

Genetic Screening

Babies with genetic diseases can be born to healthy parents who may be carriers for these genetic conditions which may be discovered through blood tests.

Plea	ase indicate whether or not you are in	nterested in testing for the following	g:	
	Spinal Muscular Atrophy (SMA)	s)	YES	\square NO
	SMA is the leading genetic cause of motor neurons of the nervous systen is a carrier of SMA.			
	Fragile X Syndrome		YES	□ №
	Fragile X syndrome is the most common inherited cause of intellectual disability, with a prevalence between 1 in 4000 and 1 in 6000 in males and about half of that in females. Of those boys affected by fragile X syndrome, about 1 out of 3 will have autism or autistic-like behavior. Nearly 1 out of 225 women are carriers.			
	Cystic Fibrosis (CF)		YES	□ NO
	CF is a genetic disease that cause severe degenerative damage to the lungs and pancreas. Between 1 in 16 to 87 people are carriers of this disease.			
	 The Counsyl Test (with Fragile 	X)	YES	□ №
	A comprehensive test for <u>100-plus</u> genetic diseases <u>which includes CF & SMA</u> . For more information, go to http://www.counsyl.com for the complete list of diseases which are tested.			
	Ashkenazi Jewish Panel		YES	□ NO
	you are of Ashkenazi Jewish (Eastern European Jewish) heritage this genetic test will assess the risk of aving a child with any of the 16 disorders which are commonly found in this population:			
	*Bloom Syndrome, *Canavan Disease, *Cystic Fibrosis, *Dihydrolipoamide Dehydrogenase Deficiency, *Familial Hyperinsul *Familial Dysautonomia, *Fanconi Anemia Group C, *Gaucher Disease, *Glycogen Storage Disease Type 1a, *Maple Syrup Disease, *Mucolipidosis IV, *Nemaline Myopathy, *Niemann-Pick Disease Types A, *Tay-Sachs Disease (Enzyme & DNA), *Syndrome type IF & III			
	am NOT interested in any gene	etic screening.		
NIAI	1 ⊑·	SIGNATI IRE:	DATE:	