MITOCHONDRIAL REPLACEMENT THERAPY (MRT) - SCIENCE & TECHNOLOGY

NEWS: Eight healthy babies were born in Britain with the help of an experimental technique called Mitochondrial Replacement Therapy (MRT).

WHAT'S IN THE NEWS?

Background and Global Significance

- **High-Risk Mothers**: The mothers who underwent the treatment carried **mutations in their mitochondrial DNA**, putting their children at high risk of **life-threatening mitochondrial disorders**.
- Successful Births: Eight healthy babies (four boys and four girls) were born to seven women using mitochondrial donation treatment, with no signs of the mitochondrial diseases they were at risk of inheriting.
- UK Leadership: The United Kingdom became the first country in the world to formally approve the use of mitochondrial donation therapy (MDT) in 2015.
- India's Legal Status: As of now, India does not permit the use of mitochondrial donation or replacement therapies.

Understanding Mitochondrial Genes

- What Are Mitochondria?
 - Mitochondria are **membrane-bound organelles** found in almost every human cell, often referred to as the "**powerhouses of the cell**" due to their role in **energy production** (via ATP).
- Mitochondrial DNA: Unlike nuclear DNA, mitochondria have their own circular DNA, with 37 genes essential for normal mitochondrial function.
- Maternal Inheritance: All mitochondrial DNA is inherited from the mother, which means any mutation in her mitochondria is passed on to all her children.

What is Mitochondria?

- About:
 - Mitochondria are membrane-bound organelles found in the cells of most eukaryotic organisms.
 - They are often referred to as the "powerhouses" of cells because they generate the majority of the cell's energy in the form of adenosine triphosphate (ATP).

Functions:

- Mitochondria carry out cellular respiration, a process that converts nutrients into ATP.
- Mitochondria convert energy from carbohydrates, fats, and proteins into a usable form for the cell.
- They metabolize glucose to produce ATP, which powers various cellular processes.

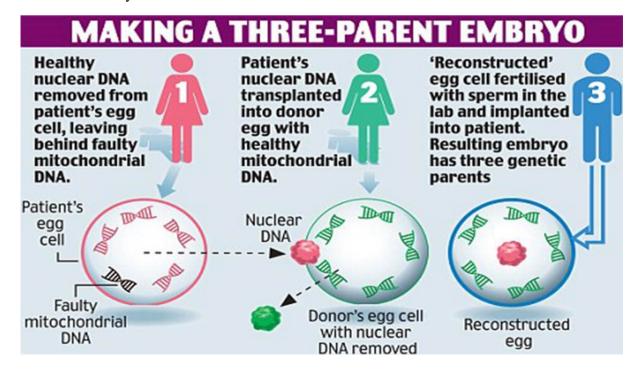
• Mitochondria participate in **cell signaling pathways**, influencing processes like **cell growth**, **differentiation**, and apoptosis.

• Inheritance:

- Mitochondria have their own DNA, known as mitochondrial DNA (mtDNA), which encodes a small number of essential proteins.
- In most animals, mtDNA is inherited solely from the mother.
- Mutations in mtDNA can lead to mitochondrial disorders and various health conditions.

• Mitochondrial Diseases:

- Certain mutations in mitochondria can lead to mitochondrial diseases, affecting
 energy production and impacting various organs, including the brain, nerves,
 muscles, kidneys, heart, and liver.
- These diseases can result in severe symptoms, such as organ failure, muscle wastage, and even brain damage. Unfortunately, there is no cure for mitochondrial diseases, but they can be managed to some extent.
- Few examples of mitochondrial diseases are **Leigh Syndrome**, **Kearns-Sayre syndrome** (**KSS**), Mitochondrial Myopathy and Mitochondrial DNA Depletion Syndrome.



Impact of Mitochondrial Diseases

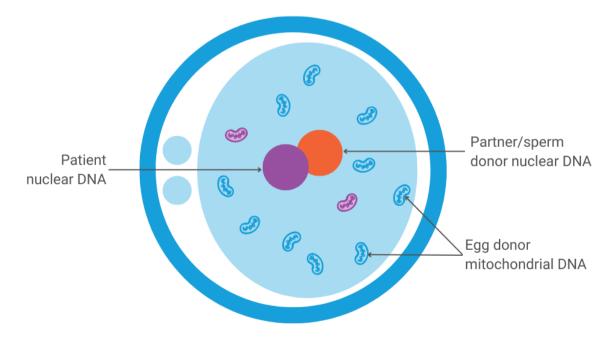
- Mode of Inheritance: Since mitochondria come only from the egg (not sperm), mutations in a mother's mitochondria are transmitted to all her biological children.
- Health Consequences:

- Mitochondrial diseases affect organs that require **high energy**, like the **brain**, **heart**, and muscles.
- Symptoms often appear in early childhood.
- Children may face **developmental delays**, **muscle weakness**, may require **wheelchairs**, and in severe cases, may **die young**.
- Approx. 1 in 5,000 births is affected by some form of mitochondrial disorder.

Mitochondrial Donation Treatment (MDT)

- Also Called:
 - Mitochondrial Replacement Therapy (MRT)
 - Three-Parent Baby Technique
 - Three-Person In Vitro Fertilisation (IVF)
- **Objective**: Prevent the **transmission of defective mitochondria** from mother to child, thereby eliminating mitochondrial diseases **before birth**.

Embryo created by mitochondrial donation



Procedure: How MDT Works

- Step-by-Step:
 - 1. **Eggs are collected** from both the affected mother and a healthy female donor.

- 2. The mother's egg is **fertilized with the father's sperm**, forming an embryo with **mutated mitochondria**.
- 3. The nuclear genetic material from the fertilized embryo is transferred into a donor egg that has had its own nucleus removed but contains healthy mitochondria.
- 4. The new embryo now contains:
 - Nuclear DNA from the biological mother and father
 - Healthy mitochondrial DNA from the donor woman
- 5. This embryo is then **implanted into the womb**, and if successful, results in a pregnancy.
- Genetic Outcome: The resulting baby carries 99.8% of DNA from the biological parents and 0.2% from the mitochondrial donor.

Ethical and Scientific Importance

- Ethical Discussions:
 - Involves **germline modification**, which is inheritable by future generations.
 - Raises questions on parental identity, regulation, and long-term effects.
- Scientific Breakthrough:
 - Represents a new frontier in genetic medicine.
 - Converts an **inherited disorder prevention technique** into a **potential reproductive option** for at-risk families.

Global Outlook and Future Prospects

- Ongoing Global Monitoring: Scientists across the world are observing the UK results for long-term safety, efficacy, and ethical acceptability.
- Potential for Wider Use:
 - With strong regulation and safety protocols, MDT may become a standard medical option for families carrying mitochondrial mutations.
 - May also influence gene therapy, IVF practices, and genomic medicine policy globally.

Source: https://www.livemint.com/news/trends/medical-breakthrough-in-uk-babies-born-after-ivf-using-dna-from-three-people-11752750527061.html