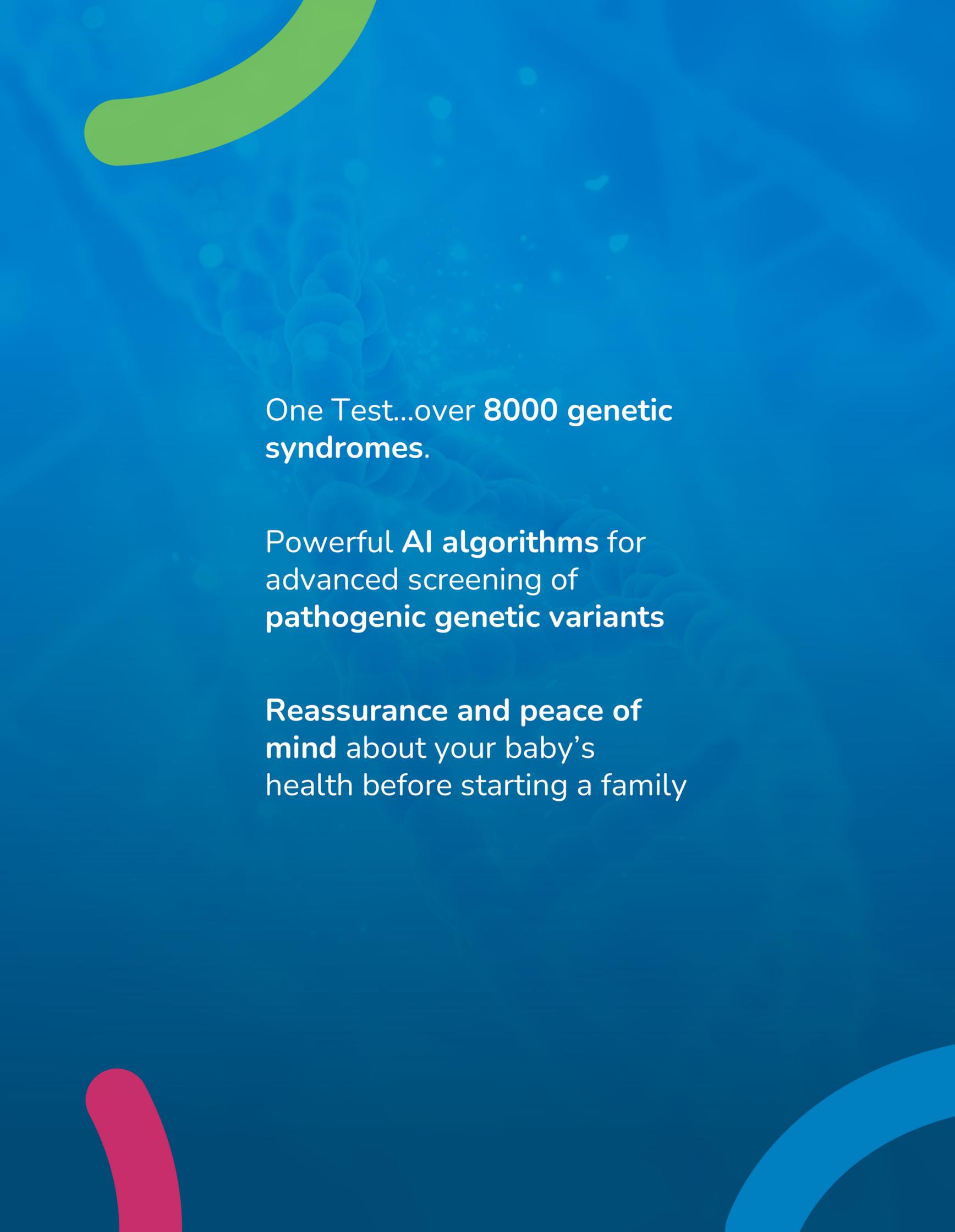




# SureMart

Empowering Families for  
the best start in life



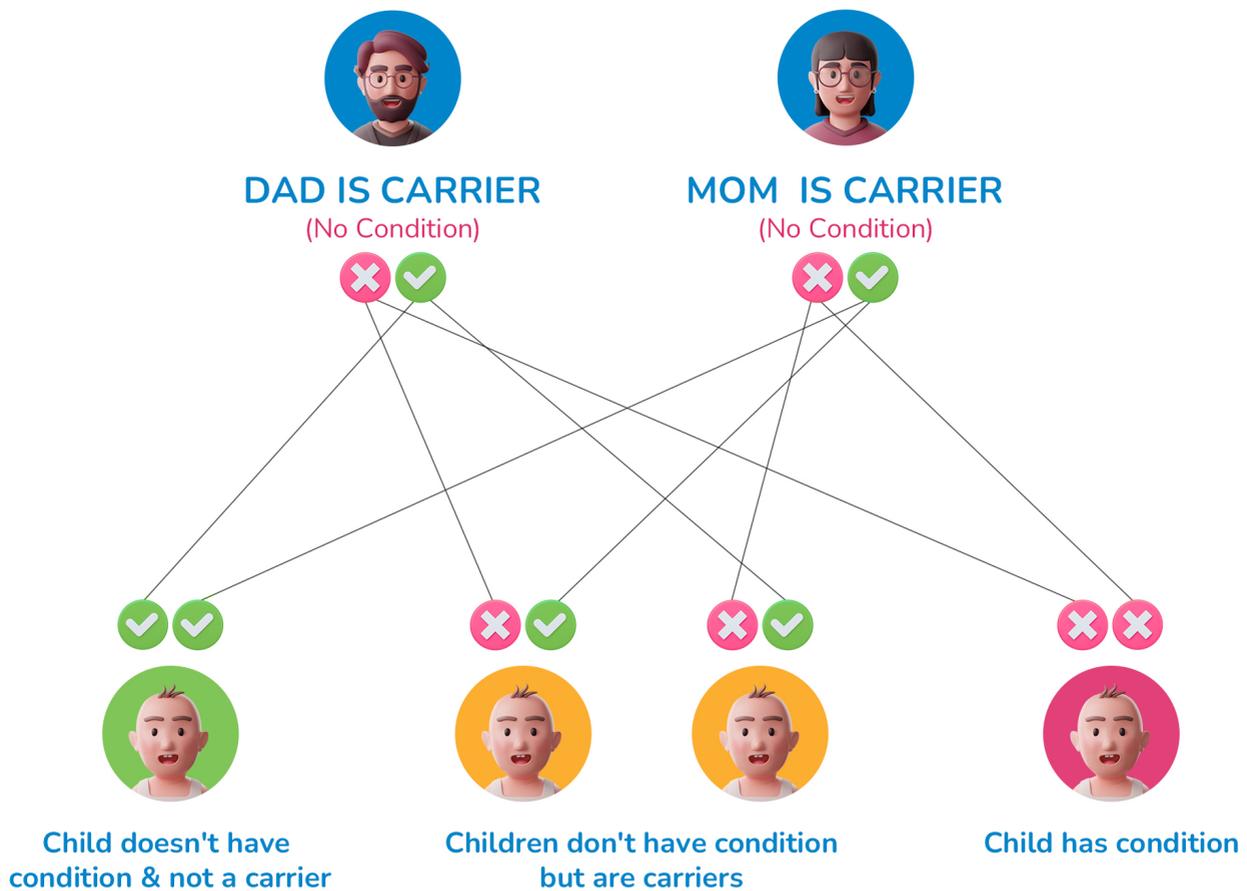
One Test...over **8000 genetic syndromes.**

Powerful **AI algorithms** for advanced screening of **pathogenic genetic variants**

**Reassurance and peace of mind** about your baby's health before starting a family

# What are the risks of not knowing

Autosomal Recessive Inheritance Pattern



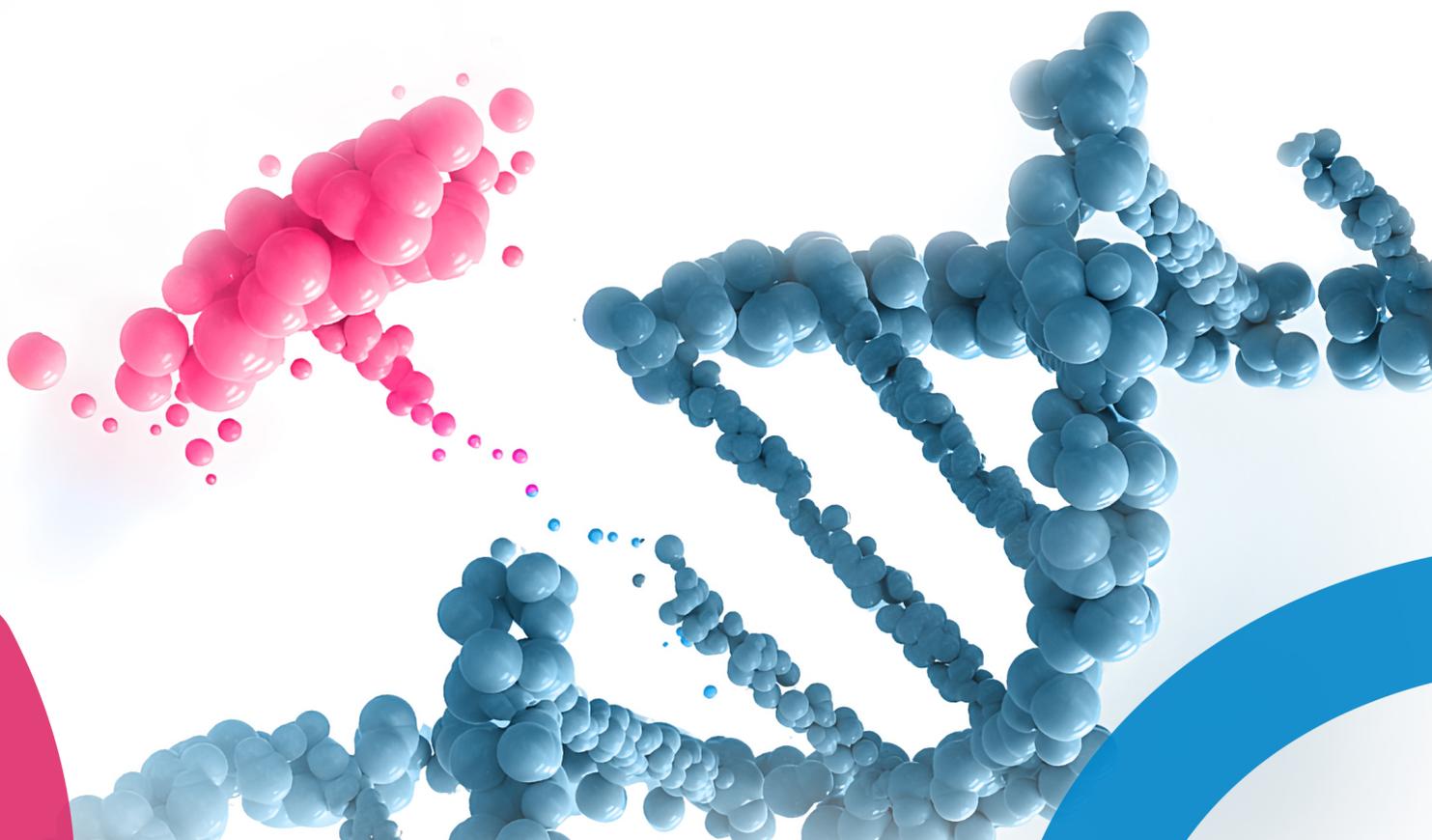
75% Children don't have the condition

25% have the condition



**Carriers of a genetic disorder** do not have any symptoms of this disorder. They do not know they are carriers. When two carriers have a child and both pass on the same genetic mutation, the child will be affected for this genetic condition. There is a **1 in 4 risk** of having an affected child when both parents are carriers of the same **genetic mutation**.

**According to data by WHO, 10 in 1000 babies are born with genetic diseases.** Some of these babies will die in the neonatal period, others will have life-limiting disabilities for the rest of their life





## Who is this test for?

Suremart is for any couple that is planning to start a family together. If the couple already has a sick child, we can recommend a course of action for the next pregnancy.

## What answers does this test give?

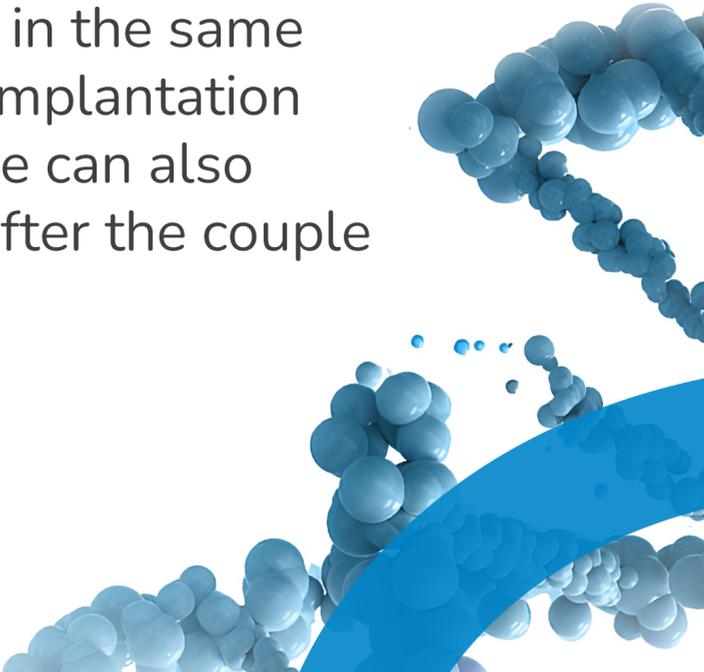
SureMart reveals and quantifies the risk of having a child affected with

1. Single-gene and complex diseases,
2. Rare diseases
3. Population-specific diseases
4. Suspected but unknown genetic solutions

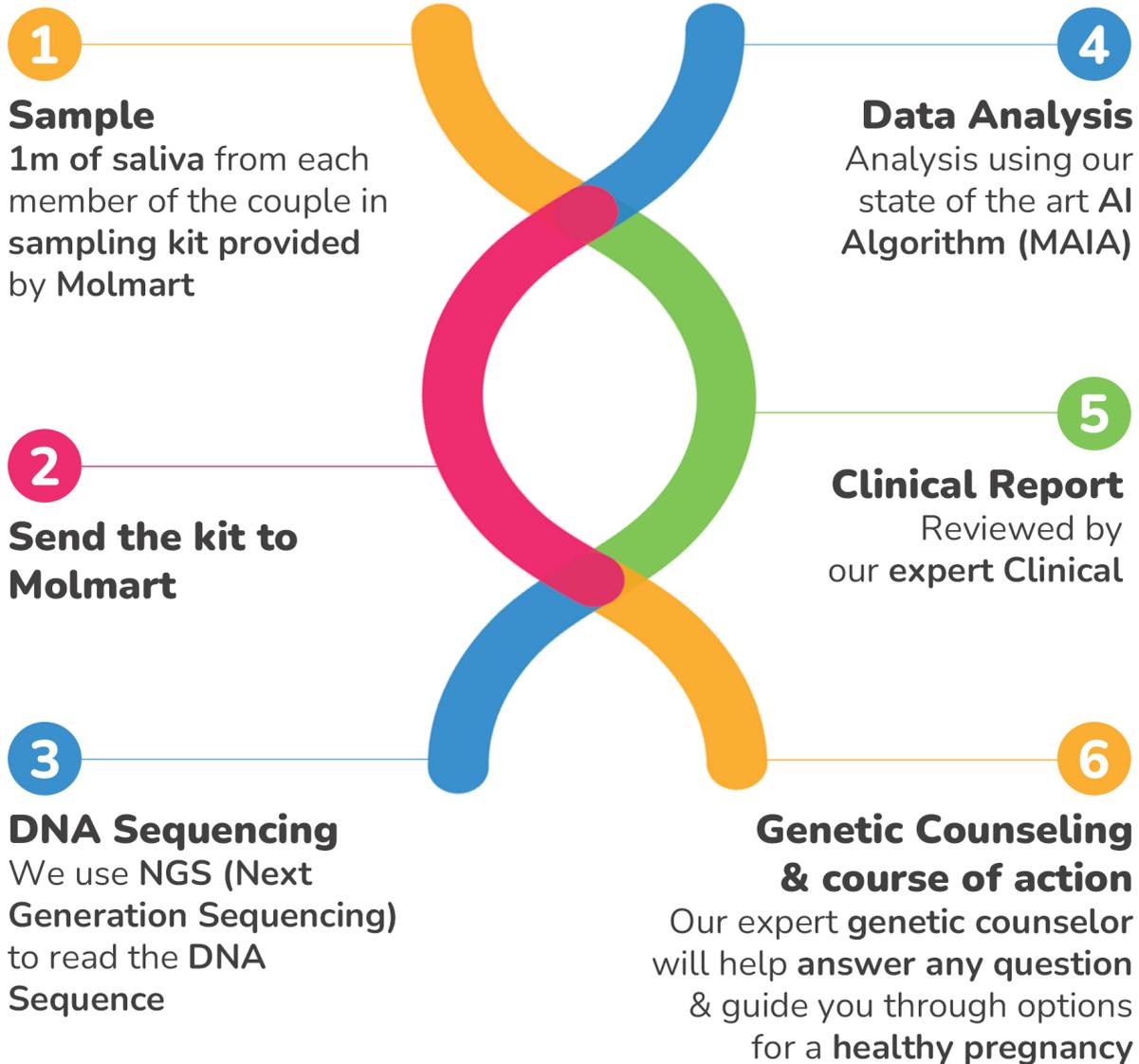
## How can the results be used?

If the couple are not carriers, no further action is necessary.

In the case where the couple are carriers of the same genetic mutation(s) in the same gene, we can assist with Preimplantation **Genetic Diagnosis (PGD)**. We can also perform Prenatal Diagnosis after the couple has conceived naturally.



# The SureMart journey



**SureMart** is the most cost-effective solution for **comprehensive genetic screening** - it substitute of single-gene tests with one test



# SureMart

Contact information

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