



**Australian Government**  
**Department of Health and Aged Care**  
 Therapeutic Goods Administration

**Public Summary**

<b>Summary for ARTG Entry:</b>	299066	ImmuneForce
<b>ARTG entry for</b>	Medicine Listed	
<b>Sponsor</b>	FIT-BioCeuticals Limited	
<b>Postal Address</b>	Care of Blackmores Ltd PO Box 1725, Warriewood, NSW, 2102 Australia	
<b>ARTG Start Date</b>	29/01/2018	
<b>Product Category</b>	Medicine	
<b>Status</b>	Active	
<b>Approval Area</b>	Listed Medicines	

**Conditions**

Colouring agents used in listed medicine for ingestion, other than those listed for export only under section 25 of the Act, shall be only those included in the list of 'Colourings permitted in medicines for oral use'.

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 Complementary Medicines Branch, Therapeutic Goods Administration, upon request.

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 . ImmuneForce**

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	24/11/2020
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**Permitted Indications**

Maintain/support immune system health  
 Maintain/support healthy immune system function

**Indication Requirements**

Product presentation must not imply or refer to serious immunological diseases.

**Standard Indications**

No Standard Indications included on Record

**Specific Indications**

No Specific Indications included on Record

**Warnings**

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

**Additional Product information**

**Pack Size/Poison information**

<b>Pack Size</b>	<b>Poison Schedule</b>
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**Components**

**1 . Formulation 1**

<b>Dosage Form</b>	Tablet, film coated
<b>Route of Administration</b>	Oral

**Visual Identification**

**Active Ingredients**

<b>Andrographis paniculata</b>	<b>142.86 mg</b>
<b>Echinacea purpurea root Extract dry concentrate standardised</b>	<b>166.67 mg</b>
Equivalent: Echinacea purpurea (Dry)	

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	1 g
<b>hydrastis canadensis root Extract dry concentrate</b>	<b>125 mg</b>
Equivalent: hydrastis canadensis (Dry)	750 mg
<b>Inula helenium root Extract dry concentrate</b>	<b>100 mg</b>
Equivalent: Inula helenium (Dry)	1 g
<b>Thymus vulgaris leaf Extract dry concentrate</b>	<b>214.29 mg</b>
Equivalent: Thymus vulgaris (Dry)	1.5 g
<b>zinc amino acid chelate</b>	<b>62.5 mg</b>
Equivalent: zinc	12.5 mg

**Other Ingredients (Excipients)**

calcium carbonate  
calcium hydrogen phosphate dihydrate  
Carnauba Wax  
chlorophyllin-copper complex  
colloidal anhydrous silica  
croscarmellose sodium  
crospovidone  
ethylcellulose  
hypromellose  
macrogol 400  
macrogol 8000  
magnesium stearate  
maltodextrin  
microcrystalline cellulose  
silicon dioxide

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