

Assessment of Uterine Disorders in Tunisian Women with Recurrent Pregnancy Loss: A Single Center Experience

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COMMENTARY

Repetitive pregnancy misfortune (RPL) is characterized by ESHRE as at least three back to back pregnancy misfortunes and influences 1-3% of ladies endeavoring to have a youngster. An assortment of potential etiologies have been portrayed. Frequently, anatomic issues, both inborn and gained; are a significant reason for clinically perceived unsuccessful labor (15%). Despite the fact that the job of uterine abnormalities in RPL is begging to be proven wrong, appraisal of uterine life structures is broadly suggested. Additionally, clinical administration of pregnancy-misfortune patients with uterine issues is questionable, and there is no decisive proof that careful treatment lessens the danger of pregnancy misfortune [1].

The point of this examination was to evaluate the pervasiveness of uterine issues in Tunisian ladies with history of repetitive pregnancy misfortune (RPL) and to outline methodologies for clinical administration. The examination included 158 couples with 2 and more pregnancy misfortune. So as to investigate issues of the uterine hole, hysterosalpingography was performed. Of an aggregate of 158 couples, 32 ladies had uterine variations from the norm with a general occurrence of 20.3 % [2].

We have discovered more procured than inherent uterine issues, essentially the intrauterine synechia. Anatomic turmoil is visit in Tunisian Women with RPL with a general recurrence of 20.3%. Appraisal of the uterine life structures is suggested in ladies with 2 pregnancy misfortunes, to explain the reasons for RPL. While hysterosalpingography is valuable as screening test, hysteroscopy remains the backbone of analysis and treatment [3].

Intermittent pregnancy misfortune is traditionally characterized in light of the fact that the event of at least three sequential pregnancy misfortune; in any case, the American Society of Reproductive Medicine (ASRM) has as of late reclassified repetitive pregnancy misfortune as at least two pregnancy misfortunes. A pregnancy misfortune is characterized as a clinically-perceived pregnancy automatically finishing before 20 weeks. A clinically-perceived pregnancy implies the pregnancy has been envisioned on a ultrasound or that pregnancy tissue was distinguished after a pregnancy misfortune [4].

Most pregnancy misfortunes result from chromosomal, or hereditary, irregularities, and are arbitrary occasions. The abnormality may come from the egg, the sperm, or the first embryo. Approximately 12-15% of all clinically recognized pregnancies end in miscarriage; however, it's estimated that a minimum of 30-60% of all conceptions will end within the primary 12 weeks of gestation. Up to 50% of the time, the lady doesn't even realize that she was ever pregnant. the danger of miscarriage increases with the amount of previous pregnancy losses, but is usually but 50%.

Advancing maternal age is related to an increased risk of miscarriage, which is assumed to flow from to poor egg quality resulting in chromosomal (genetic) abnormalities. Sometimes, the mother or father themselves may have a small irregularity in their genes, but the offspring might be more severely affected and thus end in miscarriage. Sometimes, there might be an abnormality within the uterus (the womb) that results in miscarriage [5].

The miscarriage could also be thanks to poor blood supply to the pregnancy or inflammation. Some women could also be born with an irregularly shaped uterus, and a few women may develop abnormalities with their uterus over time. A woman's system can also play a task in recurrent pregnancy loss. Hormone abnormalities can also impact pregnancy loss, including thyroid disease and diabetes. Abnormalities during a mother's blood coagulation can also affect pregnancy loss. Many early miscarriages (the ones that happen within the first 3 months of pregnancy) are thanks to genetic abnormalities within the embryo or fetus.

Normally, there are 46 chromosomes that contain the genes for normal development. Many early miscarriages happen because the fetus has an additional chromosome or one is missing. for instance, babies with mongolism have 47 chromosomes. Chromosome abnormalities occur for no known reason in up to 60% of first-trimester miscarriages. Genetic abnormalities typically don't allow development into a healthy baby. As women age, the miscarriage risk thanks to these genetic abnormalities increases – from 10%-15% in women younger than 35 years old to quite 50% in women over 40 years old.

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REFERENCES

1. Su MT, Lin SH, Chen YC. Genetic association studies of angiogenesis- and vasoconstriction-related genes in women with recurrent pregnancy loss: a systematic review and meta-analysis. *Human Reproduction Updat.* 2011;17(6):803-812
2. Bradley LA, Palomaki GE, Bienstock J, Scott JA. Can Factor V Leiden and prothrombin G20210A testing in women with recurrent pregnancy loss result in improved pregnancy outcomes?: Results from a targeted evidence-based review. *Genetics in Medicine.* 2012.
3. Sokolov DI, Mikhailova VA, Agnayeva AO, Bazhenov DO, Khokhlova EV, Bespalova ON. NK and trophoblast cells interaction: cytotoxic activity on recurrent pregnancy loss. *Gynecological Endocrinology.* 2019;35:5-10.
4. Mahjoub T, Mtiraoui N, Tamim H, Hizem S, Finan RR, Nsiri B, Almawi WY. Association between adverse pregnancy outcomes and maternal factor V G1691A (Leiden) and prothrombin G20210A genotypes in women with a history of recurrent idiopathic miscarriages. *American J hematology.* 2005;80(1):12-19.
5. Ghamdi AA, Makhshen SF. Etiology of Recurrent Pregnancy Loss in Saudi Females. *Saudi J Med Med Sci.* 2016;4(3):187-191.