



Mini Review

Volume 3 Issue 4 - August 2017
DOI: 10.19080/JPCR.2017.03.555620

J of Pharmacol & Clin Res

Copyright © All rights are reserved by Ana Maria trindade Grégio Hardy

Pyshotropic Drugs: Side Effects on Mouth



Emanuela Carla dos Santos¹, Eliana Cristina Fosquiera², Aline Cristina Batista Rodrigues Johann³, Mariana Rinaldi², Patrícia Vida Cassi Bettega² and Ana Maria Trindade Gregio*⁴

¹Master student in Stomatology, Pontifical Catholic University of Parana, Brazil

²Doctorate student in Stomatology, Pontifical Catholic University of Parana, Brazil

³Department of Oral Patology, Pontifical Catholic University of Parana, Brazil

⁴Department of Pharmacology, Pontifical Catholic University of Parana, Brazil

Submission: August 02, 2017; **Published:** August 16, 2017

***Corresponding author:** Ana Maria trindade Grégio Hardy, Department of Pharmacology, Pontifical Catholic University of Parana, Brazil, Email: anamaria.gregio@gmail.com

Mini Review

Modern life, globalization, informatization and all technical and scientific resources brought to man a better quality and longer life expectancy, but with such progress emerged diseases hitherto little studied, afflicting a significant percentage of the population. Depression is commonly accompanied by anxiety disorders such as phobias, panic disorder, generalized anxiety disorder (GAD), obsessive compulsive disorder (OCD), and acute posttraumatic stress. These are also chronic psychiatric conditions, severe and recurrent, with an estimated prevalence of around 29% that result in functional impairment, associated with significant social costs, since the treatment thereof include psychological therapy and Central nervous system (CNS) depressant drugs, called psychotropic drugs. These drugs, such as antidepressants, have been widely reported since the 1980s to date WHO 2012 [1-5].

Among these diseases, affective disorders might be highlight. These are characterized by changes in mood (depression, anxiety or mania) and thought disorders. Depression constitutes the most common form of this event, ranging from mild to severe, which is sometimes accompanied by hallucinations and delusions (psychotic depression). Many studies work with the hypothesis that depression comes from deficiency of monoamines (norepinephrine and serotonin), and the most appropriate treatment is to increase the supply of these neurotransmitters in the central nervous system. Other statistics show that depression has increased due to the increased life expectancy. It is common in the elderly population. The treatment of psychiatric disorders and affection disorders is mainly antidepressants, antipsychotics and anxiolytics [5-8].

Several authors have also reported that patients with psychiatric disorders such as depression complain of dry mouth. Depressive changes where the dry mouth is present are 20% more frequent in women than in men. The most common age is between 30 and 59 years. Drug mechanism of action on the central nervous system for treating depression consists on the reuptake of the neurotransmitter, the inhibition of neurotransmitter degradation, and increase in the bonding time with the central receptors and/or increase neurotransmitter synthesis. However, pharmacologically, these drugs promote adverse reactions and side effects to the patient that uses this medication, especially for prolonged use [9-12].

Such drugs have been investigated by the scientific community, due to the increase and continuous prescription, and obviously the reporting of undesirable effects by the patients. The study of the relation structure-activity (REA) led to the discovery that certain receptors for such monoamines - Beta1 and alpha2-adrenergic systematically undergo a down-regulation following chronic treatment with antidepressants. This has been demonstrated in experimental animals by reducing the number of binding sites as well as reduction in functional response to agonists. These findings do not explain the theory of deficiency of monoamines, but corroborate the unwanted effects of these drugs, so adding the functional loss of the theory of activity of the adrenergic and cholinergic receptors present in the salivary glands, promoting hyposalivation (dry mouth), urinary retention, constipation, bladder atony, bowel atony, xerophthalmia (dry eyes) and even insomnia and agitation at the beginning of treatment.

The appearance of patients in medical and dental office with dry mouth and dry eyes complaint has been constant, because of these medicines. It is noteworthy that hypo salivation may experience some symptoms like burning mouth, loss of taste, angular cheilitis, difficulty in speaking and swallowing, and when worsens led to the increase in the number of caries, oral candidiasis, halitosis and periodontal disease. The therapeutic approach of the patient with xerostomia (dry mouth) varies according to individual characteristics. The treatment is to relieve symptoms, prevent or correct any sequelae of salivary dysfunction and also stimulating the production and release of salivary secretion by glands. But it is an empirical therapy, because literature does not explain much about this. The use of artificial saliva, mouthwashes, increased fluid intake and citrus drinks can collaborate, but these are palliative treatment.

Drugs that stimulate the release of salivary secretion are the best therapy, that is, the drugs that activate the expulsion of the stored saliva in the salivary glands. The inconvenient of this therapy is the appearance of systemic side effects such as diarrhea, sweating, vomiting, and stomach discomfort. Studies have been conducted at the Pontifical Catholic University of Parana in Pharmacology and Pathology Laboratory - Odontology, which has investigate a medicinal plant, an alkaloid with secretagogue action (stimulating secretions) in the innovation formulation, for treatment of hyposalivation and causes no side effects, because its use is local, not promoting systemic action.

This formulation is innovative because there is no similar in the pharmaceutical market, besides this topical application, used in the oral cavity. In the USA and some countries of Europe, patients with hyposalivation and/or dry mouth are treated with Salagen, but in tablet form and oral solution. Because of its systemic use there are a number of side effects, mainly secretagogue action, that increase body secretions. Finally, we highlight the importance of these studies in the Pathology and Pharmacology Laboratory of PUCPR and also further studies at the University of Toledo in the Department of Pharmacology (Pharmacy College) - Toledo - US to clarify the new formulation and the psychotropic drugs effect on mouth. Further studies should be published in the future reporting this action mechanism.

References

1. Gregio AMT, Durscki JRC, Lima AAS, Machado MAN, Ignacio SA, et al. (2006) Association of amitryptiline and diazepam on the histomorphometry of rat parotid glands *Pharmacology online* 2: 96-108.
2. Mattioli TM, Alanis LR, Sapelli Sda S, de Lima AA, de Noronha L, et al. (2016) Effects of Benzodiazepines on Acinar and Myoepithelial Cells. *Front Pharmacol* 24(7): 173.
3. Rinaldi M, Johann AC, Rocha F, Ignacio SA, Rosa EA, et al. (2015) Histomorphometric analysis of salivary gland in wistar rats treated chronically with two benzodiazepines. *Curr Pharm Biotechnol* 16(6): 573-578.
4. Mattioli TM, Silva Sd, Grégio AM, Machado MÂ, Lima AA, et al. (2011) The effects of antidepressants and pilocarpine on rat parotid glands: an immunohistochemical study. *Clinics (Sao Paulo)* 66(9): 1605-1610.
5. Da Silva S, De Azevedo LR, De Lima AA, Ignácio SA, Machado MA, et al. (2009) Effects of fluoxetine and venlafaxine and pilocarpine on rat parotid glands. *Med Chem* 5(5): 483-490.
6. Zaclikevis MV, D'Agulham AC, Bertassoni LE, Machado MA, de Lima AA, et al. (2009) Effects of benzodiazepine and pilocarpine on rat parotid glands: histomorphometric and sialometric study. *Med Chem* 5(1): 74-78.
7. De Almeida Pdel V, Grégio AM, Machado MA, De Lima AA, Azevedo LR (2008) Saliva composition and functions: a comprehensive review. *J Contemp Dent Pract* 9(3): 72-80.
8. De Almeida Pdel V, Grégio AM, Brancher JA, Ignácio SA, Machado MA, et al. (2007) Effects of antidepressants and benzodiazepines on stimulated salivary flow rate and biochemistry composition of the saliva. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 106(1): 58-65.
9. Veiga FF, Johan ACBR, Kagy VS, Muniz LTB, Alanis LAR, et al. (2016) Action of lithium carbonate on parotid acini. *Dent Oral Craniofac Res, USA*.
10. Patrícia Del Vigna de Ameida, Aline Cristina Batista Rodrigues Johann, Luciana Reis de Azevedo Alanis, Antônio Adilson Soares de Lima and Ana Maria Trindade Grégio (2011) Antidepressants: Side Effects in the Mouth. *Antidepressants: Side Effects in the Mouth -Oral Health- Book 1*: Edited by: Prof. Dr. Mandeep Singh Viridi. (Org.) 37693.
11. Ru-Band Lu (2012) Effects of Fluoxetine and Venlafaxine on parotids glands- Experimental Study-Ana Maria Trindade Grégio Effects of Antidepressants. ISBN 978-953-51-0663-0, hard cover, 194 pages, Publisher: InTech.
12. Gregio AMT, Veiga FF, Johan ACBR, Kagy VS, Muniz LTB, et al. (2016) Action of lithium carbonate on parotid acini. *Dent Oral Craniofac Res, USA*.



This work is licensed under Creative Commons Attribution 4.0 License
DOI: 10.19080/JPCR.2017.02.555620

Your next submission with Juniper Publishers will reach you the below assets

- Quality Editorial service
- Swift Peer Review
- Reprints availability
- E-prints Service
- Manuscript Podcast for convenient understanding
- Global attainment for your research
- Manuscript accessibility in different formats
(Pdf, E-pub, Full Text, Audio)
- Unceasing customer service

Track the below URL for one-step submission

<https://juniperpublishers.com/online-submission.php>