

Please read through the information on the outside and inside of this pack before you start using this medicine.

If you have any questions, or if there is anything you do not understand, ask your pharmacist.

WHAT ARE THEY FOR?

Panadol Extra Advance Tablets provide extra effective pain relief:

- headache
- migraine
- helps reduce temperature
- feverishness
- toothache
- sore throat
- aches and pains of colds and flu
- backache
- period pain
- rheumatic pain

TAKE SPECIAL CARE

Ask your doctor before you take this medicine:

- if you have **liver or kidney disease**, including **alcoholic liver disease**.
- if you have a **severe infection**. (See Section 1, inside for more information.)

HOW TO TAKE

For oral use.

Adults and children aged 16 years and over:

Swallow 2 tablets every 4 hours as needed. Do not take more than 8 tablets in 24 hours.

Children aged 12-15 years:

Swallow 1 tablet every 4 hours as needed. Do not take more than 4 tablets in 24 hours.

Do not take more frequently than every 4 hours. Do not give to children under 12 years.

Do not take more medicine than the label tells you to. If you do not get better talk to your doctor.

Avoid too much caffeine in drinks like coffee and tea. High caffeine intake can cause **difficulty sleeping, shaking and an uncomfortable feeling in the chest.**

WARNING: Talk to a doctor at once if you take too much of this medicine, even if you feel well. This is because too much paracetamol can cause delayed, serious liver damage.

Contains paracetamol. Do not take anything else containing paracetamol while taking this medicine. If your headaches become persistent, see your doctor.

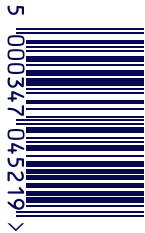
INGREDIENTS

Each tablet contains the active ingredients Paracetamol 500 mg and Caffeine 65 mg. Also includes a mixture of sodium methyl (E 219), sodium ethyl (E 215) and sodium propyl (E 217) parahydroxybenzoates.

Store below 25°C. **Keep out of the sight and reach of children.**

The marketing authorisation holder is GlaxoSmithKline Consumer Healthcare (UK) Trading Limited, Brentford, TW8 9GS, U.K. and all enquiries should be sent to this address. PL 44673/0078

NPA001934



Panadol

Extra Advance 500mg/65mg Tablets



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BN: DO NOT USE AFTER THE EXP. DATE:



14

gsk

Panadol

Extra Advance

500mg/65mg Tablets
Paracetamol + Caffeine



Extra Effective
Pain Relief

Fast Absorption

7.0

5.3

7.3

159.0

1. CHECK BEFORE YOU TAKE PANADOL EXTRA ADVANCE TABLETS



Do not take Panadol Extra Advance Tablets:

- if you have ever had an **allergic reaction** to paracetamol, caffeine or to any of the other ingredients (listed in Section 3).
- 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.

Ask your doctor before you take this medicine if you:

- have **liver or kidney problems** • are **underweight or malnourished**
- regularly **drink alcohol** • have a **severe infection** as this 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

Contact a doctor immediately if you get a combination of these symptoms. You may need to avoid using this product altogether or limit the amount of paracetamol that you take



If you are taking other medicines

Talk to your doctor or pharmacist before taking these tablets if you are taking any prescribed medicines; particularly **metoclopramide or domperidone** (for nausea [feeling sick] or vomiting [being sick]) or **colestyramine** (to lower **blood cholesterol**). If you take **blood thinning drugs** (anticoagulants e.g. **warfarin**) and you need to take a pain reliever on a daily basis, **talk to your doctor** because of the risk of bleeding. But you can still take **occasional** doses of Panadol Extra Advance Tablets at the same time as anticoagulants. This product is not recommended if you are taking **lithium**.



Pregnancy and breast feeding

Due to the caffeine content of this product it **should not be used** if you are **pregnant or breast feeding**.

2. POSSIBLE SIDE EFFECTS

Like all medicines Panadol Extra Advance Tablets can have side effects but not everybody gets them. A small number of people have had side effects.

Very rare cases of serious skin reactions have been reported.

Stop taking the medicine and **tell your doctor immediately** if you experience:

- **Allergic reactions** which may be severe such as skin rash and itching sometimes with swelling of the mouth or face or shortness of breath
- **Skin rash or peeling, or mouth ulcers**
- **Breathing problems**. These are more likely if you have experienced them before when taking other painkillers such as ibuprofen and aspirin
- **Unexplained bruising or bleeding**
- **Nausea, sudden weight loss, loss of appetite and yellowing of the eyes and skin.**

If your symptoms continue or your headache becomes persistent, **see your doctor**.

Reporting of side effects

If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via the Yellow Card Scheme at www.mhra.gov.uk/yellowcard or search for MHRA Yellow Card in the Google Play or Apple App Store. By reporting side effects, you can help provide more information on the safety of this medicine.

3. FURTHER INFORMATION

Active ingredients: Each tablet contains Paracetamol 500 mg and Caffeine 65 mg.

Other ingredients: Starch pregelatinised, povidone k-25, calcium carbonate, crospovidone, alginate acid, magnesium stearate and sodium methyl (E 219), sodium ethyl (E 215) and sodium propyl (E 217) parahydroxybenzoates.

The tablet coating contains titanium dioxide (E 171), hypromellose, macrogol, polysorbate 80 and carnauba wax.

The parahydroxybenzoates may cause allergic reactions (possibly delayed).

Packs of Panadol Extra Advance Tablets contain 8, 14, or 32 tablets.

The manufacturer is GlaxoSmithKline Dungarvan Ltd., Co. Waterford, Ireland. This information was last revised in August 2020.

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