

Generic Name: Rimegepant  
Trade Name: NURTEC  
CDS Effective Date: August 22, 2025  
Supersedes: July 21, 2022  
Approved by BPOM:

**LOCAL PRODUCT DOCUMENT  
PT. PFIZER INDONESIA**

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**1. NAME OF THE MEDICINAL PRODUCT**

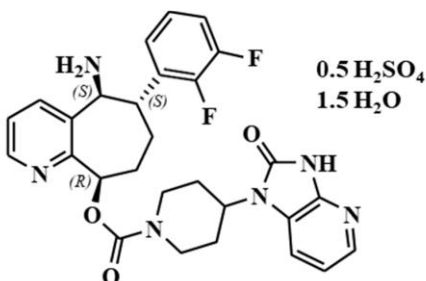
NURTEC 75 mg orally disintegrating tablet

**2. QUALITATIVE AND QUANTITATIVE COMPOSITION**

Each orally disintegrating tablet contains rimegepant sulfate, equivalent to 75 mg rimegepant free base.

For the full list of excipients, see section 6.1.

Structure of rimegepant sulfate:



**3. PHARMACEUTICAL FORM**

Orally disintegrating tablet (ODT). ODT can also be referred to as 'oral lyophilizate'.

The orally disintegrating tablet is white to off-white, circular, and debossed with the symbol

**4. CLINICAL PARTICULARS**

**4.1 Therapeutic indications**

NURTEC is indicated in adults for the:

- acute treatment of migraine with or without aura in adults;

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- preventive treatment of episodic migraine in adults who have at least 4 migraine attacks per month.

## 4.2 Posology and method of administration

### Posology

#### *Acute treatment of migraine*

The recommended dose is 75 mg rimegepant, as needed, once daily.

#### *Prophylaxis of migraine*

The recommended dose is 75 mg rimegepant every other day.

The maximum dose per day is 75 mg rimegepant.

NURTEC can be taken with or without meals.

#### *Concomitant medicinal products*

Another dose of rimegepant should be avoided within 48 hours when it is concomitantly administered with moderate inhibitors of CYP3A4 or with strong inhibitors of P-gp (see section 4.5).

### Special patient populations

#### *Elderly (aged 65 and over)*

There is limited experience with rimegepant in patients aged 65 years or older. No dose adjustment is required as the pharmacokinetics of rimegepant are not affected by age (see section 5.2).

#### *Renal impairment*

No dose adjustment is required in patients with mild, moderate, or severe renal impairment. Severe renal impairment resulted in a > 2-fold increase in unbound AUC but less than a 50% increase in total AUC (see section 5.2). Caution should be exercised during frequent use in patients with severe renal impairment. Rimegepant has not been studied in patients with end-stage renal disease and in patients on dialysis. Use of rimegepant in patients with end-stage renal disease (CL<sub>cr</sub> < 15 ml/min) should be avoided.

#### *Hepatic impairment*

No dose adjustment is required in patients with mild (Child-Pugh A) or moderate (Child-Pugh B) hepatic impairment. Plasma concentrations (unbound AUC) of rimegepant were significantly higher in subjects with severe (Child-Pugh C) hepatic impairment (see section 5.2). The use of rimegepant in patients with severe hepatic impairment should be avoided.

#### *Paediatric population*

The safety and efficacy of Nurtec in paediatric patients (< 18 years of age) have not been established. No data are available.

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### Method of administration

NURTEC is for oral use.

The orally disintegrating tablet should be placed on the tongue or under the tongue. It will disintegrate in the mouth and can be taken without liquid.

Patients should be advised to use dry hands when opening the blister and referred to the package leaflet for complete instructions.

### **4.3 Contraindications**

Hypersensitivity to the active substance or to any of the excipients listed in section 6.1.

### **4.4 Special warnings and precautions for use**

Hypersensitivity reactions, including dyspnoea and rash, have occurred in less than 1% of patients treated with NURTEC in clinical studies (see section 4.8). Hypersensitivity reactions, including serious hypersensitivity, such as anaphylactic reaction, have been reported in the clinical and post-marketing settings (see Section 4.8). Some hypersensitivity reactions can occur days after administration. If a hypersensitivity reaction occurs, rimegepant should be discontinued and appropriate therapy should be initiated (see section 4.3).

NURTEC is not recommended:

- in patients with severe hepatic impairment (see section 4.2);
- in patients with end-stage renal disease ( $CL_{Cr} < 15$  ml/min) (see section 4.2);
- for concomitant use with strong inhibitors of CYP3A4 (see section 4.5);
- for concomitant use with strong or moderate inducers of CYP3A4 (see section 4.5).

#### Medication overuse headache (MOH)

Overuse of any type of medicinal products for headaches can make them worse. If this situation is experienced or suspected, medical advice should be obtained, and treatment should be discontinued. The diagnosis of MOH should be suspected in patients who have frequent or daily headaches despite (or because of) the regular use of medicinal products for acute headache.

### **4.5 Interaction with other medicinal products and other forms of interaction**

Rimegepant is a substrate of CYP3A4, P-glycoprotein (P gp) and breast cancer resistance protein (BCRP) efflux transporters (see section 5.2).

#### CYP3A4 inhibitors

Inhibitors of CYP3A4 increase plasma concentrations of rimegepant. Concomitant administration of rimegepant with strong CYP3A4 inhibitors (e.g., clarithromycin, itraconazole, ritonavir) is not recommended (see section 4.4). Concomitant administration of rimegepant with itraconazole resulted in a significant increase in rimegepant exposure (AUC by 4-fold and  $C_{max}$  1.5-fold).

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Concomitant administration of rimegepant with medicinal products that moderately inhibit CYP3A4 (e.g., diltiazem, erythromycin, fluconazole) may increase exposure to rimegepant. Concomitant administration of rimegepant with fluconazole resulted in increased exposures of rimegepant (AUC by 1.8-fold) with no relevant effect on  $C_{max}$ . Another dose of rimegepant within 48 hours should be avoided when it is concomitantly administered with moderate inhibitors of CYP3A4 (e.g., fluconazole) (see section 4.2).

#### CYP3A4 inducers

Inducers of CYP3A4 decrease plasma concentrations of rimegepant. Concomitant administration of NURTEC with strong CYP3A4 inducers (e.g., phenobarbital, rifampicin, St John's wort (*Hypericum perforatum*)) or moderate CYP3A4 inducers (e.g., bosentan, efavirenz, modafinil) is not recommended (see section 4.4). The effect of CYP3A4 induction may last for up to 2 weeks after discontinuation of the strong or moderate CYP3A4 inducer. Concomitant administration of rimegepant with rifampicin resulted in a significant decrease (AUC reduced by 80% and  $C_{max}$  by 64%) in rimegepant exposure, which may lead to loss of efficacy.

#### P-gp and BCRP only inhibitors

Inhibitors of P-gp and BCRP efflux transporters may increase plasma concentrations of rimegepant. Another dose of NURTEC within 48 hours should be avoided when it is concomitantly administered with strong inhibitors of P-gp (e.g., cyclosporine, verapamil, quinidine) (see section 4.2). Concomitant administration of rimegepant with cyclosporine (a potent P-gp and BCRP inhibitor) or with quinidine (a selective P-gp inhibitor) resulted in a significant increase of similar magnitude in rimegepant exposure (AUC and  $C_{max}$  by > 50%, but less than two-fold).

### **4.6 Fertility, Pregnancy and Lactation**

#### Pregnancy

There are limited data from the use of rimegepant in pregnant women. Animal studies demonstrate that rimegepant is not embryocidal, and no teratogenic potential has been observed at clinically relevant exposures. Adverse effects on embryo-foetal development (decreased foetal body weight and increased skeletal variations in rats) were only observed at exposure levels associated with maternal toxicity (approximately 200 times greater than clinical exposures) following administration of rimegepant during pregnancy (see section 5.3). As a precautionary measure, it is preferable to avoid the use of NURTEC during pregnancy.

#### Lactation

In a single center study of 12 breast-feeding women treated with a single dose of rimegepant 75 mg, minimal concentrations of rimegepant were observed in breast milk. The relative percentage of a maternal dose estimated to reach the infant is less than 1%. There are no data on the effects on milk production. The developmental and health benefits of breast-feeding should be considered along with the mother's clinical need for NURTEC and any potential adverse reactions on the breastfed infant from rimegepant or from the underlying maternal condition.

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## Fertility

Animal studies showed no clinically relevant impact on female and male fertility (see section 5.3).

### **4.7 Effects on ability to drive and use machines**

NURTEC has no or negligible influence on the ability to drive and use machines.

### **4.8 Undesirable effects**

#### Summary of the safety profile

The most common adverse reaction was nausea for acute treatment (1.2%) and for migraine prophylaxis (1.4%). Most of the reactions were mild or moderate in severity. Hypersensitivity, including dyspnoea and severe rash, occurred in less than 1% of patients treated.

#### Tabulated list of adverse reactions

Adverse reactions are listed by MedDRA system organ class in Table 1. The corresponding frequency category for each drug reaction is based on the following convention (CIOMS III): very common ( $\geq 1/10$ ); common ( $\geq 1/100$  to  $< 1/10$ ); uncommon ( $\geq 1/1,000$  to  $< 1/100$ ); rare ( $\geq 1/10,000$  to  $< 1/1,000$ ); very rare ( $< 1/10,000$ ).

**Table 1 Adverse drug reactions**

System Organ Class	Adverse reaction	Frequency
<b>Acute Treatment</b>		
Immune system disorders	Anaphylactic reaction <sup>a</sup> Hypersensitivity, including dyspnoea and severe rash	Uncommon Uncommon
Gastrointestinal disorders	Nausea	Common
<b>Prophylaxis</b>		
Immune system disorders	Anaphylactic reaction <sup>a</sup> Hypersensitivity <sup>a</sup>	Not known Not known
Gastrointestinal disorders	Nausea	Common

<sup>a</sup> Adverse Drug Reactions (ADR) identified post-marketing.

#### *Long-term safety*

Long-term safety of rimegepant was assessed in two one year, open-label extensions; 1662 patients received rimegepant for at least 6 months and 740 received rimegepant for 12 months for acute or prophylactic treatment.

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### Description of selected adverse reactions

#### *Hypersensitivity reactions*

Hypersensitivity, including dyspnoea and severe rash, occurred in less than 1% of patients treated in clinical studies. Hypersensitivity reactions can occur days after administration, and delayed serious hypersensitivity has occurred.

### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via:

Pusat Farmakovigilans/MESO Nasional

Direktorat Pengawasan Keamanan, Mutu, dan Ekspor Impor Obat, Narkotika, Psikotropika,  
Prekursor dan Zat Adiktif

Badan Pengawas Obat dan Makanan

Jl. Percetakan Negara No. 23, Jakarta Pusat, 10560

Email: [pv-center@pom.go.id](mailto:pv-center@pom.go.id)

Phone: +62-21-4244691 Ext.1079

Website: <https://e-meso.pom.go.id/ADR>

PT Pfizer Indonesia

Email: [IDN.AEReporting@pfizer.com](mailto:IDN.AEReporting@pfizer.com)

Website: [www.pfizersafetyreporting.com](http://www.pfizersafetyreporting.com)

## **4.9 Overdose**

There is limited clinical experience with NURTEC overdose. No overdose symptoms have been reported. Treatment of an overdose of NURTEC should consist of general supportive measures including monitoring of vital signs and observation of the clinical status of the patient. No specific antidote for the treatment of rimegepant overdose is available. Rimegepant is unlikely to be significantly removed by dialysis because of high serum protein binding (see section 5.2).

## **5. PHARMACOLOGICAL PROPERTIES**

### **5.1 Pharmacodynamic properties**

Pharmacotherapeutic group: Analgesics, calcitonin gene-related peptide (CGRP) antagonists, ATC code: N02CD06

#### Mechanism of action

Rimegepant selectively binds with high affinity to the human calcitonin gene-related peptide (CGRP) receptor and antagonizes CGRP receptor function.

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The relationship between pharmacodynamic activity and the mechanism(s) by which rimegepant exerts its clinical effects is unknown.

#### Clinical efficacy: acute treatment

The efficacy of NURTEC for the acute treatment of migraine with and without aura in adults was studied in three randomized, double-blind, placebo-controlled trials (Studies 1-3). Patients were instructed to treat a migraine of moderate to severe headache pain intensity. Rescue medicinal products (i.e., NSAIDs, paracetamol, and/or an antiemetic) was allowed 2 hours after the initial treatment. Other forms of rescue medicinal products such as triptans were not allowed within 48 hours of initial treatment. Approximately 14% of patients were taking preventive medicinal products for migraine at baseline. None of the patients in Study 1 were on concomitant preventive medicinal products that act on the calcitonin gene-related peptide pathway.

The primary efficacy analyses were conducted in patients who treated a migraine with moderate to severe pain. Pain freedom was defined as a reduction of moderate or severe headache pain to no headache pain, and most bothersome symptom (MBS) freedom was defined as the absence of the self-identified MBS (i.e., photophobia, phonophobia, or nausea). Among patients who selected an MBS, the most commonly selected symptom was photophobia (54%), followed by nausea (28%), and phonophobia (15%).

In Study 1, the percentage of patients achieving headache pain freedom and MBS freedom at 2 hours after a single dose was statistically significantly greater in patients who received NURTEC compared to those who received placebo (Table 2). In addition, statistically significant effects of NURTEC compared to placebo were demonstrated for the additional efficacy endpoints of pain relief at 2 hours, sustained pain freedom from 2 to 48 hours, use of rescue medication within 24 hours, and ability to function normally at 2 hours after dosing. Pain relief was defined as a reduction in migraine pain from moderate or severe severity to mild or none. Pivotal single attack, double-blind, placebo-controlled studies 2 & 3 were conducted in patients with migraine who received one 75 mg rimegepant bioequivalent dosage form.

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**Table 2: Migraine Efficacy Endpoints for Acute Treatment Studies**

	Study 1		Study 2		Study 3	
	NURTEC 75 mg	Placebo	Rimegepant 75 mg	Placebo	Rimegepant 75 mg	Placebo
<b>Pain Free at 2 hours</b>						
n/N*	142/669	74/682	105/537	64/535	104/543	77/541
% Responders	21.2	10.9	19.6	12.0	19.2	14.2
Difference compared to placebo (%)	10.3		7.6		4.9	
p-value		<0.0001 <sup>a</sup>		0.0006 <sup>a</sup>		0.0298 <sup>a</sup>
<b>MBS Free at 2 hours</b>						
n/N*	235/669	183/682	202/537	135/535	199/543	150/541
% Responders	35.1	26.8	37.6	25.2	36.6	27.7
Difference compared to placebo (%)	8.3		12.4		8.9	
p-value		0.0009 <sup>a</sup>		<0.0001 <sup>a</sup>		0.0016 <sup>a</sup>
<b>Pain Relief at 2 hours</b>						
n/N*	397/669	295/682	312/537	229/535	304/543	247/541
% Responders	59.3	43.3	58.1	42.8	56.0	45.7
Difference compared to placebo	16.1		15.3		10.3	
p-value		<0.0001 <sup>a</sup>		<0.0001 <sup>a</sup>		0.0006 <sup>a</sup>
<b>Sustained Pain Freedom 2 to 48 hours</b>						
n/N*	90/669	37/682	53/537	32/535	63/543	39/541
% Responders	13.5	5.4	9.9	6.0	11.6	7.2
Difference compared to placebo (%)	8.0		3.9		4.4	
p-value		<0.0001 <sup>a</sup>		0.0181 <sup>b</sup>		0.0130 <sup>b</sup>

\*n=number of responders/N=number of patients in that treatment group

<sup>a</sup> Significant p-value in hierarchical testing

<sup>b</sup> Nominal p-value in hierarchical testing

MBS: most bothersome symptom

Figure 1 presents the percentage of patients achieving migraine pain freedom within 2 hours following treatment in Study 1.

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**Figure 1: Percentage of Patients Achieving Pain Freedom within 2 Hours in Study 1**

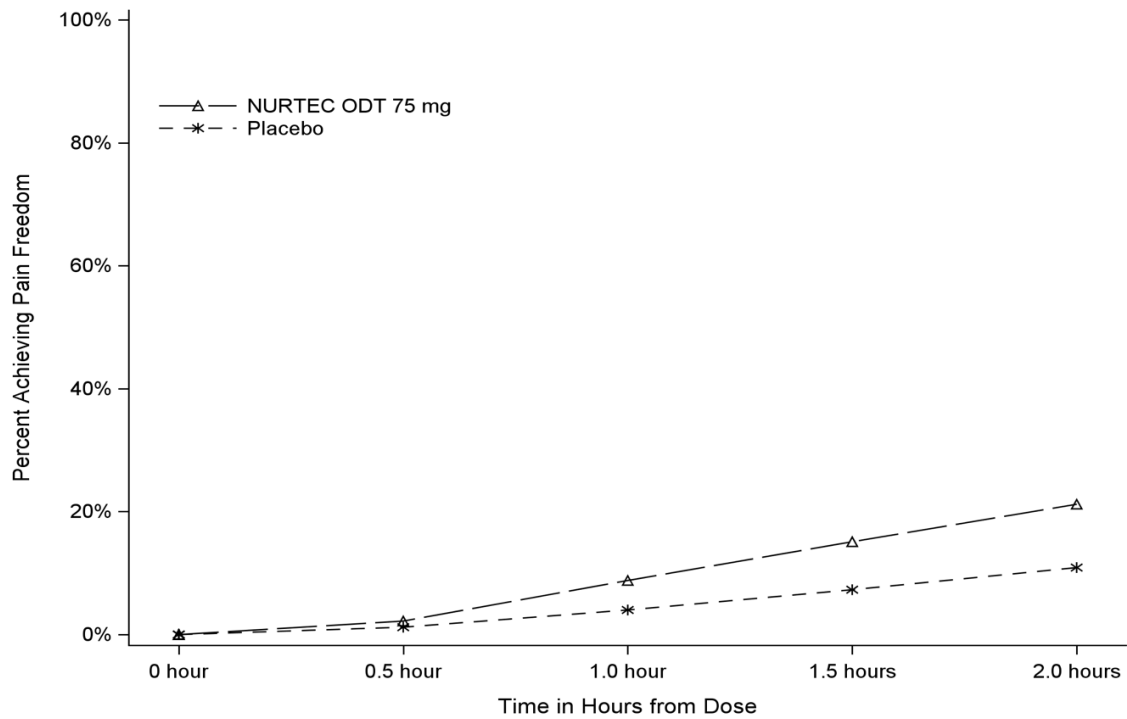
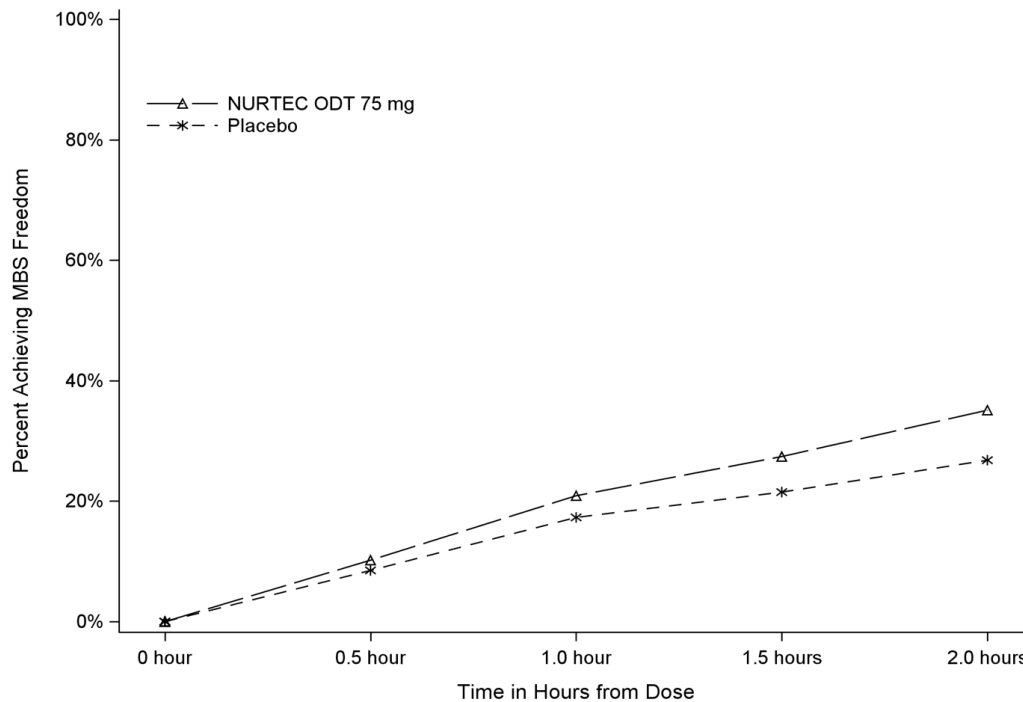


Figure 2 presents the percentage of patients achieving MBS freedom within 2 hours in Study 1.

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**Figure 2: Percentage of Patients Achieving MBS Freedom within 2 Hours in Study 1**



The incidence of photophobia and phonophobia was reduced at 2 hours following administration of NURTEC 75 mg as compared to placebo in all 3 studies.

#### Clinical efficacy: prophylaxis

The efficacy of rimegepant was evaluated as a prophylactic treatment of migraine in a randomized, double-blind, placebo-controlled study (Study 4).

Study 4 included male and female adults with at least a 1-year history of migraine (with or without aura). Patients had a history of 4 to 18 migraine attacks of moderate to severe pain intensity per 4-week period within the 12 weeks prior to the screening visit. Patients experienced an average of 10.9 headache days during the 28-day observational period, which included an average of 10.2 migraine days, prior to randomization into the study. The study randomized patients to receive rimegepant 75 mg (N=373) or placebo (N=374) for up to 12 weeks. Patients were instructed to take randomized treatment once every other day (EOD) for the 12-week treatment period. Patients were allowed to use other acute treatments for migraine (e.g., triptans, NSAIDs, paracetamol, antiemetics) as needed. Approximately 22% of patients were taking preventive medicinal products for migraine at baseline. Patients were allowed to continue in an open-label extension study for an additional 12 months.

The primary efficacy endpoint for Study 4 was the change from baseline in the mean number of monthly migraine days (MMDs) during Weeks 9 through 12 of the double-blind treatment phase. Secondary endpoints included the achievement of a  $\geq 50\%$  reduction from baseline in monthly moderate or severe migraine days.

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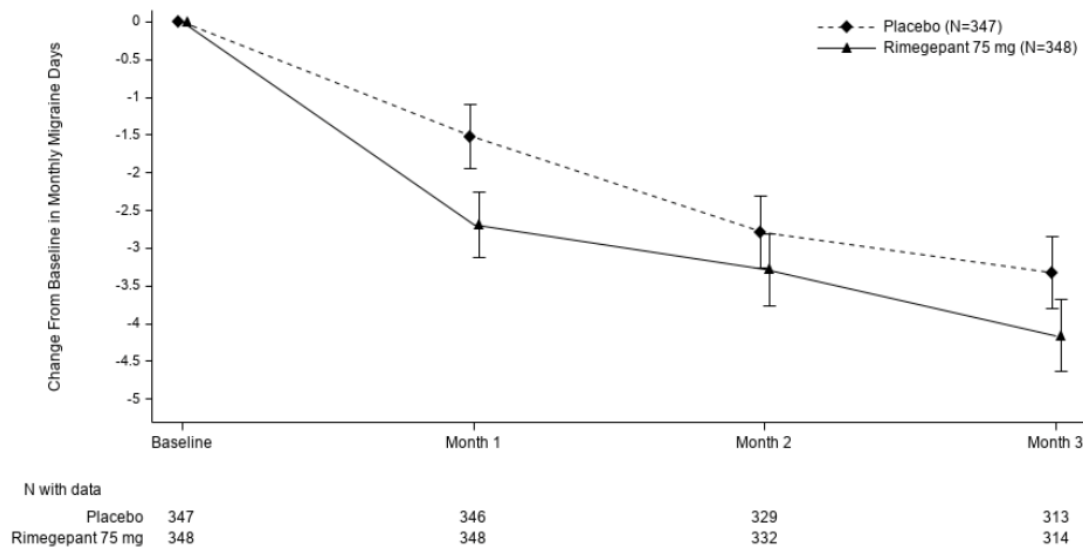
Rimegepant 75 mg dosed EOD demonstrated statistically significant improvements for key efficacy endpoints compared to placebo, as summarized in Table 3 and shown graphically in Figure 3.

**Table 3: Key Efficacy Endpoints for Study 4**

	<b>Rimegepant 75 mg EOD</b>	<b>Placebo EOD</b>
<b>Monthly Migraine Days (MMD) Weeks 9 through 12</b>	<b>N=348</b>	<b>N=347</b>
Change from baseline	-4.3	-3.5
Change compared to placebo	-0.8	
p-value	0.010 <sup>a</sup>	
<b>≥ 50% Reduction in Moderate or Severe MMDs Weeks 9 through 12</b>	<b>N=348</b>	<b>N=347</b>
% Responders	49.1	41.5
Difference compared to placebo	7.6	
p-value	0.044 <sup>a</sup>	

<sup>a</sup> Significant p-value in hierarchical testing

**Figure 3: Change from Baseline in Monthly Migraine Days in Study 4**

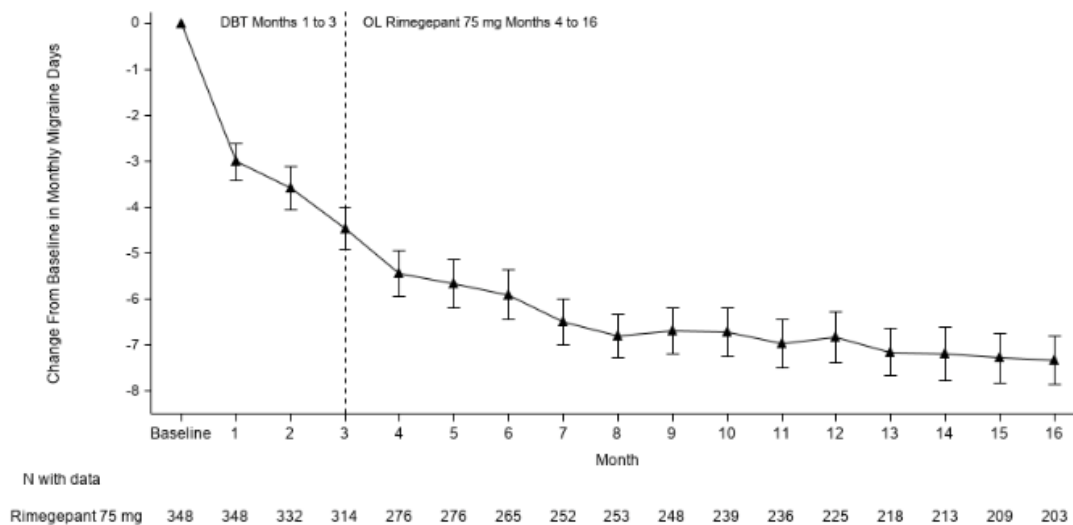


*Long-term efficacy*

Patients participating in Study 4 were allowed to continue in an open-label extension study for an additional 12 months. Efficacy was sustained for up to 1 year in an open-label study extension in which patients received rimegepant 75 mg every other day plus as needed on non-scheduled dosing days (Figure 4). A portion composed of 203 patients assigned to rimegepant completed the overall 16-month treatment period. In these patients, the overall mean reduction from baseline in the number of MMDs averaged over the 16-month treatment period was 6.2 days.

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**Figure 4: Longitudinal Plot of the Change in Mean Number of Monthly Migraine Days (MMDs) from the Observation Period Over Time during Double-Blind Treatment (Months 1 to 3) and during Treatment with Open-label Rimegepant (Months 4 to 16)**



## 5.2 Pharmacokinetic properties

### Absorption

Following oral administration, rimegepant is absorbed with the maximum concentration at 1.5 hours. Following a suprathreshold dose of 300 mg, the absolute oral bioavailability of rimegepant was approximately 64%.

### *Effects of food*

Following administration of rimegepant under fed conditions with a high-fat or low-fat meal,  $T_{max}$  was delayed by 1 to 1.5 hours. A high-fat meal reduced  $C_{max}$  by 41 to 53% and AUC by 32 to 38%. A low-fat meal reduced  $C_{max}$  by 36% and AUC by 28%. Rimegepant was administered without regard to food in clinical safety and efficacy studies.

### Distribution

The steady state volume of distribution of rimegepant is 120 L. Plasma protein binding of rimegepant is approximately 96%.

### Biotransformation

Rimegepant is primarily metabolized by CYP3A4 and to a lesser extent by CYP2C9. Rimegepant is the primary form (~77%) with no major metabolites (i.e., > 10%) detected in plasma.

Based on *in vitro* studies, rimegepant is not an inhibitor of CYP1A2, 2B6, 2C8, 2C9, 2C19, 2D6, or UGT1A1 at clinically relevant concentrations. However, rimegepant is a weak inhibitor of CYP3A4

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with time-dependent inhibition. Rimegepant is not an inducer of CYP1A2, CYP2B6, or CYP3A4 at clinically relevant concentrations.

### Elimination

The elimination half-life of rimegepant is approximately 11 hours in healthy subjects. Following oral administration of [<sup>14</sup>C]-rimegepant to healthy male subjects, 78% of the total radioactivity was recovered in feces and 24% in urine. Unchanged rimegepant is the major single component in excreted feces (42%) and urine (51%).

### *Transporters*

*In vitro*, rimegepant is a substrate of BCRP and P-gp efflux transporters. Inhibitors of P-gp and BCRP efflux transporters may increase plasma concentrations of rimegepant (see section 4.5).

Rimegepant is not a substrate of OATP1B1 or OATP1B3. Considering its low renal clearance, rimegepant was not evaluated as a substrate of the OAT1, OAT3, OCT2, MATE1, or MATE2-K.

Rimegepant is not an inhibitor of P-gp, BCRP, OAT1, or MATE2-K at clinically relevant concentrations. It is a weak inhibitor of OATP1B1 and OAT3.

Rimegepant is an inhibitor of OATP1B3, OCT2, and MATE1. Concomitant administration of rimegepant with metformin, a MATE1 transporter substrate, resulted in no clinically significant impact on either metformin pharmacokinetics or on glucose utilization. No clinical drug interactions are expected for rimegepant with OATP1B3 or OCT2, at clinically relevant concentrations.

### Linearity/non-linearity

Rimegepant exhibits greater than dose proportional increases in exposure following single oral administration, which appears to be related to a dose-dependent increase in bioavailability.

### Age, sex, weight, race, ethnicity

No clinically significant differences in the pharmacokinetics of rimegepant were observed based on age (see section 4.2), sex, race/ethnicity, body weight, migraine status, or CYP2C9 genotype.

### Renal impairment

In a dedicated clinical study comparing the pharmacokinetics of a single 75 mg dose of rimegepant in subjects with mild (estimated creatinine clearance [CL<sub>cr</sub>] 60-89 ml/min), moderate (CL<sub>cr</sub> 30-59 ml/min), and severe (CL<sub>cr</sub> 15-29 ml/min) renal impairment to that with normal subjects (healthy pooled control), a less than 50% increase in total rimegepant exposure was observed following a single 75 mg dose. The unbound AUC of rimegepant was 2.57-fold higher in subjects with severe renal impairment. NURTEC has not been studied in patients with end-stage renal disease (CL<sub>cr</sub> < 15 ml/min) (see section 4.2).

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### Hepatic impairment

In a dedicated clinical study comparing the pharmacokinetics of rimegepant in subjects with mild, moderate, and severe hepatic impairment to that with normal subjects (healthy matched control), the total exposure of rimegepant (unbound AUC) following single 75 mg dose was 3.89-fold higher in subjects with severe impairment (Child-Pugh class C). There were no clinically meaningful differences in the exposure of rimegepant in subjects with mild (Child-Pugh class A) and moderate hepatic impairment (Child-Pugh class B) compared to subjects with normal hepatic function (see section 4.2).

### **5.3 Pre-clinical safety data**

Non-clinical data reveal no special hazard for rimegepant in humans based on conventional studies of safety pharmacology, repeat-dose toxicity, genotoxicity, phototoxicity, reproduction or development, or carcinogenic potential.

Rimegepant-related effects at higher doses in repeat-dose studies included hepatic lipidosis in mice and rats, intravascular hemolysis in rats and monkeys, and emesis in monkeys. These findings were observed only at exposures considered sufficiently in excess of the maximum human exposure indicating little relevance to clinical use ( $\geq 12$  times [mice] and  $\geq 49$  times [rats] for hepatic lipidosis,  $\geq 95$  times [rats] and  $\geq 9$  times [monkeys] for intravascular hemolysis, and  $\geq 37$  times for emesis [monkeys]).

In a fertility study in rats, rimegepant-related effects were noted only at the high dose of 150 mg/kg/day (decreased fertility and increased pre-implantation loss) that produced maternal toxicity and systemic exposures  $\geq 95$  times the maximum human exposure. Oral administration of rimegepant during organogenesis resulted in foetal effects in rats but not rabbits. In rats, decreased foetal body weight and increased incidence of foetal variations were observed only at the highest dose of 300 mg/kg/day that produced maternal toxicity at exposures approximately 200 times the maximum human exposure. Additionally, rimegepant had no effects on pre- and postnatal development in rats at doses up to 60 mg/kg/day ( $\geq 24$  times the maximum human exposure) or on growth, development, or reproductive performance of juvenile rats at doses up to 45 mg/kg/day ( $\geq 14$  times the maximum human exposure).

## **6. PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

gelatin  
mannitol (E421)  
mint flavour  
sucralose

### **6.2 Incompatibilities**

Not applicable.

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### **6.3 Special precautions for storage**

Do not store above 30 °C.

Store in the original package in order to protect from moisture.

### **6.4 Nature and contents of container**

Unit dose blisters made of polyvinyl chloride (PVC), oriented polyamide (OPA) and aluminium foil and sealed with a peelable aluminum foil.

Pack sizes:

Unit dose 2 x 1 oral disintegrating tablet

Unit dose 8 x 1 oral disintegrating tablet

### **6.5 Special precautions for disposal**

No special requirements for disposal.

Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

## **7. MARKETING AUTHORISATION HOLDER**

### **Manufactured by:**

Catalent UK Swindon Zydis Limited  
Frankland Road, Blagrove, United Kingdom

### **Imported by:**

PT. Pfizer Indonesia,  
Jakarta, Indonesia

## **8. MARKETING AUTHORISATION NUMBER(S)**

DKI2346200181A1

## **9. DATE OF REVISION OF THE TEXT**

09/2025

**HARUS DENGAN RESEP DOKTER**

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## **Leaflet kemasan: Informasi untuk pengguna**

### **NURTEC 75 mg tablet orodispersibel** Rimegepant sulfat

**Baca semua bagian leaflet ini dengan cermat sebelum mulai menggunakan obat ini karena berisi informasi penting bagi Anda.**

- Simpan leaflet ini. Anda mungkin perlu membacanya kembali.
- Jika Anda memiliki pertanyaan lebih lanjut, tanyakan kepada dokter, apoteker, atau perawat Anda.
- Obat ini telah diresepkan hanya untuk Anda. Jangan memberikannya kepada orang lain. Obat ini dapat membahayakan mereka, sekalipun tanda-tanda penyakit mereka sama dengan Anda.
- Jika Anda mengalami efek samping apa pun, konsultasikan dengan dokter atau perawat Anda. Termasuk setiap kemungkinan efek samping yang tidak tercantum dalam leaflet ini. Lihat bagian 13.

#### **Isi leaflet ini:**

1. Nama obat
2. Bentuk sediaan
3. Deskripsi obat
4. Apa kandungan obat ini?
5. Kekuatan obat
6. Apa kegunaan obat ini?
7. Berapa banyak dan seberapa sering Anda seharusnya menggunakan obat ini? Apa yang harus dilakukan jika ada dosis yang terlewat?
8. Kapan seharusnya Anda tidak menggunakan obat ini?
9. Apa yang harus dipertimbangkan saat menggunakan obat ini?
10. Apa saja obat lain atau makanan yang harus dihindari selama menggunakan obat ini?
11. Apakah obat tersebut aman bagi ibu hamil dan menyusui?
12. Apakah pasien diperbolehkan mengemudi dan mengoperasikan mesin selama menggunakan obat ini?
13. Apa saja potensi efek yang tidak diinginkan dari penggunaan obat ini?
14. Tanda-tanda dan gejala-gejala overdosis
15. Apa yang harus dilakukan jika Anda menggunakan lebih dari dosis yang dianjurkan?
16. Bagaimana cara menyimpan obat ini?
17. Petunjuk penggunaan/penanganan
18. Nomor izin edar
19. Nama dan alamat pemohon dan/atau pemilik obat sesuai dengan ketentuan yang berlaku
20. Tanggal revisi
21. Peringatan khusus

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### **1. Nama obat**

NURTEC

### **2. Bentuk sediaan**

Tablet orodispersibel (ODT).

### **3. Deskripsi obat**

Tablet orodispersibel berwarna putih hingga putih pucat, bundar, dan terukir simbol .

### **4. Apa kandungan obat ini?**

Setiap NURTEC mengandung rimegepant sulfat, setara dengan 75 mg rimegepant.

Terdiri dari zat tambah:

Gelatin

Mannitol (E421)

Mint flavour

Sucralose

### **5. Kekuatan obat**

75 mg.

### **6. Apa kegunaan obat ini?**

NURTEC diindikasikan untuk:

- pengobatan migrain akut dengan atau tanpa aura pada orang dewasa;
- pengobatan preventif migrain episodik pada orang dewasa yang mengalami setidaknya 4 serangan migrain per bulan.

### **7. Berapa banyak dan seberapa sering Anda seharusnya menggunakan obat ini?**

#### **Apa yang harus dilakukan jika ada dosis yang terlewat?**

Selalu gunakan obat ini dengan tepat sesuai anjuran dokter atau apoteker Anda.

Tanyakan kepada dokter atau apoteker jika Anda merasa tidak yakin.

NURTEC digunakan secara oral dan bisa dikonsumsi dengan atau tanpa air minum.

Untuk pencegahan migrain, dosis yang disarankan adalah satu tablet (75 mg rimegepant) dua hari sekali.

Untuk pengobatan setelah serangan migrain dimulai, dosis yang dianjurkan adalah satu tablet (75 mg rimegepant) sesuai kebutuhan, tidak lebih dari sekali sehari.

Dosis maksimum hariannya adalah satu tablet (75 mg rimegepant) per hari.

#### **Jika Anda lupa meminum NURTEC**

Jika Anda mengonsumsi NURTEC untuk mencegah migrain dan melewatkan satu dosis, cukup konsumsi dosis berikutnya pada jadwal yang sesuai. **Jangan mengonsumsi dosis ganda untuk memenuhi dosis yang terlupa.**

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Jika Anda memiliki pertanyaan lebih lanjut seputar penggunaan obat ini, tanyakan kepada dokter atau apoteker Anda.

## **8. Kapan seharusnya Anda tidak menggunakan obat ini?**

### **Jangan gunakan NURTEC:**

- jika Anda alergi atau memiliki riwayat alergi terhadap rimegepant atau bahan lainnya dalam obat ini.

## **9. Apa yang harus dipertimbangkan saat menggunakan obat ini?**

Konsultasikan dengan dokter Anda sebelum menggunakan NURTEC:

- jika Anda menderita gangguan hati yang berat
- jika Anda menderita penurunan fungsi ginjal atau menjalani dialisis ginjal

Selama pengobatan menggunakan NURTEC, hentikan menggunakan obat ini dan segera beri tahu dokter Anda:

- jika Anda mengalami gejala reaksi alergi, (cth., kesulitan bernapas, ruam yang berat, pembengkakan lidah, mulut, atau wajah, kesulitan menelan, rasa sesak di tenggorokan, atau suara parau). Gejala ini dapat terjadi beberapa hari setelah pemberian.

### **Pasien anak-anak dan remaja**

NURTEC tidak boleh diberikan kepada anak-anak dan remaja di bawah usia 18 tahun karena belum ada penelitian di dalam kelompok umur ini.

## **10. Apa saja obat lain atau makanan yang harus dihindari selama menggunakan obat ini?**

Beri tahu dokter atau apoteker Anda jika Anda sedang, belum lama ini, atau akan menggunakan obat lain. Hal ini dikarenakan beberapa obat dapat memengaruhi cara kerja NURTEC atau NURTEC dapat memengaruhi cara kerja obat lain.

Berikut ini adalah daftar contoh obat-obatan yang harus dihindari saat menggunakan NURTEC:

- itrakonazol dan klaritromisin (obat-obatan yang digunakan untuk mengobati infeksi jamur atau bakteri).
- ritonavir dan evafirenz (obat-obatan yang digunakan untuk mengobati infeksi HIV).
- bosentan (obat yang digunakan untuk mengobati tekanan darah tinggi).
- St. John's wort (obat herbal yang digunakan untuk mengobati depresi).
- fenobarbital (obat yang digunakan untuk mengobati epilepsi).
- rifampisin (obat yang digunakan untuk mengobati tuberkulosis).
- modafinil (obat yang digunakan untuk mengobati narkolepsi).

Jangan mengonsumsi NURTEC lebih dari sekali setiap 48 jam bersama:

- flukonazol dan eritromisin (obat-obatan yang digunakan untuk mengobati infeksi jamur atau bakteri).

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- diltiazem, kuinidin, dan verapamil (obat-obatan yang digunakan untuk mengobati irama jantung abnormal, nyeri dada (angina), atau tekanan darah tinggi).
- siklosporin (obat yang digunakan untuk mencegah penolakan organ setelah transplantasi organ).

## **11. Apakah obat tersebut aman bagi ibu hamil dan menyusui?**

### **Kehamilan**

Jika Anda sedang hamil atau merasa diri Anda mungkin hamil, segera hubungi dokter sebelum menggunakan NURTEC. Lebih diutamakan menghindari penggunaan NURTEC selama kehamilan karena efek obat ini terhadap ibu hamil masih belum diketahui.

### **Menyusui**

Jika Anda sedang menyusui atau berencana untuk menyusui, bicarakan dengan dokter atau apoteker Anda sebelum menggunakan obat ini. Anda dan dokter Anda harus memutuskan apakah Anda akan menggunakan NURTEC saat menyusui.

## **12. Apakah pasien diperbolehkan mengemudi dan mengoperasikan mesin selama menggunakan obat ini?**

NURTEC diperkirakan tidak memengaruhi kemampuan Anda mengemudi atau menggunakan mesin.

## **13. Apa saja potensi efek yang tidak diinginkan dari penggunaan obat ini**

Seperti obat-obatan lainnya, obat ini dapat menimbulkan efek samping, sekalipun tidak semua orang mengalaminya.

**Hentikan menggunakan NURTEC dan segera hubungi dokter Anda jika Anda memiliki tanda-tanda reaksi alergi** (seperti ruam berat atau napas pendek) atau tanda-tanda reaksi alergi berat yang dikenal sebagai ‘anafilaksis’ (seperti pembengkakan lidah, mulut, atau wajah, kesulitan menelan atau bernapas, rasa sesak di tenggorokan, atau suara parau). Reaksi alergi, termasuk anafilaksis, bersama penggunaan NURTEC tidaklah umum (dapat dialami hingga 1 di antara 100 orang).

Efek samping yang umum (dapat dialami hingga 1 di antara 10 orang) adalah mual. Untuk melaporkan efek samping, hubungi [www.pfizersafetyreporting.com](http://www.pfizersafetyreporting.com) atau email di [IDN.AEReporting@pfizer.com](mailto:IDN.AEReporting@pfizer.com).

### **Melaporkan efek samping**

Jika Anda mengalami efek samping apa pun, konsultasikan dengan dokter, apoteker, atau perawat Anda. Termasuk setiap kemungkinan efek samping yang tidak tercantum dalam leaflet ini. Dengan melaporkan efek samping, Anda bisa membantu memberikan informasi lebih lanjut mengenai keamanan obat ini.

## **14. Tanda-tanda dan gejala-gejala overdosis**

Efek samping apa pun dapat terjadi.

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**15. Apa yang harus dilakukan jika Anda menggunakan lebih dari dosis yang dianjurkan?**

Konsultasikan dengan dokter atau apoteker atau segera pergi ke rumah sakit. Bawa serta kemasan obat dan leaflet ini.

**16. Bagaimana cara menyimpan obat ini?**

Jauhkan obat ini dari pandangan dan jangkauan anak-anak.

Jangan gunakan obat ini setelah melewati tanggal kedaluwarsa yang tertera pada wadahnya.

Simpan pada suhu di bawah 30 °C. Simpan pada kemasan blister asli untuk melindungi dari kelembaban.

Tidak membuang obat melewati limbah pembuangan atau limbah pembuangan rumah tangga.

Tanyakan pada apoteker bagaimana cara yang tepat untuk pembuangan obat jika tidak digunakan.

**17. Petunjuk penggunaan**

2 x 1 oral tablet orodispersibel per unit blister.

8 x 1 oral tablet orodispersibel per unit blister

NURTEC digunakan secara oral.

Letakkan tablet di atas lidah atau di bawah lidah, obat akan hancur di dalam mulut.

NURTEC dapat dikonsumsi dengan atau tanpa air minum.

Instruksi:

- Gunakan tangan yang kering saat membuka kemasan blister.
- Kupas foil penutup belakang dari satu blister dan ambil tablet orodispersibel ini dengan hati-hati. Jangan mendorong tablet agar keluar dari foil.
- Segera konsumsi tablet setelah membuka kemasan blister. Jangan simpan tablet di luar kemasan blister untuk penggunaan di masa mendatang.

**18. Nomor izin edar**

DKI2346200181A1

**19. Nama dan alamat pemohon dan/atau pemilik obat sesuai dengan ketentuan yang berlaku**

**Diproduksi oleh:**

Catalent UK Swindon Zydis Limited  
Frankland Road, Blagrove, United Kingdom

**Diimpor oleh:**

PT. Pfizer Indonesia,  
Jakarta, Indonesia

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**20. Tanggal revisi**

09/2025

**21. Peringatan khusus**

**HARUS DENGAN RESEP DOKTER**

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