



**Australian Government**  
**Department of Health**  
 Therapeutic Goods Administration

**Public Summary**

|                                |  |   |
|--------------------------------|--|---|
| <b>Summary for ARTG Entry:</b> | 20229  | BLACKMORES DUO CELLOID P.P.M.P. Tablets |
| <b>ARTG entry for</b>          | Medicine Registered  |   |
| <b>Sponsor</b>                 | FIT-BioCeuticals Limited   |   |
| <b>Postal Address</b>          | Care of Blackmores Ltd PO Box 1725, Warriewood, NSW, 2102<br>Australia |   |
| <b>ARTG Start Date</b>         | 14/10/1991   |   |
| <b>Product Category</b>        | Medicine   |   |
| <b>Status</b>                  | Active   |   |
| <b>Approval Area</b>           | 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 CELLOID P.P.M.P. Tablets**

|                     |                         |                       |            |
|---------------------|-------------------------|-----------------------|------------|
| <b>Product Type</b> | Single Medicine Product | <b>Effective Date</b> | 28/04/2021 |
|---------------------|-------------------------|-----------------------|------------|

**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<br>Protect from Light |

**Pack Size/Poison information**

| Pack Size   | Poison Schedule                                |
|-------------|--|
| 170 tablets | Not Scheduled after consideration by Committee |
| 84 tablets  | Not Scheduled after consideration by Committee |

**Components**

1 .

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

**Active Ingredients**

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|   |              |
|---|--------------|
| <b>dibasic potassium phosphate</b>      | <b>33 mg</b> |
| Equivalent: potassium                   | 14.8 mg      |
| <b>magnesium phosphate pentahydrate</b> | <b>65 mg</b> |
| Equivalent: magnesium                   | 13.4 mg      |

**Other Ingredients (Excipients)**

- Acacia
- lactose monohydrate
- magnesium stearate
- maize starch
- microcrystalline cellulose
- soy polysaccharide

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