# Advance Pain & Fever



**Description**Each tablet contains 500mg Paracetamol Ph.Eur.

## Therapeutic Indication

Introduction:
Paracetamol is an analgesic and antipyretic. Its mechanism of action is believed to include inhibition of prostaglandin synthesis, primarily within the central nervous system.

## Indication:

- Treatment of mild-to-moderate pain and treatment of fever including:

   Headache
   Migraine
- Muscle ache

- DysmenorrheaSore throatMusculoskeletal pain
- Fever and pain associated with vaccination/immunisation
  Pain after dental procedures/tooth extraction
- Toothache

- Osteoarthritis pain
   Carache/Otalgia
   Respiratory tract infections including cold and flu

## **Excipients**

Tablet core:
Pregelatinised starch, calcium carbonate, alginic acid, crospovidone, povidone (K-25), magnesium stearate, colloidal anhydrous silica, parahydroxybenzoates (sodium methyl parahydroxybenzoate (E219), sodium ethyl parahydroxybenzoate (E215) and sodium propyl parahydroxybenzoate (E217).

# **Dosage and Administration**

Oral administration only

Adults (including the elderly) and children aged 12 years and over:

500mg to 1000mg paracetamol (1 to 2 tablets), taken every 4 to 6 hours as required. Maximum daily dose: 4000mg.

Do not exceed the stated dose. Minimum dosing interval: 4 hours. The lowest dose necessary to achieve efficacy should be used for the shortest duration of treatment.

# Children, 6 to 11 years:

6-8 years: 250mg (½ tablet), 9-11 years: 500mg (1 tablet) every 4 to 6 hours as required, maximum daily dose, 60mg/kg presented in divided doses of 10 - 15 mg/kg throughout the 24 hour period. Do not exceed the stated dose. No more than four doses in any 24 hour period. Minimum dosing interval, 4 hours. Maximum duration of continued use without medical advice, 3 days. Not recommended for children under the age of 6 years. If symptoms persist, medical advise must be sought.

## Contraindications

Paracetamol is contraindicated in patients with a previous history of hypersensitivity to paracetamol or excipients

## Warnings and Precaution for use:

- Contains paracetamol.
  Do not take more than the recommended dose as it may cause serious harm to your liver.
  Do not use this medicine if you are taking any other prescription or non-prescription medicines containing paracetamol to treat pain, fever, symptoms of cold and flu, or to aid sleep.

# Check with your doctor before use if you:

- have liver or kidney problems.
  are underweight or malnourished.
  regularly drink alcohol.

You may need to avoid using this product altogether or limit the amount of paracetamol that you take if you have a severe infection, are severely malnourished, severely underweight, or are a chronic heavy alcohol user as these may increase the risk of metabolic acidosis.

# Signs of metabolic acidosis include:

- deep, rapid, difficult breathing
   feeling sick (nausea), being sick (vomiting)
   loss of appetite

# Panadol

# Advance Pain & Fever



Contact a doctor immediately if you get a combination of these symptoms

Sodium methyl-, sodium ethyl- and sodium propyl- parahydroxybenzoates (E219, E215, E217) may cause allergic reactions (possibly delayed).

# **Pregnancy and Lactation**

Talk to your health care professional before taking paracetamol if you are pregnant.

Consider taking the lowest effective dose for the shortest period of time. You can take this product whilst breastfeeding.

Before taking this medicine, make sure you consult your doctor if you are taking warfarin or similar medicines used to thin the blood.

## Side Effects/Adverse Reactions:

When taken at recommended doses, paracetamol is usually free from side effects and is unlikely to cause an effect on ability to drive and operate machines. Stop taking this medicine and tell your doctor immediately if:

• You experience allergic reactions such as skin rash or itching, sometimes with breathing problems or swelling

- You experience a skin rash or peeling, or mouth ulcers.
  You have previously experienced breathing problems with aspirin or non-steroidal anti-inflammatories, and experience a similar reaction with this product.
- You experience unexplained bruising or bleeding.

- Paracetamol overdose may cause liver failure which may require liver transplant or lead to death. Acute pancreatitis has been observed, usually with hepatic dysfunction and liver toxicity
- Seek medical advice immediately even if you do not have any symptoms because of the risk of liver failure.

Immediate medical management is required in the event of overdose, even if symptoms of overdose are not present. Administration of N-acetylcysteine or methionine may be required.

Contact the nearest Emergency Medical Centre for management and expert treatment.

# PANADOL ADVANCE is a trademark of the GlaxoSmithKline group of companies.

# THIS IS MEDICINE

- Medicines are products which affect your health, failure to follow the instruction may be harmful for you.
- Follow your doctor's advice carefully, the method of use and the instructions of the pharmacist who sold the
- Your doctor and the pharmacist are experts in medicines and their benefits and risks.
  Do not stop your course of treatment early unless advised to do so by your doctor or pharmacist.

# KEEP MEDICINE OUT OF REACH OF CHILDREN



Manufactured by GlaxoSmithKline Knockbrack, Dungaryan, Co. Waterford Ireland Packed and Released by GlaxoSmithKline Limited, Likoni Road, Industrial Area. PO. Box 78392-00507, Viwandani, Nairobi, Kenya

ke.customer-relations@gsk.com

www.gsk.com

GSK is committed to the effective collection and management of human safety information relating to our products and we encourage healthcare professionals to report adverse events to us

Full Prescribing Information is available on request from GlaxoSmithKline Limited.

Revised in May 2019

# PANADOL ADANCE LEAFLET-FRONT-REVISED

# PANADOL ADANCE LEAFLET-BACK-REVISED

Artwork Version : 2	Artwork Information Panel		
Manufacturing Site Number: XXX	Manufacturing Site(s): XXX	Technical Drawing : XXX	
Product Market Trade Name: Panadol	Approving Market(s):	Material Spec: XXX	
Print Process:	Colour Standard Reference: XXX	Material Type: XXX	XXXX
Total Colours & Varnishes: X  BLACK		Body Text Size: 7.5 pt	Smallest Text Size: 6.5 pt
		Leading: 7.8 pt	
Total Special Finishes: X		Horizontal Scale: 100%	
		Microtext: XXX	
Superseded Artwork Code: XXX	Project Name: Panadol	Additional Info (3): XXX	
Artwork copyright is the property of the GSK Group of Companies. All suppliers providing a service to GSK for printed components of any description must ensure that they have a licence for all fonts / software used in conjunction with GSK artwork. The distribution and use of fonts / software without a licence constitutes an intellectual property infringement. GSK will not accept any liability for the breach of third party intellectual property rights by printed component suppliers. The GSK certification / audit process requires suppliers to declare that they do not use unlicensed fonts / software and may require the supplier to produce evidence of such licence to GSK.			
1			i l

SCALE: 1:1



CLIENT: GlaxoSmithKline Limited, Kenya JOB TITLE: Advance NA NA NA NA Oral

FILE NAME:XXX-Panadol-Panadol-Advance-Leaflet-NA\_NANA\_NA-IND-20024006-PROJECT CO-ORDINATOR: Praiakta D DIGITAL MECH. SPECIALIST: jatin patil

SOFTWARE USED:

■ ILLUSTRATOR: CS6/CC ■ PHOTOSHOP: CS6/CC JOB NO: IND-20024006-1-V2+

This Artwork is Read and QC Approved CLIENT SAP NO\MATERIAL CODE:

MPORTANT NOTES: DIADEIS HAS TAKEN ALL PRUDENT EFFORTS TO ENSURE THAT THE CONTENT AND COLOURS ARE AS PER CLIENTS' NEEDS. EVEN THOUGH THIS INPUT HAS BEEN CHECKED BY DIADEIS FOR ACCURACY. IT IS THE RESPONSIBILITY OF THE CLIENT TO MAKE ALL FINAL APPROVALS. HOWEVER, AS AND BY WAY OF ABUNDANT CAUTION, THE CYLINDER MAKERS/PROCESSORS/PRINTERS AND/OR WHOSOEVER USING THE DIADEIS INPUTS SHOULD ALSO, BY THE CHIENTS. DIADEIS IN THE SHOULD ALSO, BY THE FINAL APPROVAL BY THE CLIENTS. DIADEIS IS NOT RESPONSIBLE AND CAN NOT BE HELD LIABLE, IF FOR ANY REASON WHATSOEVER, THE FINAL INPUTS ARE NOT THE SAME AS THAT APPROVED BY THE CLIENTS COLOUR SWATCHES/NUMBERS PROVIDED ABOVE ARE FOR REFERENCE AND NOT TO BE MATCHED WITH THE PAPER PRINTOUT. PRINTERS ARE ADVISED TO TAKE APPROVAL OF THE COLOURS BEFORE PRINT/COMMERCIAL RUN.

DIADEIS, 1st Floor, Om Sadan, Mehra Estate, LBS Marg, Vikhroli (W), Mumbai 400 079, India. T. 00 91 22 4200 8585. F. 00 91 22 4200 8688. Email: customise@diadeis.in, Website: www.diadeis.com

Attention

KLD and Dimensions are only for reference and are not to be printed.