

Available online at http://www.journalcra.com

International Journal of Current Research Vol. 9, Issue, 12, pp.63120-63122, December, 2017 INTERNATIONAL JOURNAL OF CURRENT RESEARCH

# **RESEARCH ARTICLE**

## ANTINEOPLASTIC EFFECT OF SELECTED ESSENTIAL OILS ON ETHANOL INDUCED LIVER TOXICITY IN ALBINO WISTAR RATS

## <sup>1</sup>Christina, K. J., \*,<sup>2</sup>Balwin Nambikkairaj, <sup>2</sup>Umasankar, K. and <sup>2</sup>Ramya, D. R.

<sup>1</sup>Department of Biochemistry, Chettinad Hospitals and Research Institute, Kelambakkam <sup>2</sup>P. G. and Research Department of Zoology, Voorhees College, Vellore, (T.N.), India

#### ARTICLE INFO

## ABSTRACT

Article History: Received 25<sup>th</sup> September, 2017 Received in revised form 08<sup>th</sup> October, 2017 Accepted 16<sup>th</sup> November, 2017 Published online 31<sup>st</sup> December, 2017

Key words:

Antineoplastic, Hepatoprotective, Ethanol, *Syzygium aromaticum, Rosa damascene* essential oil, Albino wistar rats. Liver is a solid glandular organ made up of hepatic lobules separated from one another by connective tissue. Each lobe consists of a mass of polyhedral hepatic cells, which are glandular in appearance containing spherical nuclei. A network of blood vessels pierces the lobules. The hepatic cells are arranged in longitudinal cords. It has a gall bladder and secretes bile. Liver is the major metabolic centre and any damage to this organ would subsequently lead to so many physiological disturbances. In the present study antineoplastic effect of *Syzygium aromaticum* and *Rosa damascena* essential oils on ethanol induced liver toxicity in male wistar rats were carried out. From the results it is evident that the administration of N-nitrosodiethylamine considerably reduced the protein, non enzyme antioxidants such as glutathione, vitamins C and E, and enzymatic antioxidants such glutathione peroxidase, superoxide dismutase and catalase. All the values are statistically significant.

*Copyright* © 2017, *Christina et al.* This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Citation: Christina, K. J., Balwin Nambikkairaj, Umasankar, K. and Ramya, D. R. 2017. "Antineoplastic effect of selected essential oils on ethanol induced liver toxicity in albino wistar rats", *International Journal of Current Research*, 9, (12), 63120-63122.

## INTRODUCTION

Liver is the first organ to metabolize all foreign compounds. The liver is a vital organ present in vertebrates and some other animals. It has a wide range of functions including detoxification and protein synthesis (Heinrich and Barnes, 2004). The liver is a necessary organ for survival. This organ plays a major role in metabolism and has a number of functions in the body, including glycogen storage, decomposition of red blood cells, plasma protein synthesis, hormone production and detoxification (Kaul et al., 2000). It lies below the diaphragm in the abdominal- pelvic region of the abdomen. It produced bile an alkaline compound which aids in digestion via the emulsification of lipids (Yassa et al., 2009). The liver highly specialized tissues regulate a wide variety of high volume biochemical reactions, including the synthesis are necessary for normal vital functions (Gholamhoseinian et al., 2009). In the present investigation antineoplastic effect of Syzygium aromaticum and Rosa damascena essential oil on ethanol induced liver toxicity in male wistar rats was carried out. The antioxidant and the herb Syzygium aromaticum and Rosa damascena essential oil used for hepatoprotective (Thomas et al., 2004).

## \*Corresponding author: Balwin Nambikkairaj,

P. G. and Research Department of Zoology, Voorhees College, Vellore, (T.N.), India.

## **MATERIALS AND METHODS**

Study on the selected herbal plants Syzygium aromaticum and Rosa damascena essential oils were orally treated in different concentration of liver toxicity in male albino wistar rats compared to control animals. Hepatoprotective activity of different factor such as total protein, glutathione peroxidase, superoxide dismutase and catalase were analyzed by sigma Diagnostic kit (Sigma chemical company catalogue, 1997). The test animal used was albino wistar rats. LD<sub>50</sub> analysis was measured using albino wistar rats treatment with Syzygium aromaticum and Rosa damascena, essential oil. The LD<sub>50</sub> analysis was orally fed with different dose of ethanol and the LD<sub>50</sub> value was calculated as per the method of Finney (1971).

## Eight groups were maintained

- 1. Control rat fed with normal diet
- 2. Albino wistar rat induced ethanol
- 3. Rat + ethanol + 100µl/ ml *Syzygium aromaticum* essential oil
- 4. Rat + ethanol + 200µl/ ml *Syzygium aromaticum* essential oil
- 5. Rat + ethanol + 300µl/ ml *Syzygium aromaticum* essential oil

- 6. Rat + ethanol + 100µl/ ml Rosa damascena essential oil
- 7. Rat + ethanol +  $200\mu$ l/ ml *Rosa damascena* essential oil
- 8. Rat + ethanol + 300µl/ ml Rosa damascena essential oil

The rats were maintained in the above condition for 10 weeks and analyzed for various biochemical factors and antioxidant enzymes in the blood sample of the chronically alcohol exposed rats.

**Statistical Analysis**: All the data were analyzed as per the method of Pillai and Sinha (1968).

## **RESULTS AND DISCUSSION**

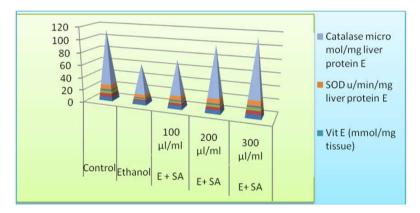
Table 1-2 and Fig 1-2 Plate 1-2 indicate the result of the factors. The plant show *Syzygium aromaticum* and *Rosa damascena* essential oil in the ethanol induced hepatotoxicity in the rats were studied. The serum biochemical, non enzymatic and enzymatic antioxidants in the ethanol hepatotoxicity induced *Syzygium aromaticum* and *Rosa damascena* essential oil ingested rats (Jirovetz *et al.*, 2006) were analyzed. The results indicate that the administration of ethanol had considerably affected the various metabolic

 Table 1. Biochemical, Non enzymatic antioxidant and enzyme antioxidant in Syzygium aromaticum treated ethanol toxicity induced rats

| Parameters  | Control         | Ethanol         | E + SA 100 µl/ml | E+ SA 200 µl/ml | E+ SA 300 µl/ml |
|---|-----------------|-----------------|------------------|-----------------|-----------------|
| Protein g/l   | 6.5±2.02        | 6.0±2.02        | 6.1±2.02         | 6.25±2.02       | 6.7±2.02        |
| Glutathione peroxidase micro g/min/mg liver protein E | 5.40±3.12       | 2.20±3.12       | 3.4±3.22         | 4.3±2.12        | 5.60±3.12       |
| GSH (mg/mg protein)                                   | 4.64±1.22       | 1.72±1.24       | 2.8±1.36         | 3.48±1.36       | 4.75±1.34       |
| Vit C (mmol/mg tissue)                                | $1.42 \pm 1.26$ | $1.01 \pm 3.00$ | $1.19\pm2.34$    | $1.36 \pm 2.30$ | $1.46 \pm 2.32$ |
| Vit E (mmol/mg tissue)                                | 0.96±2.14       | $0.67 \pm 2.24$ | 0.71±2.12        | 0.78±2.12       | $0.98 \pm 2.12$ |
| SOD u/min/mg liver protein E                          | 7.16±3.16       | 4.4±3.12        | 6.0±3.12         | 6.7±3.14        | 7.18±3.12       |
| Catalase micro mol/mg liver protein E                 | 86.5±2.14       | 47.0±2.12       | 54.0±2.22        | 78.0±2.12       | 88.24±2.12      |

Values are mean  $\pm$  SD of 6 individual observations.

Values are significant at P = 0.001.



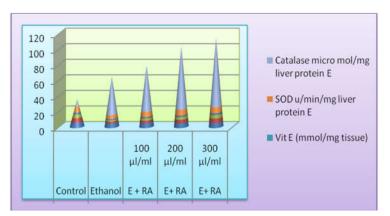
# Figure 1. Biochemical, Non enzymatic antioxidant and enzyme antioxidant in *Syzygium aromaticum* treated ethanol toxicity induced rats

Table 2. Biochemical, Non enzymatic antioxidant and enzyme antioxidant in Rosa damascena treated ethanol toxicity induced rats

| Parameters  | Control       | Ethanol         | $E + RA 100 \ \mu l/ml$ | E+ RA 200 µl/ml | E+ RA 300 µl/ml |
|---|---------------|-----------------|-------------------------|-----------------|-----------------|
| Protein g/l   | 6.5±0.14      | 6.0±0.14        | 6.1±0.14                | 6.2±0.14        | 6.5±0.14        |
| Glutathione peroxidase micro g/min/mg liver protein E | 5.40±0.16     | 2.20±0.16       | 3.2±0.16                | 4.3±0.16        | 5.50±0.16       |
| GSH (mg/mg protein)                                   | 4.64±1.12     | $1.72 \pm 1.12$ | 2.9±1.12                | 3.48±1.12       | 4.70±1.12       |
| Vit C (mmol/mg tissue)                                | $1.42\pm2.18$ | $1.01 \pm 2.20$ | 1.20±2.20               | $1.38\pm2.18$   | $1.45 \pm 2.18$ |
| Vit E (mmol/mg tissue)                                | 0.96±1.21     | 0.67±1.21       | 0.74±1.21               | 0.78±1.21       | $0.96 \pm 2.21$ |
| SOD u/min/mg liver protein E                          | 7.16±3.14     | 4.4±3.14        | 6.2±3.14                | 6.9±3.14        | 7.16±3.14       |
| Catalase micro mol/mg liver protein E                 | 8.65±2.16     | 48±2.16         | 58±2.16                 | 80±2.16         | 87.54±2.16      |

Values are mean  $\pm$  SD of 6 individual observations.

Values are significant at P = 0.001.



pathways and had reduced the biochemical and antioxidant factors (Seidemann et al., 2005). The level of Glutathione, Superoxide dismutase and Catalase have reduced from 4.64 µg/min/ml SOD liver protein, 7.16 u/min/ml Catalase liver protein E and 86.5µmol/mg Glutathione peroxidase liver protein, 2.20 µg/min/ml SOD liver protein, 4.4 u/min/ml catalase liver protein and 48 µmol /mg liver protein in ethanol exposed rats. Whereas the administration of 300 µl/min/ml of Syzygium aromaticum had improved factors such as glutathione, superoxide dismutase and catalase have reduced from 5.50 µg/min/ml SOD liver protein, 7.16 u/min/ml catalase liver protein and 87.54 µmol/mg liver protein respectively (Shahriari et al., 2007). The increase in the Syzygium aromaticum concentration had linearly increased the levels of antioxidant and biochemical factors, thus improving the liver health with ethanol induced hepatotoxicity (Lisin et al., 1999).

#### Conclusion

- It is concluded that orally treated *Syzygium aromaticum* and *Rosa damascena* essential oil can be used for the hepatoprotective activity.
- It is a complementary study that is also applied to humans.

## REFERENCES

- Finney, D.J. 1971. Probit Analysis 3<sup>rd</sup> ed. Cambridge: Cambridge university press.
- Gholamhoseinian, A., Fallah, H. and Sharififar, F. 2009. Inhibitory effect of methanol extract of *Rosa damascena* Mill. Flowers on a-glucosidase activity and postprandial hyperglycemia in normal and diabetic rats. *Phytomedicine*, 16:935-941.

- Heinrich, M. and Barnes, J. 2004. Clove, Syzygium aromaticum (L.) Merr. & L.M. Perry. Fundamentals of Pharmacognosy and Phytotherapy. Churchill Livingstone, Edinburgh-London, Great Britain, 275-276.
- Jirovetz, L. and Buchbauer, G. 2006. Chemical Composition and Antioxidant Properties of Clove Leaf Essential Oils. *Journal of Agricultural and Food Chemistry*, 54:6303-6307.
- Kaul, V.K., Singh, V. and Singh, B. 2000. Damask rose and marigold: prospective industrial crops. J Med Aromat Plant Sci., 22: 313-318.
- Lisin, G., Safiyev, S. and Craker, L.E. 1999. Antimicrobial activity of some essential oils. *Acta Horticulturae (ISHS)*, 501:283-288.
- Pillai, S.K. and Shina, H.C. 1986. In Statistical methods for biological workers pubs. Ramprasad and sons Agra, India.
- Seidemann, J. 2005. Syzygium aromaticum (L.) Merr. LM Perry. World Spice Plants. Springer-Verlag, Berlin-Heidelberg, Germany, 355-356.
- Shahriari, S., Yasa, N., Mohammadirad, A., Khorasani, R. and Abdollahi, M. 2007. *In vitro* antioxidant potential of *Rosa damascene* extract from guilan, Iran comparable to αtocopherol. *Int J Pharmacol.*, 3:187-190.
- Sigma Diagnostic kits (Sigma Chemical company catalogue, 1997).
- Thomas, P.D.R. and Montvale, N.J. 2004. Physicians' Desk Reference (PDR) for Herbal Medicines. Clove-Syzygium aromaticum. Edn 3, 204-208.
- Yassa, N., Masoomi, F., Rohani Rankouhi, S.E. and Hadjiakhoondi, A. 2009. Correspondence chemical composition and antioxidant activity of the extract and essential oil of *Rosa damascena* from Iran, Population of Guilan. Daru; 17:175-180.

\*\*\*\*\*\*