



Australian Government
Department of Health
 Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: 213033 CAPECITABINE AN capecitabine 150 mg film-coated tablet blister pack

ARTG entry for Medicine Registered
Sponsor Juno Pharmaceuticals Pty Ltd
Postal Address Level 2 6 Bond Street, South Yarra, VIC, 3141
 Australia
ARTG Start Date 22/07/2014
Product category Medicine
Status Active
Approval area Drug Safety Evaluation Branch

Conditions

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. CAPECITABINE AN capecitabine 150 mg film-coated tablet blister pack

Product Type	Single Medicine Product	Effective date	15/06/2017
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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

Colon Cancer:Capecitabine is indicated for the adjuvant treatment of patients with Duke/Es stage C and highrisk stage B, colon cancer, either as monotherapy or in combination with oxaliplatin.,Colorectal Cancer:Capecitabine is indicated for the treatment of patients with advanced or metastatic colorectal cancer.,Oesophagogastric Cancer:Capecitabine is indicated for the first-line treatment of patients with advanced oesophagogastric cancer in combination with a platinum-based regimen.,Breast Cancer:Capecitabine is indicated for the treatment of patients with locally advanced or metastatic breast cancer after failure of taxanes and an anthracycline containing chemotherapy regimen unless therapy with these and other standard agents are clinically contraindicated.,Capecitabine in combination with docetaxel is indicated for the treatment of patients with locally advanced or metastatic breast cancer after failure of prior anthracycline containing chemotherapy.

Warnings

See Product Information and Consumer Medicine Information for this product

Additional Product information

Container information

Type	Material	Life Time	Temperature	Closure	Conditions
Blister Pack	PVC/PVDC/Al	3 Years	Store below 25 degrees Celsius	Not recorded	Store in a Dry Place
Blister Pack	Al/Al	3 Years	Store below 25 degrees Celsius	Not recorded	Store in a Dry Place

Pack Size/Poison information

Pack Size	Poison Schedule
30 Tablets in blister pack	(S4) Prescription Only Medicine
120 Tablets in blister pack	(S4) Prescription Only Medicine
60 Tablets in blister pack	(S4) Prescription Only Medicine

Components

1. CAPECITABINE AN capecitabine 150 mg film-coated tablet blister pack

Dosage Form Tablet, film coated
Route of Administration Oral
Visual Identification Light peach coloured, oblong shaped, biconvex, film-coated tablets,

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debossed with 150 on one side and plain on other side.

Active Ingredients

Capecitabine

150 mg

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