Medical Family Tree

Exploring Your Roots



MEDICAL FAMILY TREE: EXPLORING YOUR ROOTS

PURPOSE: You or someone in your family has been diagnosed with Fabry disease. We know that Fabry disease is an inherited disorder, which means that others in your family may also be at-risk for having the disorder. A medical family tree can be used to determine who in the family is at-risk for Fabry disease. The purpose of this worksheet is to organize your family history information so that you can be better prepared to have a medical family tree discussion with your Patient Education Liaison (PEL).

On the next page, please list your relatives and provide as much information as you can. You may find it helpful to check with other family members for more information. This worksheet is meant to be a starting point to your medical family tree conversation, so do not worry if you cannot complete the entire worksheet. You can use the questions below to help guide your discussion.

Questions to ask your relatives:

- Does anyone complain of constant stomach aches, having to go to the bathroom a lot or frequently having diarrhea? Does anyone have irritable bowel disease?
- Is there anyone who does not sweat or sweats only a little? Is there anyone who does not like to go outside when it is hot or cold out?
- Is there anyone who says they have a history of burning pain in their hands or feet?
- Is there anyone who has described their hands or feet as feeling that they are "on fire" or as having "pins and needles" in them?
- Is anyone on dialysis?
- Does anyone see a doctor for their kidneys (nephrologist)?
- Has anyone had a kidney transplant?
- Does anyone see a doctor for their heart (cardiologist)?
- Has anyone had a stroke?
- Is there anyone who passed away at a young age (prematurely)?
- Has anyone ever been told they have whorls/swirls in their eyes, or a distinctive pattern on the cornea, called corneal whorling or corneal verticillata?
- Does anyone have difficulty hearing or complain of ringing in their ears?
- Does anyone have unexplained pain or been told they had "growing pains"?
- Has anyone been diagnosed with Rheumatoid Arthritis, Multiple Sclerosis or Fibromyalgia?
- Does anyone have red spots or dark dots on their torso or bathing suit area?

Your Parents (check all that apply)

Mother (Name):									
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY	DISE	ASE		
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls		
	Stomach/GI issues		Little/No sweat		Premature death		Pain		
Fathe	r (Name):								
	DIAGNOSED WITH FABR	Y DI	SEASE		TESTED FOR FABRY DISEASE				
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls		
	Stomach/GI issues		Little/No sweat		Premature death		Pain		
-	If you have additional details about the boxes you checked for any of your relatives, please note them here:								

Your Children (check all that apply)

Child 1 (Name):								
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY I	DISE	ASE	
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Child	2 (Name):							
	☐ DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Child	3 (Name):							
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Child	4 (Name):							
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Child	5 (Name):							
	☐ DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY I	DISE	ASE	
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	

Your Siblings (check all that apply)

Sibling	g 1 (Name):						
	DIAGNOSED WITH FABRY DISEASE			TESTED FOR FABRY	D FOR FABRY DISEASE		
	Heart issues		Kidney/Dialysis	Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat	Premature death		Pain	
Sibling	g 2 (Name):						
	DIAGNOSED WITH FABRY DISEASE			TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis	Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat	Premature death		Pain	
Sibling	g 3 (Name):						
	DIAGNOSED WITH FABRY DISEASE			TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis	Stroke □ Eye wh		Eye whorls	
	Stomach/GI issues		Little/No sweat	Premature death		Pain	
Sibling	g 4 (Name):						
	DIAGNOSED WITH FABRY DISEASE			TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis	Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat	Premature death		Pain	
Sibling	g 5 (Name):						
	DIAGNOSED WITH FABRY DISEASE			TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis	Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat	Premature death		Pain	

For patient's personal use only.

Maternal Relatives (check all that apply)

Relative 1 (Name/Relation): ☐ DIAGNOSED WITH FABRY DISEASE ☐ TESTED FOR FABRY DISEASE ☐ Kidney/Dialysis ☐ Heart issues □ Stroke □ Eye whorls ☐ Stomach/GI issues □ Little/No sweat □ Premature death □ Pain Relative 2 (Name/Relation): ☐ DIAGNOSED WITH FABRY DISEASE ☐ TESTED FOR FABRY DISEASE ☐ Heart issues ☐ Kidney/Dialysis □ Eye whorls □ Stroke ☐ Stomach/Gl issues □ Little/No sweat □ Premature death □ Pain Relative 3 (Name/Relation): ☐ DIAGNOSED WITH FABRY DISEASE ☐ TESTED FOR FABRY DISEASE ☐ Kidney/Dialysis ☐ Heart issues □ Stroke □ Eye whorls ☐ Stomach/GI issues □ Little/No sweat □ Premature death □ Pain Relative 4 (Name/Relation): ☐ DIAGNOSED WITH FABRY DISEASE ☐ TESTED FOR FABRY DISEASE ☐ Kidney/Dialysis ☐ Heart issues □ Eye whorls □ Stroke ☐ Stomach/GI issues □ Little/No sweat □ Premature death □ Pain Relative 5 (Name/Relation): ☐ DIAGNOSED WITH FABRY DISEASE ☐ TESTED FOR FABRY DISEASE ☐ Heart issues ☐ Kidney/Dialysis □ Stroke □ Eye whorls ☐ Stomach/GI issues □ Little/No sweat □ Premature death □ Pain

Paternal Relatives (check all that apply)

Relative 1 (Name/Relation):								
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY	DISE	ASE	
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Relati	ve 2 (Name/Relation):							
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Relative 3 (Name/Relation):								
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Relative 4 (Name/Relation):								
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY DISEASE			
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	
Relati	ve 5 (Name/Relation):							
	DIAGNOSED WITH FABRY DISEASE				TESTED FOR FABRY	DISE	ASE	
	Heart issues		Kidney/Dialysis		Stroke		Eye whorls	
	Stomach/GI issues		Little/No sweat		Premature death		Pain	

For patient's personal use only.

SANOFI GENZYME OFFERS SUPPORT SERVICES FOR YOU AND YOUR FAMILY

For more than 30 years, Sanofi Genzyme has been committed to helping meet the needs of people who are living with rare disorders like Fabry disease. Our commitment extends to those who have Fabry disease and their families, as well as healthcare professionals. We offer a wide spectrum of services, all personalized and tailored to you through your Case Manager and Patient Education Liaison.

Patient Education Liaisons (PELs) can help:

- Educate and support you on Fabry disease management and inheritance
- Organize, support, and coordinate patient and/or family educational meetings
- Construct a medical family tree
- Provide educational support about Fabry disease to community groups, employers and/or educators
- Provide support to certain healthcare providers/offices

GETTING STARTED IS EASY

Contact a CareConnectPSS Case Manager: 1-800-745-4447 (option 3) Monday–Friday 8am–6pm EST

Connect with us online:

careconnectpss.com



50 Binney Street Cambridge, MA 02142