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MATRIC NUMBER: 19/MHS02/126

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SPERMATOGENESIS

Spermatogenesis is the process of formation of spermatocytes from spermatogonia. Spermatogenesis is initiated at puberty, continues throughout the remainder of a man's life, and takes place in the walls of the seminiferous tubules.

The walls of the tubules are composed of two compartments separated by tight junctions between the Sertoli cells:

- The basal layer, which consists of the Leydig cells and the spermatogonia
- The adluminal layer, which is made up of Sertoli cells and spermatocytes

The initial step in the process is transformation of type A spermatogonia, which are epithelioid-like cells, to type B spermatogonia, a process involving four divisions. The type B cells embed in the Sertoli cells. In association with the Sertoli cells, the type B cells are transformed to primary spermatocytes and then, in a step involving the first meiotic division, to secondary spermatocytes. The secondary spermatocytes undergo a second meiotic division, yielding spermatids, each of which has 23 unpaired chromosomes. The steps described are stimulated by testosterone and follicle stimulating hormone (FSH).

MALE INFERTILITY

Male infertility refers to a male's inability to cause pregnancy in a fertile female. Important causes of male infertility include the following:

- Androgen dysfunction with normal sperm cell production, caused by hypothalamic-pituitary defects, Leydig cell defects, or androgen resistance
- Isolated dysfunction of sperm cell production with normal androgen levels, resulting from infection or trauma, congenital deformation of passages, or formation of nonmotile or otherwise abnormal sperm
- Combined androgen and sperm cell production defects resulting from

- (1) developmental defects, such as Klinefelter's syndrome or abnormal testicular descent,
- (2) acquired testicular defects, such as infections, autoimmune reactions, or systemic diseases such as chronic liver and kidney diseases

Unknown causes: In 50 percent of infertile males, no cause can be identified