

# The GC MS Analysis of a Rare Medicinal Plant *Aloe barbadensis*

S. Jayakumari<sup>1</sup>, Prabhu K<sup>2</sup>, Mudiganti Ram Krishna Rao<sup>3</sup>, Bhupesh<sup>4</sup>, D.Kumaran<sup>5</sup>, Aishwariya Ramesh<sup>5</sup>

<sup>1</sup> Assistant Professor, Department of Anatomy, Sree Balaji Medical College & Hospital (SBMCH), Bharath University, BIHER, Chrompet, Chennai-600044, Tamil Nadu, India.

<sup>2</sup> Associate Professor, Department of Anatomy, Sree Balaji Medical College & Hospital (SBMCH), Bharath University, BIHER, Chrompet, Chennai-600044, Tamil Nadu, India.

<sup>3</sup> Professor, Department of Industrial Biotechnology, Bharath University, Selaiyur, Chennai-600044, Tamil Nadu, India.

<sup>4</sup> Senior Scientist, Central Research Lab., Sree Balaji Medical College & Hospital (SBMCH), Bharath University, BIHER, Chrompet, Chennai-600044, Tamil Nadu, India.

<sup>5</sup> Student, Sree Balaji Medical College & Hospital (SBMCH), Bharath University, BIHER, Chrompet, Chennai-600044, Tamil Nadu, India

## Abstract

The present study deals with the GC MS analysis of Ethyl acetate extract of the plant *Aloe barbadensis*. The plant was collected from Moonaru hills, Kerala. Its color was red and the jelly inside also was blood red in color. The difference of this plant from the common *Aloe vera* has prompted us to go for the GC MS analysis to find whether any major difference exists among them. It is interesting to find that the results are different from already existing reports on *Aloe vera*. Some important constituents such as Idazoxan, Dextroamphetamine, 1-ethylpropylamine, Resepidone, P-methyl benzoic acid, 2-Thiophene carboxylic acid, Dimethyl 4-Chlorophenyl thiophosphate, Benzphetamine, Picloram and Jasmonoyl acid, Glyphosate, Mepyramine, Physostigmine, Methylphosphonic acid, Selaginine, 2,4- Dichlorophenol were shown in the GC MS analysis results. These differences could be due to the geographical and microclimatic conditions or this species of *Aloe* could be a different from the others. further studies to identify the reasons for these differences are under progress.

**Key Words** *Aloe barbadensis*, GC MS, Idazoxan, Reserpidone, 2-Thiophene carboxylic acid, Selaginine

## INTRODUCTION

*Aloe barbadensis* is a rare species of *Aloe* genus which has medicinal value similar to *Aloe vera*. The present study deals with the GC MS analysis of Ethyl acetate extract of the plant to understand the bio molecules present in it. This plant is different than the *Aloe vera* plants in having red color and with the jelly inside is also blood red in color. It was collected from Moonaru hills, Kerala and the plant material was subjected to GC MS analysis after processing it suitably. The difference of this plant from the common *Aloe vera* has prompted us to go for this test to find there is any major difference. It is interesting to find that the results are different from already existing reports on *Aloe vera* [1-7]. These differences could be due to the geographical and microclimatic conditions or this species of *Aloe* could be a different from the others. Further studies to identify the reasons for these differences are under progress. The present study is a part of our aim towards herbal standardization process. (8-27)

## MATERIALS AND METHODS

### Preparation of plant extract

Fresh leaves of *Aloe barbadensis* were collected from Moonaru hills, Kerala and the pulp material were extracted with 90% ethanol using a continuous hot percolation method in a Soxhlet apparatus for 18 hrs. The extract was concentrated in a rotary evaporator to yield a semi solid

mass. Then it was kept at -20°C in deep freezer. Further the extract was lyophilized using alpha Christ lyophilizer. The powder contents were stored at 4°C in refrigerator until use.

### Gas Chromatography Mass Spectroscopy

The Gas Chromatography and Mass Spectroscopy (GC MS) examination was done by using Perkin – Gas Chromatogram coupled to a mass identifier, Turbo mass gold – Perkin Elmer Turbomass spectrometer with an Elite - (100% Dimethyl poly siloxane), P - 265, 30m x 0.25 mm ID x 0.25µm of fine column. Injection temperature was kept up at 250°C, Helium stream rate as 1.5 ml/min and particle source temperature at 290°C. Infusion was performed in the split less mode and the volume was 1 µL. The instrument was set to an underlying temperature of 70°C, and kept up at this temperature for 3 min. Towards the end of this period, the temperature was emerged up to 300°C, at the rate of an expansion of temperature 10°C/min. The mass spectra of were attained by electron ionization (EI) at 70 eV, and the indicator worked in sweep mode from 40 – 700 m/z. The MS begin time was 3 min end time was 35 min with dissolvable cut time was around 3 min. The fundamental compound constituents were distinguished by coordinating mass spectra with spectra of reference masses in the library of the National Institute of Standards and Innovation (NIST 11). The retention values and probable type of molecules were presented.

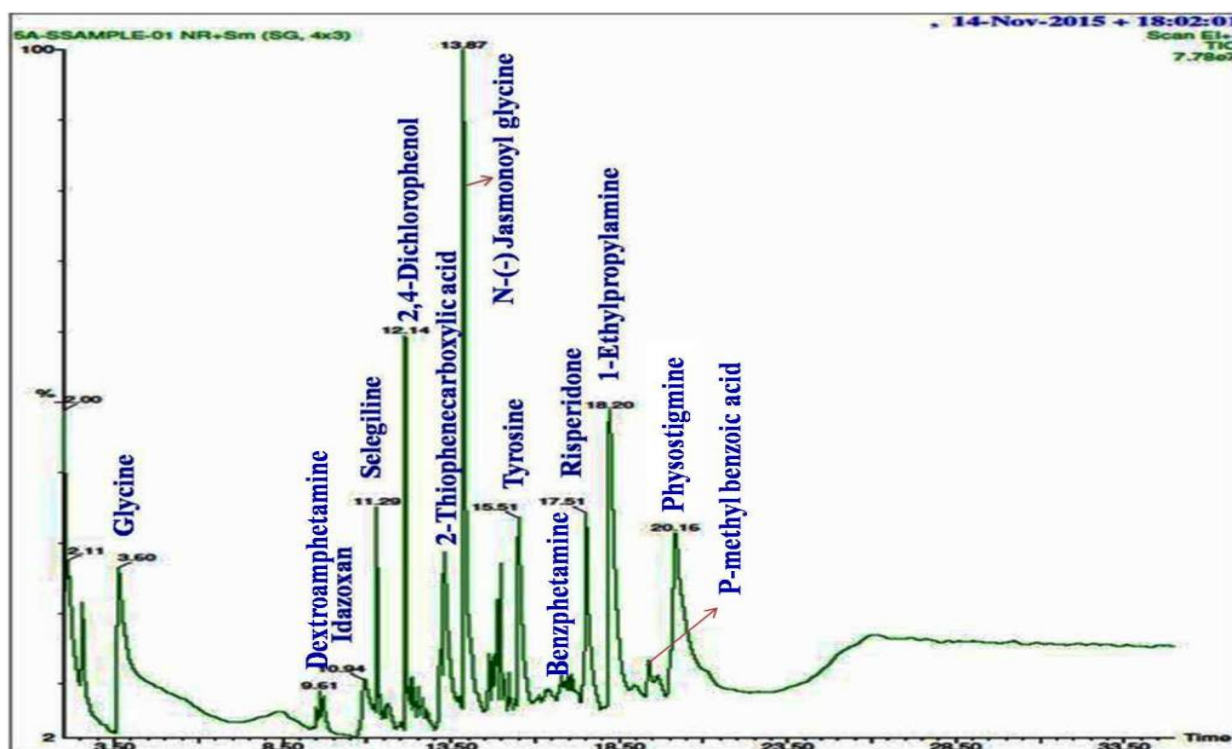


Figure 1. GC MS profile of Aloe barbadensis (Red variety).

## RESULTS AND DISCUSSION

Figure-1 indicated the identification of bioactive compounds present in the ethyl acetate extract of *Aloe barbadensis*. The extract was subjected to the Gas Chromatography Mass Spectroscopy. The GC MS spectrum was identified and elucidated for the presence of bioactive compounds from the NIST library. The Bioactive compounds are listed in Table 1. Some important constituents such as Idazoxan, Dextroamphetamine, 1-ethylpropylamine, Risperidone, P-methyl benzoic acid, 2-Thiophene carboxylic acid, Dimethyl 4-Chlorophenyl thiophosphate, Benzphetamine, Picloram and Jasmonoyl acid, Glyphosate, Mepyrmine, Physostigmine, Methylphosphonic acid, Selaginine, 2,4- Dichlorophenol was found. The medicinal roles of some of the important bioactive components is mentioned below.

### 1. Idazoxan

This chemical is a  $\alpha_2$  adrenoceptor antagonist and has been extensively used for preclinical study to support the " $\alpha_2/D_2$  receptor hypothesis" for atypical antipsychotic effects. [28-31]

### 2. Dextroamphetamine

This molecule is sold in the brand names of *Adderall*, *Adderall XR* and used for treating attention deficit hyperactivity and narcolepsy. This stimulates the brain by increasing the level of neurotransmitters like dopamine and norepinephrine.

### 3. 1-ethylpropylamine.

This bioactive compound and its derivatives are known as corticotrophin releasing factor 1 receptor antagonists.

### 4. Risperidone

This compound is used mainly for the treatment of schizophrenia, bipolar disorder and irritability in people with autism. It is taken either orally or by injection into a muscle. The injectable version is long acting and lasts for about two weeks.

### 5. P-methyl benzoic acid

This molecule, which is also present in *Aloe vera*, has been shown to have antibacterial and antioxidant activities.

### 6. 2-Thiophenecarboxylic acid.

Mishra *et al*, 2011 have reviewed the medicinal role of this compound and its derivatives.[37] The derivatives of this compound are reported to be antiallergic (Gillespie *et al*, 1985), antibacterial (Elslager *et al*, 1972), analgesic and anti-inflammatory (Santagati *et al*, 1994

### 7. Dimethyl 4-Chlorophenyl thiophosphate

This compound is a known antifungal.[41]

### 8. Benzphetamine

This compound is used for reduction of appetite in obesity management. [42]

### 9. Picloram and Jasmonoyl acid derivatives

These compounds are used in tissue culture for better callus growth. [43]

### 10. Glyphosate

This is a pesticide and can cause various diseases in humans and animals. The possible presence of this compound could reflect the use of more chemical as fertilizers where from this plant was collected.

### 11. Mepyrmine is an antidepressant and antihistaminic. [44]

**12. Physostigmine**

Physostigmine acts by interfering with the metabolism of acetylcholine. It is a covalent (reversible - bond hydrolyzed and released) inhibitor of acetylcholinesterase, the enzyme responsible for the breakdown of acetylcholine in the synaptic cleft of the neuromuscular junction. [45] Physostigmine is used to treat glaucoma, Alzheimer's disease, and delayed gastric emptying. It has been shown to improve short term memory.

**13. Methylphosphonic acid**

Aminophosphonic acid is reported to have a number of biological activities. [46-49] These compounds are defined as amino acid derivatives, in which the carboxylic acid group [-C(=O)OH] is replaced by the phosphonic acid moiety [-P(=O)(OH)<sub>2</sub>] [50]. Such modification inhibits the activity of certain enzymes by effective competition for the active site of the enzyme and by forming strong electrostatic binding [51]. Thus, the aminophosphonic acids found application as enzyme inhibitors [52, 53].

**14. Selegiline**

Selegiline is a monoamine oxidase inhibitor (MAOI). It works by prolonging the anti-Parkinson activity of levodopa, which may help to slow the progression of Parkinson disease. Selegiline has no anti-Parkinson

effects of its own and must always be given in combination with levodopa/carbidopa. [54]

**15. 2, 4- Dichlorophenol**

2, 4-Dichlorophenol (2, 4-DCP) is a chlorinated derivative of phenol with the molecular formula C<sub>6</sub>H<sub>4</sub>Cl<sub>2</sub>O. 2, 4-DCP is used primarily as an intermediate in the preparation of the herbicide 2,4-dichlorophenoxyacetic acid (2,4-D).

The plants of *Aloe* genus are grown widely all over the world for their ornamental as well as medicinal values. The jelly of *Aloe* is used mostly in skin care industries which is claimed to cause smoothness and healthy skin. Domestically the jelly is used for skin care, hair care and also as a coolant. The juice is taken in the early hours as it is supposed to stimulate good health. Although some reports on the various health benefits of *Aloe* are there, it is a long way to go to establish these claims at standardized level. The present work is one step in this direction.

**CONCLUSION**

From the above discussion it could be concluded that *Aloe barbadensis* contain some very important bio-molecules which have significant medicinal properties. Further work to understand the molecular mechanism of their action is under progress.

**Table 1. The table indicates the GC MS results of *Aloe barbadensis* showing Retention time and possible types of compounds.**

Sl. No	Retention Time (Min)	Type of Possible compound/s as Per NIST
1.	3.614	Glycine, Bromophenol. Gly-Gly
2	4.209	1-Ethylpropylamine, Pro-Tyr, L-Pro-L-Ser, Phosphoric acid, monododecyl ester, Dextroamphetamine, methylphosphonic acid pos 45 d[M+H] <sup>+</sup> 15
3	4.429	Dextroamphetamine, Benzphetamine
4	4.820	1-Ethylpropylamine, Pro-Tyr, L-Pro-L-Ser, 3-Bromophenol
5	5.630	1-Ethylpropylamine, Picloram, Glyphosate, free acid, Mepyramine, Physostigmine
6	10.967	Idazoxan, Pro-Tyr
7	11.292	Methadone, L-Deprenyl, Dextroamphetamine, Selegiline
8	12.137	2,4-Dichlorophenol
9	13.303	Idazoxan, 2-Thiophenecarboxylic acid
10	13.863	Risperidone, p-Methylbenzoic acid, Aspartic acid, N-(-)-Jasmonoyl glycine
11	15.514	Pro-Tyr, Idazoxan, 2-Thiophenecarboxylic acid
11	17.510	Idazoxan, 2-Thiophenecarboxylic acid, Dimethyl 4-chlorophenyl thiophosphate
12	18.200	1-Ethylpropylamine, Pro-Tyr, Oxalic acid, Alanine
13	20.146	1,3-Butadiene, 2,3-dimethyl-, Cyclohexene, 1,4-Hexadiene Cyclopentane, methylene

## REFERENCES

- Edward, D. S., Mariappan V. *World J Pharmacy and Pharmaceu Sci*, 2014, 3, (5),749 - 758.
- Saljooghianpour, M., Javarant T.A. *African J of Biotechnology*, 2013, 12(49), 6876 - 6880
- Manikandan, V. G., Muhammad, I. M. H. *Int. J. Bio. Pharma. And Allied. Sci.*, 2014, 3(1), 56-62
- Kumar, A. S., Muthuselvam, M. *World J of Agri Sci.*, 2009, 5 (5), 572 - 576.
- Botes, L. , Francois, H. van der Westhuizen, Du Toit Loots. *Molecules*, 2008, 13, 2169 - 2180
- Lakshmi, P. T. V. Rajalakshmi P. A. *Int Res J of Pharmacy*, 2011, 2(5), 247 - 249.
- Satyaprabha G, Kumaravel S, Ruffina D, Parveen Kumar P. *Journal of Pharmacy Res*, 2010, 3(12), 2970 - 2973.
- Rao, M. R. K., Philip, S, Kumar, M. H., Saranya, Y., Divya, D., Prabhu K. *J Chem Pharma Res*, 2015, 7(7):131 - 139.
- Ravi, A, Prabhu, J. S. P, Rao, M. R. K., Prabhu, K., Kalaiselvi, V.S., Saranya, Y. *Int. J. Pharm. Sci. Rev. Res.*, 2015, 33(2), 58 - 62
- Chandrasekar, T, Rao, M. R. K., Vijaya Kumar, R., Prabhu, K., Nandha Kumar, S, Divya, D. *J Chem Pharma Res*, 2015, 7(8), 124 - 136
- Sadhanandham, S, Narayanan, G, Rao M. R. K., Prabhu, K., Sumathi Jones, Aparna Ravi, Shruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2015, 34(2), 273 - 281
- Phillips, S, Rao, M. R. K., Prabhu, K., Minu Priya, Kalaivani, S., Aparna Ravi, Shruti Dinakar. *J Chem Pharma Res*, 2015, 7(9), 393 - 401.
- Rao, M. R. K., Nandha Kumar S., Sumathi Jones, Arul Amudha Elizabeth, Prabhu, K., Aparna Ravi, Shruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2015, 34(2), 6 - 12
- Rao, M. R. K., Aparna Ravi, Shridhar Narayanan, Prabhu, K., Kalaiselvi, V. S., Shruthi Dinakar, Guru Rajan, Kotteeswaran, N. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 36(1), 158 - 166
- Rao, M. R. K., Hassan Mohammad, Sridhar Narayanan, Prabhu, K., Kalaiselvi, V. S., Aparna Ravi, Hari Babu, Guru Rajan, Suganya, S. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 37(1), 19 - 25
- Aparna Ravi, Monika Gupta, Rao, M. R. K., Kalaivani, Kalaiselvi, V.S., Prabhu, K., Shruthi Dinakar, Rao, G. V. *Der Pharmacia Lettre*, 2015, 7 (12),45 - 52
- Aparna Ravi, Hassan Mohammad, Rao, M. R. K., Prabhu K., Hari Babu, Shridhar Narayanan, Guru Rajan, Sanjay Singh. *Int J Phama Tech.*, 2015, 7(3), 10091-10112
- Sivakumaran, G, Rao, M. R. K., Prabhu, K, Kalaiselvi V. S., Sumathi Jones, Johnson, W. M., Antony, J. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 37(1), 190 - 199
- Angeline Jessica, Rao M. R. K., Jacintha Anthony, Prabhu, K., Johnson, W. M. S., Shanthi Balasubramanian, B., Lakshmi Sundaram, Shruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 39(2), 216 - 224.
- Edel Queen, Z, Rao, M.R. K., Jecintha Anthony, Prabhu, K., Johnson, W.M. S., Shanthi Balasubramanian, B., Lakshmi Sundaram, Shruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 39(2), 169 - 172.
- Cynthia Shankari, Vathsala Venkatesan, Mudiganti Ram Krishna Rao, Saravanan, Prabhu K., Seppan Prakash. *Int. J. Pharm. Sci. Rev. Res.*, 2016, 40(1), 33 - 37.
- Rao, M.R.K., Saikumar P., Prabhu K., Arul Amutha Elizabeth, Sumathi, Lakshmi Sundaram, Sruthi Dinakar, Kumari Sangita Singh, Ayub Alam. *Int. J. Pharm. Sci. Rev. Res.*, 2017, 42(1), 15 - 19.
- Nirupa, Mudiganti Ram Krishna Rao, Prabhu K, Kaliaselvi, V.S., Kumaran, D., Sivaram E., Sruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2017, 42(1), 35 - 41.
- Sivasankar Reddy Konda, Mudiganti Ram Krishna Rao, Minu Priya, Prabhu K., V. S. Kalaivani, V.S., Kumaran, D., Ayub Alam, Kumari Sangeeta Singh, Lakshmi Sundaram. *Int. J. Pharm. Sci. Rev. Res.*, 2017, 42(1), 29 - 34.
- Muthu Lakshmi Muthiah, Mudiganti Ram Krishna Rao, Arul Amutha Elizabeth, Farhana Rahman, *Int. J. Pharm. Sci. Rev. Res.*, 2017, 42(2), 236 - 238.
- Praveen Kumar, P., Rao, M.R. K., Arul Amutha Elizabeth, Prabhu, K., Lakshmi Sundaram, R., Shruthi Dinakar. *Int. J. Pharm. Sci. Rev. Res.*, 2017, 42(1), 5 - 8.
- Jai Prabhu, Prabhu, K, Mudiganti Ram Krishna Rao, Kalaiselvi V. S., Vani Krishna, Aishwarya Ramesh. *Int. J. Pharm. Sci. Rev. Res.*, 2017, 44(1), 235 - 239.
- Adam J. Prus, Patricia A. Zornio, Candice J. Schuck, Tanya Heerts, Sarah M. Jacobson and Dominika A. Winiarski. *Drug Dev Res*, 2010, 71, 261-267.
- Fagan, G. P., Chapleo, C. B., Lane, A. C., Myers, M., Roach, A. G., Smith, C. F. C., Stillings, M. R., Welbourn, A. P. *J Med Chem*, 1998, 31(5), 944 - 948.
- Krystal, J. H., McDougale, C. J., Woods, S. W., Price, L. H., Heninger, G. R., Charney, D. S. *Psychopharmacology*, 1992. 313 - 319.
- Martire, M., Pistrutto, G., Preziosi, P. *Neuroscience Letters*, 1988, 86(3), 328 - 333
- Lakhan, S. E., Kirchgessner, A. *Brain Behaviour*, 2012, 2(5), 661 - 667.
- Kenny, J. D., Taylor, N. E., Brown, E. N., Solt, K. *PLOS*, 2015, DOI: 10.1371/journal.pone.0131914
- Tetsuji Saito, Tetsuo Obitsu, Takashi Kondo, Toshiaki Matsui, Yuuki Nagao, Kensuke Kusumi, Naoya Matsumura, Sonoko Ueno, Akihiro Kishi, Seishi Katsumata, Yoshifumi Kagamiishi, Hisao Nakai, Masaaki Toda. *Bioorganic and Medicinal Chemistry*, 2011, 9, 15.
- Canitano, R., Scandurra, V. *Neuropsychiatr Dis Treat.*, 2008, 4(4), 723 - 730.
- Fatemeh N-Borandozi. *Org Med Chem Lett*. 2013, 3, 5
- Raghav Mishra, Jha, K. K., Sachin Kumar, Isha Tomer. *Der Pharma Chemica*, 2011, 3 (4), 38 -54.
- Gillespie E., Dungan K. M., Gomol A. W., Seidehamel R. J. *Int. J. Immunopharmacol*, 1985, 7, 655.
- Elslager E. F., Jacob P., Werbel L. M. *J Hetero Chem.*, 1972, 9, 775.
- Santagati, A., Modica M., Santagati M., Garuso A., Cutuli, V. *Pharmazie*, 1994, 49, 64 -65
- Pati Bibek, Banerjee Subhasis. *Journal of Pharmacy Research*, 2012, 5(12), 5493.
- Yanvoski, S. Z., Yanovski, J. A. *JAMA*, 2014, 74 - 86.
- Abdul Bakrudeen Ali Ahmed, Adhikarla Suryanarayana Rao, Mandali Venkateswara Rao Rosna Mat Taha. *Braz Arch Biol Technol*. 2011, 54(1), 7 - 11.
- Tran, V.T, Chang, R. S., Snyder, S. H. *Proc. Nati. Acad. Sci*. 1978, 75(12), 6290-6294.
- Katzung, B. G., Masters, S. Trever, A. (2009). *Basic and Clinical Pharmacology.*, 2009, p. 110.
- Kafarski P., Lejczak B., Tyka, R., Koba L., Pliszczak E., Wiczorek P. *Journal of Plant Growth Regulation*, 1995, 14(4), 199 - 203.
- Naydenova, E. D, Topashka-Ancheva, M., Todorov, P., Tordanova T., Troev K. *Bioorganic and Medicinal Chemistry*, 2006, 14(7), 2190 - 2196
- Naydenova E. D., Todorov P. T., Troev K. D. *Amino Acids*, 2010, 38(1), 23 - 30.
- Kraszewski, A., Stawinski J. *Pure and Applied Chemistry*, 2007, 79(12), 2217 - 2227.
- Lejczak, B. Kafarski P. *Topics in Heterocyclic Chemistry*, 2009, 20, 31 - 63,
- Pawelczak, M., Nowak, K., Kafarski, P. *Phosphorus, Sulfur and Silicon and Related Elements*, 1998, 132, 65 - 71
- Tanner M. E., S. Vaganay, J. van Heijenoort, Blanot D. *Journal of Organic Chemistry*, 1996, 1(5), 1756 - 1760, 1996.
- Bubenik, M., Rej, R., N. Nguyen-Ba, G. Attardo, Ouellet F., Chan L. *Bioorganic and Medicinal Chemistry Letters*, 2002, 12(21), 3063 - 3066,
- Amsterdam, J. D. *Journal of Clinical Psychiatry*, 2003, 64 (2): 208-214.