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XOLAIR THERAPY PATIENT CONSENT

l,	am acknowledging that I will begin my Xolair treatment.
The following points regarding Xolair we	ere reviewed and discussed in great detail:
 a. The nature and purpose of X 	olair treatment program
b. The risks of the treatment, inc	cluding the possibility of an allergic reaction as well as
the risk that the treatment progra	am 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 need to evaluate my asthma by keeping records of my medication use, symptoms and need for unscheduled care
- g. Risk of anaphylaxis and epinephrine use, with a proper demonstration of an epinephrine auto-injector
- h. Office policies regarding Xolair (i.e. calling ahead for mixing and scheduled office visit required prior to administration if experiencing an increase in asthma symptoms)

I have had sufficient opportunity to discuss my condition with my allergist and all of my questions have been answered to my satisfaction. I have read and understood the Xolair treatment information form. I believe that I have adequate knowledge upon which to base an informed consent to this program.

I consent to other diagnostic and therapeutic procedures and the monitoring program that the physician decides might be necessary due to unexpected conditions (such as treatment of an allergic reaction). I have read and fully understand this form.

PATIENT	_ DATE
PARENT or LEGAL GUARDIAN	DATE
WITNESS	_DATE

Patients on Xolair should have an office visit with prescribing allergist Dr. Patel every 6 months and pulmonary function tests every 1-2 years at the minimum, or if there has been an increase in asthma symptoms or to assess response to treatment.