



Australian Government
Department of Health
Therapeutic Goods Administration

Public Summary

| | | |
|--------------------------------|--|--|
| Summary for ARTG Entry: | 20263 | BLACKMORES DUO CELLOIDS P.S.P.C. Tablets |
| ARTG entry for | Medicine Registered | |
| Sponsor | FIT-BioCeuticals Limited | |
| Postal Address | Care of Blackmores Ltd PO Box 1725, Warriewood, NSW, 2102 Australia | |
| ARTG Start Date | 14/10/1991 | |
| Product Category | Medicine | |
| Status | Active | |
| Approval Area | Registered Complementary Medicines | |

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"

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 . BLACKMORES DUO CELLOIDS P.S.P.C. Tablets

| | | | |
|---------------------|-------------------------|-----------------------|------------|
| Product Type | Single Medicine Product | Effective Date | 11/05/2021 |
|---------------------|-------------------------|-----------------------|------------|

Permitted Indications

No Permitted Indications included on Record

Indication Requirements

No Indication Requirements included on Record

Standard Indications

No Standard Indications included on Record

Specific Indications

Mineral supplement.

Warnings

No Warnings included on Record

Additional Product information

Container information

| Type | Material | Life Time | Temperature | Closure | Conditions |
|--------|--------------|-----------|--------------------------------|--------------|--|
| Bottle | Not recorded | 4 Years | Store below 30 degrees Celsius | Not recorded | Store in a Dry Place Protect from Light |

Pack Size/Poison information

| | |
|------------------|--|
| Pack Size | Poison Schedule |
| 80 | Not Scheduled after consideration by Committee |

Components

1 .

| | |
|--------------------------------|---|
| Dosage Form | Tablet, uncoated |
| Route of Administration | Oral |
| Visual Identification | A round white biconvex tablet embossed with 'b' and breakbar. |

Active Ingredients

| | |
|--------------------|-------|
| potassium chloride | 65 mg |
|--------------------|-------|

Public Summary



Australian Government
Department of Health
Therapeutic Goods Administration

| | |
|--------------------------|--------------|
| Equivalent: potassium | 34.09 mg |
| potassium sulfate | 33 mg |
| Equivalent: potassium | 14.81 mg |

Other Ingredients (Excipients)

Acacia

lactose monohydrate

magnesium stearate

maize starch

microcrystalline cellulose

soy polysaccharide

© Commonwealth of Australia. This work is copyright. You are not permitted to re-transmit, distribute or commercialise the material without obtaining prior written approval from the Commonwealth. Further details can be found at <http://www.tga.gov.au/about/website-copyright.htm>.

Public Summary