

RESEARCH ARTICLE

Significance of *Rumex Vesicarius* as Anticancer Remedy Against Hepatocellular Carcinoma: a Proposal-Based on Experimental Animal Studies

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Abstract

Rumex vesicarius is an edible herb distributed in Egypt and Saudi Arabia. The whole plant has significant value in folk medicine and it has been used to alleviate several diseases. Hepatocellular carcinoma (HCC), the major primary malignant tumor of the liver, is one of the most life-threatening human cancers. The goal of the current study was to explore the potent role of *Rumex vesicarius* extract against HCC induced in rats. Thirty adult male albino rats were divided into 3 groups: (I): Healthy animals received orally 0.9 % normal saline and served as negative control group, (II): HCC group in which rats were orally administered N-nitrosodiethylamine NDEA, (III): HCC group treated orally with *R. vesicarius* extract in a dose of 400 mg/kg b.wt daily for two months. ALT and AST, ALP and γ -GT activities were estimated. CEA, AFP, AFU, GPC-3, Gp-73 and VEGF levels were quantified. Histopathological examination of liver tissue sections was also carried out. The results of the current study showed that the treatment of the HCC group with *R. vesicarius* extract reversed the significant increase in liver enzymes activity, CEA, AFP, AFU, glypican 3, golgi 73 and VEGF levels in serum as compared to HCC-untreated counterparts. In addition, the favorable impact of *R. vesicarius* treatment was evidenced by the marked improvement in the histopathological features of the liver of the treated group. In conclusion, the present experimental setting provided evidence for the significance of *R. vesicarius* as anticancer candidate with a promising anticancer potential against HCC. The powerful hepatoprotective properties, the potent antiangiogenic activity and the effective antiproliferative capacity are responsible for the anticancer effect of this plant.

Keywords: Hepatocellular carcinoma - polygonaceae - *R. vesicarius* - angiogenesis - apoptosis - rats

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Introduction

Anticancer therapy is one of the biggest challenges in medicine. Therefore, searching for highly efficient antitumor drugs remains a hot area of research. Plants and plant-based products are gaining importance against variety of diseases due to their non-toxic and no / less side effects. Hence, we have been continuously working on the identification and characterization of medicinal plants (Khan, 2012). *Rumex vesicarius* L. of the family Polygonaceae is an edible herb, locally called hummayed, growing as wild herb throughout desert and semi-desert areas of Saudi Arabia. *R. vesicarius* is widely used as food, as a medicinal herb and as an antidote to scorpion stings (Shahat et al., 2013). The medicinal importance of *R. vesicarius* is a reflection to its chemical composition since the plant contains many bioactive substances such as flavonoids, anthraquinones, carotenoids, vitamins, lipids and organic acids, which are known to be anticancer,

antioxidant and antimicrobial agents (Barbosa et al., 2008; Imran et al., 2011). Of particular interest, It has been demonstrated that all plant parts contained quercetin in high amounts (Bakry et al., 2012). The intake of dietary antioxidants of plant origin like carotenoids, phenolic compounds and flavonoids may lead to the protection against non-communicable diseases in human beings such as cancer, cardiovascular diseases and cataract (Matkowski, 2008).

R. vesicarius has many important medicinal uses such as treatment of tumors, liver diseases, bad digestion, constipation, calculi, heart troubles, pains, diseases of the spleen, hiccup, flatulence, asthma, bronchitis, dyspepsia, piles, scabies, leucoderma, toothache and nausea. This plant is also used as laxative, stomachic, tonic, analgesic, appetizer, diuretic, astringent, purgative and antispasmodic. Moreover, can be used to reduce biliary disorders and control cholesterol levels (Lakshmi et al., 2009). The lipid constituents of *R. vesicarius*

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were examined by both liquid chromatography/mass spectrometry (LC/MS) and by gas chromatography/mass spectrometry (GC/MS). Their essential oil compositions consisted mainly of thujene, limonene, fenchon, estragole, and anethole. The crude lipid extract and the methanol extract showed strong antioxidant activity and radical quenching potential against 2, 2-diphenyl-1-picrylhydrazyl (DPPH) systems (Bakry et al., 2012; Elfotouh, et al., 2013).

Cancer is the leading cause of death around the world (Zeng et al., 2015). Cancer is, still a devastating disease, responsible for roughly one quarter of deaths and one of the most imminent health problems in the developed world and it continues to be a major disease for those in developed and developing countries. Cancer deserves a high priority of research owing to the large number of deaths, the enormous human suffering, and related health care. Hepatocellular carcinoma (HCC) is one of the major health burdens worldwide (Lyer et al., 2010) and is a leading cause of cancer death worldwide (Mitupatum et al., 2015). It is one of the most life-threatening human cancers in the world, resulting in almost one million deaths every year (Center and Jemal, 2011). HCC is the third leading cause of cancer deaths worldwide, with prevalence 16-32 times higher in developing countries. The incidence of HCC is higher in males than in females with the male: female ratio usually averaging between 2:1 and 4:1 (El-Serag and Rudolph, 2007). Also, the rising trend of HCC has been associated with increased prevalence of hepatitis C virus (HCV) infection (Serag, 2002). Lingering infection with hepatitis B virus (HBV) and hepatitis C virus (HCV) represents the major risk factor for HCC (Gao et al., 2012). In Egypt, HCC is the second most common malignancy in males and the fifth in the females. Egypt has the highest prevalence of hepatitis C virus in the world with an average of approximately 13.8% in the general population (Deuffic et al., 2006). The major risk factor of HCC is infection by HCV, which accounts for 20% of acute hepatitis, 70% of chronic hepatitis, 40% of cirrhosis and up to 90% of HCC cases (Hosny et al., 2008). Lack of effective diagnostic tools for early detection and limited treatment options for patients with advanced HCC contribute to a dismal prognosis coupled with high mortality for this disease (Thomas et al., 2010; Bishayee, 2012).

Diethylnitrosamine (NDEA) is found in a wide variety of foods and beverages such as cheese, soybeans, smoked, salted and dried fish, cured meat, alcoholic beverages as well as in ground water having a high level of nitrates (Liao et al., 2001). In rats, NDEA is a potent hepatocarcinogen influencing the initiation stage of carcinogenesis during a period of enhanced cell proliferation accompanied by hepatocellular necrosis. NDEA induces DNA carcinogen adducts, DNA-strand breaks and in turn hepatocellular carcinomas without cirrhosis through the development of putative preneoplastic focal lesions (Ahmed et al., 2010). Anticancer activities from many functional food sources have been reported in years, but correlation between cancer prevalence and types of food with anticancer activities from crop origin center in the world as well as food source with human migration are unclear (Zeng et

al., 2015). The current study was undertaken to elucidate the anticancer potential of *R. vesicarius* extract against diethylnitrosamine-induced hepatocellular carcinoma in male rats with special concern on its mechanism of action.

Materials and Methods

Chemicals

N-nitrosodiethylamine (NDEA) (CAS no. 55-18-5) was purchased from Sigma-Aldrich Chemicals Co. (St Louis, MO, USA). All other chemicals and solvents were of analytical grade.

Plant material

Rumex vesicarius was collected from the different area of the campus of the plants of King Saud University, Riyadh, Saudi Arabia in April 2012. The plant was identified by the Plant Taxonomist at the Herbarium Unit. The voucher specimens (15936) was deposited at the Herbarium of the Faculty of Pharmacy, King Saud University, Riyadh, Saudi Arabia

Plant extraction

The plant was collected and the aerial part was dried under shade. The dried samples were powdered and used for solvent extraction. For extract preparation, 1.2 kg of dried sample was extracted twice with 3 L of 80% methanol. The extracts were filtered through Whatman No.1 filter paper and concentrated using a rotary evaporator under reduced pressure at 40°C to obtain residue with yields of 13%

Experimental set up

A total number of thirty adult male rats of Wistar strain weighing 170-200g were used in the present work. The animals were obtained from the Animal House Colony of the National Research Centre, Giza, Egypt. The rats were housed in polypropylene cages in an environmentally controlled clean air room with a temperature of 25±1°C, an alternating 12h light/12h dark cycle, a relative humidity of 60±5% and free access to tap water and a standard rodent chow (Wadi El Kabda Co., Cairo, Egypt). The rats were allowed to adapt to these conditions for 2 weeks before starting the experimental set-up. The experimental protocol was approved by the Ethical Committee for Medical Research, National Research Centre, Egypt. After the acclimatization period, the animals were divided into three groups of equal average body weight and kept in well ventilated caged. They were labeled namely group *i*): Normal healthy animals received orally 0.9% normal saline and served as negative control group, group *ii*): HCC group in which the rats were orally administered with N-nitrosodiethylamine (NDEA) (dissolved in 0.9% normal saline), in a dose of 20 mg/kg b.wt. five times weekly for six weeks according to the modified method of Darwish and El-Boghdady (2011) and group *iii*): HCC-treated group in which the rats were treated orally with *R. vesicarius* extract in a dose of 400 mg/kg b.wt (Raghavendra and Reddy, 2011) daily for two months.

Samples collection

After the completion of this round, the animals were fasted overnight and the blood samples were collected, under diethyl ether anesthesia, from the retroorbital venous plexus in a dry clean centrifuge tubes and allowed to coagulate for 45 minutes at room temperature to obtain sera to be used for biochemical analyses. Clear serum samples were separated by centrifugation at 1800 xg for 15 minutes at 4°C using cooling centrifuge. Serum samples were stored at -20°C pending further biochemical analyses. After collection of the blood samples, the animals were sacrificed by cervical dislocation and the liver specimens from experimental animals were quickly excised, rinsed with saline and fixed in 10% formalin saline solution for histological examination.

Biochemical analyses

Serum alanine aminotransferase (ALT) and asparatate aminotransferase (AST) activities were estimated using colorimetric kit purchased from Salucea Co. Ltd (Netherlands) according to ECCLS (1989) method. Serum alkaline phosphatase (ALP) and gamma-glutamyl transferase (γ -GT) activities were determined using colorimetric kit purchased from Reactivos GPL Co. Ltd (Barcelona) according to Tietz (1995) method. Serum carcinoembryonic (CEA) was quantified by enzyme linked immunosorbent assay (ELISA) technique using a kit purchased from Immunospec Co., Ltd (USA), according to Schwartz (1987) method. Serum alpha-fetoprotein (AFP) level was measured by ELISA technique using ELISA kit purchased from Immunospec Co., Ltd (USA), according to the method of Hirai (1982). Serum alpha-L-fucosidase (AFU) activity was evaluated by ELISA technique using a kit purchased from Glory Science Co., Ltd (USA), according to the manufacturer's instructions provided with AFU assay kit. Serum glypican-3 (GPC-3) level was assayed by ELISA technique using a kit purchased from Glory Science Co., Ltd (USA), according to the manufacturer's instructions provided with GPC-3 assay kit. Serum golgi protein 73 (Gp-73) level was detected by ELISA technique using a kit purchased from Glory Science Co., Ltd (USA), according to the manufacturer's instructions provided with Gp-73 assay kit. Serum vascular endothelial growth factor (VEGF) level was estimated by ELISA technique using a kit purchased from Glory Science Co., Ltd (USA), according to the manufacturer's instructions provided with VEGF assay kit.

Histopathological examination

After fixation of the liver specimens in formal saline

(10%) for 24 hours, the tissues were washed in running tap water and dehydrated in series of alcohol (methyl, ethyl and absolute alcohol). The specimens were cleared in xylene and embedded in paraffin at 56 degree in hot air oven for twenty four hours. The paraffin wax tissue blocks were sectioned by sledge microtome at thickness of 4 μ m. The obtained tissue sections were collected on clean glass slides and left in the oven at 40°C for dryness before examination under the light electric microscope (Banchroft et al., 1996).

Statistical analysis

The experimental results were represented as arithmetic means with their standard errors. Data were analyzed by one way analysis of variance (ANOVA) using the Statistical Package for the Social Sciences (SPSS) program, version 14 followed by least significant difference (LSD) to compare significance between groups (Armitage and Berry, 1987). The level of significance was set at $P < 0.05$. Percentage difference representing the percent of variation with respect to the corresponding control group was also calculated using the following formula:

$$\% \text{ difference} = \frac{\text{Treated value} - \text{Control value}}{\text{Control value}} \times 100$$

Results

Biochemical data

The results in Table (1) represented the influence of treatment with *R. vesicarius* extract on serum liver enzymes (AST, ALT, ALP and γ -GT) activity in HCC bearing rats. The HCC group showed significant elevation ($P < 0.05$) in serum AST (49.79%), ALT (98.85%), ALP (133.43%) and γ -GT (173.17%) activity relative to the negative control group (Table 1). Treatment of the HCC group with *R. vesicarius* extract elicited significant reduction ($P < 0.05$) in serum AST (-22.83%), ALT (-30.9%), ALP (-38.21%) and γ -GT (-24.28%) activity with respect to the untreated HCC group (Table 1).

The results depicted in Table (2) illustrated the influence of treatment with *R. vesicarius* extract on serum tumor markers (CEA, AFP and AFU), glypican 3, golgi 73 and VEGF levels in HCC bearing rats. The HCC group revealed significant increase ($P < 0.05$) in serum CEA (500%), AFP (111.05%), AFU (275%), glypican 3 (63.68%), golgi 73 (39.393%) and VEGF (41.06%) levels in comparison with the negative control group. In contrast, treatment of the HCC group with *R. vesicarius*

Table 1. Influence of Treatment with *R. Vesicarius* Extract on Serum Liver Enzymes Activity in HCC bearing rats

Parameter	AST (U/L)	ALT (U/L)	ALP (U/L)	γ -GT (U/L)
Groups				
Negative control	84.0 \pm 1.53	44.7 \pm 1.05	109.5 \pm 7.11	14.48 \pm 1.06
HCC group	125.8 \pm 13.2 ^a (49.79%)	88.9 \pm 4.31 ^a (98.85%)	255.6 \pm 12.6 ^a (133.43%)	39.55 \pm 1.91 ^a (173.17%)
HCC+ <i>R.vesicarius</i> group	97.1 \pm 2.14 ^b (-22.83 %)	61.5 \pm 1.66 ^b (-30.9 %)	157.9 \pm 9.68 ^b (-38.21 %)	29.95 \pm 1.56 ^b (-24.28 %)

*Data were expressed as means \pm standard error (SE) for 10 animals/group.

Table 2. Influence of Treatment with *R. vesicarius* Extract on Serum Tumor Markers (CEA, AFP and AFU) Levels, Glypican 3, Golgi 73 and VEGF Levels in HCC Bearing Rats

Parameter Groups	CEA (ng/ml)	AFP (ng/ml)	AFU (pg/ml)	Glypican 3 (pg/ml)	Golgi 73 (ng/ml)	VEGF (ng/L)
Negative control	0.12±1.250×10 ⁻²	19.03±1.52	32.33±3.07	2.96±0.18	231.0±7.35	91.66±2.61
HCC group	0.60±6.0×10 ^{-2a} (500%)	40.07±2.98 ^a (111.05 %)	121.2±6.1 ^a (275 %)	4.85±0.13 ^a (63.68%)	322.0±3.74 ^a (39.393 %)	129.3±2.09 ^a (41.06%)
HCC+ <i>R.vesicarius</i> group	0.25±1.20×10 ^{-2b} (-57.5%)	30.44±0.41 ^b (-24.08 %)	65.83±2.66 ^b (-45.7%)	3.63±0.2 ^b (-25.02%)	284.1±9.62 ^b (-11.75%)	100.76±3.8 ^b (-22.08%)

*Data were expressed as means±standard error (SE) for 10 animals / group, ^a: P<0.05 vs negative control; ^b: P>0.05 vs HCC group

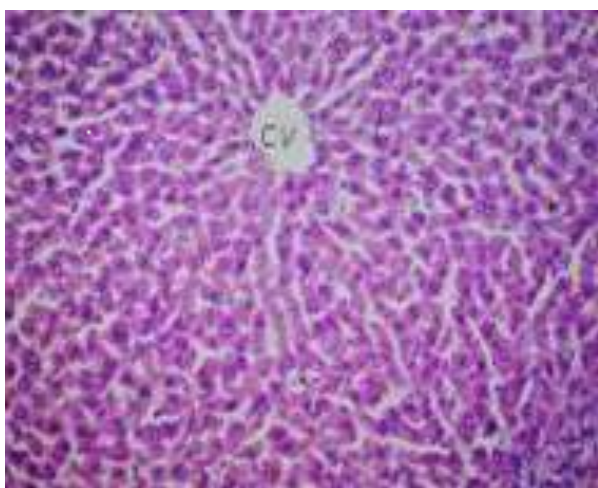


Figure 1. Photomicrograph of Liver Tissue Section of Rat in the Negative Control Group Showing Normal Histological Structure of the Central Vein and the Surrounding Hepatocytes in the Hepatic Parenchyma (H&E x 40)

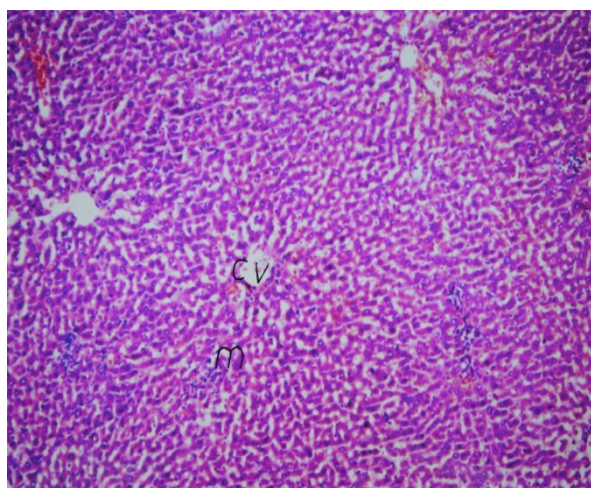


Figure 3. Photomicrograph of Liver Tissue Section of rat in HCC Group Treated with *R. vesicarius* Extract Showing Very Few Inflammatory Cells Infiltration in between the Hepatocyte (H&E x 16)

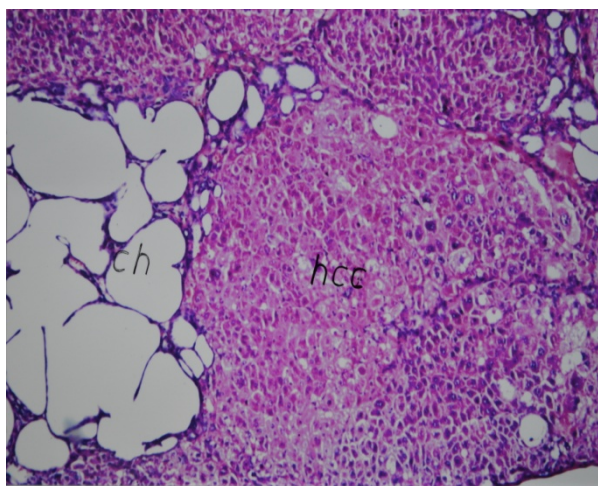


Figure 2. Photomicrograph of liver Tissue Section of rat in HCC Group Showing Foci of Anaplastic Hepatocellular Carcinoma with Other Foci of Cystic Cholangio Carcinoma (ch) (H&E x 16)

extract resulted in significant decrease (P<0.05) in serum CEA (-57.5%), AFP (-24.08%), AFU (-45.7%), glypican 3 (-25.02%), golgi 73 (-11.75%) and VEGF (-22.08%) levels as compared to the untreated HCC group (Table 2).

Histopathological findings

Photomicrograph of liver tissue section of rat in the negative control group showed no evidences for histopathological alteration and the normal structural organization of the central vein and the surrounding hepatocytes in the hepatic parenchyma is observed (Figure 1). Photomicrograph of liver tissue section of rat in HCC group revealed hepatic parenchyma with foci of anaplastic hepatocellular carcinoma as well as other foci of cystic cholangio carcinoma associated with areas of telangiectasis with haemorrhage. Also, individual hepatocellular necrosis is noted in the micrograph of a cross-sectioned liver tissue of rat in HCC group (Figure 2). Photomicrograph of liver tissue section of rat in HCC group treated with *R. vesicarius* extract showed no evidence of anaplasia but very few inflammatory cells infiltration in the hepatic parenchyma were observed (Figure 3).

Discussion

Chemical induction of liver carcinogenesis in the experimental animals initiated by the potent hepatocarcinogen diethylnitrosamine (NDEA) has been considered as one of the most accepted and widely used experimental model. This model closely mimics a subclass of human hepatocellular carcinoma (HCC) to study hepatocarcinogenesis and to allow screening of the potential anti-cancer agents on various phases of

neoplastic disease (Chakraborty et al., 2007). The present study results demonstrated that HCC group showed significant elevation in serum AST, ALT, ALP and γ -GT activity versus to the negative control group. Significant elevation of serum AST and ALT activities were seen in a variety of liver conditions, including viral infection, cirrhosis, non-alcoholic steatohepatitis (NASH), drug toxicity, liver tissue degeneration and necrosis (Yang et al., 2009). AST elevations often predominate in patients with cirrhosis and even in liver diseases that typically have increased ALT level (Green and Flamm, 2002). Concerning ALP, it has been found that ALP among liver function tests, in addition to other tumor characters is an independent factor for disease-free survival and overall survival (Tong et al., 2010). Recent studies have suggested that preoperative ALP levels could be utilized to monitor and predict recurrence in high risk HCC patients (Kim et al., 2013). Regarding γ -GT, it is well established that the elevated serum γ -GT activity is associated with diseases of the liver, biliary system, pancreas, and different types of cancers including HCC (Fentiman, 2012). Experimental studies have shown that γ -GT was strikingly activated during the course of hepatocarcinogenesis induced by several hepatocarcinogens in animals (Fiala and Fiala, 1973). Chemical carcinogens may initiate some systematic factors that induce γ -GT synthesis (Vanisree and Shyamaladevi, 1998). The elevated values of γ -GT reflect the progress of carcinogenesis, since its activity correlates with tumor growth rate, differentiation and survival of the host (Koss and Greengurd, 1982).

The current data indicated that treatment of HCC group with *R. vesicarius* extract elicited significant reduction in serum AST, ALT, ALP and γ -GT activity relative to the untreated HCC group. These results are in agreement with Chen (2010) and Saleem et al. (2014) who reported that *R. dentatus* significantly controlled the elevation of liver enzymes. This hepatoprotective action is possibly ascribed to the presence of many active constituents. *R. dentatus* active ingredients include chlorogenic acid, quercetin, myricetin, vitamin C and kaempferol. Among these isolated constituents, myricetin (Rashed et al., 2013), quercetin (Janbaz et al., 2004) and kaempferol (Song et al., 2003), have been reported to have hepatoprotective activity. Of particular interest, quercetin, the active compound of *R. vesicarius* offered a multimechanistic approach in its protective effect against liver injury induced by ethanol in rats (Chen, 2010).

In the current work, that HCC group showed significant elevation in serum CEA, AFP and AFU levels in comparison with the negative control group. CEA is an important tumor-associated antigen, and its overexpression has been used to identify or diagnose early colorectal, gastric, pancreatic, ovarian cancer and others (Ladd et al., 2009). Li et al. (2008) reported that when the rat liver tumor induced by DENA appeared, the CEA content in serum elevated. It has long been recognized that exposure of rats to certain carcinogens like NDEA causes an elevation of circulating AFP levels (Sell et al., 1983) and the reproduction of AFP in adults is associated with hepatocellular carcinoma. This finding is greatly supported by the study of Liu et al (2012), Ahmed et al.

(2013), and Song et al. (2013). The upregulation of AFP gene expression in NDEA-challenged rats might be due to necrosis of hepatocytes caused by NDEA. Hepatocyte localization within or outside the liver plate is the defining factor that regulates the activity of AFP synthesis on a cellular level (Lazarevich, 2000). Although the precise mechanism of this regulation is not fully understood but the possible explanation for the reinitiation of AFP synthesis by neoplastic hepatocytes includes either increased transcription of AFP gene or post-translational modification affecting AFP production (Motalleb et al., 2008).

Alpha L-fucosidase (AFU) activity in serum was found to be significantly increased in HCC group as shown in the present work. This result comes in line with the previous study of Chen et al. (2012). The exact mechanism behind the increased AFU in HCC is presently unknown, but the possible mechanism for that increase seems to be due to the increased synthesis of proteins by tumor with a consequent increase in fucose turnover (Deugnier, 1984). This hypothesis is supported by experimental data which revealed (i) elevated levels of fucose and fucosyltransferase activity in rat hepatoma tissue when compared to nontumor adjacent liver and (ii) disturbances in plasma membrane fucoderivative leading to an increase in fucoprotein levels (Vischer and Reutter, 1978) and (iii) the presence of abnormal fucoglycosides in hepatoma tissue (Holmes and Hakomori, 1982). One study showed the increased level of fucose in NDEA-treated rats when compared with control rats. (Balamurugan, 2011). Elevation in fucose content in the liver tissue of cancer patients have also been reported by some investigators and it has a considerable interest because of its potential application as diagnostic and/or prognostic marker. Biochemical studies have shown that changes in fucose content appears to be associated with tumor progression rather than with malignant transformation (Li et al., 1995).

In view of our data, the treatment of the HCC group with *R. vesicarius* extract led to significant decrease in serum CEA, AFP and AFU levels as compared to the untreated HCC group. Reporter assays indicated that plants belonging to genus *R. vesicarius* possessed antitumor activity for different tumor cell lines including colon, ovary, melanoma, breast, central nervous system and gastric cancer. These authors related this effect to the antiproliferative activity of these plants. The antiproliferative activity of *R. vesicarius* could be attributed to its content of polyphenols (Zhang et al., 2012) including quercetin and myricetin compounds. Quercetin is a potential anticancer, flavonoid molecule with ubiquitous nature. The suggested mechanisms for the chemopreventive action of quercetin include its capability to suppress cell proliferation and angiogenesis in cancer cells (Masuku et al., 2014). Moreover, quercetin was found to inhibit progression of human breast MCF-7 cancer cells through down regulation of proteins CDK2, cyclin A, D, E, p53 and p57 involved in cell cycle, resulting in the arrest of cell cycle. Also, quercetin has been reported to block cell cycle at G2/M through up-regulation of p21 and cyclin B to regulate cell cycle arrest at the G1 phase and G2/M phase in breast cancer cell lines (Moon et al., 2008). It was

also found that quercetin initiated apoptosis in cancer cells via the mitochondrial pathway involves the activation of caspase-3 downstream from caspase-9 (Aalinkeel et al., 2008). In addition, the antitumor efficacy of *R. vesicarius* is possibly ascribed to the synergies between quercetin and myricetin. Myricetin has been found to inhibit 7,12-dimethylbenz(a)-anthracene, benzo[a]pyrene and N-methyl-N-nitrosourea-induced tumor formation in a skin tumorigenesis model (Mukhtar et al., 1988).

In light of our data, HCC group showed significant elevation in serum glypican 3 (GPC 3) and golgi 73 (GP 73) levels in respect to the negative control group. Coston et al. (2008) demonstrated the sensitivity and specificity of GPC3 for HCC by 88% and 97%, respectively. GPC3 is a very specific marker not only for differentiating HCC from non hepatic tumors with epithelial differentiation, but also for differentiating HCC from hepatic adenoma (HA) and focal nodular hyperplasia (FNH). Golgiprotein-73 (GP73) is a resident Golgi glycoprotein expressed in epithelial human cells. GP73 serum levels are higher in early HCC patients than in cirrhotic patients. Serum GP73 is dramatically elevated in patients with HCC, and the sensitivity and specificity of GP73 for HCC might be superior to those of AFP (Willyard, 2007). It is considered as a possible tumor marker for HCC and indeed it shows a specificity of 75% and a sensitivity of 69% (Malaguarnera et al., 2010).

Treatment of the HCC group with *R. vesicarius* extract experienced significant decrease in serum GPC 3 and GP 73 serum levels relative to the untreated HCC group as shown in the present study. It is believed that oxidative stress plays critical roles in the initiation and progression of hepatocarcinogenesis. It has been hypothesized that polymorphisms that impair anti-oxidative capacity may influence HCC risk. Moreover, reactive oxygen species have been related to the aetiology of cancer as they are known to be mitogenic and therefore capable of tumour promotion. Thus, it is possible that cumulative defects in protection from oxidative stress may result in increased risk of liver cancer in the Moroccan population (Ezzikouri et al., 2010). *R. vesicarius* species have been studied as potential anticarcinogens because of their flavonoids content which stimulate apoptosis, inhibit proliferation and show antioxidant activity (Ramos, 2008).

In the present work, serum VEGF level of rats in HCC group showed significant increase which is in consistent with the results of Shahat et al. (2012). This finding indicated the high angiogenic activity in rats bearing HCC induced by NDEA. Earlier report by Jozkowicz and co-workers indicated the increased nitric oxide level in NDEA treated animals which enhances the angiogenesis by stimulating the synthesis of VEGF (Jozkowicz et al., 2001). NDEA administration has been found to increase nitric oxide synthase-2 (NOS2) activity indicating the generation of reactive oxygen species (ROS), promoting carcinogenesis and possibly leading towards angiogenesis. It seems that more advanced stage tumors actually express higher levels of VEGF protein. It has been shown that high expression levels of VEGF mRNA and protein in lung cancer patients is correlated with advanced lung cancer stage (Fontanini et al., 1997).

The present data demonstrated that the treatment of the HCC group with *R. vesicarius* extract produced significant reduction in serum VEGF level in comparison with the untreated HCC group. Accumulated evidences have suggested that some of *R. vesicarius* species could inhibit VEGF expression (Mirzoeva et al., 2008) and tumor angiogenesis *in vivo* (Fang et al., 2007). VEGF is highly stimulated by hypoxia-inducible factor 1 (HIF-1), a transcription factor that consists of two subunits, HIF-1a and HIF-1b (Powis, 2004). Several studies demonstrated that most flavonoids could inhibit HIF-1a and/or VEGF expression (Buchler et al., 2004; Fang et al., 2005; Fu et al., 2007) as well as VEGF-mediated angiogenic signalling in tumoral cells (Bagli et al., 2004). Therefore, it is assumed that flavonoids affected VEGF expression by reducing HIF-1a intracellular protein levels (Anso et al., 2010).

Photomicrograph of liver tissue section of HCC group showed hepatic parenchyma with foci of anaplastic hepatocellular carcinoma as well as other foci of cystic cholangio carcinoma associated with areas of telangiectasis with haemorrhage as well as individual hepatocellular necrosis. These remarkable features of hepatocellular carcinoma are in agreement with the studies of Abdallah and Khattab, (2004) and Seufi et al. (2009). It has been reported that histological examination of liver tissue of HCC group showed inflammatory cells infiltration and fibroblastic cells proliferation that divided the cancer and necrosed hepatocytes of the parenchyma into nodules with hyperchromatic nuclei as well as cellular pleomorphism and polarity (Seufi et al., 2009). Photomicrograph of liver tissue section of HCC group treated with *R. vesicarius* extract showed very few inflammatory cells infiltration in the hepatic parenchyma. In accordance of our results Saleem et al. (2014) reported that animals treated with aqueous methanolic extract of *R. dentatus* showed almost normal hepatocytes and normal structural organization of the liver.

The present setup provided experimental evidences that *R. vesicarius* possessed a promising anticancer potential against HCC induced in rats. This effect could be related to its hepatoprotective properties, potent antiangiogenic activity and effective antiproliferative potential as well.

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