

Antidepressants and Male Infertility: Involvement of Spermatogonial Stem Cells Self-renewal Disruption

Ali Shalizar Jalali

Histology and Embryology Research
Laboratories, Department of Basic Sciences,
Faculty of Veterinary Medicine, Urmia
University, Urmia, Iran

Received: February 20, 2017; **Accepted:** February 22, 2017; **Published:** February 27, 2017

Nearly 350 million people are affected by depression world-wide and antidepressants are the most commonly used medications [1,2]. Recently, it has been reported that the percentage of Americans on antidepressant therapies had nearly doubled, increasing from 6.8% to 13% and selective serotonin reuptake inhibitors (SSRIs) are the most commonly used antidepressant [3]. In line with that, it was also found that SSRIs treatment can lead to sexual side effects in 55% of patients [4]. Moreover, experimental evidence revealed that SSRIs administration depletes testicular germinal epithelium resulting in defective spermatogenesis and reduced male fertility [5,6].

Spermatogonial stem cells (SSCs) as self-renewing adult stem cells in the testis are responsible for regulated continual spermatogenesis throughout the most of a male's lifespan and their biological activities can be influenced by chemicals and/or radiation induced testicular damages [7,8]. Supporting these facts, it has been shown that SSRIs are associated with SSCs self-renewal suppression in adult rats confirming the possible involvement of these compounds in male germline maintenance disruption [9].

Corresponding author: Ali Shalizar Jalali

✉ a.shalizar@urmia.ac.ir

Histology and Embryology Research
Laboratories, Department of Basic Sciences,
Faculty of Veterinary Medicine, Urmia
University, Urmia, Iran.

Tel: 00984431942593

Citation: Jalali AS. Antidepressants and Male Infertility: Involvement of Spermatogonial Stem Cells Self-renewal Disruption. *J. stem cell Bio. transplant.* 2017, 1:1.

Accordingly, since antidepressants may cause severe testiculopathy, profound considerations should be given to the drastic consequences of their increased consumption.

References

- 1 Pilia M, Bairwa M, Kumar N, Khanna P, Kurana H (2013) Elderly depression in India: An emerging public health challenge. *Australas Med J* 6: 107-111.
- 2 Davey CG, Chanen AM (2016) The unfulfilled promise of the antidepressant medications. *Med J Aust* 204:348-350.
- 3 Kantor ED, Rehm CD, Haas JS, Chan AT, Giovannucci EL (2015) Trends in Prescription Drug Use among Adults in the United States from 1999-2012. *JAMA* 314: 1818-1831.
- 4 Ferguson JM (2001) SSRI antidepressant medications: adverse effects and tolerability. *Prim Care Companion J Clin Psychiatry* 3: 22-27.
- 5 Bataineh HN, Daradka T (2007) Effects of long-term use of fluoxetine on fertility parameters in adult male rats. *Neuro Endocrinol Lett* 28: 321-325.
- 6 Hajizadeh Z, Soleimani Mehranjani M, Najafi G, Shariatzadeh SMA, Shalizar Jalali A (2016) Black grape seed extract modulates fluoxetine-induced oxidative stress and cytotoxicity in mouse testis. *Jundishapur J Nat Pharm Prod*.
- 7 Grewenig A, Schuler N, Rube CE (2011) Persistent DNA damage in spermatogonial stem cells after fractionated low-dose irradiation of testicular tissue. *Int J Radiat Oncol Biol Phys* 92: 1123-1131.
- 8 Singh SR, Burnicka-Turek O, Chauhan C, Hou SX (2011) Spermatogonial stem cells, infertility and testicular cancer. *J Cell Mol Med* 15: 468-483.
- 9 Akbari H, Hasanzadeh S, Shalizar Jalali A (2016) Fluoxetine suppresses spermatogonial stem cell self-renewal in adult rats. *Int J Fertil Steril*.