

Omalizumab (Xolair[®])

Reduce direct handling to a minimum and wear appropriate personal protective equipment	
Xolair [®] dosing may be weight based; ensure accuracy of documented weight before administration	
Form & Storage	Pre-filled syringe containing 75mg/mL and 150mg/mL solution for InjectionStore in a fridge at 2°C - 8°C
Reconstitution	Already in solution
Administration	For subcutaneous administration only
	 The syringe should be taken out of the refrigerator 20 minutes before injecting to allow it to reach room temperature. Doses of more than 150 mg should be divided across two or more injection sites. The injections are administered subcutaneously in the deltoid region of the arm. Alternatively, the injections can be administered in the thigh if there is any reason precluding administration in the deltoid region.
Monitoring	 Pre and post injection vital signs Local or systemic allergic reactions, including anaphylaxis and anaphylactic shock, may occur when taking omalizumab, also with onset after a long duration of treatment. Most of these reactions occurred within 2 hours after the first and subsequent injections of Xolair but some started beyond 2 hours and even beyond 24 hours after the injection. For the first three injections, the patient is monitored in the infusion unit for two hours For subsequent injections, the monitoring period should be 20 minutes Blood tests including FBC, U/E and LFTs monthly before first 3 doses by GP/phlebotomy, thereafter every three months by GP/Phlebotomy Once the patient is established on this treatment (more than three doses), subsequent injections may be given in the asthma out patient's clinic If the patient presents to the unit and meets the criteria in 7.7*, medical review may be required prior to administration of this medication
Documentation Requirements	Document batch numbers and expiry dates of vials in medical notes.
Additional Information	*See PPG-CUH-CUH-243 <u>Policy Procedure and Guidelines for Management</u> of Patients Attending CUH Infusion Unit for Intravenous Therapy CUH for more information
Information provided relates to Xolair [®] (Novartis)	

This information has been summarised to act as a guide for those administering IV medication. The monograph should be used in conjunction with the drug data sheet and BNF for information on dose, adverse effects, cautions and contra-indications. Further information is available from Pharmacy on 22146 or 22542