

OUR UNIVERSAL APPROACH TO EXPANDED CARRIER SCREENING USING WHOLE EXOME SEQUENCING



All our CGT tests are supported by in-house genetic counselling

		CGT Bank			CGT SYNC		
	Methodology	WHOLE EXOME SEQUENCING (WES)					
	Type of panel	Exclusive Panel for Gamete Donors ACOG	Expanded Panel	Premium Expanded Panel	When a donor/partner is a known carrier		
	Genes	M: 7 genes F: 72 genes (include 65 X-linked)	M: 470 genes F: 535 genes (include 65 X-linked)	M: 1,979 genes F: 2,054 genes (include 65 X-linked)	Customised (F: option to include 65 X-linked)		
Z.S.	Disease causing variants	~3,800	~20,000	>50,000	_		
+	Numbers of diseases	Up to 75	Up to 570	More than 2,200	_		
00	Estimated carrier rate (%)*	~11%	~55%	~67%	-		
Z	Estimated mean of variants/individual**	1	1.7	2.7	_		
ii	Mean depth	150X	150X	150X	150X		
E	Complementary tests	M/F: CYP21A2, HBA1/2, SMN1 F only: DMD, FMR1, F8	M/F: CYP21A2, HBA1/2, SMN1 F only: DMD, FMR1, F8	M/F: CYP21A2, HBA1/2, SMN1 F only: DMD, FMR1, F8	***M/F: CYP21A2, HBA1/2, SMN1 ***F only: DMD, FMR1, F8		
Ū	Sample	Blood	Blood	Blood	Blood		
O	TAT	25 working days	25 working days	25 working days	25 working days		

^{*} In-house database of 30,000 tests

M: Male; F: Female

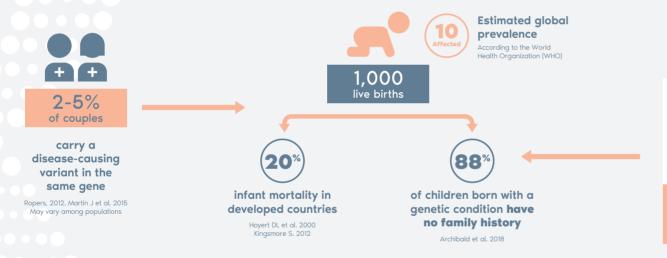
^{**} Estimated mean of positive individuals

^{***} If included in customised test



CGT is an advanced genetic test that determines the chance of having a child affected with a genetic condition.

CGT helps individuals and couples make informed reproductive decisions.



The American College of Obstetricians and Gynecologists (ACOG) makes the following recommendations:



Information about genetic carrier screening should be provided to every pregnant woman.

ACOG Committee Opinion. March

	SOME COMMON MONOGENIC CONDITIONS DETECTED WITH CGT ARE:	APPROXIMATE CARRIER FREQUENCY			
	Cystic fibrosis		in	25	
4	Spinal muscular atrophy	1	in	50	
	Autosomal recessive polycystic kidney disease	1	in	70	

