



Review Article

A REVIEW ON CURATIVE EFFECTS OF HERBAL DRUGS ON VARIOUS CHEMICAL INDUCED HEPATOTOXICITY

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ABSTRACT

A possible risk factor of drug-induced hepatotoxicity is the metabolism of drug enzyme activity. Some drugs when used for prolonged or by overdose may induce hepatotoxicity. Chronic alcohol consumption, exposure of zinc oxide nanoparticles widely used in industry & cosmetics, reveals high potency risks of hepatotoxicity. Now, herbal medicines and formulations are developing to overcome liver injuries. Hepatoprotective effect of the goat milk, *Momordica dioica Roxb.*, *Berberis aristata*. The aim of the present review is to compiling data on promising hepatotoxic as well as hepatoprotective agents.

KEY WORDS: Hepatotoxicity, Herbal plants, Hepatoprotective, Antioxidants.

INTRODUCTION

Generally, hepatotoxicity (liver dysfunction or liver damage) that occurs due to surplus of drugs or xenobiotics. Hepatotoxins or hepatotoxicants are the chemicals that produced the liver dysfunction or liver damage¹. Hepatotoxicants are exogenous compounds, may include overdoses of certain medicinal drugs, industrial chemicals, herbal remedies and dietary supplements. Usually, hepatotoxic response is expressed, when hepatocytes, biliary epithelial cells and liver vasculature affected by toxicants². More than 900 drugs have been concerned in causing liver injury and it is the most common explanation for a drug to be quiet from the market. Drug-induced liver injury is liable for 5% of all hospital admissions and 50% of all acute liver failures. More than 75 percent of cases of idiosyncratic drug reactions result in liver transplantation or death³. Liver pathology is a significant tool for identifying and characterizing liver injury. Main features of liver injury during hepatotoxicity may include zonal necrosis, hepatitis, cholestasis, steatosis, granuloma and vascular lesions diseases.

Zonal Necrosis: This kind of injury may be caused by exogenous substances like paracetamol and carbon tetrachloride. It may mark as an extremely high level of alanine aminotransferase and severe disturbance of liver function leading to acute liver failure.

Hepatitis: This type of liver injury shows hepatocellular necrosis associated with infiltration of inflammatory cells. It may be further characterized into three categories: viral, focal and chronic.

Cholestasis: This type of liver injury leads to destruction of bile flow, itching and jaundice. **Steatosis:** This type of liver injury

may manifest as triglyceride accumulation which leads to either small droplet or large droplet fatty liver.

Granuloma: Hepatic granulomas are associated with granulomas located in periportal or portal areas and show features of systemic vasculitis and hypersensitivity.

Vascular Lesions: Such condition is caused by injury to the vascular endothelium⁴.

AGENTS THAT CAUSES HEPATOTOXICITY

Drug induced hepatotoxicity is the most important cause of acute liver failure in many countries. Some of the medications causing hepatotoxicity as a potential side effect of various drugs. Aspirin, Ketoprofen, Tetracycline, Nucleoside reverse transcriptase inhibitors, Valproic acid, Acetaminophen and Methotrexate may lead to macrovesicular steatosis. Most tricyclic antidepressants are potentially hepatotoxic⁵.

HEPATOPROTECTIVE AGENTS

Silibinin extracted from *Silybum marianum* commonly called as “Milk thistle” is used against several drugs-induced hepatotoxicity by significant increase in the activities of marker enzymes (aspartate and alanine transaminase, alkaline phosphatase)⁶. Chlorogenic acid stabilizes cell membrane; prevent hepatocyte apoptosis⁷. Hepatoprotective natural products such as *Andrographis paniculata*, *Chamomile capitula*, *Coccinia grandis*, *Flacourtia indica*, *Wedelia calendulacea*, *Annona squamosa*, *Prostecheamichuacana*, are reported⁸. *Berberis* species also known as Indian Barberry⁹. All species of the *Berberis* are rich source of vitamins, minerals, antioxidants, anthocyanin etc¹⁰. *Thymus vulgaris* have protective effect on alcohol mediated hepatotoxicity. Virgin coconut oil with its

potent natural antioxidant property shows hepatoprotective effect against methotrexate induced liver toxicity and oxidative stress¹¹.

CONCLUSION

The liver plays a central role in transforming and clearing chemicals and is susceptible to the toxicity. Certain medicinal agents, when taken in overdoses and sometimes even when introduced within therapeutic ranges, may injure the organ. Other chemical agents, such as those used in laboratories and industries, natural chemicals and herbal remedies can also induce hepatotoxicity. Drug-induced hepatic injury is the most common reason cited for withdrawal of an approved drug. Knowledge of the commonly implicated agents and a high index of suspicion are essential in diagnosis.

In general, herbal medicines have been considered as safe by the general public, since they are naturally occurring and have been applied in treatment for over thousands of years. As the use of herbal medicine is rapidly increasing globally. In the present study we assume that traditional medicinal plant used in Ayurvedic, Chinese and other medicinal systems in the world for a long time may act as the potent hepatoprotective.

REFERENCES

1. Hartmut J, Gregory JG, Arthur IC, Jack AH, Dominique P, John JL. Mechanisms of Hepatotoxicity. *Toxicological Sciences*. Volume 65, Issue 2, 1 February 2002, Pages 166–176.
2. Navarro VJ, Senior JR. Drug-Related Hepatotoxicity, *New England Journal of Medicine*, Volume 354, Issue 7, 16 February 2006, Pages 731-9.
3. Aashish P, Tarun S, Pallavi B. Drug induced hepatotoxicity- A review, *Journal of Applied Pharmaceutical Science*, Volume 02, Issue 05, January 2012, Pages 233-243.
4. Anita S, Tej KB, Om PS. Clinical Biochemistry of Hepatotoxicity, *Journal of Clinical Toxicology*, S4:001. doi:10.4172/2161-0495.S4-001.
5. Voican CS, Corruble E, Naveau S, Perlemuter G. Antidepressant-Induced Liver Injury: A Review for Clinicians. *The American Journal of Psychiatry*. Volume 171, Issue 4 April 2014, Pages 404-15.
6. Ramanathan R, Sivanesan K. Evaluation of ameliorative ability of Silibinin against zidovudine and isoniazid-induced hepatotoxicity and hyperlipidaemia in rats: Role of Silibinin in Phase I and II drug metabolism. *Chemico-Biological Interactions*, Volume 273, 1 August 2017, Pages 142-153.
7. Dai C, Xinyu Z, Lihan X, Xiang L, Lihua H, Chunling W. Protective and prophylactic effects of chlorogenic acid on aluminum-induced acute hepatotoxicity and hematotoxicity in mice, *Chemico-Biological Interactions*, Volume 273, 1 August 2017, Pages 125-132.
8. Anil K. A review on hepatoprotective herbal drugs. *International Journal of Research in Pharmacy and Chemistry*, Volume 2, 2012, Pages 2231–2781.
9. Mathew G, Lincy J, Sumi J. Phytochemical and Pharmacological screening of anti-inflammatory activity of *Berberis lycium* root extract. *Pharmacophore*, Volume 7, Issue 1, 2016, Pages 01-05.
10. Purvika S, Rajni M, Monika S. *Berberis lycium* Medicinal Plant with Immense Value, *Indian Journal of Pharmaceutical & Biological Research*, Volume 1, Issue 1, January 2013, Pages 27-37.
11. Zakaria ZA, Rofiee MS, Somchit MN, Zuraini A, Sulaiman MR, Teh LK et al. Hepatoprotective Activity of Dried and Fermented Processed Virgin Coconut Oil. *Evidence Based Complement Alternative Medicine*, Volume 2011, Article ID 142739, Pages 8, doi:10.1155/2011/142739.

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