For Patients:

Do you want to understand your genetic risk and learn about genetic testing?

Do you have a family history or past medical history of a genetic disorder?

Do you find yourself lost in a diagnostic odyssey?

Have you had...

- a delayed diagnosis?
- a series of expensive tests with no definite diagnosis?
- limited access to technologies and expertise?
- a lack of information and difficulty understanding it?

Igenomix®
WITH SCIENCE ON YOUR SIDE

GPDx

Genomic Precision Diagnostic

by Igenomix®

The answers you need in a drop of blood, saliva or a buccal swab

Your genetics matter, learn about them

www.igenomix.co.uk
info.uk@igenomix.com
+44 (0)20 8068 8176

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WITH SCIENCE ON YOUR SIDE

V. 2021

Accurate diagnosis of genetic conditions must be a global health priority

- Igenomix provides expertise, support and the latest technologies to give you an accurate and personalised diagnosis.
- Igenomix's genetic specialists support you to select the best test according to clinical presentation and family history.

What are the benefits of genetic testing?

- For several disorders, this is the **only solution to make an accurate diagnosis** and help avoid additional unnecessary clinical investigations.

- **Decreases referrals between different specialties,** thus allowing high quality patient care.

- Reduces the uncertainty in a differential diagnosis and provides a directed focus for the clinician **to provide appropriate treatment.**

Our Services

We offer genetic counselling with all of our services

Preconception

- Identify if the parents are carrying a genetic mutation before pregnancy.
- Decrease the risk of genetic diseases being passed on.

Prenatal

- Identify high risk pregnancies.
- Uses advanced genetic technologies to analyse a fetal sample.

Neonatal

- Identify genetic variants that could cause disease in newborns.
- Prevent complications and increase life expectancy by initiating early treatment.

Childhood/Adulthood

- Identify or rule out a variety of genetic conditions.
- Early and accurate diagnosis to ensure high quality patient care.
- Identify risk of cancer, heart disease and other genetic conditions.

What are Rare Diseases?

Rare diseases are a diverse group of conditions; very few people are affected by them in comparison to more common conditions like diabetes or heart disease.

Although rare diseases may be individually rare, they are collectively common, with 1 in 17 people being affected by a rare disease at some point in their lives.*

~80%

Of rare diseases have a genetic origin**

~75%

Of rare diseases affect children***

* Extrapolated from www.eurordis.org/IMG/pdf/princeps_document-EN.pdf
** www.eurordis.org/IMG/pdf/princeps_document-EN.pdf
*** https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/867940/dhsc-2020-update-to-the-rare-diseases-implementation-plan-for-england.pdf