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XOLAIR THERAPY PATIENT CONSENT

am acknowledging that I will begin my Xolair treatment.

(Print name legibly)	
The following points regarding Xolair were reviewed and discussion. The nature and purpose of Xolair treatment program b. The risks of the treatment, including the possibility of an alle program may not accomplish the desired objectives c. The possible outcome of the treatment d. The available alternative medical treatment e. The prognosis if the program is not followed f. The need for regular therapy and follow up (including the new my medication use, symptoms and need for unscheduled care g. Risk of anaphylaxis and epinephrine use, with proper demonth. Office policies regarding Xolair (i.e. calling ahead for mixing a administration if experiencing increase in asthma symptoms (if	ergic reaction as well as the risk that the treatment ed to evaluate my asthma by keeping records of (if applicable)) estration of epinephrine auto injector and scheduled office visit required prior to
I have had sufficient opportunity to discuss my condition with ranswered to my satisfaction. I have read and understand the X	my allergist and all of my questions have been olair treatment information form.
I believe that I have the adequate knowledge upon which to be	ase an informed consent to this program.
I consent to other diagnostic and therapeutic procedures and t might be necessary due to unexpected conditions (such as trea	the monitoring program that the physician decides atment of an allergic reaction).
I have read and fully understand this form.	
PATIENT	DATE
PARENT or LEGAL GUARDIAN	DATE
WITNESS	DATE
Patients on Xolair for asthma should have an office visit with their prescribing allergist minimum, or if there has been an increase in asthma symptoms or to assess response	t every 6 months and pulmonary function tests every 1-2 years at to treatment.