**

Public Summary

Summary for ARTG Entry: 15584 SUISSE HAIR & NAIL Tablet bottle

ARTG entry for	Medicine
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	10/09/1991
Product category	Medicine
Status	Revoked
Approval area	

Conditions

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Products

1. Hair And Nail Tablet

Product Type Single Medicine Product **Effective date** 24/02/1995

Warnings

No Warnings included on Record

Standard Indications

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

Specific Indications

No Specific Indications included on Record

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size
85 tablets

Poison Schedule
Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet
Route of Administration	Oral
Visual Identification	Tablet, round, convex faces, diameter 11.3 mm, light green speckled

Active Ingredients

Calcium hydrogen phosphate	**424 mg**
Equisetum arvense	**230 mg**
Magnesium oxide	**8.3 mg**
Silicon dioxide	**25 mg**

Public Summary

Summary for ARTG Entry: 129846 SWISSE MEN'S ULTIVITE MULTIVITAMIN, MINERAL AND ANTI-OXIDANT FORMULA 1 WITH HERBS (CAPSULE)

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	31/07/2006
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm,as amended from time to time.

Products

1. SWISSE MEN'S ULTIVITE MULTIVITAMIN, MINERAL AND ANTI-OXIDANT FORMULA 1 WITH HERBS (CAPSULE)

Product Type Single Medicine Product **Effective date** 28/08/2009 9:44:39 AM

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

Contains caffeine [state quantity per dosage unit or per mL or per gram of product] [must be clear and legible].,

This product contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.,

Do not take while on warfarin therapy without medical advice.,

Contains lactose (or words to that effect).,

If symptoms persist consult your healthcare practitioner (or words to that effect).,

Contains [insert name of ingredient] (or words to that effect).

Standard Indications

Source of folic acid. Can assist in maintaining normal blood.,

Source of folic acid. Can assist in maintaining normal blood. Blood tonic.,

Male support. Balances and supports normal male physiology and function.,

Aids, assists or helps in the maintenance or improvement of general well-being.,

Beneficial during times of stress. [Warning S required],

Helps relieve nervous tension, stress and mild anxiety. [Warning S required],

For mineral (may state the mineral) supplementation.,

For vitamin (may state the vitamin) supplementation.,

Prevention/treatment of vitamin [XX] and/or mineral [YY] and/or nutritional deficiencies (This indication is not be to used for the treatment of iron deficiency conditions).,

May assist in the management of dietary folate deficiency.

Specific Indications

Designed to meet men's nutritional needs and general well-being. The herbal ingredients in this formulation may help men perform in peak condition and improve stress adaptation.

Additional Product Information

Pack Size/Poison Information

Pack Size	Poison Schedule

Components

1. COMPONENT ONE

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

Apium graveolens	**1 mg**
Equivalent: Apium graveolens (Dry)	10 mg
Astragalus membranaceus	**2.5 mg**
Equivalent: Astragalus membranaceus (Dry)	25 mg
Avena sativa	**25 mg**
Equivalent: Avena sativa (Fresh)	250 mg
Barosma betulina	**1.25 mg**
Equivalent: Barosma betulina (Dry)	5 mg
Berberis vulgaris	**1.25 mg**
Equivalent: Berberis vulgaris (Dry)	7.5 mg
Biotin	**25 microgram**
Calcium ascorbate dihydrate	**100 mg**
Calcium citrate	**50 mg**
Calcium pantothenate	**35 mg**
Camellia sinensis	**1.67 mg**
Equivalent: Camellia sinensis (Dry)	10 mg
Carica papaya	**5 mg**
Centella asiatica	**6.25 mg**
Equivalent: Centella asiatica (Dry)	25 mg
Cholecalciferol	**.0025 mg**
Choline bitartrate	**12.5 mg**
Chromium picolinate	**25 microgram**
Citrus bioflavonoids extract	**20 mg**
Copper gluconate	**100 microgram**
Crataegus monogyna	**12.5 mg**
Equivalent: Crataegus monogyna (Dry)	50 mg
Cyanocobalamin	**15 microgram**
Cynara scolymus	**6.25 mg**
Equivalent: Cynara scolymus (Fresh)	25 mg
d-alpha-Tocopheryl acid succinate	**20.256 mg**
Dunaliella salina	
Equisetum arvense	**3.75 mg**
Equivalent: Equisetum arvense (Dry)	15 mg
Ferrous fumarate	**4.8 mg**
Foeniculum vulgare	**1.7 mg**
Equivalent: Foeniculum vulgare (Dry)	7.5 mg
Folic acid	**250 microgram**
Ginkgo biloba	**1 mg**
Equivalent: Ginkgo biloba (Dry)	50 mg
Inositol	**12.5 mg**
Lutein	**100 microgram**
Lycopersicon esculentum	
Lysine hydrochloride	**25 mg**
Magnesium oxide - heavy	**50 mg**

Manganese amino acid chelate	**6 mg**
Nicotinamide	**15 mg**
Panax ginseng	**8.33 mg**
Equivalent: Panax ginseng (Dry)	25 mg
Petroselinum crispum	**5 mg**
Potassium iodide	**33 microgram**
Potassium sulfate	**4.46 mg**
Pyridoxine hydrochloride	**15 mg**
Riboflavine	**15 mg**
Selenomethionine	**32.5 microgram**
Serenoa repens	**10 mg**
Equivalent: Serenoa repens (Dry)	100 mg
Silybum marianum	**357 microgram**
Smilax officinalis	**6.25 mg**
Equivalent: Smilax officinalis (Dry)	25 mg
Spearmint Oil	**750 microgram**
Thiamine hydrochloride	**15 mg**
Turnera diffusa	**12 mg**
Equivalent: Turnera diffusa (Dry)	60 mg
Tyrosine	**500 microgram**
Ubidecarenone	**500 microgram**
Vaccinium myrtillus	
Vitis vinifera	**4.17 mg**
Zinc amino acid chelate	**15 mg**
Zingiber officinale	**500 microgram**
Equivalent: Zingiber officinale (Dry)	2.5 mg

Public Summary

Summary for ARTG Entry: 124329 Swisse Ultiboost Bone Dense

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/12/2005
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm as amended from time to time.

Products

1. Swisse Ultiboost Bone Dense

Product Type Single Medicine Product **Effective date** 31/05/2011 12:08:48 PM

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Vitamin D helps calcium absorption (or words of like intent) and a diet deficient in calcium can lead to osteoporosis in later life,

Source of calcium. A diet deficient in calcium can lead to osteoporosis in later life.

Specific Indications

Swisse Ultiboost Bone Dense is best taken at night because the mineral flux in your body that maintains bone growth is greatest during sleep.
It has been shown that taking excessive amounts of calcium without the necessary supplementary nutrients may hinder the rate of calcium absorption.
Swisse Ultiboost Bone Dense should be combined with weight bearing exercises, as determined by your Health Care Practitioner, to assist in increasing bone density to minimize the risk of osteoporosis in later life.
Formulated to strengthen and protect bones and teeth

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Jar/Can	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size

Poison Schedule

Components

1. Formulation 1

Dosage Form	Tablet, film coated
Route of Administration	Oral

Visual Identification

Active Ingredients	
Calcium ascorbate dihydrate	50 mg
Calcium carbonate	750 mg
Cholecalciferol	.012 mg
Chromic chloride	51 microgram
Copper gluconate	250 microgram
Magnesium aspartate	80 mg
Magnesium oxide - heavy	461.95 mg
Magnesium phosphate	150 mg
Manganese amino acid chelate	25 mg
Potassium aspartate	27.28 mg
Potassium iodide	66 microgram
Zinc sulfate monohydrate	33.34 mg

Public Summary

Summary for ARTG Entry: 122610 Swisse Men's Ultivite Multi-Vitamin Mineral & Anti-oxidant Formula 1 with herbs (tablets)

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	12/10/2005
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm as amended from time to time.,

Where the medicine is a conventional release folic acid supplement preparation, in tablet form, of strength of 100 micrograms or more per dosage unit, the additional conditions are as follows: Where the medicine contains folic acid as a single active ingredient, the dissolution characteristics for folic acid in this medicine must comply with the dissolution requirements in the United States Pharmacopoeia 24th Edition (USP24) monograph for Folic Acid Tablets. Where the medicine contains multiple active ingredients, the dissolution characteristics for folic acid in this medicine must comply with the dissolution requirements in the United States Pharmacopoeia 24th Edition (USP24) monograph for Nutritional Supplements, as amended by its First Supplement. The dissolution characteristics for this medicine must comply with the dissolution requirements of the relevant USP24 monograph over the shelf life of the product when stored under the conditions included on the product label. It is the responsibility of manufacturers to develop and validate the analytical methods for the determination of folic acid that should be used in conjunction with the USP24 dissolution criteria. This condition does not apply to chewable, effervescent or dispersible tablets or sustained release tablets.

Products

1. Swisse Men's Ultivite Multi-Vitamin Mineral & Anti-oxidant Formula 1 with herbs (tablets)

Product Type Single Medicine Product **Effective date** 28/08/2009 9:44:39 AM

Warnings

Contains [insert name of ingredient] (or words to that effect).,

Do not take while on warfarin therapy without medical advice.,

Contains lactose (or words to that effect).,

This product contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.,

Contains caffeine [state quantity per dosage unit or per mL or per gram of product] [must be clear and legible].,

If symptoms persist consult your healthcare practitioner (or words to that effect).,

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Aids, assists or helps in the maintenance or improvement of general well-being.,

May assist in the management of dietary folate deficiency.,

Source of folic acid. Can assist in maintaining normal blood.,

Source of folic acid. Can assist in maintaining normal blood. Blood tonic.,

Male support. Balances and supports normal male physiology and function.,

Helps relieve nervous tension, stress and mild anxiety. [Warning S required],

Prevention/treatment of vitamin [XX] and/or mineral [YY] and/or nutritional deficiencies (This indication is not be to used for the treatment of iron deficiency conditions).,

For vitamin (may state the vitamin) supplementation.,

For mineral (may state the mineral) supplementation.,

Beneficial during times of stress. [Warning S required]

Specific Indications

Designed to meet men's nutritional needs and general well-being. The herbal ingredients in this formulation may help men perform in peak condition and improve stress adaptation.

Additional Product information

Pack Size/Poison information

Pack Size | **Poison Schedule**

Components

1. COMPONENT ONE

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Apium graveolens	**2 mg**
Equivalent: Apium graveolens (Dry)	20 mg
Astragalus membranaceus	**5 mg**
Equivalent: Astragalus membranaceus (Dry)	50 mg
Avena sativa	**50 mg**
Equivalent: Avena sativa (Fresh)	500 mg
Barosma betulina	**2.5 mg**
Equivalent: Barosma betulina (Dry)	10 mg
Berberis vulgaris	**2.5 mg**
Equivalent: Berberis vulgaris (Dry)	15 mg
Biotin	**50 microgram**
Calcium ascorbate dihydrate	**200 mg**
Calcium citrate	**100 mg**
Calcium pantothenate	**70 mg**
Camellia sinensis	**3.34 mg**
Equivalent: Camellia sinensis (Dry)	20 mg
Carica papaya	**10 mg**
Centella asiatica	**12.5 mg**
Equivalent: Centella asiatica (Dry)	50 mg
Cholecalciferol	**.005 mg**
Choline bitartrate	**25 mg**
Chromium picolinate	**50 microgram**
Citrus bioflavonoids extract	**40 mg**
Copper gluconate	**200 microgram**
Crataegus monogyna	**25 mg**
Equivalent: Crataegus monogyna (Dry)	100 mg
Cyanocobalamin	**30 microgram**
Cynara scolymus	**12.5 mg**
Equivalent: Cynara scolymus (Fresh)	50 mg
d-alpha-Tocopheryl acid succinate	**41.33 mg**
Dunaliella salina	
Equisetum arvense	**7.5 mg**
Equivalent: Equisetum arvense (Dry)	30 mg
Ferrous fumarate	**9.61 mg**
Foeniculum vulgare	**3.41 mg**
Equivalent: Foeniculum vulgare (Dry)	15 mg
Folic acid	**500 microgram**
Ginkgo biloba	**2 mg**

Equivalent: Ginkgo biloba (Dry)	100 mg
Inositol	**25 mg**
Lutein	**.2 mg**
Lycopersicon esculentum	
Lysine hydrochloride	**50 mg**
Magnesium oxide - heavy	**100 mg**
Manganese amino acid chelate	**12 mg**
Nicotinamide	**30 mg**
Panax ginseng	**16.67 mg**
Equivalent: Panax ginseng (Dry)	50 mg
Petroselinum crispum	**10 mg**
Potassium iodide	**66 microgram**
Potassium sulfate	**8.92 mg**
Pyridoxine hydrochloride	**30 mg**
Riboflavine	**30 mg**
Selenomethionine	**65 microgram**
Serenoa repens	**20 mg**
Equivalent: Serenoa repens (Dry)	200 mg
Silybum marianum	**715 microgram**
Smilax officinalis	**12.5 mg**
Equivalent: Smilax officinalis (Dry)	50 mg
Spearmint Oil	**1.7 mg**
Thiamine hydrochloride	**30 mg**
Tocopherols concentrate - mixed (low-alpha type)	**466.2 microgram**
Turnera diffusa	**24 mg**
Equivalent: Turnera diffusa (Dry)	120 mg
Tyrosine	**1 mg**
Ubidecarenone	**1 mg**
Vaccinium myrtillus	
Vitis vinifera	**8.34 mg**
Zinc amino acid chelate	**30 mg**
Zingiber officinale	**1 mg**
Equivalent: Zingiber officinale (Dry)	5 mg

Public Summary

Summary for ARTG Entry: 123100 H BIO-JUVEN HAIR NAILS SKIN MINERAL & HERBAL FORMULA WITH ADDED VITAMINS

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	26/10/2005
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm as amended from time to time.

Products

1. H BIO-JUVEN HAIR NAILS SKIN MINERAL & HERBAL FORMULA WITH ADDED VITAMINS

Product Type Single Medicine Product **Effective date** 28/08/2009 9:44:39 AM

Warnings

If symptoms persist consult your healthcare practitioner (or words to that effect).,

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Aids, assists or helps in the maintenance or improvement of general well-being.,

Aids or assists in the maintenance of peripheral circulation.

Specific Indications

H Bio-Juven contains vitamins and minerals known to science today to assist in making hair, nails and skin look and feel better, plus herbs to help aid in the maintenance of peripheral circulation so that these nutrients can travel to their destination and fulfil the desired objectives. To PERI and healthy hair, skin and nails.

Additional Product Information

Pack Size/Poison Information

Pack Size **Poison Schedule**

Components

1. COMPONENT ONE

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

Biotin	**200 microgram**

Calcium ascorbate dihydrate	**60 mg**
Calcium citrate	**50 mg**
Calcium pantothenate	**25 mg**
Cholecalciferol	**.25 microgram**
Citrus bioflavonoids extract	**40 mg**
Cyanocobalamin	**25 microgram**
Cysteine hydrochloride	**60 mg**
Cystine	**20 mg**
d-alpha-Tocopheryl acid succinate	**10.0032 mg**
Dunaliella salina	
Equisetum arvense	**12.5 mg**
Equivalent: Equisetum arvense (Dry)	50 mg
Ferrous fumarate	**9.61 mg**
Folic acid	**50 microgram**
Ginkgo biloba	**10 mg**
Magnesium oxide - heavy	**51.82 mg**
Nicotinamide	**10 mg**
Panicum miliaceum	**25.46 mg**
Potassium iodide	**65.4 microgram**
Pyridoxine hydrochloride	**10 mg**
Riboflavine	**10 mg**
Salvia officinalis	**50 mg**
Thiamine hydrochloride	**10 mg**
Zinc amino acid chelate	**25 mg**
Zingiber officinale	**50 mg**

Public Summary

Summary for ARTG Entry: 100430 Childvite 2

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	29/03/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.

Products

1. Childvite 2

Product Type Single Medicine Product **Effective date** 4/10/2007 11:00:00 PM

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Aids, assists or helps in the maintenance or improvement of general well-being.

Specific Indications

A children's Multivitamin, Mineral & Antioxidant Formula with selected herbs to promote stamina, endurance and well-being.
Swisse Childvite 2 Multivitamin, Mineral and Antioxidant Formula in powder form has been formulated specifically for children of all ages, including those who are too young to swallow or chew tablets. Childvite 2 has been specially formulated with selected vitamins, minerals, antioxidants and herbs to address the nutritional needs of children low in stamina and endurance. The vitamin, mineral and antioxidant ingredients have been added to assist in the promotion of general well-being, while the herbs promote stamina and endurance.

Additional Product information

Pack Size/Poison Information

Pack Size **Poison Schedule**

Components

1. Medicine Component

Dosage Form	Oral Liquid, powder for
Route of Administration	Oral
Visual Identification	

Active Ingredients

Ascorbic acid	**5 mg/g**
Biotin	**3 microgram/g**
Calcium carbonate	**5.25 mg/g**
Calcium pantothenate	**200 microgram/g**
Citrus bioflavonoids extract	**200 microgram/g**

Cyanocobalamin	**300 ng/g**
Eleutherococcus senticosus	**39 microgram/g**
Equivalent: Eleutherococcus senticosus (Dry)	643.5 microgram/g
Ferrous sulfate - dried	**700 microgram/g**
Folic acid	**1.5 microgram/g**
Magnesium aspartate	**2.4 mg/g**
Manganese sulfate monohydrate	**50 microgram/g**
Nicotinamide	**1 mg/g**
Pyridoxine hydrochloride	**200 microgram/g**
Riboflavin sodium phosphate	**130 microgram/g**
Thiamine hydrochloride	**100 microgram/g**
Urtica dioica	**200 microgram/g**
Equivalent: Urtica dioica (Dry)	1 mg/g
Zinc sulfate monohydrate	**1.1 mg/g**

Public Summary

Summary for ARTG Entry: 122391 SWISSE WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI-OXIDANT FORMULA 1 WITH HERBS (CAPSULE)

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	4/10/2005
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Products

1. SWISSE WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI-OXIDANT FORMULA 1 WITH HERBS (CAPSULE)

Product Type Single Medicine Product **Effective date** 28/08/2009 9:44:39 AM

Warnings

Contains caffeine [state quantity per dosage unit or per mL or per gram of product] [must be clear and legible].,

Do not take while on warfarin therapy without medical advice.,

If symptoms persist consult your healthcare practitioner (or words to that effect).,

This product contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.,

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

Contains lactose (or words to that effect).

Standard Indications

Source of folic acid. Can assist in maintaining normal blood.,

Source of folic acid. Can assist in maintaining normal blood. Blood tonic.,

Helps relieve nervous tension, stress and mild anxiety. [Warning S required].,

Beneficial during times of stress. [Warning S required].,

For mineral (may state the mineral) supplementation.,

For vitamin (may state the vitamin) supplementation.,

Prevention/treatment of vitamin [XX] and/or mineral [YY] and/or nutritional deficiencies (This indication is not be to used for the treatment of iron deficiency conditions).,

May assist in the management of dietary folate deficiency.,

Aids, assists or helps in the maintenance or improvement of general well-being.

Specific Indications

Designed to assist in meeting women's nutritional needs and maintaining general well-being. The herbal ingredients may help women perform in peak condition and improve stress adaptation.
Assist in the relief of pre-menstrual symptoms/syndrome

Additional Product Information

Pack Size/Poison Information

Pack Size **Poison Schedule**

Components

1. COMPONENT ONE

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

Apium graveolens	**1 mg**
Equivalent: Apium graveolens (Dry)	10 mg
Arctostaphylos uva-ursi	**3.12 mg**
Equivalent: Arctostaphylos uva-ursi (Dry)	12.5 mg
Astragalus membranaceus	**2.5 mg**
Equivalent: Astragalus membranaceus (Dry)	25 mg
Avena sativa	**25 mg**

Equivalent: Avena sativa (Fresh)	250 mg
Biotin	**25 microgram**
Calcium ascorbate dihydrate	**100 mg**
Calcium citrate	**50 mg**
Calcium pantothenate	**37.5 mg**
Camellia sinensis	**1.67 mg**
Equivalent: Camellia sinensis (Dry)	10 mg
Carica papaya	**5 mg**
Centella asiatica	**1.25 mg**
Equivalent: Centella asiatica (Dry)	5 mg
Cholecalciferol	**.0025 mg**
Choline bitartrate	**12.5 mg**
Chromium picolinate	**25 microgram**
Citrus bioflavonoids extract	**20 mg**
Copper gluconate	**208.5 microgram**
Crataegus monogyna	**3.75 mg**
Equivalent: Crataegus monogyna (Dry)	15 mg
Cyanocobalamin	**25 microgram**
Cynara scolymus	**6.25 mg**
Equivalent: Cynara scolymus (Fresh)	25 mg
d-alpha-Tocopheryl acid succinate	**21.1 mg**
Dunaliella salina	
Eleutherococcus senticosus	**1.25 mg**
Equivalent: Eleutherococcus senticosus (Dry)	12.5 mg
Equisetum arvense	**3.75 mg**
Equivalent: Equisetum arvense (Dry)	15 mg
Ferrous fumarate	**7.94 mg**
Foeniculum vulgare	**1.7 mg**
Equivalent: Foeniculum vulgare (Dry)	7.5 mg
Folic acid	**250 microgram**
Ginkgo biloba	**50 microgram**
Equivalent: Ginkgo biloba (Dry)	2.5 mg
Glycyrrhiza glabra	**1.25 mg**
Equivalent: Glycyrrhiza glabra (Dry)	5 mg
Inositol	**12.5 mg**
Lutein	**100 microgram**
Lycopersicon esculentum	
Lysine hydrochloride	**25 mg**
Magnesium oxide	**42.5 mg**
Manganese amino acid chelate	**8 mg**
Matricaria recutita	**1.88 mg**
Equivalent: Matricaria recutita (Dry)	7.5 mg
Nicotinamide	**25 mg**
Petroselinum crispum	**5 mg**
Potassium iodide	**32.7 microgram**
Potassium sulfate	**2.23 mg**
Pyridoxine hydrochloride	**25 mg**
Riboflavine	**25 mg**
Selenomethionine	**32.5 microgram**
Silybum marianum	**357 microgram**
Spearmint Oil	**750 microgram**
Thiamine hydrochloride	**25 mg**
Ubidecarenone	**500 microgram**
Vaccinium myrtillus	
Vitis vinifera	**4.17 mg**
Zinc amino acid chelate	**12.5 mg**
Zingiber officinale	**1.5 mg**

Equivalent: Zingiber officinale (Dry)	7.5 mg

Public Summary

Summary for ARTG Entry: 91326 Opticq

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	15/10/2002
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Products

1. Opticq

Product Type Single Medicine Product **Effective date** 3/04/2009 8:39:00 AM

Warnings

No Warnings included on Record

Standard Indications

Eyes formula. Assists visual fatigue and eye-strain.,

Eyes formula. Support of healthy eye function.,

Aids, assists or helps in the maintenance or improvement of general well-being.,

Maintenance of healthy eyes.,

Eyes formula. Helps eyes to adapt to variations in light intensity.

Specific Indications

CLINICALS OPTICQ by Swisse has been developed by a professor and contains scientifically validated ingredients to relieve visual fatigue and eye strain. It is recommended for people who experience problems with night vison and those sensitive to glare. OPTICQ is based on the assessment of clinical studies, which showed certain ingredients in OPTICQ to assist in promoting healthy vision. Used as directed, OPTICQ may deliver positive results relatively quickly.For better results: OPTICQ may be effective when used on it's own, however we recommend you use this product in conjunction with Swisse Women's & Men's Ultivite, which act as a "foundation" to provide the bodywith the important nutrients necessary for general wellbeing.

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
SIMELOAD - missing code value	Not recorded	0 Unknown	SIMELOAD - missing code value	Not recorded	Not recorded

Pack Size/Poison information

Pack Size

Not Applicable

Poison Schedule

Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Information Not Provided

Active Ingredients

Calcium ascorbate	**250 mg**
Cupric sulfate pentahydrate	**3.93 mg**
d-alpha-Tocopheryl acid succinate	**165.29 mg**
Tagetes erecta	**60 mg**
Equivalent: Tagetes erecta (Dry)	330 mg
Tocopherols concentrate - mixed (low-alpha type)	**378 mg**
Zinc oxide	**24.89 mg**

This is not an ARTG Certificate document.

The onus is on the reader to verify the current accuracy of the information on the document subsequent to the date shown.
Visit www.tga.gov.au for contact information

Public Summary

Summary for ARTG Entry: 100431 Childvite 1

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	29/03/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.

Products

1. Childvite 1

Product Type Single Medicine Product **Effective date** 4/10/2007 11:00:00 PM

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Aids, assists or helps in the maintenance or improvement of general well-being.

Specific Indications

A children's Multivitamin, Mineral & Antioxidant Formula with selected herbs to promote calmness in very active children.
Swisse Childvite 1 Multivitamin, Mineral and Antioxidant Formula in powder form has been formulated specifically for children of all ages, including those who are too young to swallow or chew tablets. Childvite 1 has been specifically formulated with selected vitamins, minerals, antioxidants and herbs to address the nutritional needs of very active children. These ingredients are known to be important for well-being and the herbs have been selected to promote calmness while relieving stress and nervous tension in very active children. Childvite 1 may also be used to assist the child to prepare for sleep. Childvite 1 has been developed by the Swisse team of medical researchers in consultation with parents to provide a pleasant tasting effervescent liquid with natural colours and flavours and no preservatives.

Additional Product information

Pack Size/Poison information

Pack Size **Poison Schedule**

Components

1. Medicine Component

Dosage Form	Oral Liquid, powder for
Route of Administration	Oral
Visual Identification	

Active Ingredients

Ascorbic acid	**5 mg/g**
Biotin	**3 microgram/g**
Calcium carbonate	**5.25 mg/g**

Calcium pantothenate	**200 microgram/g**
Citrus bioflavonoids extract	**200 microgram/g**
Cyanocobalamin	**200 ng/g**
Ferrous sulfate - dried	**350 microgram/g**
Folic acid	**1 microgram/g**
Humulus lupulus	**57.14 microgram/g**
Equivalent: Humulus lupulus (Dry)	200 microgram/g
Magnesium aspartate	**2.4 mg/g**
Manganese sulfate monohydrate	**50 microgram/g**
Matricaria recutita	**375 microgram/g**
Equivalent: Matricaria recutita (Dry)	1.5 mg/g
Nicotinamide	**1 mg/g**
Pyridoxine hydrochloride	**300 microgram/g**
Riboflavin sodium phosphate	**160 microgram/g**
Thiamine hydrochloride	**150 microgram/g**
Zinc sulfate monohydrate	**1.1 mg/g**

Public Summary

Summary for ARTG Entry: 93095 Prosqn

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	18/02/2003
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The listing of this medicine is subject to statutory conditions of listing and additional conditions of listing imposed by the Secretary under the provisions of subsection 28(3) of the Therapeutic Goods Act 1989 by notice in writing.

Products

1. Prosqn

Product Type Single Medicine Product **Effective date** 4/10/2007 11:00:00 PM

Warnings

No Warnings included on Record

Standard Indications

For the symptomatic relief of medically diagnosed benign prostatic hypertrophy.,

Aids, assists or helps in the maintenance or improvement of general well-being.,

May assist in the management of medically diagnosed benign prostatic hypertrophy.

Specific Indications

CLINICALS PROSQN by Swisse has been developed by a professor and contains scientifically validated ingredients, to assist in normal prostate function. PROSQN is based on the assessment of clinical studies, which showed certain ingredients in PROSQN to provide relief from the symptoms of medically diagnosed benign prostatic hypertrophy (or hyperplasia commonly known as prostate enlargement) such as difficulties in micturition, a hesitant interrupted weak stream, urinary frequency, urgency and dribbling or leaking. PROSQN is based on natural herbal ingredients. Used as directed, PROSQN may deliver positive results relatively quickly. For better results, PROSQN may be effective when used on it's own, however we recommend you use this product in conjunction with CLINICALS CEREXL and SWISSE MEN'S ULTIVITE FORMULA 1, which act as a "foundation" to provide the body with the important nutrients necessary for general wellbeing.

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
SIMELOAD - missing code value	Not recorded	0 Unknown	SIMELOAD - missing code value	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size

Not Applicable

Poison Schedule

Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Information Not Provided

Active Ingredients

Cholecalciferol	**1.25 microgram**
Cucurbita pepo	**25 mg**
Equivalent: Cucurbita pepo (Dry)	500 mg
Curcuma longa	**4 mg**
Equivalent: Curcuma longa (Dry)	100 mg
Folic acid	**250 microgram**
Ganoderma lucidum	**2.75 mg**
Equivalent: Ganoderma lucidum (Dry)	55 mg
Glycyrrhiza glabra	**13.75 mg**
Equivalent: Glycyrrhiza glabra (Dry)	55 mg

Isatis tinctoria	**11 mg**
Equivalent: Isatis tinctoria (Dry)	55 mg
Lycopersicon esculentum	**10 mg**
Equivalent: Lycopersicon esculentum (Fresh)	3.5 g
Panax ginseng	**5.5 mg**
Equivalent: Panax ginseng (Dry)	55 mg
Selenomethionine	**16.25 microgram**
Serenoa repens	**100 mg**
Equivalent: Serenoa repens (Dry)	1 g
Serenoa repens	**217.5 mg**
Equivalent: Serenoa repens (Dry)	870 mg
Urtica dioica	**62.5 mg**
Equivalent: Urtica dioica (Dry)	1.13 g
Zinc amino acid chelate	**30 mg**

Public Summary

Summary for ARTG Entry: 120384 Venaxl

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	6/07/2005
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Products

1. Venaxl

Product Type Single Medicine Product **Effective date** 3/04/2009 8:39:00 AM

Warnings

Do not take while on warfarin therapy without medical advice.,

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

This product contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.

Standard Indications

Protects capillaries.,

Aids, assists or helps in the maintenance or improvement of general well-being.,

Liver tonic.,

Maintain health of capillaries.,

Formula to support the liver.,

May assist blood circulation.

Specific Indications

- Support for Healthy Heart & Cholesterol - CLINICALS VENAXL by Swisse has been developed by a professor and contains scientifically validated ingredients that help maintain a healthy cardiovascular system. VENAXL assists in the maintenance of healthy heart tissue and function as well as protecting capillaries and helping maintain normal blood circulation. VENAXL also assists in the maintenance of healthy cholesterol level, which is so important for the overall wellbeing of the cardiovascular system. The liver plays a key role in the detoxification process and fat metabolism which assists in promoting healthy heart function. Therefore, VENAXL has been designed so that it also provides support for healthy liver function. VENAXL may deliver positive results relatively quickly.For better results: VENAXL may be effective when used on it's own, however we recommend you use this product in conjunction with Swisse Women's or Men's Ultivite which act as a "foundation" to provide the body with the important nutrients necessary for general well being. Has antioxidant action via its free radical scavenging properties.

Additional Product information

Pack Size/Poison information

Pack Size **Poison Schedule**

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Calcium ascorbate	**50 mg**
Citrus bioflavonoids extract	**20 mg**
Crataegus monogyna	**150 mg**
Equivalent: Crataegus monogyna (Dry)	600 mg
Cyanocobalamin	**25 microgram**
Cynara scolymus	**16.67 mg**
Equivalent: Cynara scolymus (Fresh)	500 mg
d-alpha-Tocopheryl acid succinate	**82.6464 mg**
Folic acid	**250 microgram**
Levocarnitine	**50 mg**
Magnesium amino acid chelate	**250 mg**
Nicotinic acid	**25 mg**

Pyridoxine hydrochloride	**10 mg**
Selenomethionine	**16.25 microgram**
Silybum marianum	**21.43 mg**
Ubidecarenone	**25 mg**

Public Summary

Summary for ARTG Entry: 92813 Cerexl

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	3/02/2003
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Where the medicine is a convential release folic acid supplement preparation, in tablet form, of strength of 100 micrograms or more per dosage unit, the additional conditions are as follows: Where the medicine contains folic acid as a single active ingredient, the dissolution characteristics for folic acid in this medicine must comply with the dissolution requirements in the United States Phamacopoeia 24th Edition (USP24) monograph for Folic Acid Tablets. Where the medicine contains multiple active ingredients, the dissolution characteristics for folic acid in this medicine must compley with the dissolution requirements in the United States Phamacopoeia 24th Edition (USP24) monograph for Nutritional Supplements, as amended by its First Supplement. The dissolution characteristics for this medicine must comply with the dissolution requirements of the relevant USP24 monograph over the shelf life of the product when stored under the conditions included on the product label. It is the respo

Products

1. Cerexl

Product Type Single Medicine Product **Effective date** 4/10/2007 11:00:00 PM

Warnings

If symptoms persist consult your healthcare practitioner (or words to that effect).,

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.

Standard Indications

Aids, assists or helps in the maintenance or improvement of general well-being.,

Aids or assists in the prevention of muscular cramps and spasms.,

Eyes formula. Support of healthy eye function.,

May assist in the management of tinnitus. [Warning S required],

May assist peripheral circulation.,

To help maintain blood circulation to the peripheral areas of the body such as the legs, hands and feet.,

Assisting in the maintenance of blood flow in the hands, feet and legs.,

For the symptomatic relief of tinnitus. [Warning S required]

Specific Indications

Support of Memory and Stamina CLINICALS CEREXL by Swisse has been developed by a professor and contains scientifically validated ingredients that help maintain memory function and stamina. CEREXL is based on the assessment of clinical studies, which showed certain ingredients in CEREXL can improve short-term memory and enhance mental alertness. CEREXL helps improve brain function such as learning or concentration as well as helping relieve the symptoms of absentmindedness. Used as directed, CEREXL may deliver positive results relatively quickly.For better results: CEREXL may be effective when used on it's own, however we recommend you use this product in conjunction with Swisse Women's or Men's Ultivite, which act as a foundation to provide the body with the important nutrients for general wellbeing. Reduces muscular pain and leg cramps and improves blood circulation. Relieves symptoms of cold hands and feet.

Additional Product information

Pack Size/Poison information

Pack Size	Poison Schedule

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Calcium ascorbate dihydrate	**60 mg**
Cyanocobalamin	**25 microgram**
Folic acid	**250 microgram**
Ginkgo biloba	**30 mg**
Equivalent: Ginkgo biloba (Dry)	1.5 g
Glutamine	**125 mg**
Lecithin liquid - soy phosphatidylserine-enriched soy	**75 mg**
Levocarnitine	**50 mg**
Magnesium aspartate	**250 mg**
R,S-alpha Lipoic acid	**75 mg**
Thiamine hydrochloride	**50 mg**
Tyrosine	**50 mg**
Zinc amino acid chelate	**12.5 mg**

Public Summary

Summary for ARTG Entry: 107806 Swisse Ultiboost Prostate

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	7/09/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Products

1. Swisse Ultiboost Prostate

Product Type Single Medicine Product **Effective date** 29/06/2012

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

This product contains selenium which is toxic in high doses. A daily dose of 150 micrograms for adults of selenium from dietary supplements should not be exceeded.,

If symptoms persist consult your healthcare practitioner (or words to that effect).

Standard Indications

For the symptomatic relief of medically diagnosed benign prostatic hypertrophy.,

May assist in the management of medically diagnosed benign prostatic hypertrophy.,

Aids, assists or helps in the maintenance or improvement of general well-being.

Specific Indications

Swisse Ultiboost PROSTATE is based on 25 years of research and contains scientifically validated ingredients, to assist in normal prostate function. Swisse Ultiboost PROSTATE is based on the assessment of clinical studies, which indicated that certain ingredients in the formula provided relief from the symptoms of medically diagnosed Benign Prostatic Hypertrophy (or Hyperplasia, commonly known as prostate enlargement), including difficulties in micturition, a hesitant interrupted weak stream, urinary frequency, urgency and dribbling or leaking. Swisse Ultiboost PROSTATE is based on natural herbal ingredients. Used as directed, Swisse Ultiboost PROSTATE may deliver positive results relatively quickly.

Additional Product Information

Pack Size/Poison Information

Pack Size **Poison Schedule**

Components

1. Formulation 1

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Cholecalciferol	**1.25 microgram**
Cucurbita pepo	**25 mg**
Curcuma longa	**25 mg**
Folic acid	**250 microgram**
Ganoderma lucidum	**2.75 mg**
Glycyrrhiza glabra	**13.75 mg**
Isatis tinctoria	**11 mg**
Lycopersicon esculentum	**10 mg**
Panax ginseng	**5.5 mg**
Selenomethionine	**16.25 microgram**
Serenoa repens	**217.5 mg**
Serenoa repens	**100 mg**
Urtica dioica	**62.5 mg**
Zinc amino acid chelate	**30 mg**

Public Summary

Public Summary

Summary for ARTG Entry: 110295 SWISSE STUDENT HD

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	28/09/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm as amended from time to time.

Products

1. SWISSE STUDENT HD

Product Type Single Medicine Product **Effective date** 4/10/2007 11:00:00 PM

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

If symptoms persist consult your healthcare practitioner (or words to that effect).,

Contains [insert name of ingredient] (or words to that effect).

Standard Indications

Helps relieve nervous tension, stress and mild anxiety. [Warning S required],

Beneficial during times of stress. [Warning S required]

Specific Indications

Vitamin supplement with added minerals and herbs to help perform in peak condition. A vitamin, mineral and herbal supplement formulated to address the needs of students in today's demanding, competitive environment. The herbs in Swisse Student HD may be of benefit in times of stress, or when peak condition and stamina are desired. A special formulation designed to help improve stress adaptation. The herbs have been especially chosen for their beneficial effects in providing relief of nervous tension, mild anxiety and stress tension.

Additional Product information

Pack Size/Poison information

Pack Size **Poison Schedule**

Components

1. COMPONENT ONE

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	

Active Ingredients

Avena sativa	**50 mg**
Equivalent: Avena sativa (Fresh)	500 mg
Calcium pantothenate	**25 mg**
Calcium phosphate	**160 mg**
Cyanocobalamin	**10 microgram**
Ginkgo biloba	**5 mg**
Equivalent: Ginkgo biloba (Dry)	250 mg
Magnesium phosphate	**130 mg**
Nicotinamide	**10 mg**
Panax ginseng	**35 mg**
Equivalent: Panax ginseng (Dry)	350 mg
Potassium phosphate - dibasic	**65 mg**
Pyridoxine hydrochloride	**10 mg**
Riboflavine	**15 mg**
Smilax ornata	**12.5 mg**
Equivalent: Smilax ornata (Dry)	50 mg
Thiamine nitrate	**10 mg**
Zinc amino acid chelate	**25 mg**

Public Summary

Summary for ARTG Entry: 109294 Swisse Ultiboost Sleep

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	20/09/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm as amended from time to time.

Products

1. Swisse Ultiboost Sleep

Product Type Single Medicine Product **Effective date** 10/12/2009 10:39:18 AM

Warnings

If symptoms persist consult your healthcare practitioner (or words to that effect).

Standard Indications

Helps relieve nervous tension, stress and mild anxiety. [Warning S required],

Help reduce effects of mild anxiety and nervous tension. [Warning S required],

Aids, assists or helps in the maintenance or improvement of general well-being.

Specific Indications

Can?t sleep? Swisse Ultiboost SLEEP is based on the assessment of clinical studies and contains scientifically validated ingredients shown to act quickly and assist you to sleep like a baby and wake up refreshed.
Swisse Ultiboost SLEEP is a combination of herbs clinically proven to promote calmness and relaxation, and to assist in relieving nervous tension and promoting natural, restful sleep. Swisse Ultiboost SLEEP contains ingredients shown to decrease sleep latency (the time taken to fall asleep), assist with reducing symptoms such as mild anxiety and stress and minimise sleep challenges.

Additional Product information

Pack Size/Poison information

Pack Size **Poison Schedule**

Components

1. Formulation 1

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Anemarrhena asphodeloides	**10 mg**
Glycyrrhiza glabra	**12.5 mg**
Humulus lupulus	**26.67 mg**
Magnesium orotate	**100 mg**
Poria cocos	**10 mg**
Valeriana officinalis	**325 mg**

Public Summary

Summary for ARTG Entry: 114871 Imunxl

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	30/11/2004
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.,

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Office of Complementary Medicines, Therapeutic Goods Administration, upon request.,

The sponsor of the listed medicine must not, by any means, intentionally or recklessly advertise the medicine for an indication other than those accepted in relation to the inclusion of the medicine in the Register.,

All reports of adverse reactions or similar experiences associated with the use or administration of the listed medicine shall be notified to the Head, Office of Product Review, Therapeutic Goods Administration, as soon as practicable after the sponsor of the goods becomes aware of those reports. Sponsors of listed medicines must retain records of such reports for a period of not less than 18 months from the day the Head, Office of Product Review is notified of the report or reports.,

The sponsor shall not supply the listed medicine after the expiry date of the goods.,

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.,

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Therapeutic Goods Act 1989, shall be only those included in the list of 'Colourings permitted in medicines for oral use' available at http://www.tga.gov.au/industry/cm-colourings-oral-use.htm, as amended from time to time.

Products

1. Imunxl

Product Type Single Medicine Product **Effective date** 12/04/2005

Warnings

Vitamins can only be of assistance if the dietary vitamin intake is inadequate. OR Vitamin supplements should not replace a balanced diet.,

WARNING: May be dangerous if taken in large amounts or for a long period. OR WARNING: Contains zinc which may be dangerous if taken in large amounts or for a long period (or words to that effect).,

If coughing persists consult your doctor (or a healthcare professional) (or words to that effect).,

Adults only. OR Not to be used in children under two years of age without medical advice (or words to that effect).,

Adults only. OR Not to be used in children under two years of age without medical advice (or words to that effect).,

If symptoms persist consult your healthcare practitioner (or words to that effect).

Standard Indications

Temporary relief of coughs. [Warnings COU1 and COU2 required],

Relief of coughs. [Warnings COU1 and COU2 required],

Relief of the symptoms of influenza/flu. [Warnings S and COLD required],

Can assist in the treatment of flu by reducing the severity and duration of symptoms. [Warnings S and COLD required],

Relief of mucous congestion. [Warnings S and COLD required],

Relief of symptoms of mild upper respiratory infections. [Warnings S and COLD required],

Helps fight mild upper respiratory complaints. [Warnings S and COLD required],

For the symptomatic relief of recurrent upper respiratory tract infections. [Warnings S and COLD required],

May assist in the management of recurrent upper respiratory tract infections. [Warnings S and COLD required],

Aids, assists or helps in the maintenance or improvement of general well-being.,

Relief of the symptoms of allergies. [Warning S required],

Relief of the symptoms of colds. [Warnings S and COLD required],

May assist in the management of upper respiratory tract infections. [Warnings S and COLD required],

May help reduce the severity of the symptoms of colds.]Warnings S and COLD required],

May reduce the severity of colds. [Warning COLD required],

May reduce the duration of colds. [Warning COLD required],

May reduce the severity and duration of colds. [Warning COLD required],

For relief of mucous congestion.[Warnings S and COLD required],

For the symptomatic relief of upper respiratory tract infections. [Warnings S and COLD required]

Specific Indications

CLINICALS IMUNXL by Swisse has been developed by a professor and contains ingredients traditionally used and supported by clinical studiesto assist in immune challenging conditions such as colds or flu. IMUNXL is based on the assessment of traditional use and clinical studies,which showed that certain ingredients in IMUNXL provide relief from the symptoms associated with colds and flu such as a blocked or runny nose, dry cough, and mucous congestion. Used as directed, IMUNXL may deliver positive results relatively quickly. Best used when first symptoms occur or when your body is most vulnerable such as prior to winter or contact with sick people.For better results: IMUNXL may be effective when used on it's own, however we recommend you use this product in conjunction with Swisse Women's or Men's Ultivite, which act as a "foundation" to provide the body with the important nutrients necessary for general wellbeing.

Additional Product Information

Pack Size/Poison Information

Pack Size

Poison Schedule

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	

Active Ingredients

Calcium ascorbate	**605.34 mg**
Citrus bioflavonoids extract	**20 mg**
Cupric sulfate pentahydrate	**3.93 mg**
Magnesium aspartate	**250 mg**
Olea europaea	**55.56 mg**
Equivalent: Olea europaea (Dry)	250 mg
Zinc sulfate monohydrate	**68.61 mg**

Public Summary

Summary for ARTG Entry: 76578 WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT capsule bottle

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11

Products

1. WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet women's nutritional needs and help in the maintenance of improvement of general well-being.

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	2 Years	Store below 30 degrees Celsius	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
120 capsules,	Not scheduled. Not considered by committee,
60 capsules	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	Amber coloured powder in a clear hard gelatin capsule. Size 0 Length 21.4 mm Width 7.64 mm

Active Ingredients

Apium graveolens	**1 mg**
Equivalent: Apium graveolens (Dry)	10 mg
Arctostaphylos uva-ursi	**3.12 mg**
Equivalent: Arctostaphylos uva-ursi (Dry)	12.5 mg
Astragalus membranaceus	**2.5 mg**
Equivalent: Astragalus membranaceus (Dry)	25 mg
Avena sativa	**25 mg**

Equivalent: Avena sativa (Fresh)	250 mg
Bioflavonoids	**20 mg**
Biotin	**25 microgram**
Calcium ascorbate dihydrate	**100 mg**
Calcium citrate	**50 mg**
Calcium pantothenate	**37.5 mg**
Camellia sinensis	**1.67 mg**
Equivalent: Camellia sinensis (Dry)	10 mg
Carica papaya	**5 mg**
Centella asiatica	**1.25 mg**
Equivalent: Centella asiatica (Dry)	5 mg
Cholecalciferol	**2.5 microgram**
Equivalent: Cholecalciferol (Dry)	1 mg
Choline bitartrate	**12.5 mg**
Chromium picolinate	**25 microgram**
Copper gluconate	**208.5 microgram**
Crataegus monogyna	**3.75 mg**
Equivalent: Crataegus monogyna (Dry)	15 mg
Cyanocobalamin	**25 microgram**
Cynara scolymus	**6.25 mg**
Equivalent: Cynara scolymus (Fresh)	25 mg
d-alpha-Tocopheryl acid succinate	**20.66 mg**
Dunaliella salina	**33.33 mg**
Equivalent: Dunaliella salina (Fresh)	60 mg
Eleutherococcus senticosus	**1.25 mg**
Equivalent: Eleutherococcus senticosus (Dry)	12.5 mg
Equisetum arvense	**3.75 mg**
Equivalent: Equisetum arvense (Dry)	15 mg
Ferrous fumarate	**7.55 mg**
Foeniculum vulgare	**1.7 mg**
Equivalent: Foeniculum vulgare (Dry)	7.5 mg
Folic acid	**150 microgram**
Ginkgo biloba	**50 microgram**
Equivalent: Ginkgo biloba (Dry)	2.5 mg
Glycyrrhiza glabra	**1.25 mg**
Equivalent: Glycyrrhiza glabra (Dry)	5 mg
Inositol	**12.5 mg**
Lycopersicon esculentum	**1 mg**
Equivalent: Lycopersicon esculentum (Dry)	20 mg
Lysine hydrochloride	**25 mg**
Magnesium oxide - heavy	**42.5 mg**
Manganese amino acid chelate	**8 mg**
Matricaria recutita	**1.88 mg**
Equivalent: Matricaria recutita (Dry)	7.5 mg
Nicotinamide	**25 mg**
Petroselinum crispum	**5 g**
Potassium iodide	**32.7 microgram**
Potassium sulfate	**2.23 mg**
Pyridoxine hydrochloride	**25 mg**
Riboflavine	**25 mg**
Selenomethionine	**32.5 microgram**
Silybum marianum	**357 microgram**
Equivalent: Silybum marianum (Dry)	25 mg
Spearmint Oil	**750 microgram**
Tagetes erecta	**500 microgram**
Equivalent: Tagetes erecta (Dry)	10 mg
Thiamine hydrochloride	**25 mg**

Ubidecarenone	**500 microgram**
Vaccinium myrtillus	**125 microgram**
Equivalent: Vaccinium myrtillus (Fresh)	12.5 mg
Vitis vinifera	**4.17 mg**
Equivalent: Vitis vinifera (Dry)	500 mg
Zinc amino acid chelate	**12.5 mg**
Zingiber officinale	**1.5 mg**
Equivalent: Zingiber officinale (Dry)	7.5 mg

Public Summary

Summary for ARTG Entry: 79766 SWISSE WOMEN'S ULTIBOOST MINERAL & HERBAL PRE-EVENT FORMULA Tablet - film coated bottle

ARTG entry for	Medicine Listed
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	7/08/2001
Product category	Medicine
Status	Revoked
Approval area	Listed Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

A copy of a signed and dated certificate of analysis, which is not more than six months old, for the first batch of goods manufactured is to be provided to the Head, Listing Treaties & Export Section, Chemical & Non Prescription Drug Branch within six months of the date of supply of the goods.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995.

Products

1. SWISSE WOMEN'S ULTIBOOST MINERAL & HERBAL PRE-EVENT FORMULA

Product Type Single Medicine Product **Effective date** 4/02/2003

Warnings

Contains maltodextrin (or words to that effect).,

If symptoms persist, seek the advice of a healthcare professional.

Standard Indications

May assist in maintaining peripheral circulation and promoting general health.,

Aids or assists in the maintenance of peripheral circulation to relieve cold hands and feet. [Warning S required],

Assisting in the maintenance of blood flow in the hands, feet and legs.,

Aids or assists in the maintenance of peripheral circulation.,

To help maintain blood circulation to the peripheral areas of the body such as the legs, hands and feet.,

Beneficial during times of stress. [Warning S required],

Aids, assists or helps in the maintenance or improvement of general well-being.,

May assist blood circulation.

Specific Indications

Swisse Women's UltiBoost, to boost stamina and memory for peak perform ance, has been designed by a professor to be used as a companion with Swisse Women's Ultivite Formula 1 when an extra boost is required. Sw isse Women's UltiBoost can also prepare you for periods in your life w here you will be demanding more of your body and faculties such as wor king overtime, travelling or giving presentations. It's all about as sisting you to perform in peak condition and enhancing mental alertnes s, so Swisse Women's UlitBoost should be taken before and during physi cal and mental exertion. The difference is that unlike Swisse Women' s Ultivite Formula 1 that is used daily throughout the year for the ma intenance of general wellbeing, you only need to use Swisse Women's Ul tiBoost when you feel you need that `extra boost'.

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	2 Years	Store below 30 degrees Celsius	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size	Poison Schedule
N/A	Not scheduled. Not considered by committee

Components

1. COMPONENT ONE

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Extra large oval yellow mustard film coated tablet Length 23.20 mm Width 10.50 mm

Active Ingredients

Cimicifuga racemosa	**50 mg**
Equivalent: Cimicifuga racemosa (Dry)	200 mg
Cimicifuga racemosa	
Ginkgo biloba	
Ginkgo biloba	**60 mg**
Equivalent: Ginkgo biloba (Dry)	3 g
Glutamine	**250 mg**
Magnesium aspartate	**500 mg**
Panax ginseng	**100 mg**
Equivalent: Panax ginseng (Dry)	1 g
Panax ginseng	
Schizandra chinensis	
Schizandra chinensis	**40 mg**
Equivalent: Schizandra chinensis (Dry)	400 mg

Public Summary

Summary for ARTG Entry: 76577 WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11,

The sponsor shall hold stability data to support the claimed shelf life of the listed medicine according to the labelled storage conditions. The sponsor will provide stability data to the Head, Office of Medicines Authorisation,TGA upon request, except where the overseas importer accepts responsibility for stability studies. When an overseas importer accepts responsibility for providing stability data for this product it is the sponsor's responsibility to ensure that they have a written agreement to this effect from the overseas importer. The sponsor will provide a copy of this agreement to the Head, Office of Medicines Authorisation, TGA upon request.

Products

1. WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet Women's nutritional needs and help in the maintenance or improvement of general well-being.

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Jar/Can	Not recorded	3 Years	Not recorded	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size	Poison Schedule
30 tablets,	Not scheduled. Not considered by committee,
60 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Extra large oval shaped light pink film coated tablet Length 23.2 mm Width 10.5 mm

Active Ingredients

Apium graveolens	**2 mg**
Equivalent: Apium graveolens (Dry)	20 mg
Arctostaphylos uva-ursi	**6.25 mg**
Equivalent: Arctostaphylos uva-ursi (Dry)	25 mg

Astragalus membranaceus	**5 mg**
Equivalent: Astragalus membranaceus (Dry)	50 mg
Avena sativa	**50 mg**
Equivalent: Avena sativa (Fresh)	500 mg
Biotin	**50 microgram**
Calcium ascorbate dihydrate	**200 mg**
Calcium citrate	**200 mg**
Calcium pantothenate	**75 mg**
Camellia sinensis	**3.34 mg**
Equivalent: Camellia sinensis (Dry)	20 mg
Carica papaya	**10 mg**
Centella asiatica	**2.5 mg**
Equivalent: Centella asiatica (Dry)	10 mg
Choline bitartrate	**25 mg**
Chromium picolinate	**50 microgram**
Citrus bioflavonoids extract	**40 mg**
Copper gluconate	**417 microgram**
Crataegus monogyna	**7.5 mg**
Equivalent: Crataegus monogyna (Dry)	30 mg
Cyanocobalamin	**50 microgram**
Cynara scolymus	**12.5 mg**
Equivalent: Cynara scolymus (Fresh)	50 mg
d-alpha-Tocopheryl acid succinate	**41.33 mg**
Eleutherococcus senticosus	**2.5 mg**
Equivalent: Eleutherococcus senticosus (Dry)	25 mg
Equisetum arvense	**7.5 mg**
Equivalent: Equisetum arvense (Dry)	30 mg
Ferrous fumarate	**16.01 mg**
Foeniculum vulgare	**3.41 mg**
Equivalent: Foeniculum vulgare (Dry)	15 mg
Folic acid	**300 microgram**
Ginkgo biloba	**100 microgram**
Equivalent: Ginkgo biloba (Dry)	5 mg
Glycyrrhiza glabra	**2.5 mg**
Equivalent: Glycyrrhiza glabra (Dry)	10 mg
Inositol	**25 mg**
Lysine hydrochloride	**50 mg**
Magnesium oxide - heavy	**81.46 mg**
Manganese amino acid chelate	**16 mg**
Matricaria recutita	**3.75 mg**
Equivalent: Matricaria recutita (Dry)	15 mg
Nicotinamide	**50 mg**
Petroselinum crispum	**10 mg**
Potassium iodide	**66 microgram**
Potassium sulfate	**4.45 mg**
Pyridoxine hydrochloride	**50 mg**
Riboflavine	**50 mg**
Selenomethionine	**65 microgram**
Spearmint Oil	**1.5 mg**
Thiamine hydrochloride	**50 mg**
Ubidecarenone	**1 mg**
Zinc amino acid chelate	**25 mg**
Zingiber officinale	**3 mg**
Equivalent: Zingiber officinale (Dry)	15 mg

Public Summary

Public Summary

Summary for ARTG Entry: 76576 WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS NO IRON OR IODINE DIETARY SUPPLEMENT tablet jar

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11

Products

1. WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS NO IRON OR IODINE DIETARY SUPPL

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet nutritional needs of women who have been diagnosed with iron overload (hemochromatosis), or should not take iodine.

Additional Product Information

Container Information

Type	Material	Life Time	Temperature	Closure	Conditions
Jar/Can	Not recorded	2 Years	Store below 30 degrees Celsius	Not recorded	Not recorded

Pack Size/Poison Information

Pack Size	Poison Schedule
60 tablets	Not scheduled. Not considered by committee,
30 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Extra large oval white film coated tablet. Length 23.2 mm Width 10.5 mm

Active Ingredients

Apium graveolens	**2 mg**
Equivalent: Apium graveolens (Dry)	20 mg
Arctostaphylos uva-ursi	**6.25 mg**
Equivalent: Arctostaphylos uva-ursi (Dry)	25 mg
Astragalus membranaceus	**5 mg**
Equivalent: Astragalus membranaceus (Dry)	50 mg
Avena sativa	**50 mg**
Equivalent: Avena sativa (Fresh)	500 mg

Bioflavonoids	**40 mg**
Biotin	**50 microgram**
Calcium ascorbate dihydrate	**50 mg**
Calcium citrate	**200 mg**
Calcium pantothenate	**75 mg**
Camellia sinensis	**3.34 mg**
Equivalent: Camellia sinensis (Dry)	20 mg
Carica papaya	**5 mg**
Centella asiatica	**2.5 mg**
Equivalent: Centella asiatica (Dry)	10 mg
Cholecalciferol	**5 microgram**
Choline bitartrate	**25 mg**
Chromium picolinate	**50 microgram**
Copper gluconate	**417 microgram**
Crataegus monogyna	**7.5 mg**
Equivalent: Crataegus monogyna (Dry)	30 mg
Cyanocobalamin	**50 microgram**
Cynara scolymus	**12.5 mg**
Equivalent: Cynara scolymus (Fresh)	50 mg
d-alpha-Tocopheryl acid succinate	**37.19 mg**
Dunaliella salina	**66.67 mg**
Equivalent: Dunaliella salina (Fresh)	120 mg
Eleutherococcus senticosus	**2.5 mg**
Equivalent: Eleutherococcus senticosus (Dry)	25 mg
Equisetum arvense	**7.5 mg**
Equivalent: Equisetum arvense (Dry)	30 mg
Foeniculum vulgare	**3.41 mg**
Equivalent: Foeniculum vulgare (Dry)	15 mg
Folic acid	**300 microgram**
Ginkgo biloba	**100 microgram**
Equivalent: Ginkgo biloba (Dry)	5 mg
Glycyrrhiza glabra	**2.5 mg**
Equivalent: Glycyrrhiza glabra (Dry)	10 mg
Inositol	**25 mg**
Lycopersicon esculentum	**2 mg**
Equivalent: Lycopersicon esculentum (Dry)	40 mg
Magnesium oxide - heavy	**92 mg**
Manganese amino acid chelate	**16 mg**
Matricaria recutita	**3.75 mg**
Equivalent: Matricaria recutita (Dry)	15 mg
Nicotinamide	**50 mg**
Petroselinum crispum	**5 mg**
Potassium sulfate	**4.45 mg**
Pyridoxine hydrochloride	**50 mg**
Riboflavine	**50 mg**
Selenomethionine	**65 microgram**
Silybum marianum	**14.29 mg**
Equivalent: Silybum marianum (Dry)	1 g
Spearmint Oil	**1.7 mg**
Tagetes erecta	**1 mg**
Equivalent: Tagetes erecta (Dry)	20 mg
Thiamine hydrochloride	**50 mg**
Tocopherols concentrate - mixed (low-alpha type)	**3.36 mg**
Ubidecarenone	**1 mg**
Vaccinium myrtillus	**250 microgram**
Equivalent: Vaccinium myrtillus (Fresh)	25 mg
Vitis vinifera	**8.34 mg**

Equivalent: Vitis vinifera (Dry)	1 g
Zinc amino acid chelate	**25 mg**
Zingiber officinale	**3 mg**
Equivalent: Zingiber officinale (Dry)	15 mg

Public Summary

Summary for ARTG Entry: 76575 WOMEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS NO IRON OR IODINE DIETARY SUPPLEMENT capsule bottle

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11

Products

1. WOMEN'S ULTIVITE MULT-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS NO IRON OR IODINE DIETARY SUPPL

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet nutritional needs of women who have been diagnosed with Iron overload (hemochromatosis) or should not take Iodine.

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	2 Years	Store below 30 degrees Celsius	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
120 capsules,	Not scheduled. Not considered by committee,
60 capsules	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	Amber coloured powder in a clear hard gelatin capsule Size 0 Length 21.4 mm Width 7.64 mm

Active Ingredients

Apium graveolens	**1 mg**
Equivalent: Apium graveolens (Dry)	10 mg
Arctostaphylos uva-ursi	**3.12 mg**
Equivalent: Arctostaphylos uva-ursi (Dry)	12.5 mg
Astragalus membranaceus	**2.5 mg**
Equivalent: Astragalus membranaceus (Dry)	25 mg
Avena sativa	**25 mg**

Equivalent: Avena sativa (Fresh)	250 mg
Bioflavonoids	**20 mg**
Biotin	**25 microgram**
Calcium ascorbate dihydrate	**25 mg**
Calcium carbonate	**55.55 mg**
Calcium pantothenate	**37.5 mg**
Camellia sinensis	**1.67 mg**
Equivalent: Camellia sinensis (Dry)	10 mg
Carica papaya	**2.5 mg**
Centella asiatica	**1.25 mg**
Equivalent: Centella asiatica (Dry)	5 mg
Cholecalciferol	**2.5 microgram**
Choline bitartrate	**12.5 mg**
Chromium picolinate	**25 microgram**
Copper gluconate	**209 microgram**
Crataegus monogyna	**3.75 mg**
Equivalent: Crataegus monogyna (Dry)	15 mg
Cyanocobalamin	**25 microgram**
Cynara scolymus	**6.25 mg**
Equivalent: Cynara scolymus (Fresh)	25 mg
d-alpha-Tocopheryl acid succinate	**18.6 mg**
Dunaliella salina	**33.33 mg**
Equivalent: Dunaliella salina (Fresh)	60 mg
Eleutherococcus senticosus	**1.25 mg**
Equivalent: Eleutherococcus senticosus (Dry)	12.5 mg
Equisetum arvense	**3.75 mg**
Equivalent: Equisetum arvense (Dry)	15 mg
Foeniculum vulgare	**1.7 mg**
Equivalent: Foeniculum vulgare (Dry)	7.5 mg
Folic acid	**150 microgram**
Ginkgo biloba	**50 microgram**
Equivalent: Ginkgo biloba (Dry)	2.5 mg
Glycyrrhiza glabra	**1.25 mg**
Equivalent: Glycyrrhiza glabra (Dry)	5 mg
Inositol	**12.5 mg**
Lycopersicon esculentum	**1 mg**
Equivalent: Lycopersicon esculentum (Dry)	20 mg
Magnesium oxide - heavy	**46 mg**
Manganese amino acid chelate	**8 mg**
Matricaria recutita	**1.88 mg**
Equivalent: Matricaria recutita (Dry)	7.5 mg
Nicotinamide	**25 mg**
Petroselinum crispum	**2.5 mg**
Potassium sulfate	**2.23 mg**
Pyridoxine hydrochloride	**25 mg**
Riboflavine	**25 mg**
Selenomethionine	**32.5 microgram**
Silybum marianum	**7.14 mg**
Equivalent: Silybum marianum (Dry)	500 mg
Spearmint Oil	**750 microgram**
Tagetes erecta	**500 microgram**
Equivalent: Tagetes erecta (Dry)	10 mg
Thiamine hydrochloride	**25 mg**
Tocopherols concentrate - mixed (low-alpha type)	**1.68 mg**
Ubidecarenone	**500 microgram**
Vaccinium myrtillus	**125 microgram**
Equivalent: Vaccinium myrtillus (Fresh)	12.5 mg

Vitis vinifera	**4.17 mg**
Equivalent: Vitis vinifera (Dry)	500 mg
Zinc amino acid chelate	**12.5 mg**
Zingiber officinale	**1.5 mg**
Equivalent: Zingiber officinale (Dry)	7.5 mg

Public Summary

Summary for ARTG Entry: 76573 MEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11,

The sponsor shall hold stability data to support the claimed shelf life of the listed medicine according to the labelled storage conditions. The sponsor will provide stability data to the Head, Office of Medicines Authorisation,TGA upon request, except where the overseas importer accepts responsibility for stability studies. When an overseas importer accepts responsibility for providing stability data for this product, it is the sponsor's responsibility to ensure that they have a written agreement to this effect from the overseas importer. The sponsor will provide a copy of this agreement to the Head, Office of Medicines Authorisation, TGA upon request.

Products

1. MEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet man's nutritional needs and help in the maintenance or improvement of general well-being.

Additional Product Information

Pack Size/Poison Information

Pack Size	Poison Schedule
60 tablets	Not scheduled. Not considered by committee,
30 tablets	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Tablet, film coated
Route of Administration	Oral
Visual Identification	Extra large oval green film coated tablet Length 23.2 mm Width 10.5 mm

Active Ingredients

Apium graveolens	**2 mg**
Equivalent: Apium graveolens (Dry)	20 mg
Astragalus membranaceus	**5 mg**
Equivalent: Astragalus membranaceus (Dry)	50 mg
Avena sativa	**50 mg**
Equivalent: Avena sativa (Fresh)	500 mg
Barosma betulina	**2.5 mg**
Equivalent: Barosma betulina (Dry)	10 mg

Berberis vulgaris	**2.5 mg**
Equivalent: Berberis vulgaris (Dry)	15 mg
Biotin	**50 microgram**
Calcium ascorbate dihydrate	**200 mg**
Calcium citrate	**100 mg**
Calcium pantothenate	**70 mg**
Camellia sinensis	**3.34 mg**
Equivalent: Camellia sinensis (Dry)	20 mg
Carica papaya	**10 mg**
Centella asiatica	**12.5 mg**
Equivalent: Centella asiatica (Dry)	50 mg
Choline bitartrate	**25 mg**
Chromium picolinate	**50 microgram**
Citrus bioflavonoids extract	**40 mg**
Copper gluconate	**200 microgram**
Crataegus monogyna	**25 mg**
Equivalent: Crataegus monogyna (Dry)	100 mg
Cyanocobalamin	**30 microgram**
Cynara scolymus	**12.5 mg**
Equivalent: Cynara scolymus (Fresh)	50 mg
Equisetum arvense	**7.5 mg**
Equivalent: Equisetum arvense (Dry)	30 mg
Ferrous fumarate	**9.13 mg**
Foeniculum vulgare	**3.41 mg**
Equivalent: Foeniculum vulgare (Dry)	15 mg
Folic acid	**300 microgram**
Ginkgo biloba	**2 mg**
Equivalent: Ginkgo biloba (Dry)	100 mg
Inositol	**25 mg**
Lysine hydrochloride	**50 mg**
Magnesium oxide - heavy	**100 mg**
Manganese amino acid chelate	**12 mg**
Nicotinamide	**30 mg**
Panax ginseng	**16.67 mg**
Equivalent: Panax ginseng (Dry)	50 mg
Petroselinum crispum	**10 mg**
Potassium iodide	**66 microgram**
Potassium sulfate	**8.92 mg**
Pyridoxine hydrochloride	**30 mg**
Riboflavine	**30 mg**
Selenomethionine	**65 microgram**
Serenoa repens	**20 mg**
Equivalent: Serenoa repens (Dry)	200 mg
Smilax officinalis	**12.5 mg**
Equivalent: Smilax officinalis (Dry)	50 mg
Spearmint Oil	**1.5 mg**
Thiamine hydrochloride	**30 mg**
Turnera diffusa	**24 mg**
Equivalent: Turnera diffusa (Dry)	120 mg
Tyrosine	**1 mg**
Ubidecarenone	**1 mg**
Zinc amino acid chelate	**30 mg**
Zingiber officinale	**1 mg**
Equivalent: Zingiber officinale (Dry)	5 mg

Public Summary

Public Summary

Summary for ARTG Entry: 76574 MEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA 1 DIETARY SUPPLEMENT capsule bottle

ARTG entry for	Medicine Listed (Export Only)
Sponsor	Swisse Wellness Pty Ltd
Postal Address	36-38 Gipps Street, COLLINGWOOD, VIC, 3066 Australia
ARTG Start Date	19/10/2000
Product category	Medicine
Status	Revoked
Approval area	Export only Medicines

Conditions

Except where the sponsor has been contracted by an overseas party to manufacture the goods and that party will be responsible for placing the goods on the market in countries other than Australia, the sponsor must have and shall retain, while the goods remain listed, evidence necessary to substantiate and support the accuracy of the indications in relation to the listed goods and, upon the request of the Head, Office of Medicines Authorisation, Therapeutic Goods Administration, shall produce such evidence to the Director.,

The conditions applying to these goods when they are exported from Australia are given below:,

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No. 8 and condition No. 9.,

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 July 1995, other than condition No.11

Products

1. MEN'S ULTIVITE MULTI-VITAMIN MINERAL & ANTI- OXIDANT WITH HERBS FORMULA DIETARY SUPPLEMENT

Product Type Single Medicine Product **Effective date** 13/11/2008

Warnings

No Warnings included on Record

Standard Indications

Specific Indications

Designed to meet men's nutritional needs and help in the maintenance or improvement of general well-being.

Additional Product Information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Bottle	Not recorded	2 Years	Store below 30 degrees Celsius	Not recorded	Not recorded

Pack Size/Poison information

Pack Size	Poison Schedule
120 capsules,	Not scheduled. Not considered by committee,
60 capsules	Not scheduled. Not considered by committee

Components

1. Medicine Component

Dosage Form	Capsule, hard
Route of Administration	Oral
Visual Identification	Amber coloured powder in a clear hard gelatin capsule Size 0 Length 21.4 mm Width 7.64 mm

Active Ingredients

Apium graveolens	**1 mg**
Equivalent: Apium graveolens (Dry)	10 mg
Astragalus membranaceus	**2.5 mg**
Equivalent: Astragalus membranaceus (Dry)	.25 mg
Avena sativa	**25 mg**
Equivalent: Avena sativa (Fresh)	250 mg
Barosma betulina	**1.25 mg**

Equivalent: Barosma betulina (Dry)	5 mg
Berberis vulgaris	**1.25 mg**
Equivalent: Berberis vulgaris (Dry)	7.5 mg
Bioflavonoids	**20 mg**
Biotin	**25 microgram**
Calcium ascorbate dihydrate	**100 mg**
Calcium citrate	**50 mg**
Calcium pantothenate	**35 mg**
Camellia sinensis	**1.67 mg**
Equivalent: Camellia sinensis (Dry)	10 mg
Carica papaya	**5 mg**
Centella asiatica	**6.25 mg**
Equivalent: Centella asiatica (Dry)	25 mg
Cholecalciferol	**2.5 microgram**
Choline bitartrate	**12.5 mg**
Chromium picolinate	**25 microgram**
Copper gluconate	**100 microgram**
Crataegus monogyna	**12.5 mg**
Equivalent: Crataegus monogyna (Dry)	50 mg
Cyanocobalamin	**15 microgram**
Cynara scolymus	**6.25 mg**
Equivalent: Cynara scolymus (Fresh)	25 mg
d-alpha-Tocopheryl acid succinate	**20.66 mg**
Dunaliella salina	**33.33 mg**
Equivalent: Dunaliella salina (Fresh)	60 mg
Equisetum arvense	**3.75 mg**
Equivalent: Equisetum arvense (Dry)	15 mg
Ferrous fumarate	**4.56 mg**
Foeniculum vulgare	**1.7 mg**
Equivalent: Foeniculum vulgare (Dry)	7.5 mg
Folic acid	**150 microgram**
Ginkgo biloba	**1 mg**
Equivalent: Ginkgo biloba (Dry)	50 mg
Inositol	**12.5 mg**
Lycopersicon esculentum	**1 mg**
Equivalent: Lycopersicon esculentum (Dry)	20 mg
Lysine hydrochloride	**25 mg**
Magnesium oxide - heavy	**50 mg**
Manganese amino acid chelate	**6 mg**
Nicotinamide	**15 mg**
Panax ginseng	**8.33 mg**
Equivalent: Panax ginseng (Dry)	25 mg
Petroselinum crispum	**5 mg**
Potassium iodide	**33 microgram**
Potassium sulfate	**4.46 mg**
Pyridoxine hydrochloride	**15 mg**
Riboflavine	**15 mg**
Selenomethionine	**32.5 microgram**
Serenoa repens	**10 mg**
Equivalent: Serenoa repens (Dry)	100 mg
Silybum marianum	**357 microgram**
Equivalent: Silybum marianum (Dry)	25 mg
Smilax officinalis	**6.25 mg**
Equivalent: Smilax officinalis (Dry)	25 mg
Spearmint Oil	**750 microgram**
Tagetes erecta	**500 microgram**
Equivalent: Tagetes erecta (Dry)	10 mg

Thiamine hydrochloride	**15 mg**
Turnera diffusa	**12 mg**
Equivalent: Turnera diffusa (Dry)	60 mg
Tyrosine	**500 microgram**
Ubidecarenone	**500 microgram**
Vaccinium myrtillus	**125 microgram**
Equivalent: Vaccinium myrtillus (Fresh)	12.5 mg
Vitis vinifera	**4.17 mg**
Equivalent: Vitis vinifera (Dry)	500 mg
Zinc amino acid chelate	**15 mg**
Zingiber officinale	**500 microgram**
Equivalent: Zingiber officinale (Dry)	2.5 mg