

XOLAIR® Omalizumab

1. PRODUCT NAME

Xolair 75 mg solution for injection in pre-filled syringe

Xolair 150 mg solution for injection in pre-filled syringe

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each pre-filled syringe of 0.5 mL contains 75 mg of omalizumab.

Each pre-filled syringe of 1 mL contains 150 mg of omalizumab.

Omalizumab is a humanized monoclonal antibody manufactured from a mammalian cell line.

For a full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Clear to slightly opalescent, colorless to pale brownish-yellow solution in a pre-filled syringe.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Allergic Asthma

Xolair (omalizumab) is indicated for the reduction of asthma exacerbations and control of asthma symptoms when given as add-on therapy for adult and adolescent patients, 6 years and older, with severe persistent allergic asthma who have IgE \geq 30 IU/mL, a positive skin test or in vitro reactivity to a perennial aeroallergen and whose symptoms are inadequately controlled with inhaled corticosteroids.

Chronic rhinosinusitis with nasal polyps (CRSwNP)

Xolair is indicated as add-on treatment in adult patients (18 years of age and above) for the treatment of severe CRSwNP with inadequate response to intranasal corticosteroids. Recommended dosing is determined by serum immunoglobulin E levels and body weight corresponding to the recommended dose range in the [Data Sheet](#) (see Section 4.2 Dose and method of administration).

Chronic Idiopathic Urticaria (CIU)

Xolair is indicated for adults and adolescents (12 years of age and above) with chronic idiopathic urticaria who remain symptomatic despite H1 antihistamine treatment.

4.2 Dose and method of administration

Dosing principles for Allergic Asthma and CRSwNP

Xolair treatment for chronic rhinosinusitis with nasal polyps (CRSwNP) should be initiated by

specialists experienced in the diagnosis and treatment of CRSwNP.

Dosing for asthma and CRSwNP follows the same dosing principles. The appropriate dose and dosing frequency of Xolair is determined by baseline immunoglobulin E (IgE) (IU/mL), measured before the start of treatment, and body weight (kg). Prior to initial dosing, patients should have their IgE level determined by any commercial serum total IgE assay for their dose assignment. Based on these measurements 75 to 600 mg of Xolair in 1 to 4 injections may be needed for each administration. See Table 1 for a conversion chart and Tables 2 and 3 for the dose determination charts in children (6 years to less than 12 years of age) and in adults and adolescents (12 years of age and older). For doses of 225, 375 or 525 mg Xolair 150 mg can be used in combination with Xolair 75 mg.

Patients whose baseline IgE levels or body weight in kilograms are outside the limits of the dosing table should not be given Xolair.

Table 1 Conversion from dose to number of pre-filled syringes, number of injections and total injection volume for each administration

Dose (mg)	Number of syringes		Number of injections	Total injection volume (mL)
	75 mg	150 mg		
75	1	0	1	0.5
150	0	1	1	1.0
225	1	1	2	1.5
300	0	2	2	2.0
375	1	2	3	2.5
450	0	3	3	3.0
525	1	3	4	3.5
600	0	4	4	4.0

Measurement of serum IgE levels

Any commercial serum total IgE assay may be used for determination of serum total IgE for initial dose assignment. However, Xolair can interfere with accurate quantitation of serum IgE levels. The total IgE levels while on active treatment of Xolair increased an average of 4-fold post-dose as a result of omalizumab-IgE binding. If it is necessary to measure serum total IgE in subjects currently on Xolair treatment or who have discontinued within the last 12 months, the Abbott IMX assay has been shown to demonstrate reliable serum total IgE measurements.

Treatment duration, monitoring and dose adjustments

Doses do not need to be adjusted for variations in serum IgE over time. Data from clinical studies suggest that, in the absence of omalizumab treatment, there is no significant temporal variation in serum total IgE levels.

In clinical trials there were reductions in asthma exacerbation events and rescue medication use with improvements in symptom scores during the first 16 weeks of treatment. Xolair is intended for long-term treatment. In treatment for allergic asthma, at least 12 weeks of treatment is required to adequately assess whether or not a patient is responding to Xolair. The decision to continue Xolair should be based on whether a marked improvement in overall asthma control is seen (see Section 5.1 Pharmacodynamic properties – clinical trials).

Discontinuation generally results in a return to elevated free IgE levels and associated symptoms.

Total IgE levels are elevated during treatment and remain elevated for up to one year after the discontinuation of treatment. Therefore, re-testing of IgE levels during Xolair treatment cannot be used as a guide for dose determination. Dose determination after treatment interruptions lasting less than one year should be based on serum IgE levels obtained at the initial dose determination. Total serum IgE levels may be re-tested for dose determination if treatment with Xolair has been interrupted for one year or more.

Doses should be adjusted for significant changes in body weight (see Tables 2 and 3). No data are currently available to support dose adjustments based on changes in serum total IgE with increasing age.

In clinical trials for CRSwNP, changes in CRSwNP score (NPS) and nasal congestion score (NCS) were observed as early as the first assessment at 4 weeks. The need for continued therapy should be periodically reassessed based upon the patient's disease severity and level of symptom control.

Dosage for Allergic Asthma and CRSwNP

Xolair is administered subcutaneously every two or four weeks according to the dose determination charts below (Tables 2 and 3). Doses (mg) and dosing frequency are determined by baseline serum total IgE level (IU/mL), measured before the start of treatment, and body weight (kg).

Table 2 ADMINISTRATION EVERY 4 WEEKS. Xolair doses (milligrams per dose) administered by subcutaneous injection every 4 weeks

Baseline IgE (IU/ml)	Body weight (kg)									
	>20-25*	>25-30*	>30-40	>40-50	>90- >50-60	>125- >60-70	>70-80	>80-90	125	150
≥30-100	75	75	75	150	150	150	150	150	300	300
>100-200	150	150	150	300	300	300	300	300	450	600
>200-300	150	150	225	300	300	450	450	450	600	
>300-400	225	225	300	450	450	450	600	600		
>400-500	225	300	450	450	600	600				
>500-600	300	300	450	600	600	ADMINISTRATION EVERY 2 WEEKS				
>600-700	300		450	600		SEE TABLE 3				

*Body weights below 30 kg were not studied in the pivotal trials for CRSwNP.

Table 3 ADMINISTRATION EVERY 2 WEEKS. Xolair doses (milligrams per dose) administered by subcutaneous injection every 2 weeks

	Body weight (kg)										
Baseline IgE (IU/ml)	>20- 25*	>25-30*	>30-40	>40- 50	>50- 60	>60- 70	>70- 80	>80- 90	>90-125	>125-150	>150-200
> 30-100	ADMINISTRATION EVERY 4 WEEKS										225
> 100-200	SEE TABLE 2										375
> 200-300											375
> 300-400											525
> 400-500											600
> 500-600											600
> 600-700											600
> 600-700	225	225	300	375	450	450	525	600			
> 700-800	225	225	300	375	450	450	525	600			
> 800-900	225	300	375	450	525	600					
> 900-1000	225	300	375	450	525	600					
> 1000-1100	225	300	375	450	600	Insufficient Data to Recommend a Dose					
> 1100-1200	300	300	450	525	600						
> 1200-1300	300	375	450	525							
> 1300-1500	300	375	525	600							

*Body weights below 30 kg were not studied in the pivotal trials for CRSwNP.

In allergic asthma, safety and efficacy in paediatric patients below the age of 6 have not been established and use of Xolair in such patients is therefore not recommended.

In CRSwNP, safety and efficacy in patients below the age of 18 have not been established and use of Xolair in such patients is therefore not recommended.

There is limited data on doses greater than 375 mg every 2 weeks (750 mg per month) from the pivotal studies in CRSwNP and therefore higher doses are not recommended.

Dosage for Chronic Idiopathic Urticaria (CIU)

The recommended dose is 300 mg by subcutaneous injection every four weeks. Some patients may achieve control of their symptoms with a dose of 150 mg every four weeks.

Prescribers are advised to periodically reassess the need for continued therapy.

Clinical trial experience of long-term treatment beyond 6 months in this indication is limited. Xolair should be used as add-on therapy to H1 antihistamine treatment.

Special populations

Elderly population

There are limited data available on the use of Xolair in patients older than 65 years but there is no evidence that elderly patients require a different dosage from younger adult patients.

Renal or hepatic impairment

There have been no studies on the effect of impaired renal or hepatic function on the pharmacokinetics of omalizumab. Because omalizumab clearance at clinical doses is dominated by the reticular endothelial system (RES) it is unlikely to be altered by renal or hepatic impairment. While no particular dose adjustment is recommended, Xolair should be administered with caution in these patients.

Paediatric population

In allergic asthma, safety and efficacy in paediatric patients below the age of 6 years have not been established and use of Xolair in such patients is therefore not recommended.

In chronic idiopathic urticaria, safety and efficacy in paediatric patients below the age of 12 years have not been established.

Method of administration

For subcutaneous administration only. Xolair must not be administered by the intravenous or intramuscular route.

Doses of more than 150 mg should be divided across two or more injection sites.

After proper training in subcutaneous injection technique, patients or the caregiver may self-inject Xolair if a physician determines that it is appropriate. It is recommended that the first 3 doses be administered by or under the supervision of a healthcare professional.

Patients or caregivers should be instructed to inject the full amount of Xolair according to the instructions for use provided in Section 6.6.

Full instructions for use are provided in Section 6.6.

4.3 Contraindications

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

4.4 Special warnings and precautions for use

Abrupt discontinuation of systemic or inhaled corticosteroids after initiation of Xolair therapy in allergic asthma or CRSwNP is not recommended. Decreases in corticosteroids should be performed under the direct supervision of a physician and may need to be performed gradually.

Allergic reactions

As with any protein, local or systemic allergic reactions, including Anaphylaxis, may occur when taking omalizumab. Anaphylactic reactions were rare in clinical trials (see Section 4.8 Undesirable effects). In post-marketing experience anaphylaxis and anaphylactoid reactions have been reported following the first or subsequent administrations of Xolair. Most of these reactions occurred within 2 hours after Xolair administration, some occurred beyond 2 hours and even beyond 24 hours after

injections. The majority of anaphylactic reactions occurred within the first 3 doses of Xolair. A history of anaphylaxis unrelated to omalizumab may be a risk factor for anaphylaxis following Xolair administration. It is recommended that patients with a known history of anaphylaxis have each dose of Xolair administered in a setting where there is treatment for anaphylaxis available. Patients should be informed that such reactions are possible and prompt medical attention should be sought if allergic reactions occur.

Serum sickness and serum sickness-like reactions, which are delayed allergic type III reactions, have rarely been seen in patients treated with humanized monoclonal antibodies including omalizumab. The onset has typically been 1-5 days after administration of the first or subsequent injections, also after long duration of treatment. Symptoms suggestive of serum sickness include arthritis/arthralgia, rash (urticaria or other forms), fever and lymphadenopathy. Antihistamines and corticosteroids may be useful for preventing or treating this disorder, and patients should be advised to report any suspected symptoms.

Churg-Strauss syndrome and hypereosinophilic syndrome

Patients with severe asthma may rarely present systemic hypereosinophilic syndrome or allergic eosinophilic granulomatous vasculitis (Churg-Strauss syndrome), both of which are usually treated with systemic corticosteroids. In rare cases, patients on therapy with anti-asthma agents, including omalizumab, may present or develop systemic eosinophilia and vasculitis. A causal association between Xolair and these underlying conditions has not been established. These events are commonly associated with the reduction of oral corticosteroid therapy. In these patients, physicians should be alert to the development of marked eosinophilia, vasculitic rash, worsening pulmonary symptoms, paranasal sinus abnormalities, cardiac complications, and/or neuropathy. Discontinuation of omalizumab should be considered in all severe cases with the above mentioned immune system disorders.

Immunogenicity

As with all protein pharmaceuticals, a small percentage of patients may potentially develop antibodies to the protein (see Section 4.8 Undesirable effects).

Other IgE-associated disorders

Xolair has not been studied in patients with anaphylaxis, hyperimmunoglobulin E syndrome, allergic bronchopulmonary aspergillosis, food allergy or atopic dermatitis. Parasitic infestation may also result in elevation of serum IgE concentrations. In a study of asthmatic patients who had been treated for gut parasites, the level of reinfection did not differ significantly between omalizumab and placebo groups and there were no serious or severe infections. There is no current evidence to suggest that parasitic infections are predisposed to by omalizumab.

Interpretation of serum IgE levels, skin patch and skin prick testing

Patients treated with Xolair have a rapid reduction in free IgE in the serum but an overall increase in the total serum IgE (commonly measured in clinical laboratories), which reflects free IgE and IgE bound by Xolair. IgE measured following treatment cannot be used to guide treatment or dosing decisions.

Because Xolair reduces free IgE in the serum and tissues, results of skin prick testing, patch testing, and RAST testing for hypersensitivity to potential allergens may be affected. A positive test to a potential allergen in a patient receiving Xolair can be correctly interpreted as representing hypersensitivity to that allergen; however, a negative test in a patient receiving Xolair may not be interpretable. Physicians are urged to use caution in interpreting such tests in patients receiving Xolair.

Thrombocytopenia

At serum concentrations in excess of maximum human exposure used in pivotal clinical trials, dose-related thrombocytopenia occurred in 2 out of 4 non-human primate species studied. The thrombocytopenia was more pronounced in juvenile animals. No Xolair-related thrombocytopenia has been observed in clinical trials, but it has been reported in the post-market setting.

Xolair should be used with caution in patients with thrombocytopenia and patients with a history of thrombocytopenia. It is recommended that patients have a platelet count before commencing therapy with Xolair and then periodically during treatment with Xolair.

Arteriothrombotic Events (ATE)

In controlled clinical trials, interim and final analyses of an observational study, a numerical imbalance of ATE was observed (see Section 4.8 Undesirable effects).

Pre-filled syringe, latex-sensitive individuals

The removable needle cap of Xolair solution for injection in pre-filled syringe contains a derivative of natural rubber latex. Although no natural rubber latex is detected in the removable needle cap, the safe use of Xolair solution for injection in pre-filled syringe in latex-sensitive individuals has not been studied.

Use in hepatic and renal impairment

There have been no studies on the effect of impaired renal or hepatic function on the pharmacokinetics of Xolair. Xolair should be administered with caution in these patients.

Paediatric population

In allergic asthma, safety and efficacy in paediatric patients below the age of 6 have not been established and use of Xolair in such patients is therefore not recommended.

In CRSwNP, safety and efficacy in patients below the age of 18 years has not been established.

In chronic idiopathic urticaria, safety and efficacy in paediatric patients below the age of 12 years have not been established.

4.5 Interaction with other medicines and other forms of interaction

Cytochrome P450 enzymes, efflux pumps and protein binding mechanisms are not involved in the clearance of omalizumab; thus, there is little potential for drug-drug interactions. No formal drug or vaccine interaction studies have been performed with Xolair.

There is no pharmacological reason to expect that commonly prescribed medications used in the treatment of asthma will interact with omalizumab.

Allergic Asthma

In clinical studies Xolair was commonly used in conjunction with inhaled and oral corticosteroids, inhaled short-acting and long-acting beta2-agonists, leukotriene modifiers, theophyllines and oral antihistamines. There was no indication that the safety of Xolair was altered with these other commonly used asthma medications. Limited data are available on the use of Xolair in combination with specific immunotherapy (hypo-sensitisation therapy).

Chronic rhinosinusitis with nasal polyps (CRSwNP)

In clinical studies Xolair was used in conjunction with intranasal mometasone spray per protocol. Other commonly used concomitant medications included other intranasal corticosteroids, bronchodilators, antihistamines, leukotriene receptor antagonists, adrenergics/sympathomimetics, and local nasal anesthetics. There was no indication that the safety of Xolair was altered with these other commonly used CRSwNP medications.

Chronic Idiopathic Urticaria (CIU)

In clinical studies in CIU Xolair was used in conjunction with antihistamines (anti-H1, anti-H2) and leukotriene receptor antagonists (LTRAs). In the phase III studies Q4881g and Q4882g all patients received H1 antihistamines in addition to Xolair or placebo. In the phase III study Q4883g, all patients received one or more H1 antihistamine(s), and/or H2 antihistamines and/or LTRAs in addition to Xolair or placebo. There was no evidence that the safety of omalizumab was altered when used with these medicinal products relative to its known safety profile in allergic asthma. In addition, a population pharmacokinetic analysis showed no relevant effect of H2 antihistamines and LTRAs on omalizumab pharmacokinetics (see section 5.1 Pharmacodynamic properties).

4.6 Fertility, pregnancy and lactation

Pregnancy

Human IgG antibodies are known to cross the placental barrier; therefore, XOLAIR may be transmitted from the mother to the developing fetus. There are no well-controlled clinical studies of Xolair in pregnant women. A prospective pregnancy registry study (EXPECT) in 250 pregnant women with asthma exposed to Xolair showed the prevalence of major congenital anomalies was similar (8.1% vs 8.9%) between EXPECT and disease matched (moderate and severe asthma) patients. There was an increased rate of low birth weight among registry infants compared to infants in the other cohort. This study cannot definitively establish risk due to methodological limitations of the study, including small sample size, nonrandomised design and potential differences between the registry population and the comparator cohort.

Studies in cynomolgus monkeys do not indicate direct or indirect harmful effects with respect to pregnancy, embryonic/foetal development, parturition or neonatal growth at weekly doses up to 75 mg/kg SC (yielding about 10 times the maximum anticipated clinical exposure in adult patients, based on serum AUC).

If clinically needed, the use of Xolair may be considered during pregnancy.

Breastfeeding

Immunoglobulins G (IgGs) are present in human milk, and therefore it is expected that omalizumab will be present in human milk.

The EXPECT study, with 154 infants who had been exposed to Xolair during pregnancy and through breast-feeding did not indicate adverse effects on the breast-fed infant. The interpretation of data may be impacted due to methodological limitations of the study, including small sample size and non-randomised design.

Given orally, immunoglobulin G proteins undergo intestinal proteolysis and have poor bioavailability. No effects on the breast-fed newborns/infants are anticipated. Consequently, if clinically needed, the use of Xolair may be considered during breast-feeding.

In order to assess the effect of omalizumab on late gestation, and to evaluate the placental transfer and milk excretion of omalizumab, doses of 75 mg/kg/week were administered subcutaneously to female cynomolgus monkeys. Transport of omalizumab into maternal milk was limited. The serum

levels of omalizumab observed in dams, fetuses, and neonates are consistent with reported transport and distribution of IgG class immunoglobulins.

Fertility

There are no human fertility data for omalizumab. Studies in cynomolgus monkeys showed no adverse effects of omalizumab on fertility or general reproductive performance of males or females at weekly doses up to 75 mg/kg SC about 10 times the maximum anticipated clinical exposure in adult patients, based on serum AUC).

4.7 Effects on ability to drive and use machines

Patients receiving Xolair should be informed that if they experience dizziness, fatigue, syncope or somnolence they should not drive or use machines.

4.8 Undesirable effects

Clinical trial experience in allergic asthma

Adverse reactions with Xolair were observed (all studies with adult and adolescent patients 12 years of age and older) at a frequency of 6.6 % of patients, treated with active drug, during clinical trials. The most commonly reported adverse reactions were injection site reactions, including injection site pain, swelling, itching and redness (1.7%) and headaches (1%).

In clinical studies with patients 6 to <12 years of age, the most commonly reported adverse reactions were headache, pyrexia and upper abdominal pain. Most of the events were mild or moderate in severity.

The adverse reactions listed in Table 4 were recorded in clinical studies in the total allergic asthma safety population treated with Xolair. Adverse reactions are ranked under headings of frequency using the following convention: common (>1/100 to <1/10), uncommon (>1/1000 to <1/100), rare (<1/1000).

Table 4 Adverse reactions recorded in clinical studies in the total allergic asthma safety population treated with Xolair

Body as a whole disorders Very common: pyrexia* Common: injection site reactions Uncommon: weight increase, fatigue, swelling arms, post-injection phenomena
Cardiovascular disorders Uncommon: syncope and vasovagal syncope
Vascular disorders Uncommon: postural hypotension, flushing
Gastrointestinal disorders Common: abdominal pain upper* Uncommon: nausea, diarrhoea, dyspeptic signs and symptoms
Infections and infestations Uncommon: moniliasis
Immune system disorders Rare: anaphylactic reactions, anti-therapeutic antibody development
Nervous system disorders Common: headache** Uncommon: dizziness, somnolence, paresthesia
Respiratory system Uncommon: pharyngitis, coughing, allergic bronchospasm Rare: laryngoedema
Skin and subcutaneous tissue disorders

Uncommon: urticaria, rash, pruritus, photosensitivity Rare: angioedema
Adverse event special categories (causality not established) Uncommon: asymptomatic platelet decreases, parasitic infections

**In 6 to <12 year old children **Very common in 6 to <12 year old children*

The frequencies of adverse reactions in the active treatment group adult and adolescent patients were very similar to those observed in the placebo group. Weight increase (0.7% vs 0.2%, placebo), urticaria (0.4% vs 0.1%, placebo) and local injection site reactions (1.7% vs 1.3%, placebo) were slightly more commonly observed in the active treatment group than in placebo group patients.

Adverse Events (AEs): The most common adverse events observed (frequency of >20%) in this adult and adolescent patient population, were headaches, viral infections and upper respiratory tract infections. The frequencies of all adverse events for both treated (N = 1763) and placebo group patients (N =1278) for all studies were very similar.

Serious Adverse Events (SAEs): were reported for 2.6% of Xolair treated patients and 2.8% of placebo-treated patients. The most frequently reported SAE's were appendicitis and fractures (0.2% for both treatment groups and both events). Frequencies of all SAEs by body system were comparable for both treatment groups.

Allergic asthma (adult and adolescent study population): The most frequently observed adverse events in this population (N = 716 Xolair patients, N = 694 placebo patients) were viral infections, upper respiratory tract infections, sinusitis and headaches (frequency \geq 20%).

Adverse reactions occurred in 5.6% of Xolair-treated patients and in 5.2% of patients receiving placebo. The most frequently reported events of this type were headaches (1.3% vs 1.2%, placebo) and fatigue (0.6% vs 0%, placebo). All other suspected Xolair drug related adverse events occurred with a frequency of <0.5%.

Serious episodes of asthma related events requiring hospitalisation were observed in 0.2% of Xolair-treated patients and 1.3% of placebo-treated patients, in the asthma studies.

The most frequently observed (\geq 5%) adverse events in studies with asthma patients 12 years of age and older are provided in Table 5.

Table 5 *Most frequently reported adverse events regardless of causality in adult/adolescent asthma population (\geq 5% in either treatment group)*

Body system/preferred term	Xolair (N=716)	Placebo (N=694)
Body as a whole		
Fever	5%	5%
Pain	7%	5%
Digestive system		
Diarrhoea	6%	6%
Dyspepsia	6%	5%
Nausea	6%	6%
Gastroenteritis	6%	5%
Pain abdominal	4%	5%
Infections and infestations		
Infection, viral	37%	39%
Musculoskeletal		
Pain back	13%	12%
Arthralgia	8%	7%
Myalgia	6%	6%
Sprains and strains	6%	5%
Nervous		
Headache	27%	27%

Clinical trial experience in Chronic rhinosinusitis with nasal polyps (CRSwNP)

Summary of the safety profile

The data described below reflect data from two placebo-controlled studies in patients ≥ 18 years of age. In these studies, patients received either Xolair 150 to 600 mg every 2 or 4 weeks or placebo. All patients received background intranasal mometasone therapy. The safety profile in patients with CRSwNP was consistent with that in allergic asthma and CSU. The most frequently ($>3\%$) reported adverse drug reactions, which were higher in frequency in comparison to placebo are shown in Table 6.

Tabulated summary of adverse drug reactions from the clinical studies

Table 6 lists the adverse drug reactions recorded in clinical studies in the total nasal polyp safety population treated with Xolair by system organ class and by frequency. Frequencies are defined 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 ($< 1/10,000$).

Table 6 Adverse drug reactions from the clinical studies in CRSwNP

Adverse drug reactions (by MedDRA preferred term)	Omalizumab Nasal Polyp Study1 and Study 2 Pooled		Frequency category
	Placebo N=130	Omalizumab N =135	
Nervous system disorders			
Headache	7 (5.4%)	11 (8.1%)	Common
Dizziness	1 (0.8%)	4 (3.0%)	Common
Musculoskeletal and connective tissue disorders			
Arthralgia	2 (1.5%)	4 (3.0%)	Common
Gastrointestinal disorder			
Abdominal pain upper	1 (0.8%)	4 (3.0%)	Common
General disorders and administration site conditions			
Injection site reactions (Injection site reactions, Injection related reaction, injection site pain)	2 (1.5%)	7 (5.2%)	Common

Clinical trial experience in Chronic Idiopathic Urticaria (CIU)

Chronic Idiopathic Urticaria (CIU)

Summary of the safety profile

The safety and tolerability of omalizumab were investigated with the doses of 75 mg, 150 mg and 300 mg every four weeks in 975 CIU patients, 242 of whom received placebo. 733 patients were treated

with omalizumab for up to 12 weeks and 490 patients for up to 24 weeks. 175 and 412 patients were treated for up to 12 weeks and 87 and 333 patients were treated for up to 24 weeks at the recommended doses of 150 mg and 300 mg respectively.

During clinical studies with adult and adolescent patients (12 years of age and older) the most commonly reported adverse reactions observed were headache and nasopharyngitis.

Tabulated summary of adverse reactions from the clinical studies at the recommended doses (150 mg and 300 mg)

Adverse reactions (events occurring in $\geq 1\%$ of patients in any treatment group and $\geq 2\%$ more frequently in any omalizumab treatment group than in the placebo group after medical review) reported at the recommended doses (150mg and 300mg) in the three pooled Phase III studies are listed by MedDRA system organ class (Table 6). Within each system organ class, the adverse reactions are ranked by frequency, with the most frequent reactions listed first. The corresponding frequency category for each adverse reaction is based on the following convention (CIOMS III): very common ($\geq 1/10$); common ($\geq 1/100$ to $< 1/10$); uncommon ($\geq 1/1,000$ to $< 1/100$); rare ($\geq 1/10,000$ to $< 1/1000$); very rare ($< 1/10,000$); not known (cannot be estimated from the available data).

Table 7 Adverse reactions from the pooled CIU safety database (day 1 to week 12) at the recommended doses

Adverse reactions (by MedDRA preferred term)	Omalizumab Studies Q4881g, Q4882g and Q4883g Pooled			Frequency category
	Placebo N=242	150 mg N=175	300 mg N=412	
Infections and infestations				
Nasopharyngitis	17 (7.0%)	16 (9.1%)	27 (6.6%)	Common
Sinusitis	5 (2.1%)	2 (1.1%)	20 (4.9%)	Common
Viral upper respiratory tract infection	0	4 (2.3%)	2 (0.5%)	Common
Nervous system disorders				
Headache	7 (2.9%)	21 (12.0%)	25 (6.1%)	Very common
Musculo skeletal and connective tissue disorders				
Arthralgia	1 (0.4%)	5 (2.9%)	12 (2.9%)	Common

Additional events reported anytime during the day 1 to week 24 treatment period (studies Q4881g and Q4883g) that met the criteria of adverse reactions:

Infections and infestations: upper respiratory tract infections (placebo 3.1%, 150 mg 3.4%, 300 mg 5.7%), urinary tract infection (placebo 1.8%, 150 mg 4.6%, 300 mg 2.4%).

Nervous system disorders: sinus headache (placebo 0%, 150 mg 2.3%, 300 mg 0.3%).

Musculoskeletal and connective tissue disorders: myalgia (placebo 0%, 150 mg 2.3%, 300 mg 0.9%), pain in extremity (placebo 0%, 150 mg 3.4%, 300 mg 0.9%), musculoskeletal pain (placebo 0%, 150 mg 2.3%, 300 mg 0.9%).

General disorders and administration site conditions: pyrexia (placebo 1.2%, 150 mg 3.4%, 300 mg 0.9%).

Injection site reactions: Injection site reactions occurred during the studies in more omalizumab-treated patients than placebo patients (2.7% at 300 mg, 0.6% at 150 mg, 0.8% with placebo). They included: swelling, erythema, pain, bruising, itching, bleeding and urticaria.

Post-marketing observations

The following reactions have been identified through spontaneous reporting.

Immune system disorders

Anaphylaxis and anaphylactoid reactions have been reported following the first or subsequent administrations, serum sickness (see Section 4.4 SPECIAL WARNINGS AND PRECAUTIONS FOR USE and Section 4.8 ADVERSE EFFECTS (UNDESIRABLE EFFECTS) - 'Allergic events').

Skin and subcutaneous disorders

Alopecia.

Blood and lymphatic system disorders

Idiopathic severe thrombocytopenia.

Respiratory, thoracic and mediastinal disorders

Churg Strauss syndrome (ie. Eosinophilic Granulomatosis with Polyangitis)

Musculoskeletal and connective tissue disorders

Arthralgia, myalgia, joint swelling

Description of safety aspects of special interest

Allergic events

As with any protein, local or systemic allergic or Type I hypersensitivity events can occur. Frequencies of all allergic-type events were similar for both treatment groups of the total study population in adults and adolescents (4%, Xolair, 6%, placebo). Events such as vasovagal syncope, postural hypotension, allergic bronchospasm and photosensitivity occurred in <1% of patients in each treatment group (see section 4.4).

In post-marketing reports, the frequency of anaphylaxis in patients exposed to Xolair use was estimated to be 0.2% based on a total number of anaphylactic reactions observed from an estimated exposure of over 500,000 patient years.

Parasitic infections

In patients at chronic high risk of helminth infection, a placebo-controlled trial showed a slight numerical increase in infection rate with omalizumab that was not statistically significant. The course, severity, and response to treatment of infections were unaltered (see section 4.4).

Description of select adverse events pertinent to allergic asthma and CSU indications

The following adverse events have been reported in patients treated with Xolair, however, the causality of the adverse events and Xolair use has not been established.

Malignancies

During initial clinical trials in adults and adolescents 12 years of age and older, there was a numerical imbalance in cancers arising in the active treatment group (25 patients of 5015), compared with the control group (5 patients of 2854). The number of observed cases was uncommon (<1/100) in both the active (0.5%) and the control group, (0.2%). The observed cases in Xolair-treated patients included breast, non-melanoma skin, prostate, melanoma and parotid malignancies. The overall observed incidence rate of malignancy in the initial Xolair clinical trial programme was comparable to that reported in the general population (0.4%).

In a subsequent observational study comparing 5,007 Xolair-treated and 2,829 non-Xolair-treated patients followed for up to 5 years, the incidence rates of primary malignancies per 1,000 patient years were 16.01 (295/18,426 patient years) and 19.07 (190/9,963 patients years), respectively, which does not indicate an increased malignancy risk (rate ratio 0.84, 95% confidence interval, 0.62 to 1.13). In a further analysis of randomized, double-blind, placebo-controlled clinical trials including 4,254 patients on Xolair and 3,178 patients on placebo, Xolair treatment was not associated with an increased malignancy risk based on incidence rates per 1,000 patient years of 4.14 (14/3,382 patient years) for Xolair treated patients and 4.45 (11/2,474 patient years) for placebo patients (rate ratio 0.93, 95% confidence interval 0.39 to 2.27).

Arterial Thromboembolic Events (ATE)

In controlled clinical trials and during interim analyses of an observational study, a numerical imbalance of ATEs was observed. ATE included stroke, transient ischemic attack, myocardial infarction, unstable angina, and cardiovascular death (including death from unknown cause). In the final analysis of the observational study, the rate of ATE per 1,000 patient years was 7.52 (115/15,286 patients years) for Xolair-treated patients and 5.12 (51/9,963 patient years) for control patients. In a multivariate analysis controlling for available baseline cardiovascular risk factors, the hazard ratio was 1.32 (95% confidence interval 0.91 to 1.91). In a separate analysis of pooled clinical trials including all randomized double-blind, placebo-controlled clinical trials of 8 or more weeks duration, the rate of ATE per 1,000 patient years was 2.69 (5/1,856 patients years) for Xolair-treated patients and 2.38 (4/1,680 patient years) for placebo patients (rate ratio 1.13, 95% confidence interval 0.24 to 5.71).

Thrombocytopenia

In clinical trials 0.6 % of adult and adolescent patients experienced platelet counts below the lower limit of the normal laboratory range. None of these changes were associated with bleeding episodes or a decrease in haemoglobin. No pattern of persistent decrease in platelet counts has been reported in humans (patients greater than 6 years of age), as was observed in non-human primates (see section 5.3).

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicine is important. It allows continued monitoring of the benefit/risk balance of the medicine. Healthcare professionals are asked to report any suspected adverse reactions <https://nzphvc.otago.ac.nz/reporting/>.

4.9 Overdose

No case of overdose has been reported. Maximum tolerated dose of Xolair has not been determined. Single intravenous doses up to 4,000 mg have been administered to patients without evidence of dose-limiting toxicities. The highest cumulative dose administered to patients was 44,000 mg over a 20-week period and this dose did not result in any untoward acute effects.

For advice on the management of overdose please contact the National Poisons Centre on 0800 POISON (0800 764766).

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: other systemic drugs for obstructive airway diseases, ATC code: R03DX05

General characteristics

Omalizumab is a recombinant DNA-derived humanised monoclonal antibody that selectively binds to human immunoglobulin E (IgE). The antibody is an IgG₁ kappa that contains human framework regions with the complementary-determining regions of a murine parent antibody that binds to IgE.

Patients with Allergic Asthma and Chronic rhinosinusitis with nasal polyps (CRSwNP)

The allergic cascade is initiated when IgE bound to the high affinity IgE receptor Fc ϵ RI on the surface of mast cells and basophils is cross-linked by allergen. This results in the degranulation of these effector cells and the release of histamines, leukotrienes, cytokines and other mediators. These mediators are causally linked to the pathophysiology of allergic asthma including airway oedema, smooth muscle contraction and altered cellular activity associated with the inflammatory process. They also contribute to the signs and symptoms of allergic disease such as bronchoconstriction, mucus production, wheezing, dyspnoea, chest tightness, nasal congestion, sneezing, itchy, runny nose and itchy, watery eyes.

Mechanism of action

Omalizumab binds to IgE and prevents binding of IgE to the high-affinity Fc ϵ RI receptor, thereby reducing the amount of free IgE that is available to trigger the allergic cascade. Treatment of atopic subjects with omalizumab resulted in a marked down-regulation of Fc ϵ RI receptors on basophils. Treatment with Xolair inhibits IgE-mediated inflammation, as evidenced by reduced blood and tissue eosinophils and reduced inflammatory mediators, including IL-4, IL-5, and IL-13 by innate, adaptive and non-immune cells.

Pharmacodynamic effects

The *in vitro* histamine release from basophils isolated from Xolair treated subjects was reduced by approximately 90% following stimulation with an allergen compared to pre-treatment values.

Patients with Allergic Asthma

In clinical studies in asthma patients, free IgE levels in serum were reduced in a dose-dependent manner within one hour following the first dose and maintained between doses. Mean decrease in free IgE in serum was greater than 96% using recommended doses. Total IgE levels (i.e., bound and unbound) in serum increased after the first dose due to the formation of omalizumab: IgE complexes which have a slower elimination rate compared with free IgE. At 16 weeks after the first dose, average serum total IgE levels were five-fold higher compared with pre-treatment levels when using standard assays. After discontinuation of Xolair dosing, the Xolair-induced increase in total IgE and decrease in free IgE were reversible, with no observed rebound in IgE levels after drug washout. Total IgE levels did not return to pre-treatment levels for up to one year after discontinuation of Xolair.

Patients with Chronic rhinosinusitis with nasal polyps (CRSwNP)

In clinical studies in patients with CRSwNP, Xolair treatment led to a reduction in serum free IgE and an increase in serum total IgE levels, similar to the observations in patients with allergic asthma. After repeated dosing every 2 or 4 weeks, with dosage and frequency according to Table 2 (see section 4.2 Dose and method of administration), mean pre-dose serum free IgE levels decreased by approximately 95% and remained stable between 16 and 24 weeks of treatment. Total IgE levels in serum increased due to the formation of omalizumab-IgE complexes, which have a slower elimination rate compared with free IgE. After repeated dosing every 2 or 4 weeks, with dosage and frequency according to Table 2 (see section 4.2 Dose and method of administration), mean pre-dose serum total

IgE levels at Week 16 were 3- to 4- fold higher compared with pre-treatment levels, and remained stable between 16 and 24 weeks of treatment.

Patients with Chronic Idiopathic Urticaria (CIU)

There are several theories for the etiology of CIU, including one that suggests an autoimmune origin. Autoimmune antibodies to IgE and its receptor, FcεRI, have been isolated from the serum of some patients with CIU. These autoantibodies can activate basophils or mast cells leading to release of histamine.

Mechanism of action

One hypothesis for the mechanism of action of omalizumab in CIU is that it lowers free IgE levels in the blood and subsequently in the skin. This leads to down-regulation of surface IgE receptors, thereby decreasing downstream signaling via the FcεRI pathway, resulting in suppressed cell activation and inflammatory responses. As a consequence, the frequency and severity of symptoms of CIU are lessened. Another hypothesis is that lowering circulating free IgE levels leads to a rapid and non-specific desensitization of cutaneous mast cells. Down-regulation of FcεRI may help to sustain the response.

Pharmacodynamic effect

In clinical studies in CIU patients, omalizumab treatment led to a dose-dependent reduction of free IgE and an increase of total IgE levels in serum, similar to the observations in allergic asthma patients. Maximum suppression of free IgE was observed 3 days after the first subcutaneous dose. After repeated dosing once every 4 weeks, pre-dose serum free IgE levels remained stable between 12 and 24 weeks of treatment. Total IgE levels in serum increased after the first dose due to the formation of omalizumab:IgE complexes which have a slower elimination rate compared with free IgE. After repeated dosing once every 4 weeks at 75 mg to 300 mg, average predose serum total IgE levels at week 12 were two-to three-fold higher compared with pre-treatment levels, and remained stable between 12 and 24 weeks of treatment. After discontinuation of Xolair, free IgE levels increased and total IgE levels decreased towards pre-treatment levels over a 16-week treatment-free follow-up period.

Clinical efficacy and safety in Allergic Asthma

Adults and adolescents > 12 years of age

The safety and efficacy of Xolair were evaluated in five randomised, double-blind, placebo controlled, multi-centre trials.

In identical 16-week studies 1 and 2, the safety and efficacy of omalizumab as add-on therapy were demonstrated in 1,071 allergic asthmatics, who were symptomatic despite treatment with inhaled corticosteroids (beclomethasone dipropionate 500 to 1,200 micrograms/day).

In both trials omalizumab was superior to placebo with respect to the primary variable of asthma exacerbation (worsening of asthma requiring systemic corticosteroids or a doubling of the patient's baseline beclomethasone dose). The number of asthma exacerbations was significantly lower in the omalizumab group ($p=0.006$ and $p<0.001$ in studies 1 and 2, respectively). Fewer omalizumab-treated patients experienced asthma exacerbations (14.6% vs 23.3%, $p=0.009$ in study 1 and 12.8% vs 30.5%, $p<0.001$ in study 2).

In double-blind extension phases of both studies out to one year the reduction in the frequency of asthma exacerbations for omalizumab-treated patients compared to placebo-treated patients was maintained.

In studies 1 and 2, clinically meaningful improvement in asthma-related quality of life, measured by the validated Juniper's Asthma Quality of Life Questionnaire, was demonstrated in the Xolair group at

the end of the 28-week core trial compared to that observed in the placebo treated group (difference from placebo $p \leq 0.001$ in studies 1 and 2).

In study 3 the safety and corticosteroid-sparing effect of omalizumab was demonstrated in 246 patients with severe allergic asthma requiring daily treatment with high-dose inhaled corticosteroids (fluticasone $\geq 1,000$ micrograms/day) and in whom long-acting beta2-agonists were allowed. The study included a 16-week steroid stable phase with study medication added, followed by a 16-week steroid reduction phase. The percent reduction in inhaled corticosteroid dose at the end of the treatment phase was significantly greater in omalizumab-treated patients versus placebo patients (median 60% vs. 50%, $p=0.003$). The proportion of omalizumab patients who were able to reduce their fluticasone dose to ≤ 500 micrograms/day was 60.3% versus 45.8% in the placebo group.

In study 4 the safety and efficacy of omalizumab were demonstrated in 405 patients with co-morbid allergic asthma and perennial allergic rhinitis. Eligible patients had both symptomatic allergic asthma and perennial allergic rhinitis. Patients were treated with omalizumab or placebo for 28 weeks as add-on therapy to ≥ 400 micrograms of Budesonide Turbohaler. Inhaled long-acting beta2-agonists (39%) and nasal corticosteroids (17%) were allowed.

The co-primary endpoints for study 4 were the incidence of asthma exacerbations (worsening of asthma requiring systemic corticosteroids or a doubling of the patient's baseline budesonide dose) and the proportion of patients in each treatment group with a ≥ 1.0 improvement from baseline at the end of the treatment phase in both asthma and rhinitis specific quality of life assessments (Juniper Quality of Life Assessment).

Patients treated with omalizumab had a significantly lower incidence of asthma exacerbations than patients receiving placebo (20.6% omalizumab vs 30.1% placebo, $p=0.02$) and there was a significantly higher proportion of omalizumab-treated than placebo patients that improved by ≥ 1.0 points in both asthma and rhinitis specific quality of life assessments (57.7% omalizumab vs 40.6% placebo, $p < 0.0001$).

The reduction in exacerbations and improvements of quality of life in omalizumab-treated patients were seen in the context of statistically significant improvements in both rhinitis and asthma symptoms, and lung function, compared to placebo.

In study 5 the efficacy and safety of Xolair were demonstrated in a 28-week study involving 419 severe allergic asthmatics, ages 12 to 79 years, who had reduced lung function (Forced Expiratory Volume/1 second: FEV₁ 40 to 80% predicted) and poor asthma symptom control despite receiving $>1,000$ micrograms of beclomethasone dipropionate (or equivalent) plus long-acting beta2-agonist. Eligible patients had experienced multiple asthma exacerbations requiring systemic corticosteroid treatment or had been hospitalised or attended an emergency room due to a severe asthma exacerbation in the past year despite continuous treatment with high-dose inhaled corticosteroids and long-acting beta2-agonist. Subcutaneous Xolair or placebo were administered as add-on therapy to $>1,000$ micrograms beclomethasone dipropionate (or equivalent) plus long-acting beta2-agonist. Oral corticosteroid (22%), theophylline (27%) and anti-leukotriene (35%) maintenance therapies were allowed. In the treatment phase concomitant asthma therapy was not changed.

The rate of asthma exacerbations requiring treatment with bursts of systemic corticosteroids was the primary endpoint. Omalizumab reduced the rate of asthma exacerbations by 19% ($p = 0.153$). Further evaluations which did show statistical significance ($p < 0.05$) in favour of Xolair included reductions in severe exacerbations (where patient's lung function was reduced to below 60% of personal best and requiring systemic corticosteroids) and asthma-related emergency visits (comprised of hospitalisations, emergency room, and unscheduled doctor visits), and improvements in Physician's overall assessment of treatment effectiveness, Asthma-related Quality of Life (AQL), asthma symptoms and lung function. A physician's overall assessment was performed in the five above mentioned studies as a broad measure of asthma control performed by the treating physician. The physician was able to take into account Peak Expiratory Flow (PEF), day and night time symptoms, rescue medication use, spirometry and exacerbations. In all five studies a significantly greater

proportion of Xolair treated patients were judged to have achieved either a marked improvement or complete control of their asthma compared to placebo patients.

Children 6 to <12 years of age

The primary support for safety and efficacy of Xolair in the 6 to <12 years of age group comes from one randomised, double-blind, placebo controlled, multi-centre trial (study 6) and an additional supportive study (study 7).

Study 6 was a 52 week study that evaluated the safety and efficacy of Xolair as add-on therapy in 628 allergic asthmatics who were uncontrolled despite treatment with regular inhaled corticosteroids (fluticasone DPI ≥ 200 mcg/day or equivalent) with or without other controller asthma medications. Eligible patients were those with a diagnosis of asthma >1 year and a positive skin prick test to at least one perennial aeroallergen and a history of clinical features of moderate to severe persistent asthma including daytime and/or night-time symptoms along with a history of experiencing exacerbations within the year prior to study entry. Long-acting beta2-agonists (67.4%), anti-leukotriene (36.6%) and oral corticosteroid (1.3%) maintenance therapies were allowed. During the first 24 weeks of treatment, a patient's steroid doses remained constant from baseline and this was followed by a 28 week period during which inhaled corticosteroid adjustment was allowed.

A clinically significant exacerbation was defined as a worsening of asthma symptoms as judged clinically by the investigator, requiring doubling of the baseline inhaled corticosteroid dose for at least 3 days and/or treatment with rescue systemic (oral or iv) corticosteroids for at least 3 days.

Exacerbation rates during the 52-week double-blind treatment period in Xolair patients with FEV₁ >80% at baseline had relative decreases of 43% in asthma exacerbations compared with placebo ($p < 0.001$). Xolair patients had statistically significant reduction in the rate of asthma exacerbations irrespective of concomitant long-acting beta2-agonist use at baseline compared with placebo patients, representing a 45% decrease for long-acting beta2-agonist users and a 42% decrease for long-acting beta2-agonist non-users ($p < 0.001$ and $p = 0.011$, respectively).

Study 7 was a 28 week randomized, double blind, placebo-controlled study primarily evaluating safety in 334 children, aged 6-12 years of age, with asthma who were well controlled with inhaled corticosteroids. During the first 16 weeks of treatment, patients' steroid doses remained constant from baseline and this was followed by a 12 week steroid dose reduction period. The study assessed percent reduction in the dose of beclomethasone dipropionate (BDP) and the proportion of patients with a reduction in the dose of BDP at 28 weeks. The percent reduction in the dose of BDP at 28 weeks was higher in the Xolair group than in the placebo group (median reduction 100% vs. 66.7%, $p = 0.001$) as well as the proportion of patients with a reduction in the dose of BDP ($p = 0.002$). Frequency and incidence of asthma exacerbation episodes during the steroid dose-reduction phase were also lower in the omalizumab group (mean rate 0.42 vs. 0.72, $p < 0.001$; percent patients with exacerbations 18% vs. 39%, $p < 0.001$). A trend for superiority of omalizumab with respect to reduction of exacerbation frequency and incidence was evident during the first 16 weeks of the 24 week treatment period. 55.7% omalizumab patients had a complete (100%) reduction in corticosteroid dose at the end of the 28 week treatment period compared with a 43.2% of placebo patients. In addition, more omalizumab patients had a $\geq 50\%$ reduction in corticosteroid dose compared with placebo (80.4% vs. 69.5%. $p = 0.017$).

A physician's overall assessment was performed in the two above mentioned studies (6 and 7) as a broad measure of asthma control performed by the treating physician. The physician was able to take into account PEF, day and night time symptoms, rescue medication use, spirometry and exacerbations. In both studies a significantly greater proportion of Xolair treated patients were judged to have achieved either a marked improvement or complete control of their asthma compared to placebo patients.

Chronic rhinosinusitis with nasal polyps (CRSwNP)

The safety and efficacy of Xolair were evaluated in two randomized, multicenter, double-blind, placebo-controlled clinical trials that enrolled patients with CRSwNP (study 1, N=138; study 2, N=127). Patients received Xolair or placebo subcutaneously every 2 or 4 weeks, with dosage and frequency according to Tables 2 and 3 (see section 4.2 Dose and Method of administration). All patients received background intranasal mometasone therapy throughout the study. Prior sino-nasal surgery or prior systemic corticosteroid usage were not required for inclusion in the studies. Patients received Xolair or placebo for 24 weeks followed by a 4-week follow-up period. Demographics and baseline characteristics, including allergic comorbidities, are described in Table 8.

Table 8 Demographics and baseline characteristics of CRSwNP Studies

Parameter	Nasal Polyp Study 1 N=138	Nasal Polyp Study 2 N=127
Mean age (years) (SD)	51.0 (13.2)	50.1 (11.9)
% Male	63.8	65.4
Patients with systemic corticosteroid use in the previous year (%)	18.8	26.0
Mean bilateral endoscopic NPS* (SD), range 0-8	6.2 (1.0)	6.3 (0.9)
Mean nasal congestion (NC) score* (SD), range 0-3	2.4 (0.6)	2.3 (0.7)
Mean sense of smell score* (SD) range 0-3	2.7 (0.7)	2.7 (0.7)
Mean SNOT-22 total score* (SD) range 0-110	60.1 (17.7)	59.5 (19.3)
Mean blood eosinophils (cells/ μ l) (SD)	346.1 (284.1)	334.6 (187.6)
Mean total IgE IU/mL (SD)	160.9 (139.6)	190.2 (200.5)
Asthma (%)	53.6	60.6
Mild (%)	37.8	32.5
Moderate (%)	58.1	58.4
Severe (%)	4.1	9.1
Aspirin exacerbated respiratory disease (%)	19.6	35.4
Allergic rhinitis	43.5	42.5

SD=standard deviation; NPS=nasal polyp score; SNOT-22 = Sino-Nasal Outcome Test 22 Questionnaire; IgE = Immunoglobulin E; IU=international units. For NPS, NCS, sense of smell, post nasal drip, runny nose, and SNOT-22 scores higher scores indicate greater disease severity

The co-primary endpoints were bilateral nasal polyp score (NPS) and average daily nasal congestion score (NCS) at Week 24. NPS was measured via endoscopy at baseline and pre-specified time points and scored (range 0-4 per nostril) for a total NPS (range 0/best-8/worst). Nasal congestion was measured by a daily NCS (range 0/best-3/worst). Patients were required to have NPS \geq 5 and weekly average of NCS > 1 prior to randomization, despite use of intranasal mometasone. The mean NPS at baseline was balanced between the two treatment groups in both studies.

In both CRSwNP studies 1 and 2, patients who received Xolair had a statistically significant greater improvement from baseline at Week 24 in NPS and weekly average NCS than patients who received placebo. Results from Nasal Polyps Study 1 and 2 are shown in Table 9.

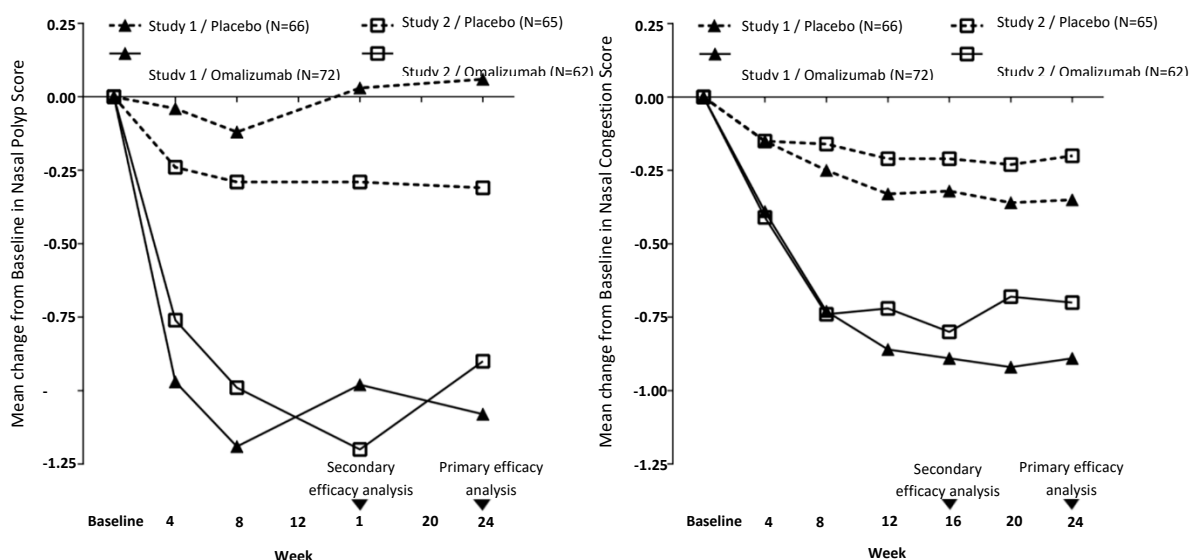
The greater improvements in NPS and NCS in the Xolair group compared to the placebo group were observed as early as the first assessment at Week 4 in both studies as seen in Figure 1. The LS mean difference in change from baseline at Week 4 in NPS in Xolair compared to placebo was -0.92 (95% CI: -1.37, -0.48) in study 1 and -0.52 (95% CI: -0.94, -0.11) in study 2. The LS mean difference in change from baseline at Week 4 in NCS in Xolair compared to placebo was -0.25 (95% CI: -0.46, -0.04) in study 1 and -0.26 (95% CI: -0.45, -0.07) in study 2. However, statistical tests at this time point were not pre-specified.

Table 9 Change from baseline at Week 24 in clinical scores from Nasal Polyp Study 1 and Nasal Polyp Study 2

	Nasal Polyp Study 1		Nasal Polyp Study 2	
	Placebo	Xolair	Placebo	Xolair
N	66	72	65	62
Nasal polyp score				
Baseline mean	6.32	6.19	6.09	6.44
LS mean change at Week 24	0.06	-1.08	-0.31	-0.90
Difference in LS means vs. placebo	-1.14		-0.59	
95% CI for difference	-1.59, -0.69		-1.05, -0.12	
p-value	<0.0001		0.0140	
7-day average of daily nasal congestion score				
Baseline mean	2.46	2.40	2.29	2.26
LS mean change at week 24	-0.35	-0.89	-0.20	-0.70
Difference in LS means vs. placebo	-0.55		-0.50	
95% CI for difference	-0.84, -0.25		-0.80, -0.19	
p-value	0.0004		0.0017	
UPSIT				
Baseline mean	13.56	12.78	13.27	12.87
LS mean change at week 24	0.63	4.44	0.44	4.31
Difference in LS means vs. placebo	3.81		3.86	
95% CI for difference	1.38, 6.24		1.57, 6.15	
p-value	0.0024		0.0011	

LS=least-square

Figure 1 Mean change from baseline in nasal congestion score and mean change from baseline in nasal polyp score by treatment group in Nasal Polyp Study 1 and Nasal Polyp Study 2



A key secondary endpoint was the assessment of the change from baseline at Week 24 of the total nasal symptom score (TNSS). Patient-reported TNSS was the sum of four equally weighted individual daily symptom scores: NCS, sense of smell score, posterior rhinorrhea score, and anterior rhinorrhea score. The TNSS ranged from 0/best-12/worst. Xolair significantly improved the average daily TNSS

compared to placebo. The LS mean difference for change from baseline to Week 24 was -1.91 points (95% CI: -2.85, -0.96; $p = 0.0001$) in study 1 and -2.09 points (95% CI: -3.00, -1.18; $p < 0.0001$) in study 2.

The SNOT-22 (Sino-Nasal Outcome Test), combines measures of sinonasal symptoms with psychological and sleep dysfunction measures and has a score that ranges from 0 to 110 (0/best-110/worst). The minimal clinically important difference has been established as 8.9. The proportion of patients who experienced an improvement from baseline at Week 24 of at least the MID (8.9 points) in SNOT-22 score for the individual studies was: Study GA39688 - 30/65 (46.2%) in the placebo arm and 53/69 (76.8%) in the omalizumab arm, OR of 4.55 (95% CI 2.07, 9.97; $p=0.0002$); Study GA39855 in and 23/63 (36.5%) in the placebo arm and 39/59 (66.1%) in the omalizumab arm, OR 3.71 (95% CI: 1.72, 8.04; $p=0.0009$).

Additional secondary endpoint analyses included Week 16 assessments of NPS and NCS. Xolair significantly improved the NPS at week 16, (range 0/best-8/worst) compared to placebo. The LS mean difference for change from baseline to Week 16 in Xolair compared to placebo was -1.01 (95% CI: -1.43, -0.60; $p < 0.0001$) in study 1 and -0.91 (95% CI: -1.39, -0.44; $p = 0.0002$) in study 2 [XX]. Xolair significantly improved the NCS at week 16, (range 0/best-3/worst) compared to placebo. The LS mean difference for change from baseline to Week 16 in average daily NCS in Xolair compared to placebo was -0.57 (95% CI: -0.83, -0.31; $p < 0.0001$) in study 1 and of -0.59 (95% CI: -0.87, -0.30; $p < 0.0001$) in study 2.

In a pre-specified pooled analysis of rescue treatment (systemic corticosteroids for ≥ 3 consecutive days or nasal polypectomy) during the 24-week treatment period, the proportion of patients requiring rescue treatment was lower in Xolair compared to placebo (2.3% versus 6.2%, respectively).

The treatment arm difference was not statistically significant (OR: 0.38; 95% CI: 0.10, 1.49; nominal $p=0.1639$). There were no sino-nasal surgeries reported in either study.

Clinical efficacy and safety in Chronic Idiopathic Urticaria (CIU)

The efficacy and safety of Xolair were demonstrated in two randomised, placebo-controlled phase III studies (study Q4881g, Q4882g) in patients with CIU who remained symptomatic despite H1 antihistamine therapy at the approved dose. All patients received Xolair or placebo in addition to H1 antihistamines. A third study (Q4883g) primarily evaluated the safety of Xolair in patients with CIU who remained symptomatic despite treatment with H1 antihistamines at increased doses and H2 antihistamine and/or leukotriene receptor antagonist (LTRA) treatment. All patients received Xolair or placebo in addition to H1 antihistamines up to 4 times the approved dose, and/or H2 antihistamines and/or LTRAs.

CIU patients with external triggers were excluded from these trials. The three studies enrolled 975 patients aged between 12 and 75 years (mean age 42.3 years; 39 patients 12-17 years, 54 patients ≥ 65 years; 259 males and 716 females). All patients were required to have inadequate symptom control, as assessed by a weekly urticaria activity score (UAS7, range 0 to 42) of ≥ 16 , and a weekly itch severity score (which is a component of the UAS7; range 0 to 21) of ≥ 8 for the 7 days prior to randomisation, despite having used an antihistamine for at least 2 weeks beforehand.

In studies Q4481g and Q4882g, patients had a mean weekly itch severity score of between 13.7 and 14.5 at baseline and a mean UAS7 score of 29.5 and 31.7 respectively. Patients in safety study Q4883g had a mean weekly itch severity score of 13.8 and a mean UAS7 score of 31.2 at baseline. Across all three studies, patients reported receiving on average 4 to 6 medications (including H1 antihistamines) for CIU symptoms prior to study enrolment. Patients received Xolair at 75 mg, 150 mg or 300 mg or placebo by subcutaneous injection every 4 weeks for 24 and 12 weeks in studies 1 and 2, respectively, and 300 mg or placebo by subcutaneous injection every 4 weeks for 24 weeks in study 3. All studies had a 16 week treatment-free follow-up period.

Table 10 Efficacy endpoints

Change from baseline to week 12 in weekly Itch Severity Score (ISS, range 0-21)	Primary endpoint in studies Q4881g and Q4882g Secondary endpoint in safety study Q4883g
Time to MID ^a response (decrease from baseline of ≥5 points) in weekly ISS up to week 12	Secondary endpoints in all three studies Q4881g, Q4882g and Q4883g
Change from Baseline to week 12 in Urticaria Activity score during a 7 day period (UAS7 ^b , range 0-42)	
Proportion of patients with Urticaria Activity Score during a 7-Day Period ≤ 6 (UAS7 ^b ≤ 6) at week 12	
Proportion of patients with Urticaria Activity Score during a 7-Day Period = 0 (UAS7 ^b = 0) at week 12 ^c	
Changes from baseline in the weekly number of hives score at week 12	
Change from baseline to week 12 in overall Dermatology Life Quality Index (DLQI)	
Proportion of patients with angioedema-free days from week 4 to week 12 ^d	

^a MID: Minimally Important Difference

^b UAS7: Composite of itch severity and number of hives measured daily and totalled over one week

^c Post hoc analysis for study Q4882g

^d Mean proportion of angioedema-free days from week 4 to week 12 was calculated for the entire study population, including patients asymptomatic for angioedema.

In studies Q4881g and Q4882g the 75 mg dose did not consistently meet either the primary efficacy endpoint or a number of secondary endpoints. It was deemed not efficacious and therefore not further presented.

Change from baseline to week 12 in weekly itch severity score

The primary efficacy endpoint, change from baseline to week 12 in weekly itch severity score was met by both the 150 mg and 300 mg doses in studies Q4881g and Q4882g and by the 300 mg dose in Q4883g (secondary endpoint; see Table 11).

Table 11 Change from baseline to week 12 in weekly itch severity score, Studies Q4881g, Q4882g and Q4883g (mITT population*)

	Placebo	Omalizumab 150mg	Omalizumab 300mg
Study Q4881g			
N	80	80	81
Mean (SD)	-3.63 (5.22)	-6.66 (6.28)	-9.40 (5.73)
Difference in LS means vs. placebo ¹	-	-2.95	-5.80
95% CI for difference	-	-4.72, -1.18	-7.49, -4.10
P-value vs. placebo ²	-	0.0012	<0.0001
Study Q4882g			
N	79	82	79
Mean (SD)	-5.14 (5.58)	-8.14 (6.44)	-9.77 (5.95)
Difference in LS means vs. placebo ¹	-	-3.04	-4.81

	Placebo	Omalizumab 150mg	Omalizumab 300mg
95% CI for difference	-	-4.85, -1.24	-6.49, -3.13
P-value vs. placebo ²	-	0.0011	<0.0001
Study Q4883g			
n	83	-	252
Mean (SD)	-4.01 (5.87)	-	-8.55 (6.01)
Difference in LS means vs. placebo ¹	-	-	-4.52
95% CI for difference	-	-	-5.97, -3.08
P-value vs. placebo ²	-	-	<0.0001

*Modified intent-to-treat (mITT) population: Included all patients who were randomized and received at least one dose of study medication.

BOCF (Baseline Observation Carried Forward) was used to impute missing data.

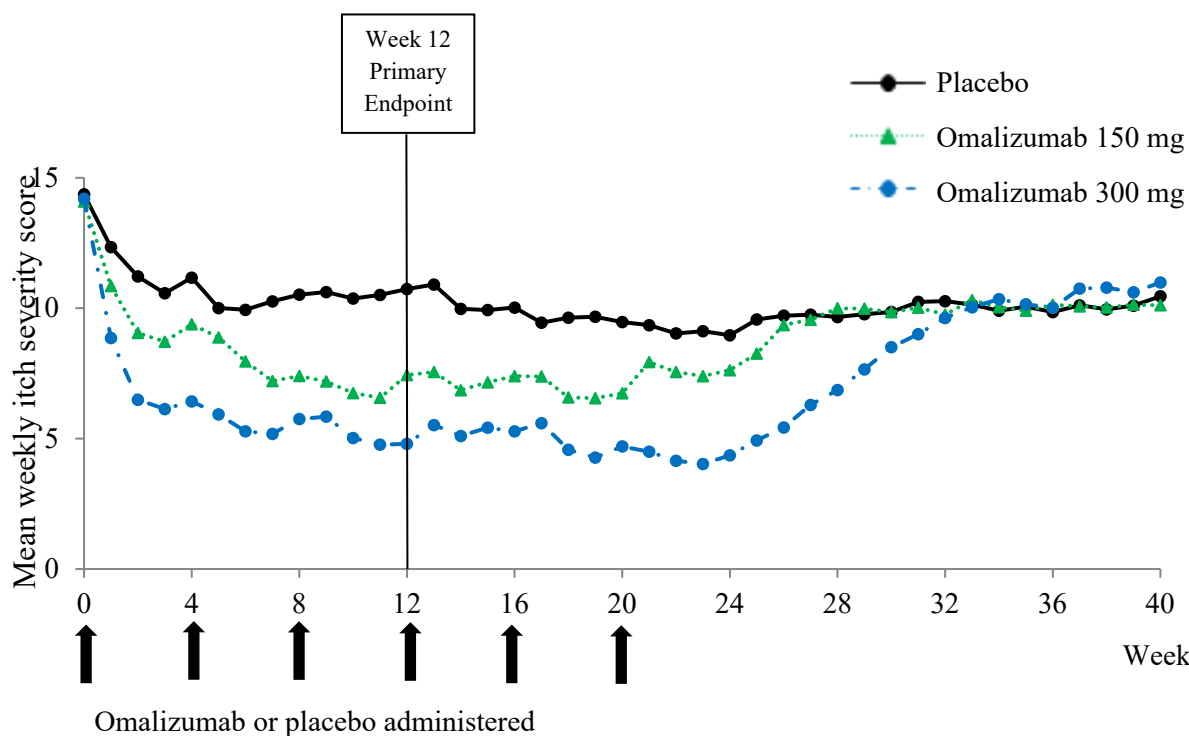
¹ The LS mean was estimated using an ANCOVA model. The strata were baseline weekly itch severity score (< 13 vs. ≥13) and baseline weight (< 80 kg vs. ≥ 80 kg).

² p-value is derived from ANCOVA t-test

Figure 2 shows the mean weekly itch severity score over time in study Q4881g. The mean weekly itch severity scores significantly decreased in both treatment groups with a maximum effect around week 12 that was sustained over the 24-week treatment period. In studies Q4883g (300 mg over the 24-week treatment period) and Q4882g (150 mg and 300 mg over the 12-week treatment period) the results were similar to those of study Q4881g.

In all three studies (see Figure 2 for study Q4881g), the mean weekly itch severity score for both doses increased gradually during the 16-week treatment-free follow-up period, consistent with symptom re-occurrence. Mean values at the end of the follow-up period were similar to the placebo group, but lower than respective mean baseline values.

Figure 2 Mean weekly itch severity score over time, Study Q4881g (BOCF, mITT population)



BOCF=baseline observation carried forward; mITT=modified intention-to-treat population

Secondary endpoints for studies Q4881g, Q4882g and Q4883g are presented in Table 12.

Table 12 Secondary efficacy endpoints in Studies Q4881g, Q4882g and Q4883g (mITT population*)

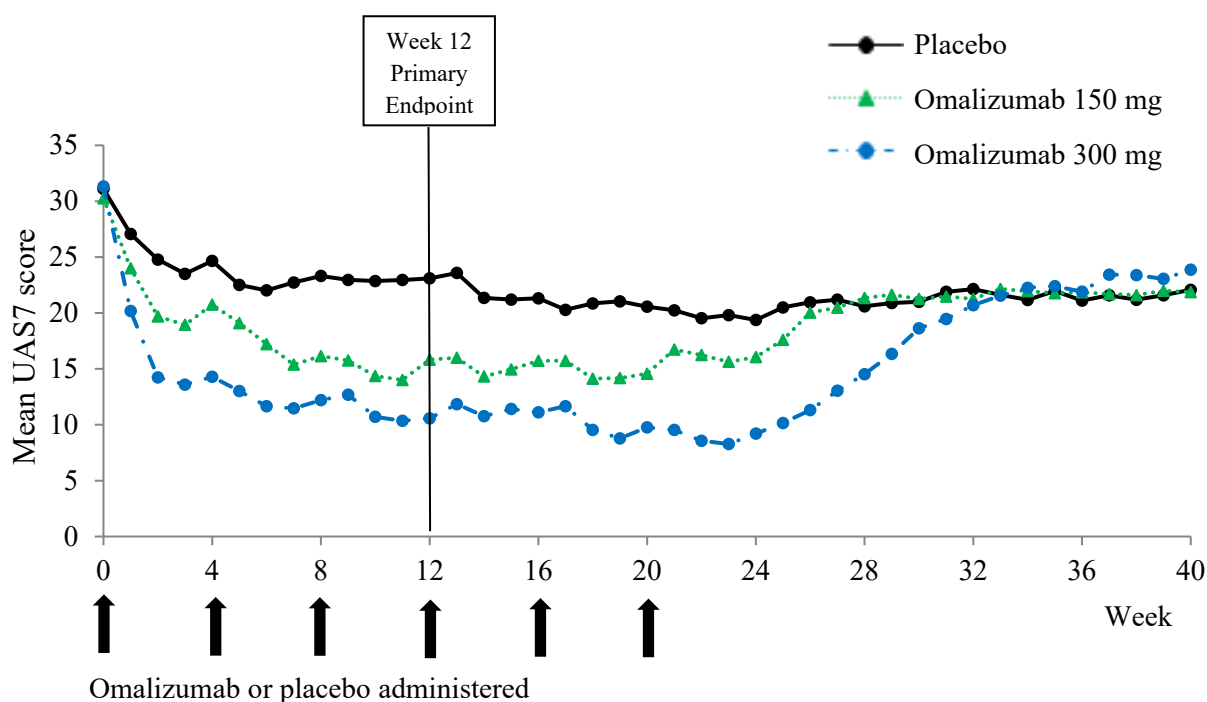
	Placebo	Omalizumab 150mg	Omalizumab 300mg
Time to MID response in weekly ISS up to week 12 (median weeks)			
Q4881g	4	2 (p=0.0301)	1 (p<0.0001)
Q4882g	4	2 (p=0.0101)	1 (p<0.0001)
Q4883g	5	NA	2 (p<0.0001)
Change from baseline to week 12 in UAS7 (mean)			
Q4881g	-8.01	-14.44 (p = 0.0008)	-20.75 (p<0.0001)
Q4882g	-10.36	-17.89 (p = 0.0001)	-21.74 (p<0.0001)
Q4883g	-8.50	NA	-19.01 (p<0.0001)
Changes from baseline in the weekly number of hives score at week 12 (mean)			
Q4881g	-4.37	-7.78 (p=0.0017)	-11.35 (p<0.0001)
Q4882g	-5.22	-9.75 (p<0.0001)	-11.97 (p<0.0001)
Q4883g	-4.49	NA	-10.46 (p<0.0001)
Change from baseline to week 12 in overall Dermatology Life Quality Index (DLQI) (mean)			
Q4881g	-6.13	-8.00 (p=0.2286)	-10.29 (p<0.0001)
Q4882g	-6.09	-8.29 (p=0.0215)	-10.15 (p=0.0004)
Q4883g	-5.11	NA	-9.69 (p<0.0001)
Proportion of angioedema-free days from week 4 to week 12 (mean)			
Q4881g	88.2%	89.6% (p=0.1747)	96.1% (p<0.0001)
Q4882g	89.2%	91.6% (p=0.0905)	95.5% (p<0.0001)
Q4883g	88.1%	NA	91.0% (p<0.0006)
Proportion of patients with UAS7 ≤ 6 at week 12 (% patients)			
Q4881g	11.3	40.0 (p<0.0001)	51.9 (p<0.0001)
Q4882g	19.0	42.7 (p=0.0010)	65.8 (p<0.0001)
Q4883g	12.0	NA	52.4 (p<0.0001)
Proportion of patients with UAS7 = 0 at week 12 (% patients)			
Q4881g	8.8	15.0 (p=0.2087)	35.8 (p<0.0001)
Q4882g	5.1	22.0 (p=0.0019)	44.3 (p<0.0001)
Q4883g	4.8	NA	33.7 (p<0.0001)

Placebo	Omalizumab 150mg	Omalizumab 300mg
*Modified intent-to-treat (mITT) population: included all patients who were randomised and received at least one dose of study medication		
p-value was derived using Cox proportional hazard model, ANCOVA, stratified Wilcoxon, and stratified CMH, as appropriate, comparing between active treatment and placebo		
NA: Not applicable.		
BOCF: Baseline Observation Carried Forward		

Figure 3 shows mean UAS7 over time in study Q4881g, displaying a significant decrease from baseline in both treatment groups with a maximum effect around week 12. The magnitude of the effect was maintained during the 24-week treatment period. In studies Q4882g (150 mg and 300 mg over the 12-week treatment period) and Q4883g (300 mg over 24-week treatment period) the results were similar to those of study Q4881g.

In all three studies (see Figure 3 for study Q4881g), the UAS7 for both omalizumab treatment groups increased gradually during the 16-week treatment-free follow-up period, consistent with symptom re-occurrence. Mean values at the end of the follow-up period were similar to the placebo group but lower than respective mean baseline values.

Figure 3 Mean UAS7 over time, Study Q4881g (BOCF, mITT population)



BOCF=baseline observation carried forward; mITT=modified intention-to-treat population; UAS7= urticaria activity score over 7 days

Efficacy after 24 weeks of treatment

Table 13 shows the results after 24 weeks of treatment. Similar magnitudes of response are seen as at 12 weeks.

Table 13 Efficacy results after 24 weeks of treatment, Studies Q4881g and Q4883g (mITT population*)

Parameter			Omalizumab	Omalizumab
Study	Week	Placebo	150 mg	300 mg
Change from baseline in weekly itch severity score (BOCF), mean				
Study Q4881g	Week 24	-5.41	-6.47	-9.84**
Study Q4883g	Week 24	-4.03	NA	-8.60**
Change from baseline in UAS7 (BOCF), mean				
Study Q4881g	Week 24	-11.73	-14.21	-22.11**
Study Q4883g	Week 24	-8.85	NA	-19.15**
Proportion of patients with UAS7 ≤ 6, % patients				
Study Q4881g	Week 24	25.0	36.3	61.7**
Study Q4883g	Week 24	16.9	NA	55.6**
Proportion of patients with UAS7 = 0, % patients				
Study Q4881g	Week 24	12.5	20.0	48.1**
Study Q4883g	Week 24	3.6	NA	42.5**

*Modified intent-to-treat (mITT) population: included all patients who were randomised and received at least one dose of study medication

** p -value ≤ 0.0001 for the corresponding test statistics between the treatment and the placebo

NA: Not applicable.

BOCF: Baseline Observation Carried Forward

The treatment duration in the studies were 12 weeks (Q4882g) and 24 weeks, respectively (Q4881g and Q4883g), therefore the Clinical trial experience of long-term treatment beyond 6 months in this indication is limited.

5.2 Pharmacokinetic properties

General characteristics

Absorption

After subcutaneous administration, omalizumab is absorbed with an average absolute bioavailability of 62%. The pharmacokinetics of omalizumab are linear at doses greater than 0.5 mg/kg.

Administration of Xolair manufactured as a lyophilized or liquid formulation resulted in similar serum concentration-time profiles of omalizumab.

Distribution

In vitro, omalizumab forms complexes of limited size with IgE. Precipitating complexes and complexes larger than one million Daltons in molecular weight are not observed *in vitro* or *in vivo*.

Tissue distribution studies in cynomolgus monkeys showed no specific uptake of ¹²⁵I-omalizumab by any organ or tissue.

Elimination

Clearance of omalizumab involves IgG clearance processes as well as clearance via specific binding and complex formation with its target ligand, IgE. Liver elimination of IgG includes degradation in the liver reticuloendothelial system (RES) and endothelial cells. Intact IgG is also excreted in bile. In studies with mice and monkeys, omalizumab: IgE complexes were eliminated by interactions with Fc γ receptors within the RES at rates that were generally faster than IgG clearance.

Patients with Allergic Asthma

Absorption

Following a single subcutaneous dose in adult and adolescent patients with asthma, omalizumab was absorbed slowly, reaching peak serum concentrations after an average of 7 to 8 days. Following multiple doses of omalizumab, areas under the serum concentration-time curve from Day 0 to Day 14 at steady state were up to 6-fold of those after the first dose.

Distribution

The apparent volume of distribution of omalizumab in patients with asthma following subcutaneous administration was 78 ± 32 mL/kg.

Elimination

In asthma patients omalizumab serum elimination half-life averaged 26 days, with apparent clearance averaging 2.4 ± 1.1 mL/kg/day. Doubling of body weight approximately doubled apparent clearance.

Age, Race/Ethnicity, Gender, Body Mass Index

The population pharmacokinetics of omalizumab were analysed to evaluate the effects of demographic characteristics. Analyses of these data suggest that no dose adjustments are necessary in asthma patients for age (6 to 76 years), race, ethnicity, gender or body mass index.

Patients with Chronic rhinosinusitis with nasal polyps (CRSwNP)

The population pharmacokinetics analyses of omalizumab suggested that pharmacokinetics of omalizumab in CRSwNP were consistent with that in asthma. Graphical covariate analyses were performed to evaluate the effects of demographic characteristics and other factors on omalizumab exposure and clinical responses. These analyses demonstrate that no dose adjustments are necessary for age (18 to 75 years) or gender. Race and ethnicity data are too limited in CRSwNP to inform dose adjustment.

Patients with Chronic Idiopathic Urticaria (CIU)

Absorption

Following a single subcutaneous dose in adult and adolescent patients with CIU, omalizumab was absorbed slowly, reaching peak serum concentrations after an average of 6 to 8 days.

In patients with CIU, omalizumab exhibited linear pharmacokinetics across the dose range of 75 mg to 600 mg given as a single subcutaneous dose. Following doses of 75 mg, 150 mg or 300 mg every 4 weeks, trough serum concentrations of omalizumab increased proportionally with the dose level.

Distribution

Based on population pharmacokinetic, distribution of omalizumab in CIU patients was similar to that in patients with allergic asthma.

Elimination

In patients with CIU, based on population pharmacokinetic simulations, omalizumab serum elimination half-life at steady state averaged 24 days and apparent clearance at steady state averaged 240 mL/day (corresponding to 3.0 mL/kg/day for an 80 kg patient).

Age, Race/Ethnicity, Gender, Body Mass Index, Baseline IgE, anti-FcεRI autoantibodies, co-mediations

The effects of demographic covariates and other factors on omalizumab exposure were evaluated using population pharmacokinetics. In addition, covariate effects were evaluated by analyzing the relationship between omalizumab concentrations and clinical responses. These analyses suggest that no dose adjustments are necessary in patients with CIU for age (12 to 75 years), race/ethnicity, gender, body weight, body mass index, baseline IgE, anti-FcεRI autoantibodies or concomitant use of H2 antihistamines or leukotriene receptor antagonists (LTRAs).

Patients with renal and hepatic impairment

There are no pharmacokinetic or pharmacodynamic data in patients with renal or hepatic impairment (see section 4.4).

5.3 Preclinical safety data

There was no evidence of a systemic anaphylactic response due to mast-cell degranulation in either adult or juvenile cynomolgus monkeys. Circulating omalizumab: IgE antibody complexes were present in all monkey studies, however there was no evidence of immune complex-mediated disease in any organ (including the kidney) following omalizumab administration. Omalizumab: IgE complexes do not fix complement or mediate complement-dependent cytotoxicity.

Chronic administration of omalizumab was well tolerated in non-human primates (both adult and juvenile animals), with the exception of a dose-related decrease in platelet counts that occurred in several non-human primate species, at serum concentrations generally in excess of maximum human exposure in pivotal clinical trials. Juvenile monkeys were more sensitive to the platelet effects than adult monkeys. In addition, acute haemorrhage and inflammation were observed at injection sites in cynomolgus monkeys, consistent with a localised immune response to repeated subcutaneous administration of a heterologous protein. Formal carcinogenicity studies have not been conducted with omalizumab.

Antibodies to omalizumab were detected in some monkeys following subcutaneous or intravenous administration. This was not unexpected based on administration of a heterologous protein. Some animals could not be evaluated because of high serum omalizumab concentrations, high IgE levels, or both. However, the animals maintained high serum omalizumab concentrations throughout the treatment periods of the studies, and there was no apparent toxicity due to the presence of anti-omalizumab antibodies.

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

L-arginine hydrochloride, L-histidine hydrochloride monohydrate, L-histidine, polysorbate 20, water for injection.

6.2 Incompatibilities

This medicinal product must not be mixed with other medicinal products.

6.3 Shelf life

18 months

6.4 Special precautions for storage

Solution for injection in pre-filled syringe: Store at 2°C to 8°C. Do not freeze. Store in original package and protect from light. The product may be kept for a total of 48 hours at 25°C.

6.5 Nature and contents of container

Pre-filled syringe comprising a type I glass syringe barrel with staked needle (stainless steel), latex-free rubber plunger stopper, and rigid needle shield composed of a latex-free rubber needle shield covered by a rigid shell. Each pre-filled syringe is mounted with a plastic safety device (needle guard) to prevent from needle stick injury.

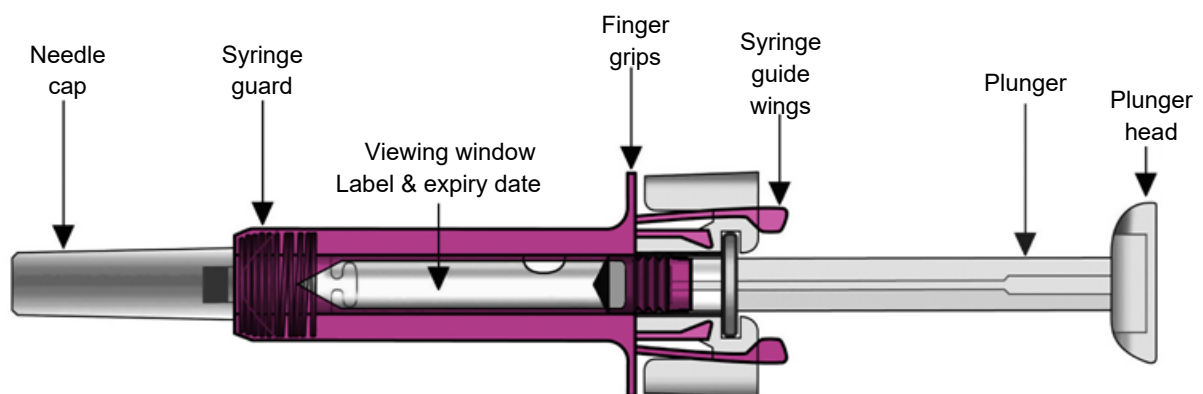
Available as a single packaged pre-filled syringe and in multipacks containing either 4 or 10 individually packaged pre-filled syringes. Not all pack sizes may be marketed.

6.6 Special precautions for disposal and other handling

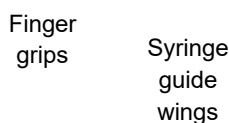
Read ALL the way through these instructions before injecting. If your doctor decides that you or a caregiver may be able to give your injections of Xolair at home, you need to be trained by your doctor, nurse or pharmacist before you inject yourself or others. Children (6 to less than 12 years of age) are not expected to inject Xolair themselves, however, if deemed appropriate by their doctor, a caregiver may give them their Xolair injection after proper training. The box contains a Xolair pre-filled syringe individually sealed in a plastic tray.

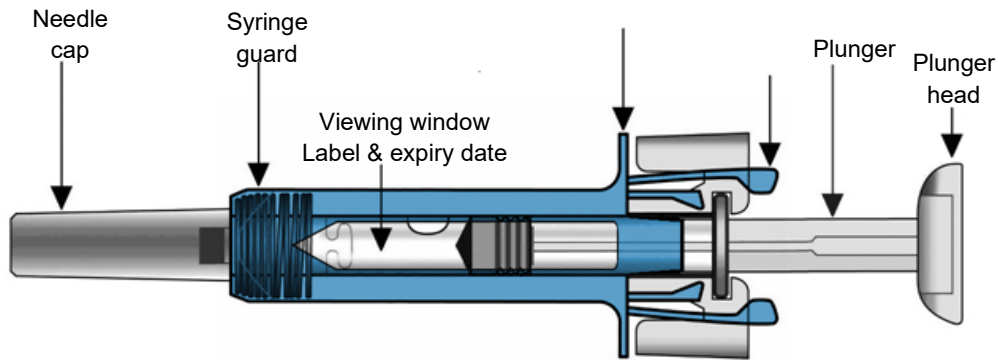
Xolair pre-filled syringe is available in two strengths, 75 mg and 150 mg. You may receive one or both strengths from the pharmacy.

Your Xolair 150 mg pre-filled syringe



Your Xolair 75 mg pre-filled syringe





After the medicine has been injected the syringe guard will be activated to cover the needle. This is intended to protect against accidental needle stick injuries.

Other items you need for your injection:

- Alcohol swab.
- Cotton ball or gauze.
- Sharps disposal container.
- Adhesive plaster



Important safety information

Caution: Keep the syringe out of the sight and reach of children.

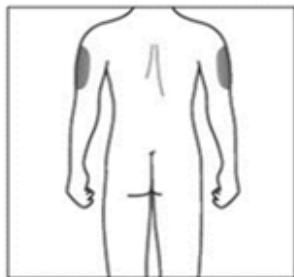
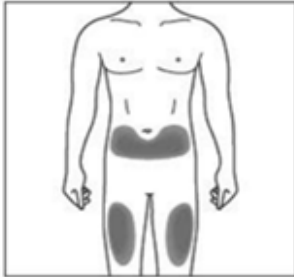
- The needle cap of the syringe may contain dry rubber (latex), which should not be handled by anyone who is sensitive to this substance.
- Do not open the sealed outer box until you are ready to use this medicine.
- Do not use this medicine if either the seal on the outer box or the seal of the plastic tray is broken, as it may not be safe for you to use.
- Do not use if the syringe has been dropped onto a hard surface or dropped after removing the needle cap.
- Never leave the syringe where others might tamper with it.
- Do not shake the syringe.
- Be careful not to touch the syringe guard wings before use. If the wings are touched, the syringe guard may be activated too early.
- Do not remove the needle cap until just before you give the injection.
- The syringe cannot be re-used. Dispose of the used syringe immediately after use in a sharps container.

Storage of the Xolair pre-filled syringe

- Store this medicine sealed in its outer box to protect it from light. Store in the refrigerator between 2°C and 8°C. DO NOT FREEZE.
- Prior to use, allow the syringe to reach room temperature (25°C) before preparing it for injection (it will take about 30 minutes). If necessary, the product may be returned to the refrigerator for later use. The total time that the syringe is kept at room temperature must not exceed 48 hours.

- **Do not** try to warm the syringe using an external heat source.
- Do not use the syringe after the expiry date which is stated on the outer box and syringe label. If it has expired, return the entire pack to the pharmacy.

The injection site



The injection site is the place on the body where you are going to use the syringe.









- The recommended site is the front of the thighs. You may also use the lower abdomen, but **not** the area 5 centimeters around the navel (belly button).
- If you need to give more than one injection for the full dose, choose a different injection site each time you inject.
- If injecting more than once into the same area, ensure a distance of at least 4 cm between injection sites.
- Do not inject into areas where the skin is tender, bruised, red, or hard. Avoid areas with scars or stretch marks.

If a caregiver is giving the injection, the outer upper arms may also be used.

Preparing the Xolair pre-filled syringe for use

Note: Depending on the dose prescribed to you by your doctor, you may need to prepare one or more pre-filled syringes, and inject the contents of them all.

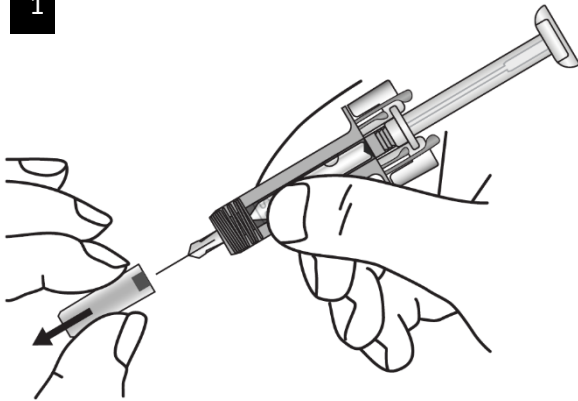
The following table give EXAMPLES of how many injections of each strength you need for a given dose. Please discuss with your doctor how frequent (4 weekly or 2 weekly) you need to administer each dose.

Dose	Syringes needed for the dose		
75 mg	1 blue (75 mg)		
150 mg	1 purple (150 mg)		
225 mg	1 blue (75 mg)	+ 1 purple (150 mg)	
300 mg	2 purple (150 mg)		
375 mg	1 blue (75 mg)	+ 2 purple (150 mg)	
450 mg	3 purple (150 mg)		
525 mg	1 blue (75 mg)	+ 3 purple (150 mg)	
600 mg	4 purple (150 mg)		

1. Take the box containing the syringe out of the refrigerator and leave it **unopened** for about 20 minutes so that it reaches room temperature (leave the syringe in the box to protect it from light).
2. When you are ready to use the syringe, wash your hands thoroughly with soap and water.
3. Clean the injection site with an alcohol swab.
4. Remove the plastic tray from the box and peel back the paper cover. Gripping the middle of the syringe guard, lift the syringe out of the tray.
5. Inspect the syringe. The liquid should be clear to slightly cloudy. Its color may vary from colorless to pale brownish-yellow. You may see an air bubble, which is normal. **DO NOT USE** if the syringe is broken or if the liquid looks distinctly cloudy or distinctly brown, or contains particles. In all these cases, return the entire pack to the pharmacy.
6. Holding the syringe horizontally, look into the viewing window to check the expiry date printed on the label. (Note: It is possible to rotate the inner part of the syringe assembly so that the label can be read in the viewing window. **DO NOT USE** if the product has expired. If expired, return the entire pack to the pharmacy.)

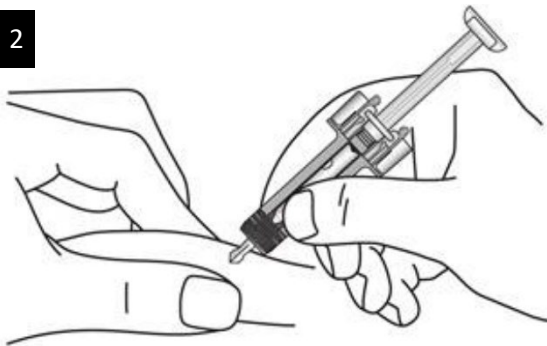
How to use the Xolair pre-filled syringe

1



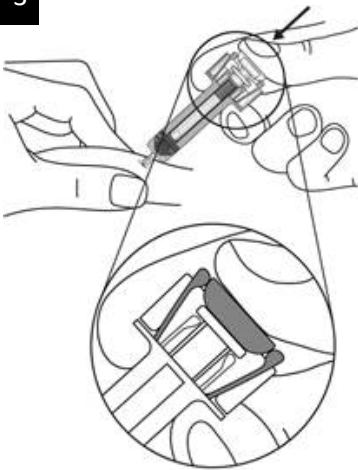
Carefully remove the needle cap from the syringe. Discard the needle cap. You may see a drop of liquid at the end of the needle. This is normal.

2



Gently pinch the skin at the injection site and insert the needle as shown. Push the needle all the way in to ensure that the medicine can be fully administered.

3



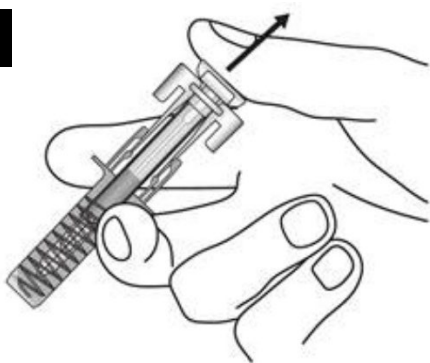
Hold the syringe as shown. **Slowly** depress the plunger **as far as it will go** so that the plunger head is completely between the syringe guard wings.

4



Keep the plunger fully depressed while you carefully lift the needle straight out from the injection site.

5



Slowly release the plunger and allow the syringe guard to automatically cover the exposed needle.

There may be a small amount of blood at the injection site. You can press a cotton ball or gauze over the injection site and hold it for 30 seconds. Do not rub the injection site. You may cover the injection site with a small adhesive bandage, if needed.

Disposal instructions



Dispose of the used syringe immediately in a sharps container (closable, puncture resistant container). For the safety and health of you and others, needles and used syringes **must never** be re-used. Any unused medicinal product or waste material should be disposed of in accordance with local requirements. Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help protect the environment.

7. MEDICINE SCHEDULE

Prescription Medicine

8. SPONSOR

Novartis New Zealand Limited

PO Box 99102
Newmarket
Auckland 1149
New Zealand

Telephone: 0800 354 335

® = Registered Trademark

9. DATE OF FIRST APPROVAL

Xolair solution for injection, pre-filled syringe: 10 August 2017

10. DATE OF REVISION OF THE TEXT

5 July 2024

SUMMARY TABLE OF CHANGES

Section changed	Summary of new information
All	Removal of reference to Xolair powder vial and diluent.

Internal document code: xol090724iNZ, CDS 26-Oct-2022