

**ANNEX I**

**SUMMARY OF PRODUCT CHARACTERISTICS**

## 1. NAME OF THE MEDICINAL PRODUCT

Rotarix powder and solvent for **oral** suspension  
Rotavirus vaccine, live

## 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

After reconstitution, 1 dose (1 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated)\* not less than  $10^{6.0}$  CCID<sub>50</sub>

\*Produced on Vero cells

### Excipients with known effect:

This product contains 13.5 mg of sorbitol, 9 mg of sucrose, 10 micrograms of glucose and 0.15 microgram of phenylalanine per dose (see section 4.4).

For the full list of excipients, see section 6.1.

## 3. PHARMACEUTICAL FORM

Powder and solvent for **oral** suspension.

The powder is white.

The solvent is a turbid liquid with a slow settling white deposit and a colourless supernatant.

## 4. CLINICAL PARTICULARS

### 4.1 Therapeutic indications

Rotarix is indicated for the active immunisation of infants aged 6 to 24 weeks for prevention of gastroenteritis due to rotavirus infection (see sections 4.2, 4.4 and 5.1).

The use of Rotarix should be based on official recommendations.

### 4.2 Posology and method of administration

#### Posology

The vaccination course consists of two doses. The first dose may be administered from the age of 6 weeks. There should be an interval of at least 4 weeks between doses. The vaccination course should preferably be given before 16 weeks of age, but must be completed by the age of 24 weeks.

Rotarix may be given with the same posology to preterm infants born after at least 27 weeks of gestational age (see sections 4.8 and 5.1).

In clinical trials, spitting or regurgitation of the vaccine has rarely been observed and, under such circumstances, a replacement dose was not given. However, in the unlikely event that an infant spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

It is recommended that infants who receive a first dose of Rotarix complete the 2-dose regimen with Rotarix. There are no data on safety, immunogenicity or efficacy when Rotarix is administered for the first dose and another rotavirus vaccine is administered for the second dose or vice versa.

#### *Paediatric population*

Rotarix should not be used in children over 24 weeks of age.

## Method of administration

Rotarix is for **oral** use only.

### **Rotarix should under no circumstances be injected.**

For instructions for the preparation or reconstitution of the medicinal product before administration, see section 6.6.

## **4.3 Contraindications**

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

Hypersensitivity after previous administration of rotavirus vaccines.

History of intussusception.

Subjects with uncorrected congenital malformation of the gastrointestinal tract that would predispose for intussusception.

Subjects with Severe Combined Immunodeficiency (SCID) disorder (see section 4.8).

Administration of Rotarix should be postponed in subjects suffering from acute severe febrile illness. The presence of a minor infection is not a contra-indication for immunisation.

The administration of Rotarix should be postponed in subjects suffering from diarrhoea or vomiting.

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

It is good clinical practice that vaccination should be preceded by a review of the medical history especially with regard to the contraindications and by a clinical examination.

There are no data on the safety and efficacy of Rotarix in infants with gastrointestinal illnesses or growth retardation. Administration of Rotarix may be considered with caution in such infants when, in the opinion of the physician, withholding the vaccine entails a greater risk.

As a precaution, healthcare professionals should follow-up on any symptoms indicative of intussusception (severe abdominal pain, persistent vomiting, bloody stools, abdominal bloating and/or high fever) since data from observational safety studies indicate an increased risk of intussusception, mostly within 7 days after rotavirus vaccination (see section 4.8). Parents/guardians should be advised to promptly report such symptoms to their healthcare provider.

For subjects with a predisposition for intussusception, see section 4.3.

Asymptomatic and mildly symptomatic HIV infections are not expected to affect the safety or efficacy of Rotarix. A clinical study in a limited number of asymptomatic or mildly symptomatic HIV positive infants showed no apparent safety problems (see section 4.8).

Administration of Rotarix to infants who have known or suspected immunodeficiency, including *in utero* exposure to an immunosuppressive treatment, should be based on careful consideration of potential benefits and risks.

Excretion of the vaccine virus in the stools is known to occur after vaccination with peak excretion around the 7th day. Viral antigen particles detected by ELISA were found in 50% of stools after the first dose and 4% of stools after the second dose. When these stools were tested for the presence of live vaccine strain, only 17% were positive.

Cases of transmission of this excreted vaccine virus to seronegative contacts of vaccinees have been observed without causing any clinical symptom.

Rotarix should be administered with caution to individuals with immunodeficient close contacts, such as individuals with malignancies, or who are otherwise immunocompromised or individuals receiving immunosuppressive therapy.

Contacts of recent vaccinees should observe personal hygiene (e.g. wash their hands after changing child's nappies).

The potential risk of apnoea and the need for respiratory monitoring for 48-72h should be considered when administering the primary immunisation series to very premature infants (born  $\leq$  28 weeks of gestation) and particularly for those with a previous history of respiratory immaturity.

As the benefit of the vaccination is high in this group of infants, vaccination should not be withheld or delayed.

A protective immune response may not be elicited in all vaccinees (see section 5.1).

The extent of protection that Rotarix might provide against other rotavirus strains that have not been circulating in clinical trials is currently unknown. Clinical studies from which efficacy data were derived were conducted in Europe, Central and South America, Africa and Asia (see section 5.1).

Rotarix does not protect against gastro-enteritis due to other pathogens than rotavirus.

No data are available on the use of Rotarix for post-exposure prophylaxis.

### **Rotarix should under no circumstances be injected.**

#### Excipients

This vaccine contains sucrose, glucose and sorbitol as excipients. Patients with rare hereditary problems of fructose intolerance, glucose-galactose malabsorption or sucrase-isomaltase insufficiency should not take this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful for patients with phenylketonuria (PKU).

This vaccine contains less than 1 mmol sodium (23 mg) in each dose, that is to say essentially 'sodium-free'.

#### Traceability

In order to improve the traceability of biological medicinal products, the name and the batch number of the administered product should be clearly recorded.

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

Rotarix can be given concomitantly with any of the following monovalent or combination vaccines [including hexavalent vaccines (DTPa-HBV-IPV/Hib)]: diphtheria-tetanus-whole cell pertussis vaccine (DTPw), diphtheria-tetanus-acellular pertussis vaccine (DTPa), *Haemophilus influenzae* type b vaccine (Hib), inactivated polio vaccine (IPV), hepatitis B vaccine (HBV), pneumococcal conjugate vaccine and meningococcal serogroup C conjugate vaccine. Clinical studies demonstrated that the immune responses and the safety profiles of the administered vaccines were unaffected.

Concomitant administration of Rotarix and oral polio vaccine (OPV) does not affect the immune response to the polio antigens. Although concomitant administration of OPV may slightly reduce the

immune response to rotavirus vaccine, clinical protection against severe rotavirus gastro-enteritis was shown to be maintained in a clinical trial involving more than 4 200 subjects who received Rotarix concomitantly with OPV.

There are no restrictions on the infant's consumption of food or liquid, either before or after vaccination.

#### **4.6 Fertility, pregnancy and lactation**

Rotarix is not intended for use in adults. There are no data on the use of Rotarix during pregnancy and lactation.

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

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

Not relevant.

#### **4.8 Undesirable effects**

##### **Summary of the safety profile**

The safety profile presented below is based on data from clinical trials conducted with either the lyophilised or the liquid formulation of Rotarix.

In a total of four clinical trials, approximately 3 800 doses of Rotarix liquid formulation were administered to approximately 1 900 infants. Those trials have shown that the safety profile of the liquid formulation is comparable to the lyophilised formulation.

In a total of twenty-three clinical trials, approximately 106 000 doses of Rotarix (lyophilised or liquid formulation) were administered to approximately 51 000 infants.

In three placebo-controlled clinical trials (Finland, India and Bangladesh), in which Rotarix was administered alone (administration of routine paediatric vaccines was staggered), the incidence and severity of the solicited events (collected 8 days post-vaccination), diarrhoea, vomiting, loss of appetite, fever, irritability and cough/runny nose were not significantly different in the group receiving Rotarix when compared to the group receiving placebo. No increase in the incidence or severity of these events was seen with the second dose.

In a pooled analysis from seventeen placebo-controlled clinical trials (Europe, North America, Latin America, Asia, Africa) including trials in which Rotarix was co-administered with routine paediatric vaccines (see section 4.5), the following adverse reactions (collected 31 days post-vaccination) were considered as possibly related to vaccination.

##### **Tabulated list of adverse reactions**

Adverse reactions reported are listed according to the following frequency:

Frequencies are reported as:

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	( $< 1/10\ 000$ )

<b>System Organ Class</b>	<b>Frequency</b>	<b>Adverse reactions</b>
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Gastrointestinal disorders	Common	Diarrhoea
	Uncommon	Abdominal pain, flatulence
	Very rare	Intussusception (see section 4.4)
	Not known*	Haematochezia
	Not known*	Gastroenteritis with vaccine viral shedding in infants with Severe Combined Immunodeficiency (SCID) disorder
Skin and subcutaneous tissue disorders	Uncommon	Dermatitis
	Very rare	Urticaria
General disorders and administration site conditions	Common	Irritability
Respiratory, thoracic and mediastinal disorders	Not known*	Apnoea in very premature infants ( $\leq 28$ weeks of gestation) (see section 4.4)

\* Because these events were reported spontaneously, it is not possible to reliably estimate their frequency.

## Description of selected adverse reactions

### Intussusception

Data from observational safety studies performed in several countries indicate that rotavirus vaccines carry an increased risk of intussusception, mostly within 7 days of vaccination. Up to 6 additional cases per 100 000 infants have been observed in these countries against a background incidence of 25 to 101 per 100 000 infants (less than one year of age) per year, respectively.

There is limited evidence of a smaller increased risk following the second dose.

It remains unclear whether rotavirus vaccines affect the overall incidence of intussusception based on longer periods of follow-up (see section 4.4).

## Other special populations

### Safety in preterm infants

In a clinical study, 670 pre-term infants from 27 to 36 weeks of gestational age were administered Rotarix and 339 received placebo. The first dose was administered from 6 weeks after birth. Serious adverse events were observed in 5.1% of recipients of Rotarix as compared with 6.8% of placebo recipients. Similar rates of other adverse events were observed in Rotarix and placebo recipients. No cases of intussusception were reported.

### Safety in infants with human immunodeficiency (HIV) infection

In a clinical study, 100 infants with HIV infection were administered Rotarix or placebo. The safety profile was similar between Rotarix and placebo recipients.

### 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 the national reporting system listed in [Appendix V](#).

## 4.9 Overdose

Some cases of overdose have been reported. In general, the adverse event profile reported in these cases was similar to that observed after administration of the recommended dose of Rotarix.

## 5. PHARMACOLOGICAL PROPERTIES

### 5.1 Pharmacodynamic properties

Pharmaco-therapeutic group: rotavirus diarrhoea vaccines, ATC code: J07BH01

#### Protective efficacy

In clinical trials, efficacy was demonstrated against gastro-enteritis due to rotavirus of the most common genotypes G1P[8], G2P[4], G3P[8], G4P[8] and G9P[8]. In addition, efficacy against uncommon rotavirus genotypes G8P[4] (severe gastro-enteritis) and G12P[6] (any gastro-enteritis) has been demonstrated. These strains are circulating worldwide.

Clinical studies have been conducted in Europe, Latin America, Africa and Asia to evaluate the protective efficacy of Rotarix against any and severe rotavirus gastro-enteritis (RVGE).

Severity of gastro-enteritis was defined according to two different criteria:

- the Vesikari 20-point scale, which evaluates the full clinical picture of rotavirus gastro-enteritis by taking into account the severity and duration of diarrhoea and vomiting, the severity of fever and dehydration as well as the need for treatment

or

- the clinical case definition based on World Health Organization (WHO) criteria

Clinical protection was assessed in the ATP cohort for efficacy, which includes all subjects from the ATP cohort for safety who entered into the concerned efficacy follow-up period.

#### Protective efficacy in Europe

A clinical study performed in Europe evaluated Rotarix given according to different European schedules (2, 3 months; 2, 4 months; 3, 4 months; 3, 5 months) in 4 000 subjects.

After two doses of Rotarix, the protective vaccine efficacy observed during the first and second year of life is presented in the following table:

	1 <sup>st</sup> year of life Rotarix N=2 572 Placebo N=1 302		2 <sup>nd</sup> year of life Rotarix N=2 554 Placebo N=1 294	
Vaccine efficacy (%) against any and severe rotavirus gastro-enteritis [95% CI]				
Genotype	Any severity	Severe <sup>†</sup>	Any severity	Severe <sup>†</sup>
G1P[8]	95.6 [87.9;98.8]	96.4 [85.7;99.6]	82.7 [67.8;91.3]	96.5 [86.2;99.6]
G2P[4]	62.0* [<0.0;94.4]	74.7* [<0.0;99.6]	57.1 [<0.0;82.6]	89.9 [9.4;99.8]
G3P[8]	89.9 [9.5;99.8]	100 [44.8;100]	79.7 [<0.0;98.1]	83.1* [<0.0;99.7]
G4P[8]	88.3 [57.5;97.9]	100 [64.9;100]	69.6* [<0.0;95.3]	87.3 [<0.0;99.7]
G9P[8]	75.6 [51.1;88.5]	94.7 [77.9;99.4]	70.5 [50.7;82.8]	76.8 [50.8;89.7]
Strains with P[8] genotype	88.2 [80.8;93.0]	96.5 [90.6;99.1]	75.7 [65.0;83.4]	87.5 [77.8;93.4]

Circulating rotavirus strains	87.1 [79.6;92.1]	95.8 [89.6;98.7]	71.9 [61.2;79.8]	85.6 [75.8;91.9]
<b>Vaccine efficacy (%) against rotavirus gastro-enteritis requiring medical attention [95% CI]</b>				
Circulating rotavirus strains	91.8 [84;96.3]		76.2 [63.0;85.0]	
<b>Vaccine efficacy (%) against hospitalisation due to rotavirus gastro-enteritis [95% CI]</b>				
Circulating rotavirus strains	100 [81.8;100]		92.2 [65.6;99.1]	

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution

Vaccine efficacy during the first year of life progressively increased with increasing disease severity, reaching 100% (95% CI: 84.7;100) for Vesikari scores  $\geq 17$ .

#### Protective efficacy in Latin America

A clinical study performed in Latin America evaluated Rotarix in more than 20 000 subjects. Severity of gastro-enteritis (GE) was defined according to WHO criteria. The protective vaccine efficacy against severe rotavirus (RV) gastro-enteritis requiring hospitalisation and/or rehydration therapy in a medical facility and the genotype specific vaccine efficacy after two doses of Rotarix are presented in the table below:

<b>Genotype</b>	<b>Severe rotavirus gastro-enteritis† (1<sup>st</sup> year of life)</b> Rotarix N=9 009 Placebo N=8 858	<b>Severe rotavirus gastro-enteritis† (2<sup>nd</sup> year of life)</b> Rotarix N=7 175 Placebo N=7 062
	<b>Efficacy (%)</b> <b>[95% CI]</b>	<b>Efficacy (%)</b> <b>[95% CI]</b>
All RVGE	84.7 [71.7;92.4]	79.0 [66.4;87.4]
G1P[8]	91.8 [74.1;98.4]	72.4 [34.5;89.9]
G3P[8]	87.7 [8.3;99.7]	71.9* [<0.0;97.1]
G4P[8]	50.8#* [<0.0;99.2]	63.1 [0.7;88.2]
G9P[8]	90.6 [61.7;98.9]	87.7 [72.9;95.3]
Strains with P[8] genotype	90.9 [79.2;96.8]	79.5 [67.0;87.9]

† Severe rotavirus gastro-enteritis was defined as an episode of diarrhoea with or without vomiting that required hospitalisation and/or re-hydration therapy in a medical facility (WHO criteria)

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution

# The numbers of cases, on which the estimates of efficacy against G4P[8] were based, were very small (1 case in the Rotarix group and 2 cases in the placebo group)

A pooled analysis of five efficacy studies\*, showed a 71.4% (95% CI: 20.1;91.1) efficacy against severe rotavirus gastro-enteritis (Vesikari score  $\geq 11$ ) caused by rotavirus G2P[4] genotype during the first year of life.

\* In these studies, the point estimates and confidence intervals were respectively: 100% (95% CI: -1 858.0;100), 100% (95% CI: 21.1;100), 45.4% (95% CI: -81.5;86.6), 74.7% (95% CI: -386.2;99.6). No point estimate was available for the remaining study.



### Protective efficacy in Africa

A clinical study performed in Africa (Rotarix: N = 2 974; placebo: N = 1 443) evaluated Rotarix given at approximately 10 and 14 weeks of age (2 doses) or 6, 10 and 14 weeks of age (3 doses). The vaccine efficacy against severe rotavirus gastro-enteritis during the first year of life was 61.2% (95% CI: 44.0;73.2). The protective vaccine efficacy (pooled doses) observed against any and severe rotavirus gastro-enteritis is presented in the following table:

<b>Genotype</b>	<b>Any rotavirus gastro-enteritis</b> Rotarix N=2 974 Placebo N=1 443	<b>Severe rotavirus gastro-enteritis†</b> Rotarix N=2 974 Placebo N=1 443
	<b>Efficacy (%)</b> <b>[95% CI]</b>	<b>Efficacy (%)</b> <b>[95% CI]</b>
G1P[8]	68.3 [53.6;78.5]	56.6 [11.8;78.8]
G2P[4]	49.3 [4.6;73.0]	83.8 [9.6;98.4]
G3P[8]	43.4* [<0.0;83.7]	51.5* [<0.0;96.5]
G8P[4]	38.7* [<0.0;67.8]	63.6 [5.9;86.5]
G9P[8]	41.8* [<0.0;72.3]	56.9* [<0.0;85.5]
G12P[6]	48.0 [9.7;70.0]	55.5* [<0.0;82.2]
Strains with P[4] genotype	39.3 [7.7;59.9]	70.9 [37.5;87.0]
Strains with P[6] genotype	46.6 [9.4;68.4]	55.2* [<0.0;81.3]
Strains with P[8] genotype	61.0 [47.3;71.2]	59.1 [32.8;75.3]

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

### Sustained efficacy up to 3 years of age in Asia

A clinical study conducted in Asia (Hong Kong, Singapore and Taiwan) (Total vaccinated cohort : Rotarix: N = 5 359; placebo: N = 5 349) evaluated Rotarix given according to different schedules (2, 4 months of age; 3, 4 months of age).

During the first year, significantly fewer subjects in the Rotarix group reported severe rotavirus gastro-enteritis caused by the circulating wild-type RV compared to the placebo group from 2 weeks after Dose 2 up to one year of age (0.0% versus 0.3%), with a vaccine efficacy of 100% (95% CI: 72.2; 100).

The protective vaccine efficacy after two doses of Rotarix observed against severe rotavirus gastro-enteritis up to 2 years of age is presented in the following table:

	<b>Efficacy up to 2 years of age</b> Rotarix N= 5 263 Placebo N= 5 256
<b>Vaccine efficacy (%) against severe rotavirus gastro-enteritis [95% CI]</b>	
<b>Genotype</b>	<b>Severe†</b>
G1P[8]	100 [80.8;100]
G2P[4]	100* [<0.0;100]

G3P[8]	94.5 [64.9;99.9]
G9P[8]	91.7 [43.8;99.8]
Strains with P[8] genotype	95.8 [83.8;99.5]
Circulating rotavirus strains	96.1 [85.1;99.5]
Vaccine efficacy (%) against rotavirus gastro-enteritis requiring hospitalisation and/or rehydration therapy in a medical facility [95% CI]	
Circulating rotavirus strains	94.2 [82.2;98.8]

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution

During the third year of life, there were no cases of severe RV gastro-enteritis in the Rotarix group (N=4 222) versus 13 (0.3%) in the placebo group (N=4 185). Vaccine efficacy was 100% (95% CI: 67.5; 100). The severe RV gastro-enteritis cases were due to RV strains G1P[8], G2P[4], G3P[8] and G9P[8]. The incidence of severe RV gastro-enteritis associated with the individual genotypes was too small to allow calculation of efficacy. The efficacy against severe RV gastro-enteritis requiring hospitalisation was 100% (95% CI: 72.4; 100).

### Immune response

The immunologic mechanism by which Rotarix protects against rotavirus gastro-enteritis is not completely understood. A relationship between antibody responses to rotavirus vaccination and protection against rotavirus gastro-enteritis has not been established.

The following table shows the percentage of subjects initially seronegative for rotavirus (IgA antibody titres  $< 20$  U/mL) (by ELISA) with serum anti-rotavirus IgA antibody titres  $\geq 20$  U/mL one to two months after the second dose of vaccine or placebo as observed in different studies.

Schedule	Studies conducted in	Vaccine		Placebo	
		N	% $\geq 20$ U/mL [95% CI]	N	% $\geq 20$ U/mL [95% CI]
<b>2, 3 months</b>	France, Germany	239	82.8 [77.5;87.4]	127	8.7 [4.4;15.0]
<b>2, 4 months</b>	Spain	186	85.5 [79.6;90.2]	89	12.4 [6.3;21.0]
<b>3, 5 months</b>	Finland, Italy	180	94.4 [90.0;97.3]	114	3.5 [1.0;8.7]
<b>3, 4 months</b>	Czech Republic	182	84.6 [78.5;89.5]	90	2.2 [0.3;7.8]
<b>2, 3 to 4 months</b>	Latin America; 11 countries	393	77.9% [73.8;81.6]	341	15.1% [11.7;19.0]
<b>10, 14 weeks and 6, 10, 14 weeks (Pooled)</b>	South Africa, Malawi	221	58.4 [51.6;64.9]	111	22.5 [15.1;31.4]

### Immune response in preterm infants

In a clinical study conducted in preterm infants, born after at least 27 weeks of gestational age, the immunogenicity of Rotarix was assessed in a subset of 147 subjects and showed that Rotarix is immunogenic in this population; 85.7% (95% CI: 79.0;90.9) of subjects achieved serum anti-rotavirus IgA antibody titres  $\geq 20$  U/mL (by ELISA) one month after the second dose of vaccine.

### Effectiveness

In observational studies, vaccine effectiveness was demonstrated against severe gastro-enteritis leading to hospitalisation due to rotavirus of common genotypes G1P[8], G2P[4], G3P[8], G4P[8] and G9P[8] as well as the less common rotavirus genotypes G9P[4] and G9P[6]. All of these strains are circulating worldwide.

#### Effectiveness after 2 doses in preventing RVGE leading to hospitalisation

Countries Period	Age range	N <sup>(1)</sup> (cases/controls)	Strains	Effectiveness % [95% CI]			
High Income countries							
Belgium 2008-2010 <sup>(2)</sup>	< 4 yrs 3-11 m	160/198	All	90 [81;95] 91 [75;97]			
	< 4 yrs	41/53	G1P[8]	95 [78;99]			
	< 4 yrs 3-11 m	80/103	G2P[4]	85 [64;94] 83 [22;96] <sup>(3)</sup>			
	< 4 yrs	12/13	G3P[8]	87* [ $<0$ ;98] <sup>(3)</sup>			
	< 4 yrs	16/17	G4P[8]	90 [19;99] <sup>(3)</sup>			
Singapore 2008-2010 <sup>(2)</sup>	< 5 yrs	136/272	All	84 [32;96]			
		89/89	G1P[8]	91 [30;99]			
Taiwan 2009-2011	< 3 yrs	184/1 623 <sup>(4)</sup>	All G1P[8]	92 [75;98] 95 [69;100]			
US 2010-2011	< 2 yrs	85/1 062 <sup>(5)</sup>	All G1P[8] G2P[4]	85 [73;92] 88 [68;95] 88 [68;95]			
	8-11 m		All	89 [48;98]			
US 2009-2011	< 5 yrs	74/255 <sup>(4)</sup>	All	68 [34;85]			
Middle Income Countries							
Bolivia 2010-2011	< 3 yrs 6-11 m	300/974	All	77 [65;84] <sup>(6)</sup> 77 [51;89]			
	< 3 yrs 6-11 m		G9P[8]	85 [69;93] 90 [65;97]			
	< 3 yrs		G3P[8]	93 [70;98]			
			G2P[4]	69 [14;89]			
			G9P[6]	87 [19;98]			
Brazil 2008-2011	< 2 yrs	115/1 481	All G1P[8] G2P[4]	72 [44;85] <sup>(6)</sup> 89 [78;95] 76 [64;84]			
			Brazil 2008-2009 <sup>(2)</sup>	< 3 yrs 3-11 m	249/249 <sup>(5)</sup>	All	76 [58;86] 96 [68;99]
				< 3 yrs 3-11 m	222/222 <sup>(5)</sup>	G2P[4]	75 [57;86] 95 [66;99] <sup>(3)</sup>
El Salvador 2007-2009	< 2 yrs 6-11 m	251/770 <sup>(5)</sup>	All	76 [64;84] <sup>(6)</sup> 83 [68;91]			
Guatemala 2012-2013	< 4 yrs	NA <sup>(7)</sup>	All	63 [23;82]			
Mexico 2010	< 2 yrs	9/17 <sup>(5)</sup>	G9P[4]	94 [16;100]			
Low Income Countries							
Malawi 2012-2014	< 2 yrs	81/286 <sup>(5)</sup>	All	63 [23;83]			

m: months

yrs: years

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

- (1) The number of fully vaccinated (2 doses) and unvaccinated cases and controls is given.
- (2) GSK sponsored studies
- (3) Data from a post-hoc analysis
- (4) Vaccine effectiveness was calculated using rotavirus-negative hospital control participants (estimates from Taiwan were calculated using combined rotavirus-negative hospital control and non-diarrhoea hospital control participants).
- (5) Vaccine effectiveness was calculated using neighbourhood controls.
- (6) In subjects who did not receive the full course of vaccination, the effectiveness after one dose ranged from 51% (95% CI: 26;67, El Salvador) to 60% (95% CI: 37;75, Brazil).
- (7) NA: not available. Vaccine effectiveness estimate is based on 41 fully vaccinated cases and 175 fully vaccinated controls.

#### Impact on mortality<sup>§</sup>

Impact studies with Rotarix conducted in Panama, Brazil and Mexico showed a decrease in all-cause diarrhoea mortality ranging from 17% to 73% in children less than 5 years of age, within 2 to 4 years after vaccine introduction.

#### Impact on hospitalisation<sup>§</sup>

In a retrospective database study in Belgium conducted in children 5 years of age and younger, the direct and indirect impact of Rotarix vaccination on rotavirus-related hospitalisation ranged from 64% (95% CI: 49;76) to 80% (95% CI: 77;83) two years after vaccine introduction. Similar studies in Armenia, Australia, Brazil, Canada, El Salvador and Zambia showed a reduction of 45 to 93% between 2 and 4 years after vaccine introduction.

In addition, nine impact studies on all-cause diarrhoea hospitalisation conducted in Africa and Latin America showed a reduction of 14% to 57% between 2 and 5 years after vaccine introduction.

<sup>§</sup>NOTE : Impact studies are meant to establish a temporal relationship but not a causal relationship between the disease and vaccination. Natural fluctuations of the incidence of the disease may also influence the observed temporal effect.

## **5.2 Pharmacokinetic properties**

Not applicable.

## **5.3 Preclinical safety data**

Non-clinical data reveal no special hazard for humans based on conventional studies of repeated dose toxicity.

# **6. PHARMACEUTICAL PARTICULARS**

## **6.1 List of excipients**

### Powder

Sucrose

Dextran

Sorbitol (E420)

Amino acids (including phenylalanine)

Dulbecco's Modified Eagle Medium (DMEM) (containing phenylalanine, sodium, glucose and other substances)

### Solvent

Calcium carbonate

Xanthan gum

Sterile water

## 6.2 Incompatibilities

In the absence of compatibility studies, this medicinal product must not be mixed with other medicinal products.

## 6.3 Shelf life

3 years.

### After reconstitution:

After the reconstitution, the vaccine should be administered immediately. If not used immediately, in-use storage should not be longer than 24 hours and at a temperature between 2-25 °C.

## 6.4 Special precautions for storage

Store in a refrigerator (2 °C – 8 °C).  
Do not freeze.

Store in the original package, in order to protect from light.

For storage conditions after reconstitution of the medicinal product, see section 6.3.

## 6.5 Nature and contents of container

1 dose of powder in a glass container (type I glass) with a stopper (rubber butyl)  
1 mL of solvent in an **oral** applicator (type I glass) with a plunger stopper and a protective tip cap (rubber butyl).

Transfer adapter for reconstitution (1/dose) in the following pack sizes:

- pack size of 1 glass container of powder plus 1 **oral** applicator of solvent
- pack size of 5 glass containers of powder plus 5 **oral** applicators of solvent
- pack size of 10 glass containers of powder plus 10 **oral** applicators of solvent
- pack size of 25 glass containers of powder plus 25 **oral** applicators of solvent

Not all pack sizes may be marketed.

## 6.6 Special precautions for disposal and other handling

### Before reconstitution:

A white deposit and clear supernatant is observed upon storage of the **oral** applicator containing the solvent. The solvent should be inspected visually for any foreign particulate matter and/or abnormal physical appearance prior to reconstitution.

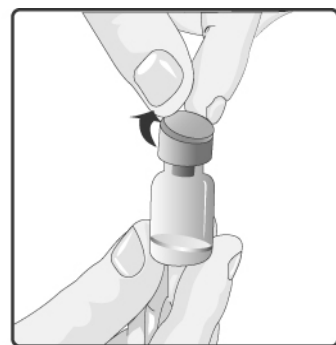
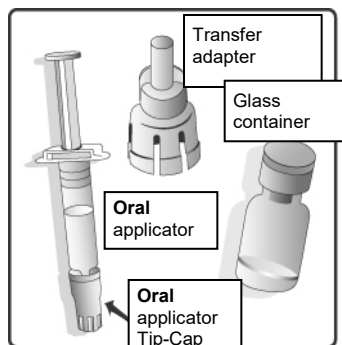
### After reconstitution:

The reconstituted vaccine is slightly more turbid than the solvent and is milky white in appearance.

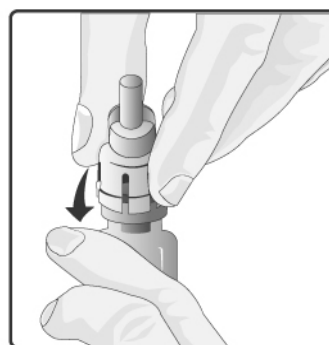
The reconstituted vaccine should also be inspected visually for any foreign particulate matter and/or abnormal physical appearance prior to administration. In the event of either being observed, discard the vaccine.

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

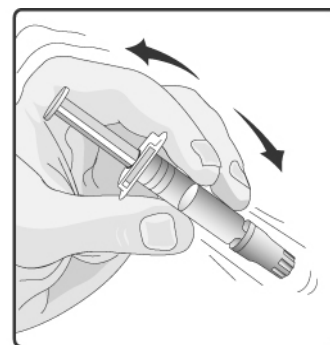
### Instructions for reconstitution and administration of the vaccine:



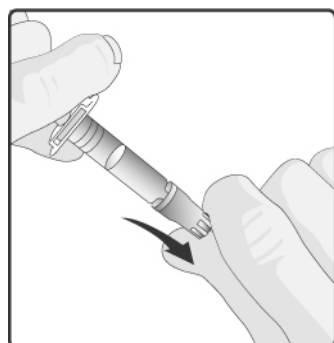
1. Remove the plastic cover from the glass container containing the powder



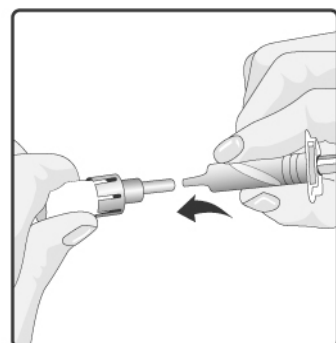
2. Connect the transfer adapter onto the glass container by pushing it downwards until the transfer adapter is properly and securely placed



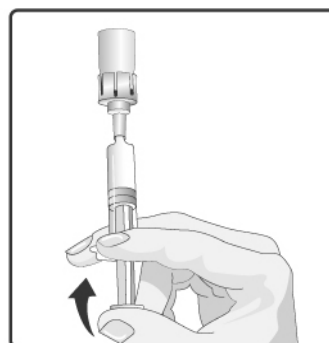
3. Shake the **oral** applicator containing the solvent vigorously. The shaken suspension will appear as a turbid liquid with a slow settling white deposit



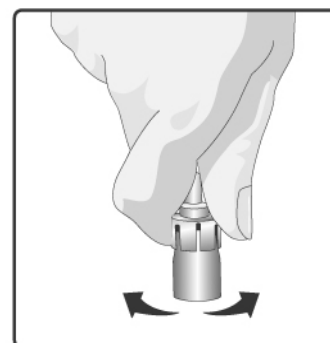
4. Remove the protective tip cap from the **oral** applicator



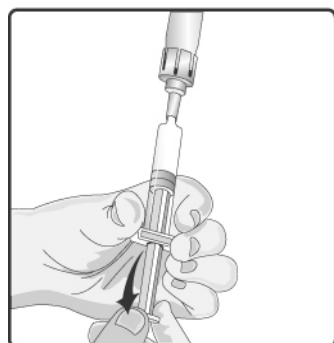
5. Connect the **oral** applicator into the transfer adapter by pushing it firmly on this device



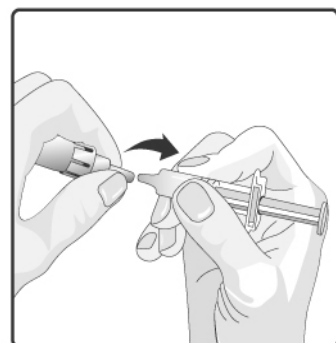
6. Transfer the entire content of the **oral** applicator into the glass container containing the powder



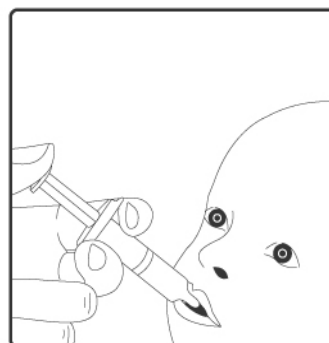
7. With the **oral** applicator still attached, shake the glass container and examine it for complete suspension of the powder. The reconstituted vaccine will appear more turbid than the solvent alone. This appearance is normal



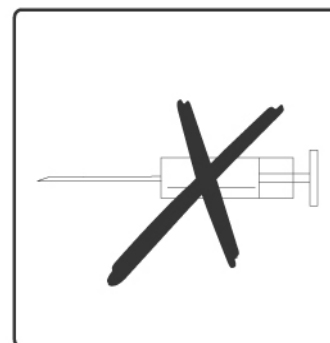
8. Withdraw the entire mixture back into the **oral** applicator



9. Remove the **oral** applicator from the transfer adapter



10. This vaccine is for **oral administration only**. The child should be seated in a reclining position. Administer the entire content of the **oral** applicator **orally** (by administering the entire content of the **oral** applicator on the inside of the cheek)



11. **Do not inject.**

If the reconstituted vaccine is to be stored temporarily before administration, replace the protective tip cap on the **oral** applicator. The **oral** applicator containing the reconstituted vaccine should be shaken gently again before **oral** administration. **Do not inject.**

## 7. MARKETING AUTHORISATION HOLDER

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

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

EU/1/05/330/001

EU/1/05/330/002

EU/1/05/330/003

EU/1/05/330/004

**9.     DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation: 21 February 2006

Date of latest renewal: 14 January 2016

**10.    DATE OF REVISION OF THE TEXT**

Detailed information on this medicinal product is available on the website of the European Medicines Agency <http://www.ema.europa.eu>.

## 1. NAME OF THE MEDICINAL PRODUCT

Rotarix **oral** suspension in pre-filled **oral** applicator

Rotarix **oral** suspension in squeezable tube

Rotarix **oral** suspension in multi-monodose (5 single dose) squeezable tube presentation connected by a bar

Rotavirus vaccine, live

## 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

1 dose (1.5 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated)\* not less than  $10^{6.0}$  CCID<sub>50</sub>

\*Produced on Vero cells

### Excipients with known effect:

This product contains 1 073 mg of sucrose, 32 mg of sodium, 10 micrograms of glucose and 0.15 microgram of phenylalanine per dose (see section 4.4).

For the full list of excipients, see section 6.1.

## 3. PHARMACEUTICAL FORM

**Oral** suspension.

Rotarix is a clear and colourless liquid.

## 4. CLINICAL PARTICULARS

### 4.1 Therapeutic indications

Rotarix is indicated for the active immunisation of infants aged 6 to 24 weeks for prevention of gastroenteritis due to rotavirus infection (see sections 4.2, 4.4 and 5.1).

The use of Rotarix should be based on official recommendations.

### 4.2 Posology and method of administration

#### Posology

The vaccination course consists of two doses. The first dose may be administered from the age of 6 weeks. There should be an interval of at least 4 weeks between doses. The vaccination course should preferably be given before 16 weeks of age, but must be completed by the age of 24 weeks.

Rotarix may be given with the same posology to preterm infants born after at least 27 weeks of gestational age (see sections 4.8 and 5.1).

In clinical trials, spitting or regurgitation of the vaccine has rarely been observed and, under such circumstances, a replacement dose was not given. However, in the unlikely event that an infant spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

It is recommended that infants who receive a first dose of Rotarix complete the 2-dose regimen with Rotarix. There are no data on safety, immunogenicity or efficacy when Rotarix is administered for the first dose and another rotavirus vaccine is administered for the second dose or vice versa.

#### *Paediatric population*



Rotarix should not be used in children over 24 weeks of age.

#### Method of administration

Rotarix is for **oral** use only.

### **Rotarix should under no circumstances be injected.**

For instructions for administration, see section 6.6.

#### **4.3 Contraindications**

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

Hypersensitivity after previous administration of rotavirus vaccines.

History of intussusception.

Subjects with uncorrected congenital malformation of the gastrointestinal tract that would predispose for intussusception.

Subjects with Severe Combined Immunodeficiency (SCID) disorder (see section 4.8).

Administration of Rotarix should be postponed in subjects suffering from acute severe febrile illness. The presence of a minor infection is not a contra-indication for immunisation.

The administration of Rotarix should be postponed in subjects suffering from diarrhoea or vomiting.

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

It is good clinical practice that vaccination should be preceded by a review of the medical history especially with regard to the contraindications and by a clinical examination.

There are no data on the safety and efficacy of Rotarix in infants with gastrointestinal illnesses or growth retardation. Administration of Rotarix may be considered with caution in such infants when, in the opinion of the physician, withholding the vaccine entails a greater risk.

As a precaution, healthcare professionals should follow-up on any symptoms indicative of intussusception (severe abdominal pain, persistent vomiting, bloody stools, abdominal bloating and/or high fever) since data from observational safety studies indicate an increased risk of intussusception, mostly within 7 days after rotavirus vaccination (see section 4.8). Parents/guardians should be advised to promptly report such symptoms to their healthcare provider.

For subjects with a predisposition for intussusception, see section 4.3.

Asymptomatic and mildly symptomatic HIV infections are not expected to affect the safety or efficacy of Rotarix. A clinical study in a limited number of asymptomatic or mildly symptomatic HIV positive infants showed no apparent safety problems (see section 4.8).

Administration of Rotarix to infants who have known or suspected immunodeficiency, including *in utero* exposure to an immunosuppressive treatment, should be based on careful consideration of potential benefits and risks.

Excretion of the vaccine virus in the stools is known to occur after vaccination with peak excretion around the 7th day. Viral antigen particles detected by ELISA were found in 50% of stools after the first dose of Rotarix lyophilised formulation and 4% of stools after the second dose. When these stools were tested for the presence of live vaccine strain, only 17% were positive. In two comparative

controlled trials, vaccine shedding after vaccination with Rotarix liquid formulation was comparable to that observed after vaccination with Rotarix lyophilised formulation.

Cases of transmission of this excreted vaccine virus to seronegative contacts of vaccinees have been observed without causing any clinical symptom.

Rotarix should be administered with caution to individuals with immunodeficient close contacts, such as individuals with malignancies, or who are otherwise immunocompromised or individuals receiving immunosuppressive therapy.

Contacts of recent vaccinees should observe personal hygiene (e.g. wash their hands after changing child's nappies).

The potential risk of apnoea and the need for respiratory monitoring for 48-72h should be considered when administering the primary immunisation series to very premature infants (born  $\leq$  28 weeks of gestation) and particularly for those with a previous history of respiratory immaturity.

As the benefit of the vaccination is high in this group of infants, vaccination should not be withheld or delayed.

A protective immune response may not be elicited in all vaccinees (see section 5.1).

The extent of protection that Rotarix might provide against other rotavirus strains that have not been circulating in clinical trials is currently unknown. Clinical studies from which efficacy data were derived were conducted in Europe, Central and South America, Africa and Asia (see section 5.1).

Rotarix does not protect against gastro-enteritis due to other pathogens than rotavirus.

No data are available on the use of Rotarix for post-exposure prophylaxis.

### **Rotarix should under no circumstances be injected.**

#### Excipients

This vaccine contains sucrose and glucose as excipients. Patients with rare hereditary problems of fructose intolerance, glucose-galactose malabsorption or sucrase-isomaltase insufficiency should not take this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful for patients with phenylketonuria (PKU).

This vaccine contains 32 mg sodium in each dose.

#### Traceability

In order to improve the traceability of biological medicinal products, the name and the batch number of the administered product should be clearly recorded.

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

Rotarix can be given concomitantly with any of the following monovalent or combination vaccines [including hexavalent vaccines (DTPa-HBV-IPV/Hib)]: diphtheria-tetanus-whole cell pertussis vaccine (DTPw), diphtheria-tetanus-acellular pertussis vaccine (DTPa), *Haemophilus influenzae* type b vaccine (Hib), inactivated polio vaccine (IPV), hepatitis B vaccine (HBV), pneumococcal conjugate vaccine and meningococcal serogroup C conjugate vaccine. Clinical studies demonstrated that the immune responses and the safety profiles of the administered vaccines were unaffected.

Concomitant administration of Rotarix and oral polio vaccine (OPV) does not affect the immune response to the polio antigens. Although concomitant administration of OPV may slightly reduce the immune response to rotavirus vaccine, clinical protection against severe rotavirus gastro-enteritis was shown to be maintained in a clinical trial involving more than 4 200 subjects who received Rotarix concomitantly with OPV.

There are no restrictions on the infant's consumption of food or liquid, either before or after vaccination.

#### **4.6 Fertility, pregnancy and lactation**

Rotarix is not intended for use in adults. There are no data on the use of Rotarix during pregnancy and lactation.

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

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

Not relevant.

#### **4.8 Undesirable effects**

##### **Summary of the safety profile**

The safety profile presented below is based on data from clinical trials conducted with either the lyophilised or the liquid formulation of Rotarix.

In a total of four clinical trials, approximately 3 800 doses of Rotarix liquid formulation were administered to approximately 1 900 infants. Those trials have shown that the safety profile of the liquid formulation is comparable to the lyophilised formulation.

In a total of twenty-three clinical trials, approximately 106 000 doses of Rotarix (lyophilised or liquid formulation) were administered to approximately 51 000 infants.

In three placebo-controlled clinical trials (Finland, India and Bangladesh), in which Rotarix was administered alone (administration of routine paediatric vaccines was staggered), the incidence and severity of the solicited events (collected 8 days post-vaccination), diarrhoea, vomiting, loss of appetite, fever, irritability and cough/runny nose were not significantly different in the group receiving Rotarix when compared to the group receiving placebo. No increase in the incidence or severity of these events was seen with the second dose.

In a pooled analysis from seventeen placebo-controlled clinical trials (Europe, North America, Latin America, Asia, Africa) including trials in which Rotarix was co-administered with routine paediatric vaccines (see section 4.5), the following adverse reactions (collected 31 days post-vaccination) were considered as possibly related to vaccination.

##### **Tabulated list of adverse reactions**

Adverse reactions reported are listed according to the following frequency:

Frequencies are reported as:

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	( $< 1/10\ 000$ )

System Organ Class	Frequency	Adverse reactions
Gastrointestinal disorders	Common	Diarrhoea
	Uncommon	Abdominal pain, flatulence
	Very rare	Intussusception (see section 4.4)
	Not known*	Haematochezia
	Not known*	Gastroenteritis with vaccine viral shedding in infants with Severe Combined Immunodeficiency (SCID) disorder
Skin and subcutaneous tissue disorders	Uncommon	Dermatitis
	Very rare	Urticaria
General disorders and administration site conditions	Common	Irritability
Respiratory, thoracic and mediastinal disorders	Not known*	Apnoea in very premature infants ( $\leq 28$ weeks of gestation) (see section 4.4)

\* Because these events were reported spontaneously, it is not possible to reliably estimate their frequency.

## Description of selected adverse reactions

### Intussusception

Data from observational safety studies performed in several countries indicate that rotavirus vaccines carry an increased risk of intussusception, mostly within 7 days of vaccination. Up to 6 additional cases per 100 000 infants have been observed in these countries against a background incidence of 25 to 101 per 100 000 infants (less than one year of age) per year, respectively.

There is limited evidence of a smaller increased risk following the second dose.

It remains unclear whether rotavirus vaccines affect the overall incidence of intussusception based on longer periods of follow-up (see section 4.4).

## Other special populations

### Safety in preterm infants

In a clinical study, 670 pre-term infants from 27 to 36 weeks of gestational age were administered Rotarix lyophilised formulation and 339 received placebo. The first dose was administered from 6 weeks after birth. Serious adverse events were observed in 5.1% of recipients of Rotarix as compared with 6.8% of placebo recipients. Similar rates of other adverse events were observed in Rotarix and placebo recipients. No cases of intussusception were reported.

### Safety in infants with human immunodeficiency (HIV) infection

In a clinical study, 100 infants with HIV infection were administered Rotarix lyophilised formulation or placebo. The safety profile was similar between Rotarix and placebo recipients.

### 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 the national reporting system listed in [Appendix V](#).

## 4.9 Overdose

Some cases of overdose have been reported. In general, the adverse event profile reported in these cases was similar to that observed after administration of the recommended dose of Rotarix.

## 5. PHARMACOLOGICAL PROPERTIES

### 5.1 Pharmacodynamic properties

Pharmaco-therapeutic group: rotavirus diarrhoea vaccines, ATC code: J07BH01

#### Protective efficacy of the lyophilised formulation

In clinical trials, efficacy was demonstrated against gastro-enteritis due to rotavirus of the most common genotypes G1P[8], G2P[4], G3P[8], G4P[8] and G9P[8]. In addition, efficacy against uncommon rotavirus genotypes G8P[4] (severe gastro-enteritis) and G12P[6] (any gastro-enteritis) has been demonstrated. These strains are circulating worldwide.

Clinical studies have been conducted in Europe, Latin America, Africa and Asia to evaluate the protective efficacy of Rotarix against any and severe rotavirus gastro-enteritis (RVGE).

Severity of gastro-enteritis was defined according to two different criteria:

- the Vesikari 20-point scale, which evaluates the full clinical picture of rotavirus gastro-enteritis by taking into account the severity and duration of diarrhoea and vomiting, the severity of fever and dehydration as well as the need for treatment

or

- the clinical case definition based on World Health Organization (WHO) criteria

Clinical protection was assessed in the ATP cohort for efficacy, which includes all subjects from the ATP cohort for safety who entered into the concerned efficacy follow-up period.

#### Protective efficacy in Europe

A clinical study performed in Europe evaluated Rotarix given according to different European schedules (2, 3 months; 2, 4 months; 3, 4 months; 3, 5 months) in 4 000 subjects.

After two doses of Rotarix, the protective vaccine efficacy observed during the first and second year of life is presented in the following table:

		1 <sup>st</sup> year of life Rotarix N=2 572 Placebo N=1 302		2 <sup>nd</sup> year of life Rotarix N=2 554 Placebo N=1 294	
Vaccine efficacy (%) against any and severe rotavirus gastro-enteritis [95% CI]					
Genotype	Any severity	Severe <sup>†</sup>	Any severity	Severe <sup>†</sup>	
G1P[8]	95.6 [87.9;98.8]	96.4 [85.7;99.6]	82.7 [67.8;91.3]	96.5 [86.2;99.6]	
G2P[4]	62.0* [<0.0;94.4]	74.7* [<0.0;99.6]	57.1 [<0.0;82.6]	89.9 [9.4;99.8]	
G3P[8]	89.9 [9.5;99.8]	100 [44.8;100]	79.7 [<0.0;98.1]	83.1* [<0.0;99.7]	
G4P[8]	88.3 [57.5;97.9]	100 [64.9;100]	69.6* [<0.0;95.3]	87.3 [<0.0;99.7]	
G9P[8]	75.6 [51.1;88.5]	94.7 [77.9;99.4]	70.5 [50.7;82.8]	76.8 [50.8;89.7]	
Strains with P[8] genotype	88.2 [80.8;93.0]	96.5 [90.6;99.1]	75.7 [65.0;83.4]	87.5 [77.8;93.4]	

Circulating rotavirus strains	87.1 [79.6;92.1]	95.8 [89.6;98.7]	71.9 [61.2;79.8]	85.6 [75.8;91.9]
<b>Vaccine efficacy (%) against rotavirus gastro-enteritis requiring medical attention [95% CI]</b>				
Circulating rotavirus strains	91.8 [84;96.3]		76.2 [63.0;85.0]	
<b>Vaccine efficacy (%) against hospitalisation due to rotavirus gastro-enteritis [95% CI]</b>				
Circulating rotavirus strains	100 [81.8;100]		92.2 [65.6;99.1]	

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

Vaccine efficacy during the first year of life progressively increased with increasing disease severity, reaching 100% (95% CI: 84.7;100) for Vesikari scores  $\geq 17$ .

#### Protective efficacy in Latin America

A clinical study performed in Latin America evaluated Rotarix in more than 20 000 subjects. Severity of gastro-enteritis (GE) was defined according to WHO criteria. The protective vaccine efficacy against severe rotavirus (RV) gastro-enteritis requiring hospitalisation and/or rehydration therapy in a medical facility and the genotype specific vaccine efficacy after two doses of Rotarix are presented in the table below:

<b>Genotype</b>	<b>Severe rotavirus gastro-enteritis† (1<sup>st</sup> year of life)</b> Rotarix N=9 009 Placebo N=8 858	<b>Severe rotavirus gastro-enteritis† (2<sup>nd</sup> year of life)</b> Rotarix N=7 175 Placebo N=7 062
	<b>Efficacy (%)</b> <b>[95% CI]</b>	<b>Efficacy (%)</b> <b>[95% CI]</b>
All RVGE	84.7 [71.7;92.4]	79.0 [66.4;87.4]
G1P[8]	91.8 [74.1;98.4]	72.4 [34.5;89.9]
G3P[8]	87.7 [8.3;99.7]	71.9* [<0.0;97.1]
G4P[8]	50.8#* [<0.0;99.2]	63.1 [0.7;88.2]
G9P[8]	90.6 [61.7;98.9]	87.7 [72.9;95.3]
Strains with P[8] genotype	90.9 [79.2;96.8]	79.5 [67.0;87.9]

† Severe rotavirus gastro-enteritis was defined as an episode of diarrhoea with or without vomiting that required hospitalisation and/or re-hydration therapy in a medical facility (WHO criteria)

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution

# The numbers of cases, on which the estimates of efficacy against G4P[8] were based, were very small (1 case in the Rotarix group and 2 cases in the placebo group)

A pooled analysis of five efficacy studies\*, showed a 71.4% (95% CI: 20.1;91.1) efficacy against severe rotavirus gastro-enteritis (Vesikari score  $\geq 11$ ) caused by rotavirus G2P[4] genotype during the first year of life.

\* In these studies, the point estimates and confidence intervals were respectively: 100% (95% CI: -1 858.0;100), 100% (95% CI: 21.1;100), 45.4% (95% CI: -81.5;86.6), 74.7% (95% CI: -386.2;99.6). No point estimate was available for the remaining study.

### Protective efficacy in Africa

A clinical study performed in Africa (Rotarix: N = 2 974; placebo: N = 1 443) evaluated Rotarix given at approximately 10 and 14 weeks of age (2 doses) or 6, 10 and 14 weeks of age (3 doses). The vaccine efficacy against severe rotavirus gastro-enteritis during the first year of life was 61.2% (95% CI: 44.0;73.2). The protective vaccine efficacy (pooled doses) observed against any and severe rotavirus gastro-enteritis is presented in the following table:

<b>Genotype</b>	<b>Any rotavirus gastro-enteritis</b> Rotarix N=2 974 Placebo N=1 443	<b>Severe rotavirus gastro-enteritis†</b> Rotarix N=2 974 Placebo N=1 443
	<b>Efficacy (%)</b> <b>[95% CI]</b>	<b>Efficacy (%)</b> <b>[95% CI]</b>
G1P[8]	68.3 [53.6;78.5]	56.6 [11.8;78.8]
G2P[4]	49.3 [4.6;73.0]	83.8 [9.6;98.4]
G3P[8]	43.4* [<0.0;83.7]	51.5* [<0.0;96.5]
G8P[4]	38.7* [<0.0;67.8]	63.6 [5.9;86.5]
G9P[8]	41.8* [<0.0;72.3]	56.9* [<0.0;85.5]
G12P[6]	48.0 [9.7;70.0]	55.5* [<0.0; 82.2]
Strains with P[4] genotype	39.3 [7.7;59.9]	70.9 [37.5;87.0]
Strains with P[6] genotype	46.6 [9.4;68.4]	55.2* [<0.0;81.3]
Strains with P[8] genotype	61.0 [47.3;71.2]	59.1 [32.8;75.3]

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

### Sustained efficacy up to 3 years of age in Asia

A clinical study conducted in Asia (Hong Kong, Singapore and Taiwan) (Total vaccinated cohort: Rotarix: N = 5 359; placebo: N = 5 349) evaluated Rotarix given according to different schedules (2, 4 months of age; 3, 4 months of age).

During the first year, significantly fewer subjects in the Rotarix group reported severe rotavirus gastro-enteritis caused by the circulating wild-type RV compared to the placebo group from 2 weeks after Dose 2 up to one year of age (0.0% versus 0.3%), with a vaccine efficacy of 100% (95% CI: 72.2;100).

The protective vaccine efficacy after two doses of Rotarix observed against severe rotavirus gastro-enteritis up to 2 years of age is presented in the following table:

Efficacy up to 2 years of age Rotarix N= 5 263 Placebo N= 5 256	
Vaccine efficacy (%) against severe rotavirus gastro-enteritis [95% CI]	
Genotype	Severe†
G1P[8]	100 [80.8;100]
G2P[4]	100* [<0.0;100]
G3P[8]	94.5 [64.9;99.9]
G9P[8]	91.7 [43.8;99.8]
Strains with P[8] genotype	95.8 [83.8;99.5]
Circulating rotavirus strains	96.1 [85.1;99.5]
Vaccine efficacy (%) against rotavirus gastro-enteritis requiring hospitalisation and/or rehydration therapy in a medical facility [95% CI]	
Circulating rotavirus strains	94.2 [82.2;98.8]

† Severe gastro-enteritis was defined as a score  $\geq 11$  on the Vesikari scale

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

During the third year of life, there were no cases of severe RV gastro-enteritis in the Rotarix group (N=4 222) versus 13 (0.3%) in the placebo group (N=4 185). Vaccine efficacy was 100% (95% CI: 67.5; 100). The severe RV gastro-enteritis cases were due to RV strains G1P[8], G2P[4], G3P[8] and G9P[8]. The incidence of severe RV gastro-enteritis associated with the individual genotypes was too small to allow calculation of efficacy. The efficacy against severe RV gastro-enteritis requiring hospitalisation was 100% (95% CI: 72.4;100).

#### Protective efficacy of the liquid formulation

Since the immune response observed after 2 doses of Rotarix liquid formulation was comparable to the immune response observed after 2 doses of Rotarix lyophilised formulation, the levels of vaccine efficacy observed with the lyophilised formulation can be extrapolated to the liquid formulation.

#### Immune response

The immunologic mechanism by which Rotarix protects against rotavirus gastro-enteritis is not completely understood. A relationship between antibody responses to rotavirus vaccination and protection against rotavirus gastro-enteritis has not been established.

The following table shows the percentage of subjects initially seronegative for rotavirus (IgA antibody titres < 20 U/mL) (by ELISA) with serum anti-rotavirus IgA antibody titres  $\geq 20$  U/mL one to two months after the second dose of vaccine or placebo as observed in different studies with Rotarix lyophilised formulation.

Schedule	Studies conducted in	Vaccine		Placebo	
		N	% $\geq 20$ U/mL [95% CI]	N	% $\geq 20$ U/mL [95% CI]
2, 3 months	France, Germany	239	82.8 [77.5;87.4]	127	8.7 [4.4;15.0]
2, 4 months	Spain	186	85.5 [79.6;90.2]	89	12.4 [6.3;21.0]
3, 5 months	Finland, Italy	180	94.4 [90.0;97.3]	114	3.5 [1.0;8.7]
3, 4 months	Czech Republic	182	84.6 [78.5;89.5]	90	2.2 [0.3;7.8]



<b>2, 3 to 4 months</b>	Latin America; 11 countries	393	77.9% [73.8;81.6]	341	15.1% [11.7;19.0]
<b>10, 14 weeks and 6, 10, 14 weeks (Pooled)</b>	South Africa, Malawi	221	58.4 [51.6;64.9]	111	22.5 [15.1;31.4]

In three comparative controlled trials, the immune response elicited by Rotarix liquid formulation was comparable to the one elicited by Rotarix lyophilised formulation.

#### Immune response in preterm infants

In a clinical study conducted in preterm infants, born after at least 27 weeks of gestational age, the immunogenicity of Rotarix was assessed in a subset of 147 subjects and showed that Rotarix is immunogenic in this population; 85.7% (95% CI: 79.0;90.9) of subjects achieved serum anti-rotavirus IgA antibody titres  $\geq 20$  U/mL (by ELISA) one month after the second dose of vaccine.

#### Effectiveness

In observational studies, vaccine effectiveness was demonstrated against severe gastro-enteritis leading to hospitalisation due to rotavirus of common genotypes G1P[8], G2P[4], G3P[8], G4P[8] and G9P[8] as well as the less common rotavirus genotypes G9P[4] and G9P[6]. All of these strains are circulating worldwide.

#### Effectiveness after 2 doses in preventing RVGE leading to hospitalisation

Countries Period	Age range	N <sup>(1)</sup> (cases/controls)	Strains	Effectiveness % [95% CI]
High Income countries				
Belgium 2008-2010 <sup>(2)</sup>	< 4 yrs 3-11 m	160/198	All	90 [81;95] 91 [75;97]
	< 4 yrs	41/53	G1P[8]	95 [78;99]
	< 4 yrs 3-11 m	80/103	G2P[4]	85 [64;94] 83 [22;96] <sup>(3)</sup>
	< 4 yrs	12/13	G3P[8]	87* [ $<0$ ;98] <sup>(3)</sup>
	< 4 yrs	16/17	G4P[8]	90 [19;99] <sup>(3)</sup>
Singapore 2008-2010 <sup>(2)</sup>	< 5 yrs	136/272 89/89	All G1P[8]	84 [32;96] 91 [30;99]
	< 3 yrs	184/1 623 <sup>(4)</sup>	All G1P[8]	92 [75;98] 95 [69;100]
US 2010-2011	< 2 yrs	85/1 062 <sup>(5)</sup>	All G1P[8] G2P[4]	85 [73;92] 88 [68;95] 88 [68;95]
	8-11 m		All	89 [48;98]
	US 2009-2011		< 5 yrs	74/255 <sup>(4)</sup>
Middle Income Countries				
Bolivia 2010-2011	< 3 yrs 6-11 m	300/974	All	77 [65;84] <sup>(6)</sup> 77 [51;89]
	< 3 yrs 6-11 m		G9P[8]	85 [69;93] 90 [65;97]
	< 3 yrs		G3P[8]	93 [70;98]
			G2P[4]	69 [14;89]
			G9P[6]	87 [19;98]

Brazil 2008-2011	< 2 yrs	115/1 481	All	72 [44;85] <sup>(6)</sup>
			G1P[8]	89 [78;95]
			G2P[4]	76 [64;84]
Brazil 2008-2009 <sup>(2)</sup>	< 3 yrs 3-11 m	249/249 <sup>(5)</sup>	All	76 [58;86] 96 [68;99]
	< 3 yrs 3-11 m	222/222 <sup>(5)</sup>	G2P[4]	75 [57;86] 95 [66;99] <sup>(3)</sup>
El Salvador 2007-2009	< 2 yrs 6-11 m	251/770 <sup>(5)</sup>	All	76 [64;84] <sup>(6)</sup> 83 [68;91]
Guatemala 2012-2013	< 4 yrs	NA <sup>(7)</sup>	All	63 [23;82]
Mexico 2010	< 2 yrs	9/17 <sup>(5)</sup>	G9P[4]	94 [16;100]
<b>Low Income Countries</b>				
Malawi 2012-2014	< 2 yrs	81/286 <sup>(5)</sup>	All	63 [23;83]

m: months

yrs: years

\* Not statistically significant ( $p \geq 0.05$ ). These data should be interpreted with caution.

(1) The number of fully vaccinated (2 doses) and unvaccinated cases and controls is given.

(2) GSK sponsored studies

(3) Data from a post-hoc analysis

(4) Vaccine effectiveness was calculated using rotavirus-negative hospital control participants (estimates from Taiwan were calculated using combined rotavirus-negative hospital control and non-diarrhoea hospital control participants).

(5) Vaccine effectiveness was calculated using neighbourhood controls.

(6) In subjects who did not receive the full course of vaccination, the effectiveness after one dose ranged from 51% (95% CI: 26;67, El Salvador) to 60% (95% CI: 37;75, Brazil).

(7) NA: Not available. Vaccine effectiveness estimate is based on 41 fully vaccinated cases and 175 fully vaccinated controls.

### Impact on mortality<sup>§</sup>

Impact studies with Rotarix conducted in Panama, Brazil and Mexico showed a decrease in all-cause diarrhoea mortality ranging from 17% to 73% in children less than 5 years of age, within 2 to 4 years after vaccine introduction.

### Impact on hospitalisation<sup>§</sup>

In a retrospective database study in Belgium conducted in children 5 years of age and younger, the direct and indirect impact of Rotarix vaccination on rotavirus-related hospitalisation ranged from 64% (95% CI: 49;76) to 80% (95% CI: 77;83) two years after vaccine introduction. Similar studies in Armenia, Australia, Brazil, Canada, El Salvador and Zambia showed a reduction of 45 to 93% between 2 and 4 years after vaccine introduction.

In addition, nine impact studies on all-cause diarrhoea hospitalisation conducted in Africa and Latin America showed a reduction of 14% to 57% between 2 and 5 years after vaccine introduction.

<sup>§</sup>NOTE: Impact studies are meant to establish a temporal relationship but not a causal relationship between the disease and vaccination. Natural fluctuations of the incidence of the disease may also influence the observed temporal effect.

## **5.2 Pharmacokinetic properties**

Not applicable.

### 5.3 Preclinical safety data

Non-clinical data reveal no special hazard for humans based on conventional studies of repeated dose toxicity.

## 6. PHARMACEUTICAL PARTICULARS

### 6.1 List of excipients

Sucrose

Di-sodium Adipate

Dulbecco's Modified Eagle Medium (DMEM) (containing phenylalanine, sodium, glucose, and other substances)

Sterile water

### 6.2 Incompatibilities

In the absence of compatibility studies, this medicinal product must not be mixed with other medicinal products.

### 6.3 Shelf life

- Pre-filled oral applicator: 3 years
- Squeezable tube fitted with a membrane and a tube cap: 3 years
- Multi-monodose (5 single dose) squeezable tube presentation connected by a bar: 2 years

The vaccine should be used immediately after opening.

### 6.4 Special precautions for storage

Store in a refrigerator (2°C – 8°C).

Do not freeze.

Store in the original package, in order to protect from light.

### 6.5 Nature and contents of container

#### Pre-filled oral applicator

1.5 mL of **oral** suspension in a pre-filled **oral** applicator (type I glass) with a plunger stopper (rubber butyl) and a protective tip cap (rubber butyl), in pack sizes of 1, 5, 10 or 25.

#### Squeezable tube

1.5 mL of **oral** suspension in a squeezable tube (polyethylene) fitted with a membrane and a tube cap (polypropylene), in pack sizes of 1, 10 or 50.

#### Multi-monodose (5-single dose) squeezable tube presentation connected by a bar

1.5 mL of **oral** suspension in a squeezable tube (polyethylene) presented in a multi-monodose (5-single dose) squeezable tube presentation connected by a bar, in pack size of 50 tubes.

Not all pack sizes may be marketed.

### 6.6 Special precautions for disposal and other handling

The vaccine is presented as a clear, colourless liquid, free of visible particles, for **oral** administration.

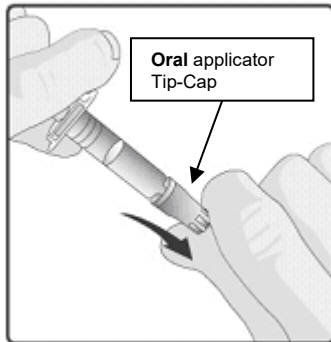
The vaccine is ready to use (no reconstitution or dilution is required).

The vaccine is to be administered **orally** without mixing with any other vaccines or solutions.

The vaccine should be inspected visually for any foreign particulate matter and/or abnormal physical appearance. In the event of either being observed, discard the vaccine.

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

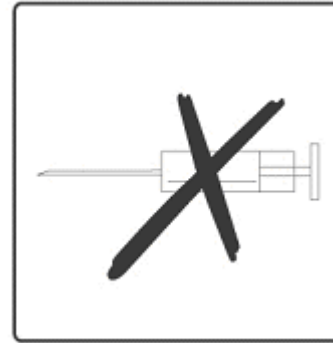
Instructions for administration of the vaccine in a pre-filled oral applicator:



1. Remove the protective tip cap from the oral applicator.



2. This vaccine is for **oral administration only**. The child should be seated in a reclining position. Administer **orally** (i.e. into the child's mouth, towards the inner cheek) the entire content of the **oral** applicator.



3. **Do not inject.**

Discard the empty **oral** applicator and tip cap in approved biological waste containers according to local regulations.

Instructions for administration of the vaccine in a squeezable tube:

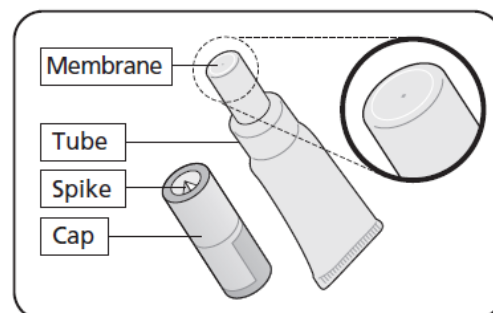
Please read the instructions for use all the way through before starting to give the vaccine.

**A What you need to do before giving Rotarix**

- Check the expiry date.
- Check the tube has not been damaged nor is already open.
- Check the liquid is clear and colourless, without any particles in it.

If you notice anything abnormal, do not use the vaccine.

- This vaccine is given orally - straight from the tube.
- It is ready to use - you do not need to mix it with anything.

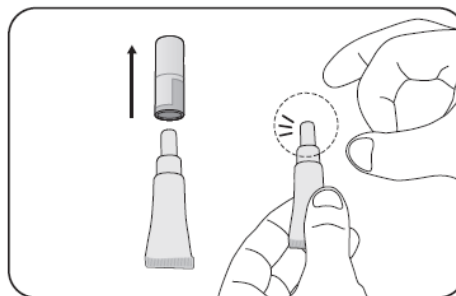


## **B Get the tube ready**

---

### **1. Pull off the cap**

- Keep the cap – you need this to pierce the membrane.
- Hold the tube upright.

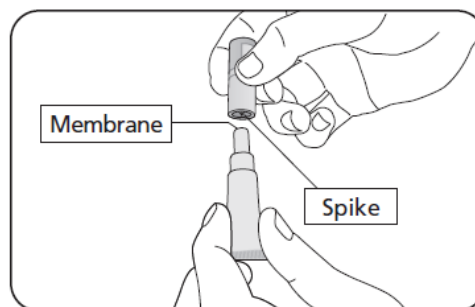


### **2. Repeatedly flick the top of the tube until it is clear of any liquid**

- Clear any liquid from the thinnest section of the tube by flicking just below the membrane.

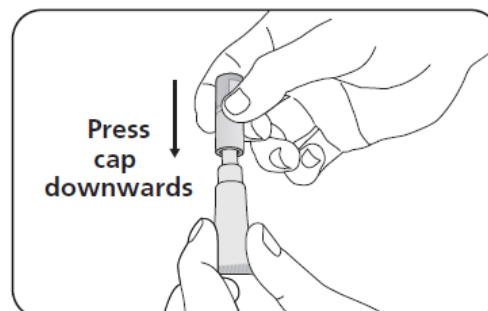
### **3. Position the cap to open the tube**

- Keep the tube held upright.
- Hold the side of tube
- There is a small spike inside the top of the cap - in the centre.
- Turn the cap upside down (180°).



### **4. To open the tube**

- You do not need to twist. Press the cap down to pierce the membrane.
- Then lift off the cap.



## **C Check the tube has opened correctly**

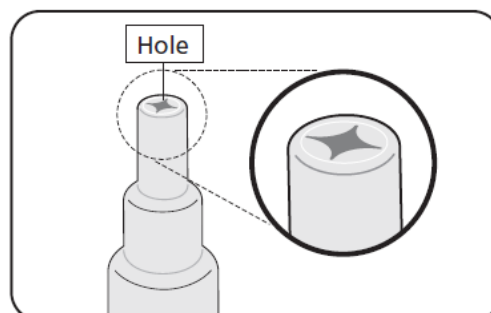
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### **1. Check the membrane has been pierced**

- There should be a hole at the top of the tube.

### **2. What to do if the membrane has not been pierced**

- If the membrane has not been pierced return to section B and repeat steps 2, 3 and 4.



#### D Give the vaccine

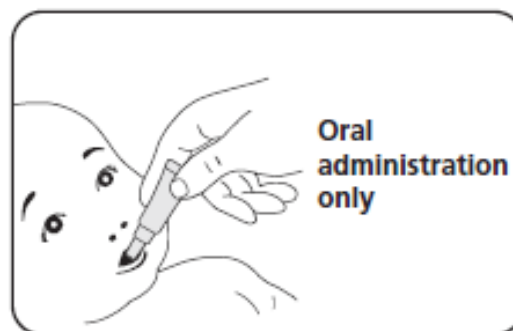
- Once the tube is open check the liquid is clear, without any particles in it.  
If you notice anything abnormal, do not use the vaccine.
- Give the vaccine straight away.

##### 1. Position the child to give the vaccine

- Seat the child leaning slightly backwards.

##### 2. Administer the vaccine

- Squeeze the liquid gently into the side of the child's mouth - towards the inside of their cheek.
- You may need to squeeze the tube a few times to get all of the vaccine out - it is okay if a drop remains in the tip of the tube.



Discard the empty tube and cap in approved biological waste containers according to local regulations.

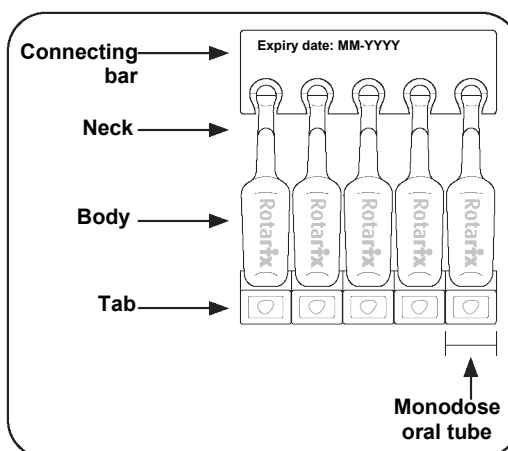
#### Instructions for administration of the vaccine in a multi-monodose (5 single dose) squeezable tube presentation connected by a bar:

Please read the instructions for use all the way through before starting to give the vaccine.

- This vaccine is given orally straight from an individual tube.
- One oral tube delivers one dose of vaccine.
- This vaccine is ready to use - do not mix it with anything else.

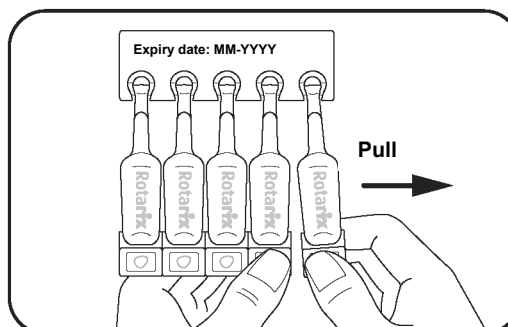
#### A. What you need to do before giving Rotarix

- Check the expiry date on the connecting bar.
- Check the liquid in the oral tubes is clear, colourless and free from any particles.
  - Do not use any of the oral tubes on the connecting bar if you notice anything unusual.**
- Check that each individual oral tube is not damaged and is still sealed.
  - Do not use the affected oral tube if you notice anything unusual.**

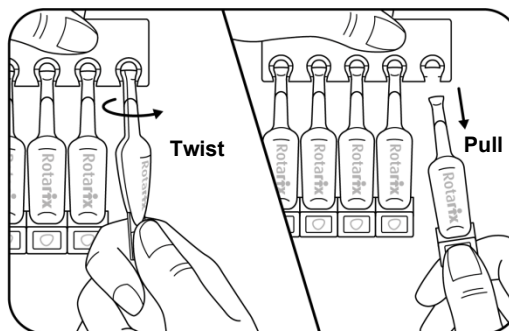


#### B. Get the oral tube ready

- To separate one oral tube from the others starting at one end:
  - Hold the tab of one of the end oral tubes to separate it from the others.
  - With your other hand, hold the tab of the oral tube next to it.
  - Pull the tab and tear it away from the oral tube next to it.

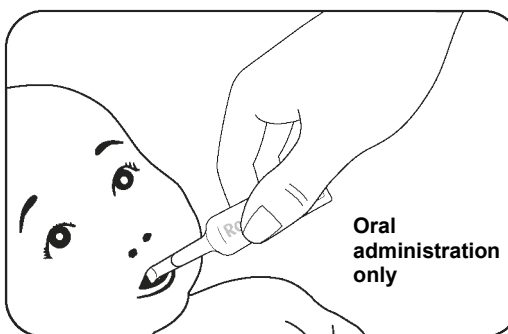


2. To open the separated oral tube:
  - d) Keep the separated oral tube held upright.
  - e) Hold the tab of the separated oral tube in one hand and the connecting bar in the other hand. **Do not hold the body of the oral tube, you may squeeze out some of the vaccine.**
  - f) Twist the separated oral tube.
  - g) Pull it from the connecting bar.



### **C. Give the vaccine orally immediately after opening**

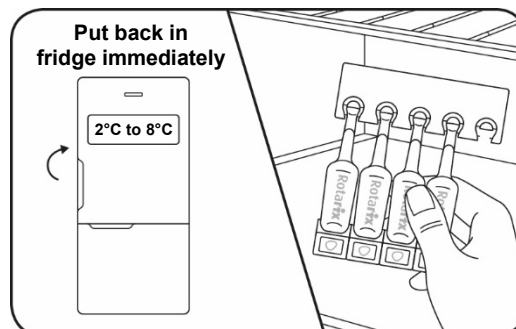
1. To position the child to receive the vaccine:
  - Seat the child leaning slightly backwards.
2. To administer the vaccine orally:
  - Squeeze the liquid gently into the side of the child's mouth, towards the inside of their cheek.
  - You may need to squeeze the oral tube a few times to get all of the vaccine out - it is okay if a drop stays in the oral tube.



### **D. Store remaining doses in the fridge immediately**

Unused oral tubes that are still attached to the connecting bar must be put back in the fridge immediately after an oral tube has been used. This is so that the unused oral tubes can be used for the next vaccination.

Discard the used oral tubes in approved biological waste containers according to local regulations.



## **7. MARKETING AUTHORISATION HOLDER**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

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

### **Pre-filled oral applicator**

EU/1/05/330/005  
EU/1/05/330/006  
EU/1/05/330/007  
EU/1/05/330/008

### **Squeezable tube**

EU/1/05/330/009  
EU/1/05/330/010  
EU/1/05/330/011

Multi-monodose (5-single dose) squeezable tube presentation connected by a bar  
EU/1/05/330/012

**9. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation: 21 February 2006

Date of latest renewal: 14 January 2016

**10. DATE OF REVISION OF THE TEXT**

Detailed information on this medicinal product is available on the website of the European Medicines Agency <http://www.ema.europa.eu>.



## **ANNEX II**

- A. MANUFACTURER(S) OF THE BIOLOGICAL ACTIVE  
SUBSTANCE(S) AND MANUFACTURER(S) RESPONSIBLE  
FOR BATCH RELEASE**
- B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY  
AND USE**
- C. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING  
AUTHORISATION**
- D. CONDITIONS OR RESTRICTIONS WITH REGARD TO THE  
SAFE AND EFFECTIVE USE OF THE MEDICINAL  
PRODUCT**

**A. MANUFACTURER(S) OF THE BIOLOGICAL ACTIVE SUBSTANCE(S) AND MANUFACTURER(S) RESPONSIBLE FOR BATCH RELEASE**

Name and address of the manufacturer(s) of the biological active substance(s)

GlaxoSmithKline Biologicals SA  
Parc de la Noire Epine  
Rue Fleming, 20  
1300 Wavre Belgium

Name and address of the manufacturer(s) responsible for batch release

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
1330 Rixensart  
Belgium

**B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE**

Medicinal product subject to medical prescription.

**Official batch release:**

In accordance with Article 114 of Directive 2001/83/EC, the official batch release will be undertaken by a state laboratory or a laboratory designated for that purpose.

**C. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING AUTHORISATION**

- **Periodic safety update reports (PSURs)**

The requirements for submission of PSURs for this medicinal product are set out in the list of Union reference dates (EURD list) provided for under Article 107c(7) of Directive 2001/83/EC and any subsequent updates published on the European medicines web-portal.

**D. CONDITIONS OR RESTRICTIONS WITH REGARD TO THE SAFE AND EFFECTIVE USE OF THE MEDICINAL PRODUCT**

- **Risk management plan (RMP)**

The marketing authorisation holder (MAH) shall perform the required pharmacovigilance activities and interventions detailed in the agreed RMP presented in Module 1.8.2. of the marketing authorisation and any agreed subsequent updates of the RMP.

An updated RMP should be submitted:

- At the request of the European Medicines Agency;
- Whenever the risk management system is modified, especially as the result of new information being received that may lead to a significant change to the benefit/risk profile or as the result of an important (pharmacovigilance or risk minimisation) milestone being reached.

**ANNEX III**  
**LABELLING AND PACKAGE LEAFLET**

## **A. LABELLING**

**PARTICULARS TO APPEAR ON THE OUTER PACKAGING**  
**GLASS CONTAINER WITH ORAL APPLICATOR AND TRANSFER ADAPTER, PACK**  
**SIZE OF 1, 5, 10 OR 25**

**1. NAME OF THE MEDICINAL PRODUCT**

Rotarix powder and solvent for **oral** suspension  
rotavirus vaccine, live

**2. STATEMENT OF ACTIVE SUBSTANCE(S)**

After reconstitution, 1 dose (1 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated)\* not less than  $10^{6.0}$  CCID<sub>50</sub>

\*Produced on Vero cells

**3. LIST OF EXCIPIENTS**

Powder: sucrose, sorbitol, glucose, phenylalanine  
See the package leaflet for further information

**4. PHARMACEUTICAL FORM AND CONTENTS**

Powder and solvent for **oral** suspension

1 glass container: powder  
1 **oral** applicator: solvent  
1 transfer adapter  
1 dose (1 mL)

5 glass containers: powder  
5 **oral** applicators: solvent  
5 transfer adapters  
5 x 1 dose (1 mL)

10 glass containers: powder  
10 **oral** applicators: solvent  
10 transfer adapters  
10 x 1 dose (1 mL)

25 glass containers: powder  
25 **oral** applicators: solvent  
25 transfer adapters  
25 x 1 dose (1 mL)

**5. METHOD AND ROUTE(S) OF ADMINISTRATION**

**Oral** use  
**Do not inject!**  
Shake before use

Read the package leaflet before use

**6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN**

Keep out of the sight and reach of children

**7. OTHER SPECIAL WARNING(S), IF NECESSARY**

**8. EXPIRY DATE**

Read the leaflet for the shelf-life of the reconstituted medicine.  
EXP {MM/YYYY}

**9. SPECIAL STORAGE CONDITIONS**

Store in a refrigerator  
Do not freeze  
Store in the original package in order to protect from light

**10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE**

**11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

**12. MARKETING AUTHORISATION NUMBER(S)**

EU/1/05/330/001 – pack of 1 (glass container + **oral** applicator + transfer adapter)  
EU/1/05/330/002 – pack of 5 (glass container + **oral** applicator + transfer adapter)  
EU/1/05/330/003 – pack of 10 (glass container + **oral** applicator + transfer adapter)  
EU/1/05/330/004 – pack of 25 (glass container + **oral** applicator + transfer adapter)

**13. BATCH NUMBER**

Lot

**14. GENERAL CLASSIFICATION FOR SUPPLY**

**15. INSTRUCTIONS ON USE**

<b>16. INFORMATION IN BRAILLE</b>
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Justification for not including Braille accepted

<b>17. UNIQUE IDENTIFIER – 2D BARCODE</b>
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2D barcode carrying the unique identifier included.

<b>18. UNIQUE IDENTIFIER - HUMAN READABLE DATA</b>
--

PC  
SN  
NN

**MINIMUM PARTICULARS TO APPEAR ON BLISTERS  
ORAL APPLICATOR WITH SOLVENT FOR RECONSTITUTION WITH POWDER**

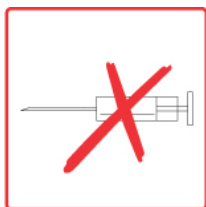
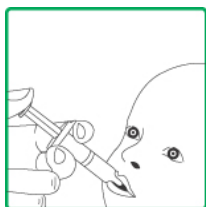
**1. NAME OF THE MEDICINAL PRODUCT**

**2. NAME OF THE MARKETING AUTHORISATION HOLDER**

**3. EXPIRY DATE**

**4. BATCH NUMBER**

**5. OTHER**





<b>MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS GLASS CONTAINER WITH POWDER TO BE RECONSTITUTED WITH SOLVENT</b>
--

<b>1. NAME OF THE MEDICINAL PRODUCT AND ROUTE(S) OF ADMINISTRATION</b>
--

Rotarix  
Powder for **oral** suspension  
rotavirus vaccine, live  
**Oral** use

<b>2. METHOD OF ADMINISTRATION</b>
------------------------------------

<b>3. EXPIRY DATE</b>
-----------------------

EXP

<b>4. BATCH NUMBER</b>
------------------------

Lot

<b>5. CONTENTS BY WEIGHT, BY VOLUME OR BY UNIT</b>
--

1 dose

<b>6. OTHER</b>
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<b>MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS ORAL APPLICATOR WITH SOLVENT FOR RECONSTITUTION WITH POWDER</b>
---

<b>1. NAME OF THE MEDICINAL PRODUCT AND ROUTE(S) OF ADMINISTRATION</b>
--

Solvent for Rotarix  
**Oral** use

<b>2. METHOD OF ADMINISTRATION</b>
------------------------------------

<b>3. EXPIRY DATE</b>
-----------------------

EXP

<b>4. BATCH NUMBER</b>
------------------------

Lot

<b>5. CONTENTS BY WEIGHT, BY VOLUME OR BY UNIT</b>
--

1 dose (1 mL)

<b>6. OTHER</b>
-----------------

**PARTICULARS TO APPEAR ON THE OUTER PACKAGING  
PRE-FILLED ORAL APPLICATOR, PACK SIZE OF 1, 5, 10 OR 25**

**1. NAME OF THE MEDICINAL PRODUCT**

Rotarix **oral** suspension in pre-filled **oral** applicator  
rotavirus vaccine, live

**2. STATEMENT OF ACTIVE SUBSTANCE(S)**

1 dose (1.5 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated) not less than  $10^{6.0}$  CCID<sub>50</sub>

**3. LIST OF EXCIPIENTS**

Sucrose, sodium, glucose, phenylalanine

See the package leaflet for further information

**4. PHARMACEUTICAL FORM AND CONTENTS**

**Oral** suspension in pre-filled **oral** applicator  
1 pre-filled **oral** applicator  
1 dose (1.5 mL)

5 pre-filled **oral** applicators  
5 x 1 dose (1.5 mL)

10 pre-filled **oral** applicators  
10 x 1 dose (1.5 mL)

25 pre-filled **oral** applicators  
25 x 1 dose (1.5 mL)

**5. METHOD AND ROUTE(S) OF ADMINISTRATION**

**Oral** use  
**Do not inject!**  
Read the package leaflet before use

**6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN**

Keep out of the sight and reach of children

**7. OTHER SPECIAL WARNING(S), IF NECESSARY**

Ready to use  
No reconstitution required

**8. EXPIRY DATE**

EXP {MM/YYYY}

**9. SPECIAL STORAGE CONDITIONS**

Store in a refrigerator  
Do not freeze  
Store in the original package in order to protect from light

**10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE**

**11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

**12. MARKETING AUTHORISATION NUMBER(S)**

EU/1/05/330/005 – pack of 1 pre-filled **oral** applicator  
EU/1/05/330/006 – pack of 5 pre-filled **oral** applicators  
EU/1/05/330/007 – pack of 10 pre-filled **oral** applicators  
EU/1/05/330/008 – pack of 25 pre-filled **oral** applicators

**13. BATCH NUMBER**

Lot

**14. GENERAL CLASSIFICATION FOR SUPPLY**

**15. INSTRUCTIONS ON USE**

**16. INFORMATION IN BRAILLE**

Justification for not including Braille accepted

**17. UNIQUE IDENTIFIER – 2D BARCODE**

2D barcode carrying the unique identifier included.

<b>18.    UNIQUE IDENTIFIER - HUMAN READABLE DATA</b>
---

PC  
SN  
NN

**PARTICULARS TO APPEAR ON THE OUTER PACKAGING  
TUBE, PACK SIZES OF 1, 10 OR 50**

**1. NAME OF THE MEDICINAL PRODUCT**

Rotarix **oral** suspension  
rotavirus vaccine, live

**2. STATEMENT OF ACTIVE SUBSTANCE(S)**

1 dose (1.5 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated) not less than  $10^{6.0}$  CCID<sub>50</sub>

**3. LIST OF EXCIPIENTS**

Sucrose, sodium, glucose, phenylalanine  
See the package leaflet for further information

**4. PHARMACEUTICAL FORM AND CONTENTS**

**Oral** suspension  
1 tube  
1 dose (1.5 mL)

10 tubes  
10 x 1 dose (1.5 mL)

50 tubes  
50 x 1 dose (1.5 mL)

**5. METHOD AND ROUTE(S) OF ADMINISTRATION**

**Oral** use  
**Do not inject!**  
Read the package leaflet before use

**6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN**

Keep out of the sight and reach of children

**7. OTHER SPECIAL WARNING(S), IF NECESSARY**



**Read the instructions before administration of the vaccine.**



This vaccine is intended for **oral administration only**.

**8. EXPIRY DATE**

EXP {MM/YYYY}

**9. SPECIAL STORAGE CONDITIONS**

Store in a refrigerator

Do not freeze

Store in the original package in order to protect from light

**10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE**

**11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

**12. MARKETING AUTHORISATION NUMBER(S)**

EU/1/05/330/009 – pack of 1 tube  
EU/1/05/330/010 – pack of 10 tubes  
EU/1/05/330/011 – pack of 50 tubes

**13. BATCH NUMBER**

Lot

**14. GENERAL CLASSIFICATION FOR SUPPLY**

**15. INSTRUCTIONS ON USE**

**16. INFORMATION IN BRAILLE**

Justification for not including Braille accepted

**17. UNIQUE IDENTIFIER – 2D BARCODE**

2D barcode carrying the unique identifier included.

**18. UNIQUE IDENTIFIER - HUMAN READABLE DATA**

PC  
SN  
NN



**PARTICULARS TO APPEAR ON THE OUTER PACKAGING  
MULTI-MONODOSE (5 SINGLE DOSE) SQUEEZABLE TUBE PRESENTATION  
CONNECTED BY A BAR, PACK SIZE OF 50 TUBES**

**1. NAME OF THE MEDICINAL PRODUCT**

Rotarix **oral** suspension  
rotavirus vaccine, live

**2. STATEMENT OF ACTIVE SUBSTANCE(S)**

1 dose (1.5 mL) contains:

Human rotavirus RIX4414 strain (live, attenuated) not less than  $10^{6.0}$  CCID<sub>50</sub>

**3. LIST OF EXCIPIENTS**

Sucrose, sodium, glucose, phenylalanine  
See the package leaflet for further information

**4. PHARMACEUTICAL FORM AND CONTENTS**

**Oral** suspension

50 tubes  
10 x 5 single dose tubes connected by a bar  
1 dose (1.5 mL)

**5. METHOD AND ROUTE(S) OF ADMINISTRATION**

**Oral** use  
**Do not inject!**  
Read the package leaflet before use.

**6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN**

Keep out of the sight and reach of children.

**7. OTHER SPECIAL WARNING(S), IF NECESSARY**



**Read the instructions before administration of the vaccine.**



This vaccine is intended for **oral administration only**.

**8. EXPIRY DATE**

EXP {MM/YYYY}

**9. SPECIAL STORAGE CONDITIONS**

Store in a refrigerator.

Do not freeze.

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

**10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE**

Dispose of in accordance with local regulations.

**11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart, Belgium

**12. MARKETING AUTHORISATION NUMBER(S)**

EU/1/05/330/012 – pack of 50 tubes

**13. BATCH NUMBER**

Lot

**14. GENERAL CLASSIFICATION FOR SUPPLY**

**15. INSTRUCTIONS ON USE**

**16. INFORMATION IN BRAILLE**

Justification for not including Braille accepted

<b>17. UNIQUE IDENTIFIER – 2D BARCODE</b>
---

2D barcode carrying the unique identifier included.

<b>18. UNIQUE IDENTIFIER - HUMAN READABLE DATA</b>
--

PC  
SN  
NN

**MINIMUM PARTICULARS TO APPEAR ON BLISTERS  
PRE-FILLED ORAL APPLICATOR**

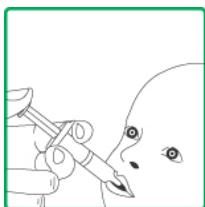
**1. NAME OF THE MEDICINAL PRODUCT**

**2. NAME OF THE MARKETING AUTHORISATION HOLDER**

**3. EXPIRY DATE**

**4. BATCH NUMBER**

**5. OTHER**



<b>MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS PRE-FILLED ORAL APPLICATOR</b>
--

<b>1. NAME OF THE MEDICINAL PRODUCT AND ROUTE(S) OF ADMINISTRATION</b>
--

Rotarix  
**Oral** suspension  
rotavirus vaccine, live  
**Oral** use

<b>2. METHOD OF ADMINISTRATION</b>
------------------------------------

<b>3. EXPIRY DATE</b>
-----------------------

EXP

<b>4. BATCH NUMBER</b>
------------------------

Lot

<b>5. CONTENTS BY WEIGHT, BY VOLUME OR BY UNIT</b>
--

1 dose (1.5 mL)

<b>6. OTHER</b>
-----------------

<b>MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS TUBE</b>
--

<b>1. NAME OF THE MEDICINAL PRODUCT AND ROUTE(S) OF ADMINISTRATION</b>
--

Rotarix  
**Oral** suspension  
rotavirus vaccine, live  
**Oral** use

<b>2. METHOD OF ADMINISTRATION</b>
------------------------------------

<b>3. EXPIRY DATE</b>
-----------------------

EXP

<b>4. BATCH NUMBER</b>
------------------------

Lot

<b>5. CONTENTS BY WEIGHT, BY VOLUME OR BY UNIT</b>
--

1 dose (1.5 mL)

<b>6. OTHER</b>
-----------------

**MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS  
MULTI-MONODOSE (5 SINGLE DOSE) SQUEEZABLE TUBE PRESENTATION  
CONNECTED BY A BAR**

**1. NAME OF THE MEDICINAL PRODUCT AND ROUTE(S) OF ADMINISTRATION**

Rotarix  
**Oral** suspension  
rotavirus vaccine, live  
**Oral** use

**2. METHOD OF ADMINISTRATION**

**3. EXPIRY DATE**

EXP

**4. BATCH NUMBER**

Lot

**5. CONTENTS BY WEIGHT, BY VOLUME OR BY UNIT**

5 single dose tubes  
1 dose (1.5 mL)

**6. OTHER**

## **B. PACKAGE LEAFLET**



## **Package leaflet: Information for the user**

### **Rotarix powder and solvent for oral suspension** rotavirus vaccine, live

**Read all of this leaflet carefully before your child receives this vaccine because it contains important information for you.**

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This vaccine has been prescribed for your child only. Do not pass it on to others.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

#### **What is in this leaflet**

1. What Rotarix is and what it is used for
2. What you need to know before your child receives Rotarix
3. How Rotarix is given
4. Possible side effects
5. How to store Rotarix
6. Contents of the pack and other information

#### **1. What Rotarix is and what it is used for**

Rotarix is a viral vaccine, containing live, attenuated human rotavirus, that helps to protect your child, from the age of 6 weeks, against gastro-enteritis (diarrhoea and vomiting) caused by rotavirus infection.

#### **How Rotarix works**

Rotavirus infection is the most common cause of severe diarrhoea in infants and young children. Rotavirus is easily spread from hand-to-mouth due to contact with stools from an infected person. Most children with rotavirus diarrhoea recover on their own. However, some children become very ill with severe vomiting, diarrhoea and life-threatening loss of fluids that requires hospitalisation.

When a person is given the vaccine, the immune system (the body's natural defences) will make antibodies against the most commonly occurring types of rotavirus. These antibodies protect against disease caused by these types of rotavirus.

As with all vaccines, Rotarix may not completely protect all people who are vaccinated against the rotavirus infections it is intended to prevent.

#### **2. What you need to know before your child receives Rotarix**

##### **Rotarix should not be given**

- if your child has previously had any allergic reaction to rotavirus vaccines or any of the other ingredients of this vaccine (listed in section 6). Signs of an allergic reaction may include itchy skin rash, shortness of breath and swelling of the face or tongue.
- if your child has previously had intussusception (a bowel obstruction in which one segment of bowel becomes enfolded within another segment).
- if your child was born with a malformation of the gut that could lead to intussusception.
- if your child has a rare inherited illness which affects their immune system called Severe Combined Immunodeficiency (SCID).
- if your child has a severe infection with a high temperature. It might be necessary to postpone the vaccination until recovery. A minor infection such as a cold should not be a problem, but talk to your doctor first.

- if your child has diarrhoea or is vomiting. It might be necessary to postpone the vaccination until recovery.

### **Warnings and precautions**

Talk to your doctor/health care professional before your child receives Rotarix if:

- he/she has a close contact such as a household member who has a weakened immune system, e.g., a person with cancer or who is taking medicines that may weaken the immune system.
- he/she has any disorder of the gastrointestinal system.
- he/she has not been gaining weight and growing as expected.
- he/she has any disease or is taking any medicine which reduces his/her resistance to infection or if his/her mother has taken during pregnancy any medicine that may weaken the immune system.

After your child has received Rotarix, contact a doctor/health care professional right away if your child experiences severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever (see also section 4 “Possible side effects”).

As always, please take care to wash your hands thoroughly after changing soiled nappies.

### **Other medicines and Rotarix**

Tell your doctor if your child is taking, has recently taken or might take any other medicines or has recently received any other vaccine.

Rotarix may be given at the same time your child receives other normally recommended vaccines, such as diphtheria, tetanus, pertussis (whooping cough), *Haemophilus influenzae* type b, oral or inactivated polio, hepatitis B vaccines as well as pneumococcal and meningococcal serogroup C conjugate vaccines.

### **Rotarix with food and drink**

There are no restrictions on your child’s consumption of food or liquids, either before or after vaccination.

### **Breast-feeding**

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

### **Rotarix contains sorbitol, sucrose, glucose, phenylalanine and sodium**

This vaccine contains 13.5 mg sorbitol in each dose.

If you have been told by your doctor that the child being vaccinated has an intolerance to some sugars, contact your doctor before your child receives this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful if your child has phenylketonuria (PKU), a rare genetic disorder in which phenylalanine builds up because the body cannot remove it properly.

This vaccine contains less than 1 mmol sodium (23 mg) in each dose, that is to say essentially ‘sodium-free’.

## **3. How Rotarix is given**

The doctor or nurse will administer the recommended dose of Rotarix to your child. The vaccine (1 mL liquid) will be given **orally**. Under no circumstance should this vaccine be administered by injection.

Your child will receive two doses of the vaccine. Each dose will be given on a separate occasion with an interval of at least 4 weeks between the two doses. The first dose may be given from the age of 6

weeks. The two doses of the vaccine must have been given by the age of 24 weeks, although they should preferably have been given before 16 weeks of age.

Rotarix may be given according to the same vaccination course to infants who were born prematurely, provided that the pregnancy had lasted at least 27 weeks.

In case your child spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

When Rotarix is given to your child for the first dose, it is recommended that your child also receives Rotarix (and not another rotavirus vaccine) for the second dose.

It is important that you follow the instructions of your doctor or nurse regarding return visits. If you forget to go back to your doctor at the scheduled time, ask your doctor for advice.

#### **4. Possible side effects**

Like all medicines, this vaccine can cause side effects, although not everybody gets them.

The following side effects may happen with this vaccine:

- ◆ Common (These may occur with up to 1 in 10 doses of the vaccine):
  - diarrhoea
  - irritability
- ◆ Uncommon (These may occur with up to 1 in 100 doses of the vaccine):
  - abdominal pain (see also below for signs of very rare side effects of intussusception)
  - flatulence
  - inflammation of the skin

Side effects that have been reported during marketed use of Rotarix include:

- Very rare: hives (urticaria)
- Very rare: intussusception (part of the intestine gets blocked or twisted). The signs may include severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever. **Contact a doctor/health care professional right away if your child experiences one of these symptoms.**
- blood in stools
- in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2-3 days after vaccination.
- children with a rare inherited illness called Severe Combined Immunodeficiency (SCID) may have an inflamed stomach or gut (gastroenteritis) and pass the vaccine virus in their stools. The signs of gastroenteritis may include feeling sick, being sick, stomach cramps or diarrhoea.

#### **Reporting of side effects**

If your child gets any of the side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via [the national reporting system listed in Appendix V](#). By reporting side effects you can help provide more information on the safety of this medicine.

#### **5. How to store Rotarix**

Keep this vaccine out of the sight and reach of children.

Do not use this vaccine after the expiry date which is stated on the carton. The expiry date refers to the last day of that month.

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

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines your child no longer uses. These measures will help to protect the environment.

## What Rotarix contains

- \*Produced on Vero cells

- Solvent: calcium carbonate, xanthan gum, sterile water

Not all pack sizes may be marketed.

**Lietuva**  
GlaxoSmithKline Biologicals SA  
Tel. +370 800003334

**България**

GlaxoSmithKline Biologicals SA  
Тел. +359 80018205

**Česká republika**

GlaxoSmithKline s.r.o.  
Tel: + 420 2 22 00 11 11  
cz.info@gsk.com

**Danmark**

GlaxoSmithKline Pharma A/S  
Tlf: + 45 36 35 91 00  
dk-info@gsk.com

**Deutschland**

GlaxoSmithKline GmbH & Co. KG  
Tel: + 49 (0)89 360448701  
produkt.info@gsk.com

**Eesti**

GlaxoSmithKline Biologicals SA  
Tel: +372 8002640

**Ελλάδα**

GlaxoSmithKline Μονοπρόσωπη Α.Ε.Β.Ε.  
Τηλ: + 30 210 68 82 100

**España**

GlaxoSmithKline, S.A.  
Tel: + 34 900 202 700  
es-ci@gsk.com

**France**

Laboratoire GlaxoSmithKline  
Tél: + 33 (0) 1 39 17 84 44  
diam@gsk.com

**Hrvatska**

GlaxoSmithKline Biologicals SA  
Tel.: +385 800787089

**Ireland**

GlaxoSmithKline (Ireland) Ltd  
Tel: + 353 (0)1 495 5000

**Ísland**

Vistor hf.  
Sími : +354 535 7000

**Italia****Luxembourg/Luxemburg**

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

**Magyarország**

GlaxoSmithKline Biologicals SA  
Tel.: +36 80088309

**Malta**

GlaxoSmithKline Biologicals SA  
Tel: +356 80065004

**Nederland**

GlaxoSmithKline BV  
Tel: + 31 (0)33 2081100

**Norge**

GlaxoSmithKline AS  
Tlf: + 47 22 70 20 00

**Österreich**

GlaxoSmithKline Pharma GmbH.  
Tel: + 43 (0)1 97075 0  
at.info@gsk.com

**Polska**

GSK Services Sp. z o.o.  
Tel.: + 48 (22) 576 9000

**Portugal**

Smith Kline & French Portuguesa - Produtos  
Farmacêuticos, Lda.  
Tel: + 351 21 412 95 00  
FI.PT@gsk.com

**România**

GlaxoSmithKline Biologicals SA  
Tel: +40 800672524

**Slovenija**

GlaxoSmithKline Biologicals SA  
Tel: +386 80688869

**Slovenská republika**

GlaxoSmithKline Biologicals SA  
Tel: +421 800500589

**Suomi/Finland**

GlaxoSmithKline S.p.A.  
Tel: + 39 (0)45 7741 111

GlaxoSmithKline Oy  
Puh/Tel: + 358 10 30 30 30

**Κύπρος**  
GlaxoSmithKline Biologicals SA  
Τηλ: +357 80070017

**Sverige**  
GlaxoSmithKline AB  
Tel: + 46 (0)8 638 93 00  
info.produkt@gsk.com

**Latvija**  
GlaxoSmithKline Biologicals SA  
Tel: +371 80205045

**United Kingdom (Northern Ireland)**  
GlaxoSmithKline Biologicals SA  
Tel: + 44 (0)800 221 441  
customercontactuk@gsk.com

**This leaflet was last revised in.**

### **Other sources of information**

Detailed information on this medicine is available on the European Medicines Agency web site:  
<http://www.ema.europa.eu>.

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The following information is intended for healthcare professionals only:

Before reconstitution:

A white deposit and clear supernatant is observed upon storage of the **oral** applicator containing the solvent.

The solvent should be inspected visually for any foreign particulate matter and/or abnormal physical appearance prior to reconstitution.

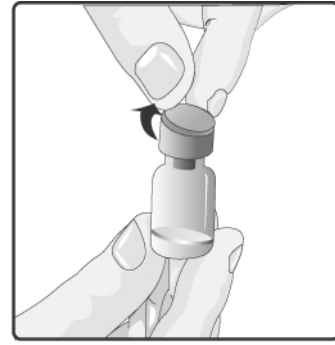
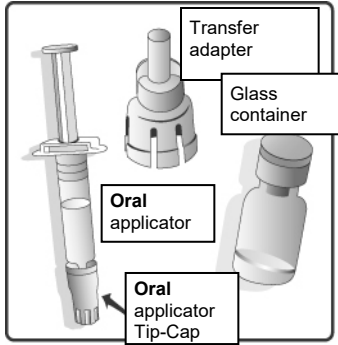
After reconstitution:

The reconstituted vaccine is slightly more turbid than the solvent and is milky white in appearance.

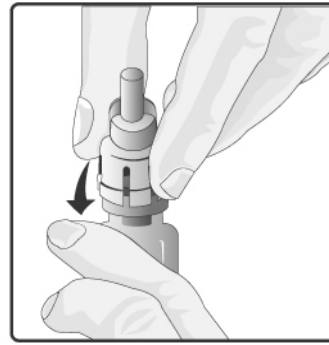
The reconstituted vaccine should also be inspected visually for any foreign particulate matter and/or abnormal physical appearance prior to administration. In the event of either being observed, discard the vaccine.

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

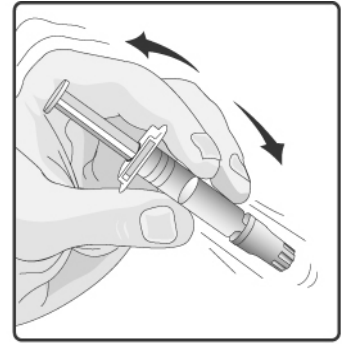
Instructions for reconstitution and administration of the vaccine:



1. Remove the plastic cover from the glass container containing the powder



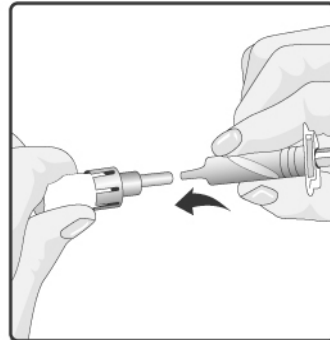
2. Connect the transfer adapter onto the glass container by pushing it downwards until the transfer adapter is properly and securely placed



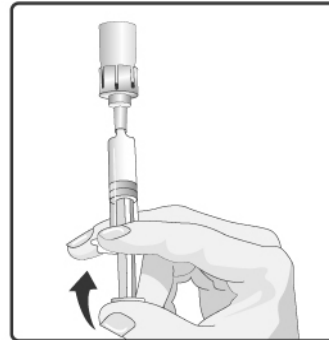
3. Shake the **oral** applicator containing the solvent vigorously. The shaken suspension will appear as a turbid liquid with a slow settling white deposit



4. Remove the protective tip cap from the **oral** applicator



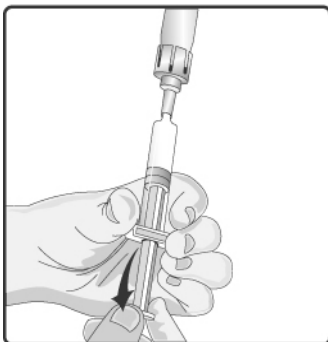
5. Connect the **oral** applicator into the transfer adapter by pushing it firmly on this device



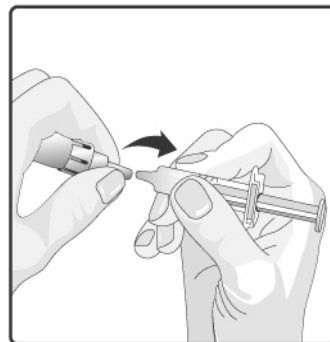
6. Transfer the entire content of the **oral** applicator into the glass container containing the powder



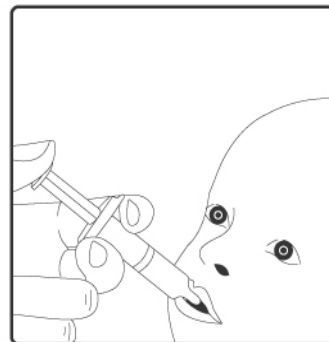
7. With the **oral** applicator still attached, shake the glass container and examine it for complete suspension of the powder. The reconstituted vaccine will appear more turbid than the solvent alone. This appearance is normal



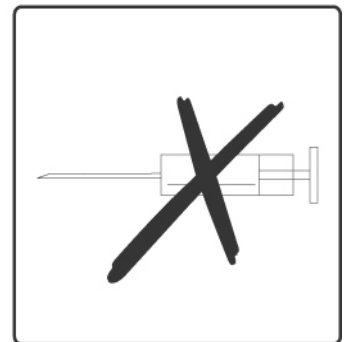
8. Withdraw the entire mixture back into the **oral** applicator



9. Remove the **oral** applicator from the transfer adapter



10. This vaccine is for **oral administration only**. The child should be seated in a reclining position. Administer the entire content of the **oral** applicator **orally** (by administering the entire content of the **oral** applicator on the inside of the cheek)



11. **Do not inject.**

If the reconstituted vaccine is to be stored temporarily before administration, replace the protective tip cap on the **oral** applicator. The **oral** applicator containing the reconstituted vaccine should be shaken gently again before **oral** administration. **Do not inject.**

## **Package leaflet: Information for the user**

### **Rotarix oral suspension in pre-filled oral applicator** rotavirus vaccine, live

**Read all of this leaflet carefully before your child receives this vaccine because it contains important information for you.**

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This vaccine has been prescribed for your child only. Do not pass it on to others.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

#### **What is in this leaflet**

1. What Rotarix is and what it is used for
2. What you need to know before your child receives Rotarix
3. How Rotarix is given
4. Possible side effects
5. How to store Rotarix
6. Contents of the pack and other information

#### **1. What Rotarix is and what it is used for**

Rotarix is a viral vaccine, containing live, attenuated human rotavirus, that helps to protect your child, from the age of 6 weeks, against gastro-enteritis (diarrhoea and vomiting) caused by rotavirus infection.

#### **How Rotarix works**

Rotavirus infection is the most common cause of severe diarrhoea in infants and young children. Rotavirus is easily spread from hand-to-mouth due to contact with stools from an infected person. Most children with rotavirus diarrhoea recover on their own. However, some children become very ill with severe vomiting, diarrhoea and life-threatening loss of fluids that requires hospitalisation.

When a person is given the vaccine, the immune system (the body's natural defences) will make antibodies against the most commonly occurring types of rotavirus. These antibodies protect against disease caused by these types of rotavirus.

As with all vaccines, Rotarix may not completely protect all people who are vaccinated against the rotavirus infections it is intended to prevent.

#### **2. What you need to know before your child receives Rotarix**

##### **Rotarix should not be given**

- if your child has previously had any allergic reaction to rotavirus vaccines or any of the other ingredients of this vaccine (listed in section 6). Signs of an allergic reaction may include itchy skin rash, shortness of breath and swelling of the face or tongue.
- if your child has previously had intussusception (a bowel obstruction in which one segment of bowel becomes enfolded within another segment).
- if your child was born with a malformation of the gut that could lead to intussusception.
- if your child has a rare inherited illness which affects their immune system called Severe Combined Immunodeficiency (SCID).
- if your child has a severe infection with a high temperature. It might be necessary to postpone the vaccination until recovery. A minor infection such as a cold should not be a problem, but talk to your doctor first.



- if your child has diarrhoea or is vomiting. It might be necessary to postpone the vaccination until recovery.

### **Warnings and precautions**

Talk to your doctor/health care professional before your child receives Rotarix if:

- he/she has a close contact such as a household member who has a weakened immune system, e.g., a person with cancer or who is taking medicines that may weaken the immune system.
- he/she has any disorder of the gastrointestinal system.
- he/she has not been gaining weight and growing as expected.
- he/she has any disease or is taking any medicine which reduces his/her resistance to infection or if his/her mother has taken during pregnancy any medicine that may weaken the immune system.

After your child has received Rotarix, contact a doctor/health care professional right away if your child experiences severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever (see also section 4 “Possible side effects”).

As always, please take care to wash your hands thoroughly after changing soiled nappies.

### **Other medicines and Rotarix**

Tell your doctor if your child is taking, has recently taken or might take any other medicines or has recently received any other vaccine.

Rotarix may be given at the same time your child receives other normally recommended vaccines, such as diphtheria, tetanus, pertussis (whooping cough), *Haemophilus influenzae* type b, oral or inactivated polio, hepatitis B vaccines as well as pneumococcal and meningococcal serogroup C conjugate vaccines.

### **Rotarix with food and drink**

There are no restrictions on your child’s consumption of food or liquids, either before or after vaccination.

### **Breast-feeding**

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

### **Rotarix contains sucrose, glucose, phenylalanine and sodium**

If you have been told by your doctor that the child being vaccinated has an intolerance to some sugars, contact your doctor before your child receives this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful if your child has phenylketonuria (PKU), a rare genetic disorder in which phenylalanine builds up because the body cannot remove it properly.

This vaccine contains 32 mg sodium (main component of cooking/table salt) in each dose.

## **3. How Rotarix is given**

The doctor or nurse will administer the recommended dose of Rotarix to your child. The vaccine (1.5 mL liquid) will be given **orally**. Under no circumstance should this vaccine be administered by injection.

Your child will receive two doses of the vaccine. Each dose will be given on a separate occasion with an interval of at least 4 weeks between the two doses. The first dose may be given from the age of 6 weeks. The two doses of the vaccine must have been given by the age of 24 weeks, although they should preferably have been given before 16 weeks of age.

Rotarix may be given according to the same vaccination course to infants who were born prematurely, provided that the pregnancy had lasted at least 27 weeks.

In case your child spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

When Rotarix is given to your child for the first dose, it is recommended that your child also receives Rotarix (and not another rotavirus vaccine) for the second dose.

It is important that you follow the instructions of your doctor or nurse regarding return visits. If you forget to go back to your doctor at the scheduled time, ask your doctor for advice.

#### **4. Possible side effects**

Like all medicines, this vaccine can cause side effects, although not everybody gets them.

The following side effects may happen with this vaccine:

- ◆ Common (These may occur with up to 1 in 10 doses of the vaccine):
  - diarrhoea
  - irritability
- ◆ Uncommon (These may occur with up to 1 in 100 doses of the vaccine):
  - abdominal pain (see also below for signs of very rare side effects of intussusception)
  - flatulence
  - inflammation of the skin

Side effects that have been reported during marketed use of Rotarix include:

- Very rare: hives (urticaria)
- Very rare: intussusception (part of the intestine gets blocked or twisted). The signs may include severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever. **Contact a doctor/health care professional right away if your child experiences one of these symptoms.**
- blood in stools
- in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2-3 days after vaccination.
- children with a rare inherited illness called Severe Combined Immunodeficiency (SCID) may have an inflamed stomach or gut (gastroenteritis) and pass the vaccine virus in their stools. The signs of gastroenteritis may include feeling sick, being sick, stomach cramps or diarrhoea.

#### **Reporting of side effects**

If your child gets any of the side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via **the national reporting system listed in Appendix V**. By reporting side effects you can help provide more information on the safety of this medicine.

#### **5. How to store Rotarix**

Keep this vaccine out of the sight and reach of children.

Do not use this vaccine after the expiry date which is stated on the carton. The expiry date refers to the last day of that month.

Store in a refrigerator (2 °C – 8 °C).

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

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines your child no longer uses. These measures will help to protect the environment.

## What Rotarix contains

- \*Produced on Vero cells

- The other ingredients in Rotarix are: sucrose , Di-sodium Adipate, Dulbecco's Modified Eagle Medium (DMEM) (containing phenylalanine, sodium, glucose, and other substances), sterile water, (see also section 2, "Rotarix contains sucrose, glucose, phenylalanine and sodium")

Not all pack sizes may be marketed.

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart  
Belgium

For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

GlaxoSmithKline Biologicals SA  
Tel. +370 80000334

GlaxoSmithKline Biologics SA  
Тел. +359 80018205

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

GlaxoSmithKline s.r.o.  
Tel: + 420 2 22 00 11 11  
cz.info@gsk.com

GlaxoSmithKline Biologicals SA  
Tel.: +36 80088309

## Malta

GlaxoSmithKline Pharma A/S  
Tlf: + 45 36 35 91 00  
dk-info@gsk.com

**Deutschland**

GlaxoSmithKline GmbH & Co. KG  
Tel: + 49 (0)89 360448701  
produkt.info@gsk.com

**Eesti**

GlaxoSmithKline Biologicals SA  
Tel: +372 8002640

**Ελλάδα**

GlaxoSmithKline Μονοπρόσωπη Α.Ε.Β.Ε.  
Τηλ: + 30 210 68 82 100

**España**

GlaxoSmithKline, S.A.  
Tel: + 34 900 202 700  
es-ci@gsk.com

**France**

Laboratoire GlaxoSmithKline  
Tél: + 33 (0) 1 39 17 84 44  
diam@gsk.com

**Hrvatska**

GlaxoSmithKline Biologicals SA  
Tel.: +385 800787089

**Ireland**

GlaxoSmithKline (Ireland) Ltd  
Tel: + 353 (0)1 495 5000

**Ísland**

Vistor hf.  
Sími : +354 535 7000

**Italia**

GlaxoSmithKline S.p.A.  
Tel:+ 39 (0)45 7741 111

**Κύπρος**

GlaxoSmithKline Biologicals SA  
Τηλ: +357 80070017

**Latvija**

GlaxoSmithKline Biologicals SA  
Tel: +371 80205045

GlaxoSmithKline Biologicals SA  
Tel: +356 80065004

**Nederland**

GlaxoSmithKline BV  
Tel: + 31 (0)33 2081100

**Norge**

GlaxoSmithKline AS  
Tlf: + 47 22 70 20 00

**Österreich**

GlaxoSmithKline Pharma GmbH  
Tel: + 43 (0)1 97075 0  
at.info@gsk.com

**Polska**

GSK Services Sp. z o.o.  
Tel.: + 48 (22) 576 9000

**Portugal**

Smith Kline & French Portuguesa - Produtos  
Farmacêuticos, Lda.  
Tel: + 351 21 412 95 00  
FI.PT@gsk.com

**România**

GlaxoSmithKline Biologicals SA  
Tel: +40 800672524

**Slovenija**

GlaxoSmithKline Biologicals SA  
Tel: +386 80688869

**Slovenská republika**

GlaxoSmithKline Biologicals SA  
Tel: +421 800500589

**Suomi/Finland**

GlaxoSmithKline Oy  
Puh/Tel: + 358 10 30 30 30

**Sverige**

GlaxoSmithKline AB  
Tel: + 46 (0)8 638 93 00  
info.produkt@gsk.com

**United Kingdom (Northern Ireland)**

GlaxoSmithKline Biologicals SA  
Tel: + 44 (0)800 221 441

**This leaflet was last revised in.**

### **Other sources of information**

Detailed information on this medicine is available on the European Medicines Agency web site:  
<http://www.ema.europa.eu>.

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The following information is intended for healthcare professionals only:

The vaccine is presented as a clear, colourless liquid, free of visible particles, for **oral** administration.

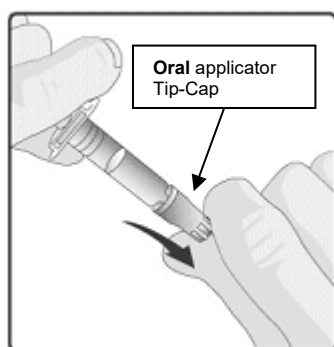
The vaccine is ready to use (no reconstitution or dilution is required).

The vaccine is to be administered **orally** without mixing with any other vaccines or solutions.

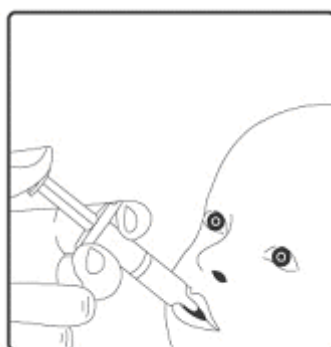
The vaccine should be inspected visually for any foreign particulate matter and/or abnormal physical appearance. In the event of either being observed, discard the vaccine.

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

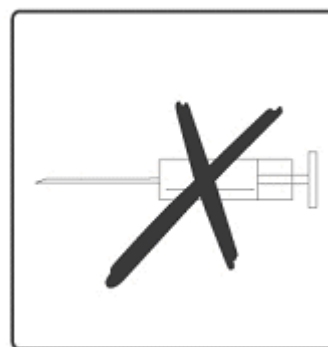
### Instructions for administration of the vaccine:



1. Remove the protective tip cap from the **oral** applicator.



2. This vaccine is for **oral administration only**. The child should be seated in a reclining position. Administer **orally** (i.e. into the child's mouth, towards the inner cheek) the entire content of the **oral** applicator.



3. **Do not inject.**

Discard the empty **oral** applicator and tip cap in approved biological waste containers according to local regulations.

## **Package leaflet: Information for the user**

### **Rotarix oral suspension in squeezable tube** rotavirus vaccine, live

**Read all of this leaflet carefully before your child receives this vaccine because it contains important information for you.**

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This vaccine has been prescribed for your child only. Do not pass it on to others.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

#### **What is in this leaflet**

1. What Rotarix is and what it is used for
2. What you need to know before your child receives Rotarix
3. How Rotarix is given
4. Possible side effects
5. How to store Rotarix
6. Contents of the pack and other information

#### **1. What Rotarix is and what it is used for**

Rotarix is a viral vaccine, containing live, attenuated human rotavirus, that helps to protect your child, from the age of 6 weeks, against gastro-enteritis (diarrhoea and vomiting) caused by rotavirus infection.

#### **How Rotarix works**

Rotavirus infection is the most common cause of severe diarrhoea in infants and young children. Rotavirus is easily spread from hand-to-mouth due to contact with stools from an infected person. Most children with rotavirus diarrhoea recover on their own. However, some children become very ill with severe vomiting, diarrhoea and life-threatening loss of fluids that requires hospitalisation.

When a person is given the vaccine, the immune system (the body's natural defences) will make antibodies against the most commonly occurring types of rotavirus. These antibodies protect against disease caused by these types of rotavirus.

As with all vaccines, Rotarix may not completely protect all people who are vaccinated against the rotavirus infections it is intended to prevent.

#### **2. What you need to know before your child receives Rotarix**

##### **Rotarix should not be given**

- if your child has previously had any allergic reaction to rotavirus vaccines or any of the other ingredients of this vaccine (listed in section 6). Signs of an allergic reaction may include itchy skin rash, shortness of breath and swelling of the face or tongue.
- if your child has previously had intussusception (a bowel obstruction in which one segment of bowel becomes enfolded within another segment).
- if your child was born with a malformation of the gut that could lead to intussusception.
- if your child has a rare inherited illness which affects their immune system called Severe Combined Immunodeficiency (SCID).
- if your child has a severe infection with a high temperature. It might be necessary to postpone the vaccination until recovery. A minor infection such as a cold should not be a problem, but talk to your doctor first.

- if your child has diarrhoea or is vomiting. It might be necessary to postpone the vaccination until recovery.

### **Warnings and precautions**

Talk to your doctor/health care professional before your child receives Rotarix if:

- he/she has a close contact such as a household member who has a weakened immune system, e.g., a person with cancer or who is taking medicines that may weaken the immune system.
- he/she has any disorder of the gastrointestinal system.
- he/she has not been gaining weight and growing as expected.
- he/she has any disease or is taking any medicine which reduces his/her resistance to infection or if his/her mother has taken during pregnancy any medicine that may weaken the immune system.

After your child has received Rotarix, contact a doctor/health care professional right away if your child experiences severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever (see also section 4 “Possible side effects”).

As always, please take care to wash your hands thoroughly after changing soiled nappies.

### **Other medicines and Rotarix**

Tell your doctor if your child is taking, has recently taken or might take any other medicines or has recently received any other vaccine.

Rotarix may be given at the same time your child receives other normally recommended vaccines, such as diphtheria, tetanus, pertussis (whooping cough), *Haemophilus influenzae* type b, oral or inactivated polio, hepatitis B vaccines as well as pneumococcal and meningococcal serogroup C conjugate vaccines.

### **Rotarix with food and drink**

There are no restrictions on your child’s consumption of food or liquids, either before or after vaccination.

### **Breast-feeding**

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

### **Rotarix contains sucrose, glucose, phenylalanine and sodium**

If you have been told by your doctor that the child being vaccinated has an intolerance to some sugars, contact your doctor before your child receives this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful if your child has phenylketonuria (PKU), a rare genetic disorder in which phenylalanine builds up because the body cannot remove it properly.

This vaccine contains 32 mg sodium (main component of cooking/table salt) in each dose.

## **3. How Rotarix is given**

The doctor or nurse will administer the recommended dose of Rotarix to your child. The vaccine (1.5 mL liquid) will be given **orally**. Under no circumstance should this vaccine be administered by injection.

Your child will receive two doses of the vaccine. Each dose will be given on a separate occasion with an interval of at least 4 weeks between the two doses. The first dose may be given from the age of 6 weeks. The two doses of the vaccine must have been given by the age of 24 weeks, although they should preferably have been given before 16 weeks of age.

Rotarix may be given according to the same vaccination course to infants who were born prematurely, provided that the pregnancy had lasted at least 27 weeks.

In case your child spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

When Rotarix is given to your child for the first dose, it is recommended that your child also receives Rotarix (and not another rotavirus vaccine) for the second dose.

It is important that you follow the instructions of your doctor or nurse regarding return visits. If you forget to go back to your doctor at the scheduled time, ask your doctor for advice.

#### 4. Possible side effects

Like all medicines, this vaccine can cause side effects, although not everybody gets them.

The following side effects may happen with this vaccine:

- ◆ Common (These may occur with up to 1 in 10 doses of the vaccine):
  - diarrhoea
  - irritability
- ◆ Uncommon (These may occur with up to 1 in 100 doses of the vaccine):
  - abdominal pain (see also below for signs of very rare side effects of intussusception)
  - flatulence
  - inflammation of the skin

Side effects that have been reported during marketed use of Rotarix include:

- Very rare: hives (urticaria)
- Very rare: intussusception (part of the intestine gets blocked or twisted). The signs may include severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever. **Contact a doctor/health care professional right away if your child experiences one of these symptoms.**
- blood in stools
- in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2-3 days after vaccination.
- children with a rare inherited illness called Severe Combined Immunodeficiency (SCID) may have an inflamed stomach or gut (gastroenteritis) and pass the vaccine virus in their stools. The signs of gastroenteritis may include feeling sick, being sick, stomach cramps or diarrhoea.

#### Reporting of side effects

If your child gets any of the side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via [the national reporting system listed in Appendix V](#). By reporting side effects you can help provide more information on the safety of this medicine.

#### 5. How to store Rotarix

Keep this vaccine out of the sight and reach of children.

Do not use this vaccine after the expiry date which is stated on the carton. The expiry date refers to the last day of that month.

Store in a refrigerator (2 °C – 8 °C).

Do not freeze.



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

The vaccine should be used immediately after opening.

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines your child no longer uses. These measures will help to protect the environment.

## **6. Contents of the pack and other information**

### **What Rotarix contains**

- The active substances are:

Human rotavirus RIX4414 strain (live, attenuated)\* not less than  $10^{6.0}$  CCID<sub>50</sub>

\*Produced on Vero cells

- The other ingredients in Rotarix are: sucrose, Di-sodium Adipate, Dulbecco's Modified Eagle Medium (DMEM) (containing phenylalanine, sodium, glucose, and other substances), sterile water, (see also section 2, "Rotarix contains sucrose, glucose, phenylalanine and sodium")

### **What Rotarix looks like and contents of the pack**

**Oral** suspension.

Rotarix is supplied as clear and colourless liquid in a single dose squeezable tube (1.5 mL).

Rotarix is available in a pack of 1, 10 or 50.

Not all pack sizes may be marketed.

### **Marketing Authorisation Holder and Manufacturer**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart  
Belgium

For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

#### **België/Belgique/Belgien**

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

#### **Lietuva**

GlaxoSmithKline Biologicals SA  
Tel. +370 80000334

#### **България**

GlaxoSmithKline Biologicals SA  
Тел. +359 80018205

#### **Luxembourg/Luxemburg**

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

#### **Česká republika**

GlaxoSmithKline s.r.o.  
Tel: + 420 2 22 00 11 11  
cz.info@gsk.com

#### **Magyarország**

GlaxoSmithKline Biologicals SA  
Tel.: +36 80088309

**Danmark**

GlaxoSmithKline Pharma A/S  
Tlf: + 45 36 35 91 00  
dk-info@gsk.com

**Deutschland**

GlaxoSmithKline GmbH & Co. KG  
Tel: + 49 (0)89 360448701  
produkt.info@gsk.com

**Eesti**

GlaxoSmithKline Biologicals SA  
Tel: +372 8002640

**Ελλάδα**

GlaxoSmithKline Μονοπρόσωπη Α.Ε.Β.Ε.  
Τηλ: + 30 210 68 82 100

**España**

GlaxoSmithKline, S.A.  
Tel: + 34 900 202 700  
es-ci@gsk.com

**France**

Laboratoire GlaxoSmithKline  
Tél: + 33 (0) 1 39 17 84 44  
diam@gsk.com

**Hrvatska**

GlaxoSmithKline Biologicals SA  
Tel.: +385 800787089

**Ireland**

GlaxoSmithKline (Ireland) Ltd  
Tel: + 353 (0)1 495 5000

**Ísland**

Vistor hf.  
Sími : +354 535 7000

**Italia**

GlaxoSmithKline S.p.A.  
Tel: + 39 (0)45 7741 111

**Κύπρος**

GlaxoSmithKline Biologicals SA  
Τηλ: +357 80070017

**Malta**

GlaxoSmithKline Biologicals SA  
Tel: +356 80065004

**Nederland**

GlaxoSmithKline BV  
Tel: + 31 (0)33 2081100

**Norge**

GlaxoSmithKline AS  
Tlf: + 47 22 70 20 00

**Österreich**

GlaxoSmithKline Pharma GmbH  
Tel: + 43 (0)1 97075 0  
at.info@gsk.com

**Polska**

GSK Services Sp. z o.o.  
Tel.: + 48 (22) 576 9000

**Portugal**

Smith Kline & French Portuguesa - Produtos  
Farmacêuticos, Lda.  
Tel: + 351 21 412 95 00  
FI.PT@gsk.com

**România**

GlaxoSmithKline Biologicals SA  
Tel: +40 800672524

**Slovenija**

GlaxoSmithKline Biologicals SA  
Tel: +386 80688869

**Slovenská republika**

GlaxoSmithKline Biologicals SA  
Tel: +421 800500589

**Suomi/Finland**

GlaxoSmithKline Oy  
Puh/Tel: + 358 10 30 30 30

**Sverige**

GlaxoSmithKline AB  
Tel: + 46 (0)8 638 93 00  
info.produkt@gsk.com

**Latvija**  
GlaxoSmithKline Biologicals SA  
Tel: +371 80205045

**United Kingdom (Northern Ireland)**  
GlaxoSmithKline Biologicals SA  
Tel: + 44 (0)800 221 441  
customercontactuk@gsk.com

**This leaflet was last revised in**

### **Other sources of information**

Detailed information on this medicine is available on the European Medicines Agency web site:  
<http://www.ema.europa.eu>.

-----  
The following information is intended for healthcare professionals only:

The vaccine is presented as a clear, colourless liquid, free of visible particles, for **oral** administration.

The vaccine is ready to use (no reconstitution or dilution is required).

The vaccine is to be administered **orally** without mixing with any other vaccines or solutions.

The vaccine should be inspected visually for any foreign particulate matter and/or abnormal physical appearance. In the event of either being observed, discard the vaccine.

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

### Instructions for administration of the vaccine:

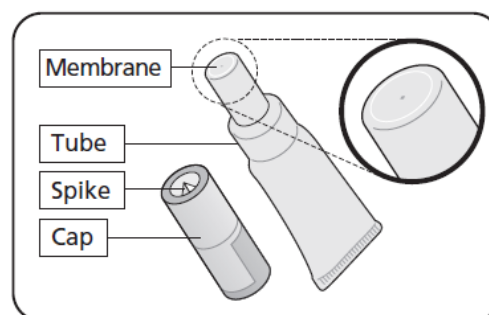
Please read the instructions for use all the way through before starting to give the vaccine.

#### **A What you need to do before giving Rotarix**

- Check the expiry date.
- Check the tube has not been damaged nor is already open.
- Check the liquid is clear and colourless, without any particles in it.

If you notice anything abnormal, do not use the vaccine.

- This vaccine is given orally - straight from the tube.
- It is ready to use - you do not need to mix it with anything.



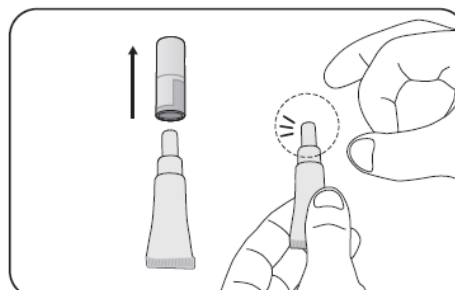
#### **B Get the tube ready**

##### **1. Pull off the cap**

- Keep the cap – you need this to pierce the membrane.
- Hold the tube upright.

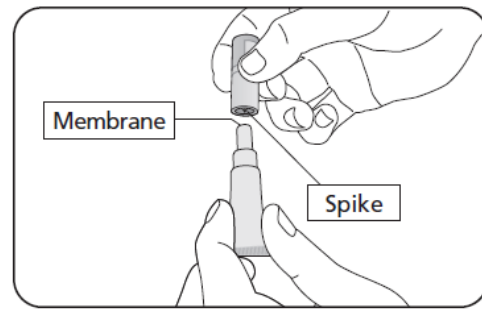
##### **2. Repeatedly flick the top of the tube until it is clear of any liquid**

- Clear any liquid from the thinnest section of the tube by flicking just below the membrane.



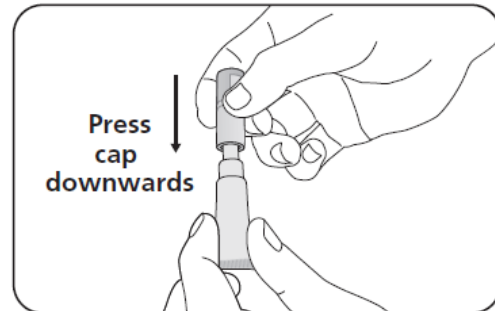
### 3. Position the cap to open the tube

- Keep the tube held upright.
- Hold the side of tube
- There is a small spike inside the top of the cap - in the centre.
- Turn the cap upside down (180°).



### 4. To open the tube

- You do not need to twist. Press the cap down to pierce the membrane.
- Then lift off the cap.



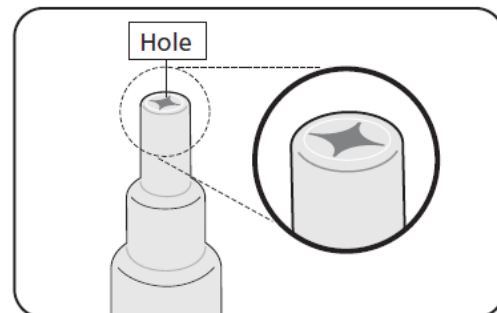
### C Check the tube has opened correctly

#### 1. Check the membrane has been pierced

- There should be a hole at the top of the tube.

#### 2. What to do if the membrane has not been pierced

- If the membrane has not been pierced return to section B and repeat steps 2, 3 and 4.



### D Give the vaccine

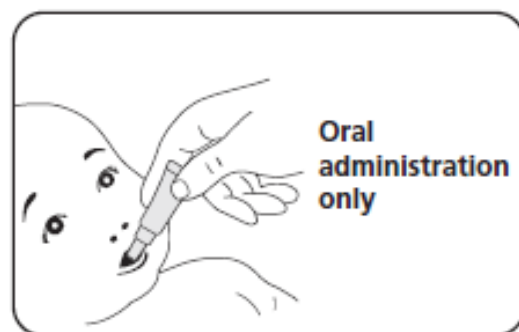
- Once the tube is open check the liquid is clear, without any particles in it.  
If you notice anything abnormal, do not use the vaccine.
- Give the vaccine straight away.

#### 1. Position the child to give the vaccine

- Seat the child leaning slightly backwards.

#### 2. Administer the vaccine

- Squeeze the liquid gently into the side of the child's mouth - towards the inside of their cheek.
- You may need to squeeze the tube a few times to get all of the vaccine out - it is okay if a drop remains in the tip of the tube.



Discard the empty tube and cap in approved biological waste containers according to local regulations.

## **Package leaflet: Information for the user**

### **Rotarix oral suspension in multi-monodose (5 single dose) squeezable tube presentation connected by a bar rotavirus vaccine, live**

**Read all of this leaflet carefully before your child receives this vaccine because it contains important information for you.**

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This vaccine has been prescribed for your child only. Do not pass it on to others.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

#### **What is in this leaflet**

1. What Rotarix is and what it is used for
2. What you need to know before your child receives Rotarix
3. How Rotarix is given
4. Possible side effects
5. How to store Rotarix
6. Contents of the pack and other information

#### **1. What Rotarix is and what it is used for**

Rotarix is a viral vaccine, containing live, attenuated human rotavirus, that helps to protect your child, from the age of 6 weeks, against gastro-enteritis (diarrhoea and vomiting) caused by rotavirus infection.

#### **How Rotarix works**

Rotavirus infection is the most common cause of severe diarrhoea in infants and young children. Rotavirus is easily spread from hand-to-mouth due to contact with stools from an infected person. Most children with rotavirus diarrhoea recover on their own. However, some children become very ill with severe vomiting, diarrhoea and life-threatening loss of fluids that requires hospitalisation.

When a person is given the vaccine, the immune system (the body's natural defences) will make antibodies against the most commonly occurring types of rotavirus. These antibodies protect against disease caused by these types of rotavirus.

As with all vaccines, Rotarix may not completely protect all people who are vaccinated against the rotavirus infections it is intended to prevent.

#### **2. What you need to know before your child receives Rotarix**

##### **Rotarix should not be given**

- if your child has previously had any allergic reaction to rotavirus vaccines or any of the other ingredients of this vaccine (listed in section 6). Signs of an allergic reaction may include itchy skin rash, shortness of breath and swelling of the face or tongue.
- if your child has previously had intussusception (a bowel obstruction in which one segment of bowel becomes enfolded within another segment).
- if your child was born with a malformation of the gut that could lead to intussusception.
- if your child has a rare inherited illness which affects their immune system called Severe Combined Immunodeficiency (SCID).
- if your child has a severe infection with a high temperature. It might be necessary to postpone the vaccination until recovery. A minor infection such as a cold should not be a problem, but talk to your doctor first.

- if your child has diarrhoea or is vomiting. It might be necessary to postpone the vaccination until recovery.

### **Warnings and precautions**

Talk to your doctor/health care professional before your child receives Rotarix if:

- he/she has a close contact such as a household member who has a weakened immune system, e.g., a person with cancer or who is taking medicines that may weaken the immune system.
- he/she has any disorder of the gastrointestinal system.
- he/she has not been gaining weight and growing as expected.
- he/she has any disease or is taking any medicine which reduces his/her resistance to infection or if his/her mother has taken during pregnancy any medicine that may weaken the immune system.

After your child has received Rotarix, contact a doctor/health care professional right away if your child experiences severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever (see also section 4 “Possible side effects”).

As always, please take care to wash your hands thoroughly after changing soiled nappies.

### **Other medicines and Rotarix**

Tell your doctor if your child is taking, has recently taken or might take any other medicines or has recently received any other vaccine.

Rotarix may be given at the same time your child receives other normally recommended vaccines, such as diphtheria, tetanus, pertussis (whooping cough), *Haemophilus influenzae* type b, oral or inactivated polio, hepatitis B vaccines as well as pneumococcal and meningococcal serogroup C conjugate vaccines.

### **Rotarix with food and drink**

There are no restrictions on your child’s consumption of food or liquids, either before or after vaccination.

### **Breast-feeding**

Based on evidence generated in clinical trials, breast-feeding does not reduce the protection against rotavirus gastro-enteritis afforded by Rotarix. Therefore, breast-feeding may be continued during the vaccination schedule.

### **Rotarix contains sucrose, glucose, phenylalanine and sodium**

If you have been told by your doctor that the child being vaccinated has an intolerance to some sugars, contact your doctor before your child receives this vaccine.

This vaccine contains 0.15 microgram phenylalanine in each dose. Phenylalanine may be harmful if your child has phenylketonuria (PKU), a rare genetic disorder in which phenylalanine builds up because the body cannot remove it properly.

This vaccine contains 32 mg sodium (main component of cooking/table salt) in each dose.

## **3. How Rotarix is given**

The doctor or nurse will administer the recommended dose of Rotarix to your child. The vaccine (1.5 mL liquid) will be given **orally**. Under no circumstance should this vaccine be administered by injection.

Your child will receive two doses of the vaccine. Each dose will be given on a separate occasion with an interval of at least 4 weeks between the two doses. The first dose may be given from the age of 6 weeks. The two doses of the vaccine must have been given by the age of 24 weeks, although they should preferably have been given before 16 weeks of age.

Rotarix may be given according to the same vaccination course to infants who were born prematurely, provided that the pregnancy had lasted at least 27 weeks.

In case your child spits out or regurgitates most of the vaccine dose, a single replacement dose may be given at the same vaccination visit.

When Rotarix is given to your child for the first dose, it is recommended that your child also receives Rotarix (and not another rotavirus vaccine) for the second dose.

It is important that you follow the instructions of your doctor or nurse regarding return visits. If you forget to go back to your doctor at the scheduled time, ask your doctor for advice.

#### 4. Possible side effects

Like all medicines, this vaccine can cause side effects, although not everybody gets them.

The following side effects may happen with this vaccine:

- ◆ Common (These may occur with up to 1 in 10 doses of the vaccine):
  - diarrhoea
  - irritability
- ◆ Uncommon (These may occur with up to 1 in 100 doses of the vaccine):
  - abdominal pain (see also below for signs of very rare side effects of intussusception)
  - flatulence
  - inflammation of the skin

Side effects that have been reported during marketed use of Rotarix include:

- Very rare: hives (urticaria)
- Very rare: intussusception (part of the intestine gets blocked or twisted). The signs may include severe stomach pain, persistent vomiting, blood in stools, a swollen belly and/or high fever. **Contact a doctor/health care professional right away if your child experiences one of these symptoms.**
- blood in stools
- in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2-3 days after vaccination.
- children with a rare inherited illness called Severe Combined Immunodeficiency (SCID) may have an inflamed stomach or gut (gastroenteritis) and pass the vaccine virus in their stools. The signs of gastroenteritis may include feeling sick, being sick, stomach cramps or diarrhoea.

#### Reporting of side effects

If your child gets any of the side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via [the national reporting system listed in Appendix V](#). By reporting side effects you can help provide more information on the safety of this medicine.

#### 5. How to store Rotarix

Keep this vaccine out of the sight and reach of children.

Do not use this vaccine after the expiry date which is stated on the carton. The expiry date refers to the last day of that month.

Store in a refrigerator (2 °C – 8 °C).

Do not freeze.

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

The vaccine should be used immediately after opening.

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines your child no longer uses. These measures will help to protect the environment.

## **6. Contents of the pack and other information**

### **What Rotarix contains**

- The active substances are:

Human rotavirus RIX4414 strain (live, attenuated)\* not less than  $10^{6.0}$  CCID<sub>50</sub>

\*Produced on Vero cells

- The other ingredients in Rotarix are: sucrose, Di-sodium Adipate, Dulbecco's Modified Eagle Medium (DMEM) (containing phenylalanine, sodium, glucose and other substances), sterile water, (see also section 2, "Rotarix contains sucrose, glucose, phenylalanine and sodium")

### **What Rotarix looks like and contents of the pack**

**Oral** suspension.

Rotarix is supplied as clear and colourless liquid in 5 single dose squeezable tubes (5 x 1.5 mL) connected by a bar.

Rotarix is available in a pack of 50 tubes.

Not all pack sizes may be marketed.

### **Marketing Authorisation Holder and Manufacturer**

GlaxoSmithKline Biologicals SA  
Rue de l'Institut 89  
B-1330 Rixensart  
Belgium

For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

#### **België/Belgique/Belgien**

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

#### **Lietuva**

GlaxoSmithKline Biologicals SA  
Tel. +370 80000334

#### **България**

GlaxoSmithKline Biologicals SA  
Тел. +359 80018205

#### **Luxembourg/Luxemburg**

GlaxoSmithKline Pharmaceuticals SA/NV  
Tél/Tel: + 32 10 85 52 00

#### **Česká republika**

GlaxoSmithKline s.r.o.  
Tel: + 420 2 22 00 11 11  
cz.info@gsk.com

#### **Magyarország**

GlaxoSmithKline Biologicals SA  
Tel.: +36 80088309



**Danmark**

GlaxoSmithKline Pharma A/S  
Tlf: + 45 36 35 91 00  
dk-info@gsk.com

**Deutschland**

GlaxoSmithKline GmbH & Co. KG  
Tel: + 49 (0)89 360448701  
produkt.info@gsk.com

**Eesti**

GlaxoSmithKline Biologicals SA  
Tel: +372 8002640

**Ελλάδα**

GlaxoSmithKline Μονοπρόσωπη Α.Ε.Β.Ε.  
Τηλ: + 30 210 68 82 100

**España**

GlaxoSmithKline, S.A.  
Tel: + 34 900 202 700  
es-ci@gsk.com

**France**

Laboratoire GlaxoSmithKline  
Tél: + 33 (0) 1 39 17 84 44  
diam@gsk.com

**Hrvatska**

GlaxoSmithKline Biologicals SA  
Tel.: +385 800787089

**Ireland**

GlaxoSmithKline (Ireland) Ltd  
Tel: + 353 (0)1 495 5000

**Ísland**

Vistor hf.  
Sími: +354 535 7000

**Italia**

GlaxoSmithKline S.p.A.  
Tel: + 39 (0)45 7741 111

**Κύπρος**

GlaxoSmithKline Biologicals SA  
Τηλ: +357 80070017

**Malta**

GlaxoSmithKline Biologicals SA  
Tel: +356 80065004

**Nederland**

GlaxoSmithKline BV  
Tel: + 31 (0)33 2081100

**Norge**

GlaxoSmithKline AS  
Tlf: + 47 22 70 20 00

**Österreich**

GlaxoSmithKline Pharma GmbH  
Tel: + 43 (0)1 97075 0  
at.info@gsk.com

**Polska**

GSK Services Sp. z o.o.  
Tel.: + 48 (22) 576 9000

**Portugal**

Smith Kline & French Portuguesa - Produtos  
Farmacêuticos, Lda.  
Tel: + 351 21 412 95 00  
FI.PT@gsk.com

**România**

GlaxoSmithKline Biologicals SA  
Tel: +40 800672524

**Slovenija**

GlaxoSmithKline Biologicals SA  
Tel: +386 80688869

**Slovenská republika**

GlaxoSmithKline Biologicals SA  
Tel.: +421 800500589

**Suomi/Finland**

GlaxoSmithKline Oy  
Puh/Tel: + 358 10 30 30 30

**Sverige**

GlaxoSmithKline AB  
Tel: + 46 (0)8 638 93 00  
info.produkt@gsk.com

**Latvija**  
GlaxoSmithKline Biologicals SA  
Tel: +371 80205045

**United Kingdom (Northern Ireland)**  
GlaxoSmithKline Biologicals SA  
Tel: +44 (0)800 221 441  
customercontactuk@gsk.com

**This leaflet was last revised in.**

### Other sources of information

Detailed information on this medicine is available on the European Medicines Agency web site:  
<http://www.ema.europa.eu>.

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The following information is intended for healthcare professionals only:

The vaccine is presented as a clear, colourless liquid, free of visible particles, for **oral** administration.

The vaccine is ready to use (no reconstitution or dilution is required).

The vaccine is to be administered **orally** without mixing with any other vaccines or solutions.

The vaccine should be inspected visually for any foreign particulate matter and/or abnormal physical appearance. In the event of either being observed, discard the vaccine.

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

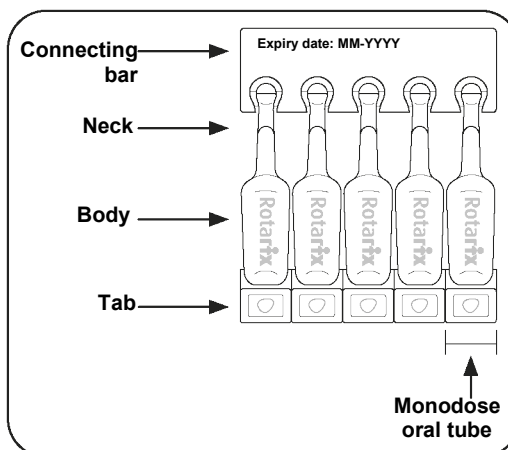
### Instructions for administration of the vaccine:

Please read the instructions for use all the way through before starting to give the vaccine.

- This vaccine is given orally straight from an individual tube.
- One oral tube delivers one dose of vaccine.
- This vaccine is ready to use - do not mix it with anything else.

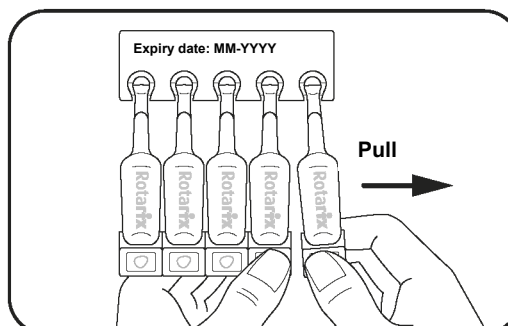
#### **A. What you need to do before giving Rotarix**

1. Check the expiry date on the connecting bar.
2. Check the liquid in the oral tubes is clear, colourless and free from any particles.
  - **Do not use any of the oral tubes on the connecting bar if you notice anything unusual.**
3. Check that each individual oral tube is not damaged and is still sealed.
  - **Do not use the affected oral tube if you notice anything unusual.**

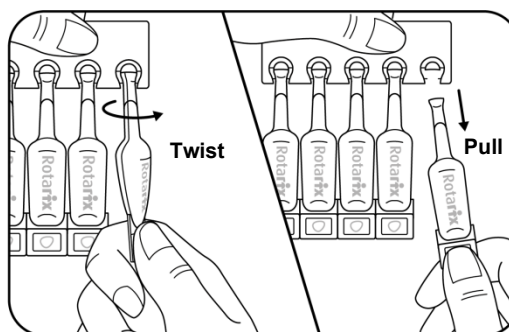


#### **B. Get the oral tube ready**

1. To separate one oral tube from the others starting at one end:
  - a) Hold the tab of one of the end oral tubes to separate it from the others.
  - b) With your other hand, hold the tab of the oral tube next to it.
  - c) Pull the tab and tear it away from the oral tube next to it.

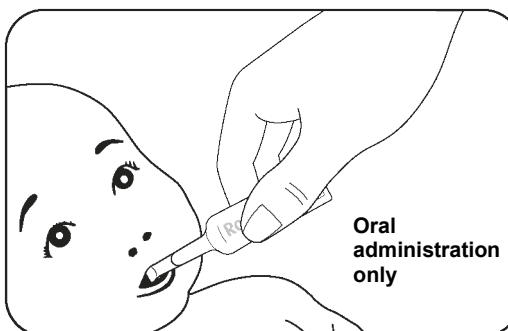


2. To open the separated oral tube:
  - d) Keep the separated oral tube held upright.
  - e) Hold the tab of the separated oral tube in one hand and the connecting bar in the other hand. **Do not hold the body of the oral tube, you may squeeze out some of the vaccine.**
  - f) Twist the separated oral tube.
  - g) Pull it from the connecting bar.



### **C. Give the vaccine orally immediately after opening**

1. To position the child to receive the vaccine:
  - Seat the child leaning slightly backwards.
2. To administer the vaccine orally:
  - Squeeze the liquid gently into the side of the child's mouth, towards the inside of their cheek.
  - You may need to squeeze the oral tube a few times to get all of the vaccine out - it is okay if a drop stays in the oral tube.



### **D. Store remaining doses in the fridge immediately**

Unused oral tubes that are still attached to the connecting bar must be put back in the fridge immediately after an oral tube has been used. This is so that the unused oral tubes can be used for the next vaccination.

Discard the used oral tubes in approved biological waste containers according to local regulations.

