

SEROTONERGIC-LIKE PROFILE OF 4-PROPYL-2H-BENZO[H]-CHROMEN-2-ONE (FCS-304) IN MICE AND RATS

PERFIL DE TIPO SEROTONINÉRGICO DE 4-PROPIL-2H-BENZO[H]-CROMEN-2-ONA (FCS-304) EN RATONES Y RATAS

Laura M. MORENO¹, Luis E. CUCA², Mario F. GUERRERO^{1*}

Received: September 21, 2018. Approved: April 24, 2019

ABSTRACT

Background: 4-propil-2H-benzo[h]-cromen-2-ona (FCS-304) is a semisynthetic coumarin with MAO-A inhibitory activity and positive results in forced swimming and tail suspension test in mice, but until now, it has not been studied in other screening antidepressant models in mice and rats. **Objectives:** The aim of this work was to assess the serotonin like effect of FCS-304 in the 5-hydroxytryptophan (5-HTP) test in mice, in the behavioral despair test in rats, and in the reserpine test in rats. **Methods:** Potentiation of 5-HTP (100 mg/kg, i.p.), induced head twitches were assessed in mice, previously treated with FCS-304 (50-75-150 mg/kg, p.o.). The behavioral despair test was performed in rats treated with FCS-304, recording the immobility time attained by the animals subjected to forced swimming. Antagonism of reserpine-induced ptosis was examined in rats, assessing the level of palpebral closure. Imipramine (30 mg/kg, p.o.) and vehicle (canola oil) served as positive and negative controls, respectively. **Results:** FCS-304 significantly potentiated 5-HTP induced head twitches in mice, in a dose dependent manner. In rats, FCS-304 significantly decreased the immobility time in the behavioral despair test and antagonized reserpine induced ptosis. **Conclusions:** These results add support to propose that FCS-304 could elicit antidepressant effects related to MAO-A inhibitory activity.

Keywords: Antidepressant, Coumarin, 5-Hydroxytryptophan, MAO-A, Reserpine, Drug screening.

RESUMEN

Antecedentes: 4-propil-2H-benzo[h]-cromen-2-ona (FCS-304) es una cumarina semisintética inhibidora de MAO-A con efectos positivos en las pruebas de nado forzado y suspensión por la cola en ratones, sin embargo, hasta ahora no se había estudiado en otros modelos de tamizado antidepressivo en ratones y ratas. **Objetivos:** el objetivo de este trabajo fue evaluar el efecto de tipo serotoninérgico de FCS-304 en la prueba de potenciación de 5-hidroxitriptofano (5-HTP) en ratones, y su respuesta en la prueba de desesperanza conductual en ratas y en la prueba de reserpina en ratas. **Métodos:** se evaluó la potenciación de las sacudidas de cabeza inducidas por 5-HTP (100 mg/kg, i.p.), en ratones tratados con FCS-304 (50-75-150 mg/Kg, v.o.). La prueba de desesperanza conductual se realizó en ratas tratadas con FCS-304, expuestas a nado forzado. El antagonismo de la ptosis palpebral inducida por reserpina se

¹ Pharmacy Department, Faculty of Sciences, Universidad Nacional de Colombia, Bogotá, D.C., Colombia.

² Chemical Department, Faculty of Sciences, Universidad Nacional de Colombia, Bogotá, D.C., Colombia.

* Author of Correspondence: mfguerrerop@unal.edu.co

examinó en ratas determinando el grado de apertura ocular. Imipramina (30 mg/kg, v.o.) y el vehículo (aceite de canola, 0,1 mL/10 g), sirvieron como controles positivo y negativo, respectivamente. **Resultados:** FCS-304 incrementó significativamente el recuento de sacudidas de cabeza inducidas por 5-HTP en ratones, en función de la dosis. En ratas, FCS-304 fue efectiva para disminuir el tiempo de inmovilidad en la prueba de desesperanza inducida por nado forzado y el grado de ptosis palpebral inducida por reserpina. **Conclusiones:** estos resultados dan soporte para proponer que FCS-304 ejercería efectos de tipo antidepressivo relacionados con la inhibición de MAO-A.

Palabras clave: Antidepressivo, Cumarina, 5-hidroxitriptófano, MAO-A, Reserpina, Evaluación de fármacos.

INTRODUCTION

Major depression is a potential serious disorder with a high impact in worldwide public health, affecting up to 20% of the population, and contributes to consequential disabling in personal development, social relations, and higher morbidity and mortality (1). In Colombia, mental disorders reach up to 40% and major depression is the most prevalent (2). Given that this disorder results from the conjunction of a variable proportion of factors of endogenous nature, added to exogenous triggers, the appropriate psychological approach combined with pharmacological treatment are the backbone of therapy to achieve a greater improvement in patient response (3, 4).

Pharmacological treatment for major depression includes: selective serotonin reuptake inhibitors (SSRIs), serotonin/norepinephrine reuptake inhibitors (SNRIs), atypical antidepressants, serotonin-dopamine activity modulators (SDAMs), tricyclic antidepressants (TCAs), monoamine oxidase inhibitors (MAOIs), and St. John's wort. Although evidence for St. John's wort's effectiveness is modest, overall, effectiveness of the different antidepressant groups appears to be similar, albeit, there are differences with respect to onset of action and adverse events profile (5).

Reducing neurotransmitter breakdown through pharmacological inhibition of central monoamine oxidase (MAO), has been an effective strategy for treatment of a broad range of affective and anxiety disorders. However, the risk of hypertensive crisis discouraged the use of non-selective MAO inhibitors and led to the search for selective and non-reversible agents, such as selegiline, a MAO-B inhibitor, usually prescribed for Parkinson

disease, but also effective for major depression in a transdermal system (6).

MAO-A selective inhibition has been another approach to get an antidepressant effect with lower risks of potentiation of cardiovascular effects of dietary amines (7). Moclobemide is the prototype agent of this group that attained clinical use in several countries, but newer MAO-A inhibitors could represent an alternative treatment (8).

Coumarin is an active metabolite, isolated from many medicinal plants, among them, *Hygrophila tyttia* ("amansamachos"), used popularly in Colombia as a tranquilizer (9, 10, 11). This natural metabolite has served as the base for obtaining derivatives directed to different pharmacological targets, among which, MAO inhibition is one of the main targets (12). Several coumarin derivatives with a variety of ring substitutions, including 3-phenyl (13, 14), 3-aryl (15, 16), and 7-benzylpiperidin coumarins (17, 18) have led to potent MAO-B inhibition, whereas 6-amino 3-pyrrolidin (19) and 7-oxy-coumarins (20) have led to MAO-A inhibition. However, the correlation of the biological activity between coumarin derivatives *in vitro* and *in vivo* is a matter that requires further optimization (21).

Previous work showed that the coumarin derivative, coded as FCS-304 (4-propyl-2H-benzo[h]-chromen-2-one) displays MAO-A inhibition at the micromolar range, with no activity against MAO-B, as well as positive results in forced swimming and tail suspension test in mice (22). This work advances the antidepressant-like profile of this compound, assessing its activity in the 5-HTP test in mice to explore its pro-serotonergic activity, as well as its response in rats; in reserpine induce ptosis and in the behavioral despair test.

MATERIALS AND METHODS

FCS-304 synthesis

FCS-304 was obtained according to previously described, following Pechman condensation (22, 23), starting from 1-naphthol and ethyl 3-oxohexanoate (yield of 80%). FCS-304 melting point values were 103 - 105 °C and infrared and ¹H, ¹³C magnetic resonance spectra leading to the 4-propyl-7,8-benzocoumarine structure (Figure 1, C₁₆H₁₄O₂, M.W: 238).

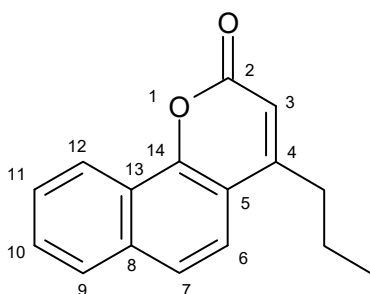


Figure 1. Structure of 4-propyl-2H-benzo[h]-chromen-2-one (FCS-304).

Experimental protocol

Animals were supplied by the Bioterium of the pharmacy department, Faculty of Sciences, Universidad Nacional de Colombia. Male ICR mice and female Wistar rats of 9-11 weeks old were kept under a 12 h dark/light cycle, 22°C temperature and 70% moisture with free access to water and food, except during the experiment day. They were randomly assigned to the following treatment groups (n=6-7 per group): FCS-304, three doses: 50, 75, and 150 mg/kg, p.o., according to a previous study (22), imipramine, as reference agent, 30 mg/kg, p.o. and vehicle: canola oil, as control group (0.1 mL/10 g, p.o.). These treatments were administered 1 h before each test. The experimental procedure was approved by the institutional ethics committee (Act 4, Jun 9, 2014, Faculty of Sciences, at the Universidad Nacional de Colombia).

5-HTP-induced head twitches in mice

This test was performed, based on Koe et al., (24, 25), administering 5-HTP, 100 mg/kg, i.p. in mice treated 1h prior with FCS-304, imipramine, or vehicle, according to the experimental protocol described above. Number of head twitches per animal was registered during 60 min.

Behavioral despair test in rats

This test was performed, based on Porsolt et al., (26, 27). Rats previously treated with FCS-304, imipramine, or vehicle were placed in a tank containing water (55 cm deep, 25°C) and left there for 5 min, measuring the total duration of immobility when the animals made only movements necessary to keep their heads above water.

Reserpine induced ptosis in rats

This test was performed based on Worms et al., (25, 28). Rats previously treated with FCS-304, imipramine or vehicle were injected with reserpine, 3.6 mg/kg, i.p. (according to preliminary assays) and 2 h later, the degree of ptosis was evaluated as follows: 0, eyes open; 1, eyes one-quarter closed; 2, eyes half closed; 3, eyes three-quarters closed; 4, eyes completely closed.

Drugs and solutions

Imipramine, L-5-hydroxytryptophan (5-HTP) and reserpine were obtained from Sigma®. Imipramine was dissolved in SSN, 0.9% (3 mg/mL), FCS-304 (4-propyl-2H-benzo[h]-chromen-2-one) was suspended in a canola oil vehicle and reserpine was prepared in a vehicle formed by a mixture of polysorbate-80, polypropyleneglycol-400, glycerin, and distilled water, in proportions of 2, 10, 10, and 78.

Statistical and data analysis

Results are expressed as mean ± standard error mean (S.E.M.). Given the small number of samples, Kolmogorov Smirnov test was applied to assess Gaussian distribution. Then, one-way ANOVA or Kruskal Wallis analysis, followed by multiple comparisons test were applied (p<0.05 against control). *GraphPad-Prism*® software, version 6, was used for data analysis.

RESULTS

Effects of FCS-304 in 5-HTP-induced twitches in mice

FCS-304, administered as a dose of 75, 150, and 300 mg/kg, p.o., significantly increased the number of mice head twitches in a dose dependent manner. Imipramine showed a slight increase (Figure 2).

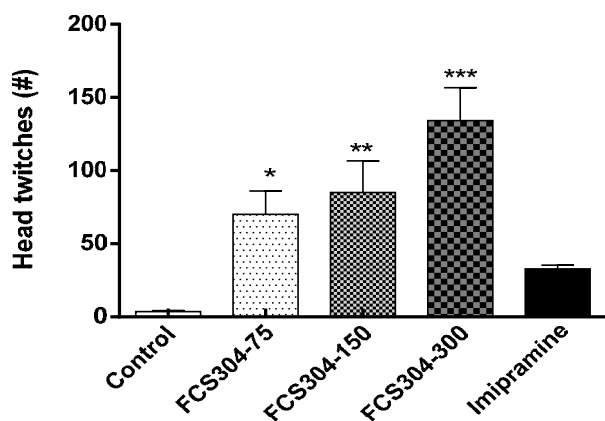


Figure 2. Effect of FCS304 (75, 150 and 300 mg/kg, v.o.), imipramine (30 mg/kg, v.o.) and control (vehicle, 0.1 mL/10 g, v.o.) on number of head twitches induced by 5-HTP (100 mg/kg, v.o.) in ICR mice. Results are expressed as means \pm S.E.M. * p <0.05, *** p <0.001 against control.

Effects of FCS-304 in behavioral despair in rats

Behavioral despair in rats, under induced forced swimming, showed a significant decrease in immobility time in rats treated with FCS-304, 300 mg/kg and the reference agent, imipramine, 30 mg/kg. Medium and low dose of FCS-304 (75, 50 mg/kg) did not achieve any difference from the control (Figure 3).

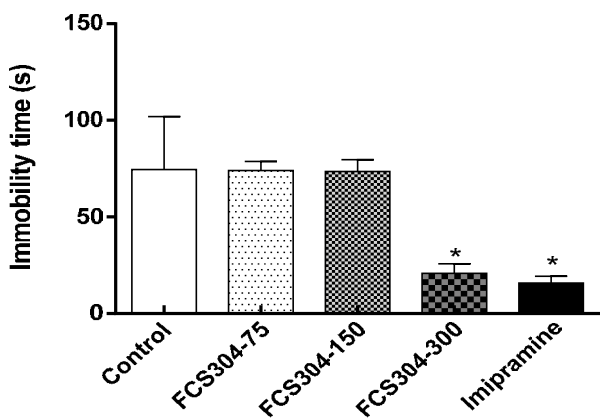


Figure 3. Effect of FCS-304 (75, 150 and 300 mg/kg, v.o.), imipramine (30 mg/kg, v.o.) and control (vehicle, 0.1 mL/10 g, v.o.) in immobility time of rats in behavioral despair when under induced forced swimming. Results are expressed as means \pm S.E.M. * p <0.05, against control.

Effects of FCS-304 in reserpinized rats

FCS-304 significantly decreased the palpebral ptosis and akinesia induced by reserpine in a dose dependent manner. Imipramine showed a greater response (Figure 4).

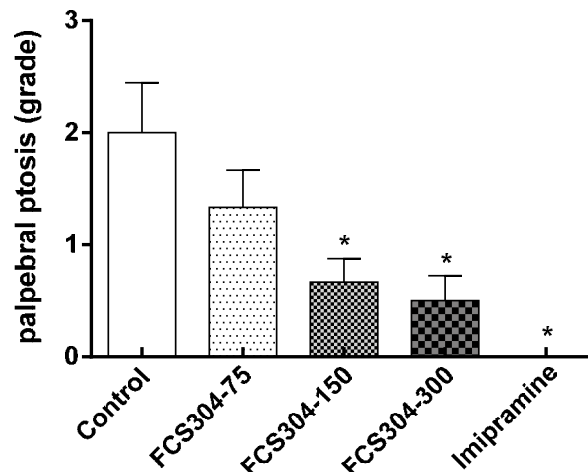


Figure 4. Effect of FCS-304 (75, 150, and 300 mg/kg, v.o.), imipramine (30 mg/kg, v.o.) and control (vehicle, 0.1 mL/10 g, v.o.) in the grade of palpebral ptosis induced by reserpine (3.6 mg/kg, i.p.) in rats. Values are represented as: 0, eyes open; 1, eyes one-quarter closed; 2, eyes half closed; 3, eyes three-quarters closed; 4, eyes completely closed * p <0.05, against control.

DISCUSSION

This study shows that FCS-304 increased the pro-serotonergic mechanism linked to the 5-OH-triptamine precursor 5-HTP, manifested by exacerbation of head twitches in mice, whereas in rats, it reduced the immobility time in forced swimming and decreased palpebral ptosis induced by reserpine. Globally considered, these results support the idea that FCS-304 follows an antidepressant like profile.

A previous work showed that FCS-304 displays an IC_{50} value of micromolar range against hMAO- whereas it lacks activity against hMAO-B (22). This selective inhibition of MAO-A isozyme by FCS-304 would represent the advantage of a less adverse effect profile related to cardiovascular risk, given that tyramine, present in foods like cheese and wine, shows similar affinity for each enzyme form, and hence, could be metabolized by MAO-B (7, 29).

Previously FCS-304 had showed activity in forced swimming and tail suspension test in mice,

therefore, it was useful to explore the activity of FCS-304 in other experimental models of depression and in another species, in this case, the rat. 5-HTP, as the immediate precursor in the biosynthesis of serotonin from tryptophan, is useful to study pro-serotonergic mechanisms of potential drugs and it is especially useful to test MAO-A inhibitors (30, 31).

Results of the 5HTP test support that FCS-304 could act through a pro-serotonergic mechanism at the central level. In addition, given that forced swimming and reserpine tests are high sensitive to non-selective agents, among them, tricyclic antidepressants (32, 33), it is reasonable to assume that imipramine shows a greater response in both assays, whereas FCS-304 is more effective against 5-HTP.

FCS-304 would share the favorable profile of selective MAO-A inhibitors: an anti-depressive profile linked to an increase in the activity of central serotonin, noradrenalin, and dopamine, neurotransmitters that participate in the regulation of the mood, whereas at the same time, have less risk of hypertensive events related to foods containing tyramine (7). Furthermore, MAO-A inhibition, due to its capacity to lower the generation of free radicals and toxins, remains as a target in the search for new drugs for the treatment of neurologic and neurodegenerative disorders (34, 35). This fact increases the pharmacological interest of FCS-304.

Coumarin derivatives have been extensively studied as sources of compounds with potential applicability in a wide range of disorders, including those of neurological nature: depression, anxiety, chronic pain, and Alzheimer disease, among them. Potent MAO-B and MAO-A coumarin inhibitors have been developed, but correlation between *in silico*, *in vitro*, and *in vivo* requires further optimization (12, 21).

Regarding FCS-304 as a MAO-A selectivity inhibitor, it was suggested that 4-hydrogen substitution and 7 aromatic radical presence, which could lead to covalent interactions with the flavin ring of the FAD cofactor required by the isozyme, would contribute the mechanism of action of this compound, but more research is needed in order to determine its structure-activity relationship (22).

CONCLUSION

In conclusion, these results support the proposal that FCS-304 is a coumarin derivative with MAO-A selective properties and antidepressant-like effects in mice and rats. More studies, including those of safety profile, are required to advance the pharmacological interest of this compound.

CONFLICTS OF INTEREST

There have been no conflicts of interest in carrying out this work.

ACKNOWLEDGMENTS

This work was supported by the Universidad Nacional de Colombia, (VRI/DIB, code: 31335). Thanks to the Animalarium of the Pharmacy Department (Prof. LF Ospina).

AUTHORS CONTRIBUTION

Laura M. Moreno (MSc) performed the experiments of this work, Luis E. Cuca (PhD) directed the chemical synthesis and Mario F. Guerrero (PhD) directed the pharmacological procedures.

REFERENCES

1. Kessler RC, Bromet EJ. The epidemiology of depression across cultures. *Annu Rev Public Health*. 2013;34:119-38.
2. Chaskel R, Gaviria SL, Espinel Z, Taborda E, Vanegas R, Shultz JM. Mental health in Colombia. *BJPsych Int*. 2015 Nov 1;12(4):95-97.
3. Pampallona S, Bollini P, Tibaldi G, Kupelnick B, Munizza C. Combined pharmacotherapy and psychological treatment for depression: a systematic review. *Arch Gen Psychiatry*. 2004 Jul;61(7):714-9.
4. Pemberton R, Fuller Tyszkiewicz MD. Factors contributing to depressive mood states in everyday life: A systematic review. *J Affect Disord*. 2016 Aug;200:103-10.
5. Cipriani A, Furukawa TA, Salanti G, Chaimani A, Atkinson LZ, Ogawa Y, Leucht S, Ruhe HG, Turner EH, Higgins JPT, Egger M, Takeshima N, Hayasaka Y, Imai H, Shinohara K, Tajika A, Ioannidis JPA, Geddes JR. Comparative efficacy and acceptability of 21 antidepressant drugs for the acute treatment of adults with major depressive disorder: a systematic review and network meta-analysis. *Lancet*. 2018 Apr 7;391(10128):1357-1366.
6. Bied AM, Kim J, Schwartz TL. A critical appraisal of the selegiline transdermal system for major depressive disorder. *Expert Rev Clin Pharmacol*. 2015;8(6):673-81.
7. Finberg JP, Rabey JM. Inhibitors of MAO-A and MAO-B in Psychiatry and Neurology. *Front Pharmacol*. 2016 Oct 18;7:340.
8. Chiucciariello L, Cooke RG, Miler L, Levitan RD, Baker GB, Kish SJ, Kolla NJ, Rusjan PM, Houle S, Wilson AA, Meyer JH. Monoamine Oxidase-A Occupancy by Moclobemide and Phenelzine: Implications for the Development of Monoamine

- Oxidase Inhibitors. *Int J Neuropsychopharmacol.* 2015 Aug 27;19(1).
- Pereira TM, Franco DP, Vitorio F, Kummerle AE. Coumarin Compounds in Medicinal Chemistry: Some Important Examples from the Last Years. *Curr Top Med Chem.* 2018;18(2):124-148.
 - Ariza SY, Rincón J, Guerrero MF. Efectos sobre el sistema nervioso central del extracto etanólico y fracciones de *Hygrophila tytttha* Leonard. *Rev Colomb Cienc Quim Farm.* 2006; 35(1):106-119.
 - Ariza SY, Rueda DC, Rincón J, Linares EL, Guerrero MF. Efectos farmacológicos sobre el sistema nervioso central inducidos por cumarina, aislada de *Hygrophila tytttha* Leonard. *VITAE.* 2007;14(2), 51-58.
 - Stefanachi A, Leonetti F, Pisani L, Catto M, Carotti A. Coumarin: A Natural, Privileged and Versatile Scaffold for Bioactive Compounds. *Molecules.* 2018 Jan 27;23(2). pii: E250.
 - Rauhämäki S, Postila PA, Niinivähmsä S, Kortet S, Schildt E, Pasanen M, Manivannan E, Ahinko M, Koskimies P, Nyberg N, Huuskonen P, Mutamäki E, Pasanen M, Juvonen RO, Raunio H, Huuskonen J, Pentikäinen OT. Structure-Activity Relationship Analysis of 3-Phenylcoumarin-Based Monoamine Oxidase B Inhibitors. *Front Chem.* 2018 Mar 2;6:41.
 - Delogu GL, Serra S, Quezada E, Uriarte E, Vilar S, Tatonetti NP, Viña D. Monoamine oxidase (MAO) inhibitory activity: 3-phenylcoumarins versus 4-hydroxy-3-phenylcoumarins. *ChemMedChem.* 2014 Aug;9(8):1672-6.
 - Costas-Lago MC, Besada P, Rodríguez-Enríquez F, Viña D, Vilar S, Uriarte E, Borges F, Terán C. Synthesis and structure-activity relationship study of novel 3-heteroaryl coumarins based on pyridazine scaffold as selective MAO-B inhibitors. *Eur J Med Chem.* 2017 Oct 20;139:1-11.
 - Matos MJ, Vilar S, García-Morales V, Tatonetti NP, Uriarte E, Santana L, Viña D. Insight into the functional and structural properties of 3-arylcoumarin as an interesting scaffold in monoamine oxidase B inhibition. *ChemMedChem.* 2014 Jul;9(7):1488-500.
 - Carotti A, Altomare C, Catto M, Gnerre C, Summo L, De Marco A, Rose S, Jenner P, Testa B. Lipophilicity plays a major role in modulating the inhibition of monoamine oxidase B by 7-substituted coumarins. *Chem Biodivers.* 2006 Feb;3(2):134-49.
 - Joubert J, Foka GB, Repsold BP, Oliver DW, Kapp E, Malan SF. Synthesis and evaluation of 7-substituted coumarin derivatives as multimodal monoamine oxidase-B and cholinesterase inhibitors for the treatment of Alzheimer's disease. *Eur J Med Chem.* 2017 Jan 5;125:853-864.
 - Mattsson C, Svensson P, Sonesson C. A novel series of 6-substituted 3-(pyrrolidin-1-ylmethyl)chromen-2-ones as selective monoamine oxidase (MAO) A inhibitors. *Eur J Med Chem.* 2014 Feb 12;73:177-86.
 - Abdelhafez OM, Amin KM, Ali HI, Abdalla MM, Batran RZ. Monoamine oxidase A and B inhibiting effect and molecular modeling of some synthesized coumarin derivatives. *Neurochem Int.* 2013 Jan;62(2):198-209.
 - Pisani L, Catto M, Leonetti F, Nicolotti O, Stefanachi A, Campagna F, Carotti A. Targeting monoamine oxidases with multipotent ligands: an emerging strategy in the search of new drugs against neurodegenerative diseases. *Curr Med Chem.* 2011;18(30):4568-87.
 - Vergel NE, López JL, Orallo F, Viña D, Buitrago DM, del Olmo E, Mico JA, Guerrero MF. Antidepressant-like profile and MAO-A inhibitory activity of 4-propyl-2H-benzo[h]-chromen-2-one. *Life Sci.* 2010 May 22;86(21-22):819-24.
 - Potdar MK, Mohile SS, Salunkhe MM. Coumarin syntheses via Pechmann condensation in Lewis acidic chloroaluminate ionic liquid. *Tetrahedron Letters.* 2001;42(52):9285-9287.
 - Koe BK, Weissman A, Welch WM, Browne RG. Sertraline, 1S,4S-N-methyl-4-(3,4-dichlorophenyl)-1,2,3,4-tetrahydro-1-naphthylamine, a new uptake inhibitor with selectivity for serotonin. *J Pharmacol Exp Ther.* 1983 Sep;226(3):686-700.
 - Kato M, Katayama T, Iwata H, Yamamura M, Matsuoka Y, Narita H. In vivo characterization of T-794, a novel reversible inhibitor of monoamine oxidase-A, as an antidepressant with a wide safety margin. *J Pharmacol Exp Ther.* 1998 Mar;284(3):983-90.
 - Porsolt RD, Bertin A, Jalfre M. Behavioral despair in mice: a primary screening test for antidepressants. *Arch Int Pharmacodyn Ther.* 1977 Oct;229(2):327-36.
 - Lapa AJ, Souccar C, Lima MT, Lima TCM. Métodos farmacológicos para el estudio de actividad sobre el sistema nervioso central. In: CYTED/CNPq (Ed), Métodos de evaluación de la actividad farmacológica de plantas medicinales. Florianópolis, Santa Catarina, pp 70-90, 2002.
 - Worms P, Kan JP, Wermuth CG, Roncucci R, Bizière K. SR 95191, a selective inhibitor of type A monoamine oxidase with dopaminergic properties. I. Psychopharmacological profile in rodents. *J Pharmacol Exp Ther.* 1987 Jan;240(1):241-50.
 - Nowakowska E1, Chodera A. Inhibitory monoamine oxidases of the new generation. *Pol Merkur Lekarski.* 1997 Jul;3(13):1-4.
 - Corne SJ, Pickering RW, Warner BT. A method for assessing the effects of drugs on the central actions of 5-hydroxytryptamine. *Brit. J. Pharmacol.* 1963;20:106-120.
 - Mahesh R, Jindal A, Gautam B, Bhatt S, Pandey D. Evaluation of anti-depressant-like activity of linezolid, an oxazolidinone class derivative - an investigation using behavioral tests battery of depression. *Biochem Biophys Res Commun.* 2011 Jun 17;409(4):723-6.
 - Castagné V, Moser P, Roux S, Porsolt RD. Rodent models of depression: forced swim and tail suspension behavioral despair tests in rats and mice. *Curr Protoc Neurosci.* 2011 Apr;Chapter 8:Unit 8.10A.
 - Ozerov AA, Bagmetova VV, Chernysheva YV, Tyurenkov IN. Comparison of the Efficiency of Adeprophen and Antidepressants of Various Groups on the Model of Reserpine-Induced Depression in Rats. *Bull Exp Biol Med.* 2016 Mar;160(5):649-52.
 - Naoi M, Maruyama W, Inaba-Hasegawa K. Type A and B monoamine oxidase in age-related neurodegenerative disorders: their distinct roles in neuronal death and survival. *Curr Top Med Chem.* 2012;12(20):2177-88.
 - Pathak A, Srivastava AK, Singour PK, Gouda P. Synthetic and Natural Monoamine Oxidase Inhibitors as Potential Lead Compounds for Effective Therapeutics. *Cent Nerv Syst Agents Med Chem.* 2016;16(2):81-97.