Comparative analysis of gene expression in granulosa cells: Implications for ovarian function and reproductive health.

Sofia Reschini*

Department of Family Medicine, Clalit Health Services, Israel

Introduction

Granulosa cells, the somatic cells surrounding developing oocytes in the ovaries, play a critical role in regulating ovarian function and female reproductive health [1]. These cells are responsible for secreting estrogen and progesterone, facilitating follicular development, and supporting oocyte maturation [2]. Recent research has focused on the gene expression profiles of granulosa cells to better understand their contribution to fertility and ovarian disorders, such as polycystic ovary syndrome (PCOS) and premature ovarian failure (POF) [3].

Studies comparing gene expression in granulosa cells from different ovarian conditions have revealed significant variations that affect ovarian function [4]. In women with PCOS, granulosa cells show altered expression of genes related to steroidogenesis, follicular development, and cell proliferation. These changes are often linked to insulin resistance and hormonal imbalances [5].

Specifically, granulosa cells in PCOS exhibit an upregulation of genes involved in androgen production, such as CYP17A1, which plays a role in the synthesis of testosterone [6]. This contributes to the elevated androgen levels commonly observed in PCOS and disrupts normal ovarian function. Conversely, in women with premature ovarian failure, granulosa cell function is impaired, leading to reduced estrogen production and the early cessation of ovarian activity [7].

Comparative gene expression analysis has also been used to explore the effects of hormonal treatments, such as those used in assisted reproductive technologies (ART) [8]. Research shows that granulosa cells from women undergoing in vitro fertilization (IVF) exhibit altered gene expression compared to natural cycles, particularly in genes related to oxidative stress and apoptosis, which may affect oocyte quality and embryo development [9]. Furthermore, identifying specific gene markers that can predict granulosa cell function and oocyte quality has become a significant area of focus in reproductive medicine, with the aim of improving ART outcomes [10].

Conclusion

The comparative analysis of gene expression in granulosa cells provides valuable insights into the mechanisms underlying

ovarian function and reproductive health. By identifying the genetic signatures associated with various ovarian conditions and treatments, researchers aim to develop more targeted and effective therapeutic strategies for women experiencing infertility or hormonal imbalances.

References

- Hsueh AJ, Adashi EY, Jones PB, et al. Hormonal regulation of the differentiation of cultured ovarian granulosa cells. Endocr Rev. 1984;5(1):76-127.
- 2. Khristi V, Chakravarthi VP, Singh P, et al. ESR2 regulates granulosa cell genes essential for follicle maturation and ovulation. Mol Cell Endocrinol. 2018;474:214-26.
- Johnson AL, Woods DC. Dynamics of avian ovarian follicle development: cellular mechanisms of granulosa cell differentiation. Gen Comp Endocrinol. 2009;163(1-2):12-7.
- Liu J, Yang Y, Yang Y, et al. Disrupting effects of bifenthrin on ovulatory gene expression and prostaglandin synthesis in rat ovarian granulosa cells. Toxicology. 2011;282(1-2):47-55.
- Racine C, Genêt C, Bourgneuf C, et al. New anti-Müllerian hormone target genes involved in granulosa cell survival in women with polycystic ovary syndrome. J Clin Endocrinol Metab. 2021;106(3):1271-89.
- 6. Havelock JC, Rainey WE, Carr BR. Ovarian granulosa cell lines. Mol Cell Endocrinol. 2004;228(1-2):67-78.
- Höfner M, Eubler K, Herrmann C, et al. Reduced oxygen concentrations regulate the phenotype and function of human granulosa cells in vitro and cause a diminished steroidogenic but increased inflammatory cellular reaction. Mol Hum Reprod. 2024;30(1):gaad049.
- Zhou W, Liu J, Liao L, et al. Effect of bisphenol A on steroid hormone production in rat ovarian theca-interstitial and granulosa cells. Mol Cell Endocrinol. 2008;283(1-2):12-8.
- 9. Huang P, Zhou Y, Tang W, et al. Long-term treatment of Nicotinamide mononucleotide improved age-

Citation: Reschini S. Comparative analysis of gene expression in granulosa cells: Implications for ovarian function and reproductive health. Gynecol Reprod Endocrinol.2024;8(5):224

^{*}Correspondence to: Sofia Reschini, epartment of Family Medicine, Clalit Health Services, Israel. E-mail: reschini@chs.isr.co

Received: 21-Aug-2024, Manuscript No. AAGGS-24-154856; *Editor assigned*: 22-Aug-2024, Pre QC No. AAGGS-24-154856(PQ); *Reviewed*: 05-Sep-2024, QC No. AAGGS-24-154856; *Revised*: 10-Sep-2024, Manuscript No. AAGGS-24-154856(R); *Published*: 17-Sep-2024, DOI: 10.35841/aajnnr-8.5.224

related diminished ovary reserve through enhancing the mitophagy level of granulosa cells in mice. J Nutr Biochem. 2022;101:108911. Lerner A, Owens LA, Coates M, et al. Expression of genes controlling steroid metabolism and action in granulosalutein cells of women with polycystic ovaries. Mol Cell Endocrinol y. 2019;486:47-54.

Citation: Reschini S. Comparative analysis of gene expression in granulosa cells: Implications for ovarian function and reproductive health. Gynecol Reprod Endocrinol.2024;8(5):224