Appraisal of circulating biochemical markers and antioxidative activity in interferon intolerant HCV patients

Abstract

Hepatitis C is small, single stranded, enveloped positive sense RNA viruses. It is member of Hepacivirus genus, family is Flaviviridae. Transmission route is intravenous drugs, blood transfusion etc. circulating biochemical marker are indicators of pathological utter in Hepatitis C virus. Decreased antioxidants and increased oxidants expose clear picture of pathology. Here this research work we will investigated the inter relationship between the HCV patients with and without interferon therapy. Interferon is a protein released animal cells in response to entrance of virus.

Keywords: biochemical markers, interferon, hepatitis C, detoxification, plasma protein production, hormone synthesis

Introduction

The liver is a vital organ which is present in vertebrates and some animals. It works as a wide range of functions, including detoxification, synthesis of protein, glycogen store room, breakdown of red blood cells, detoxification, plasma protein production, hormone synthesis and manufacture of biochemical’s required for digestion. Hepatitis C is the infectious disease which affects the causative agent is hepatitis C virus. The infection may be in some time asymptomatic but chronic infection ultimately changed to cirrhosis, which is commonly parent apparent after some years. In some cases cirrhosis changed to liver cancer, liver failure, esophageal, gastric varices.1 Liver cirrhosis caused different complications which are jaundice, bleeding, as cites, portal hypertension, varices, and hepatic encephalopathy. These are the most common causes of the liver transplant.2 Hepatitis C is commonly associated with the diabetes mellitus, thrombocytopenia, lymphoproliferative disorders, lichen planus and sjogren syndrome (an autoimmune syndrome). It is estimated that thrombocytopenia is present in 0.16- 45.4% of the people in hepatitis C virus. Hepatitis C is also linked with the cryoglobulinemia which is the inflammation of small and medium size blood vessels caused the deposition of immune complexes in which cry globulins are involved. Hepatitis C is the small, single-stranded, enveloped, positive sense RNA virus. Its family is Flaviviridae. Hepatitis C has seven major genotypes which represented as one to seven. Approximately 70% cases are caused by genotype 1, 20% genotype 2, 1% each of the other genotypes. Genotype 1 is most commonly found in Europe.3 Treatment consists of pegylated interferon and ribivir for the period of 24-48 weeks, depending upon the HCV genotype.4 Cirrhosis is the result of damaging of the liver cells from toxins, inflammation metabolic derangement and others chronic liver diseases, such as hemochromatosis, caused the inability of liver to perform its biochemical function. Its infection occur due to the long -lasting use of the alcohol, hepatitis B, hepatitis C Piekarska & Matusiak5 hepatocellular carcinoma is a type of cancer which damage the hepatocytes that is major cell type of the liver. Throughout the world it is considered the number one or number two cause of cancer death. Chronic infection which are HBV and HCV are increased the risk of hepatocellular carcinoma. About 80% people who suffer in hepatocellular carcinoma have cirrhosis.6

Endogenous oxidants are generated by multiple intracellular pathways and are important class of naturally occurring carcinogens. During aerobic metabolism ROS and endogenous oxygen containing molecules formed as normal product during aerobic metabolism.7 ROS include the number of species such as superoxide, hydroxyl, and peroxyl radicals and certain non radical such as single oxygen and hydrogen peroxide that may be easily converted into radicals. ROS caused the chromosomal alteration and genetic mutation and contribute to cancer development in multi step carcinogenesis.8 A study demonstrated ROS at submicromolar level act as novel intra and intercellular secondary messengers and modulates various aspects of cellular function including gene expression, apoptosis and proliferation.9 Mostly markers of oxidative injury used reflect free radicals attack on polyunsaturated fatty acids with the classical route of attack involving lipid peroxidation which generates end peroxides and hydro peroxides.

Moreover it has been showed that patient with chronic hepatitis C virus increased the production of the tumor necrosis factor-Alpha .Tumor necrosis factor-Alpha is a cytokine that can produced the oxidative stress by stimulating the generation of reactive oxygen species such as hydrogen peroxide ion(H2O2) and super oxide ion(O2-). ROS can damage cells and oxidative damage of DNA, proteins, and by depleting ATP stores. Metals such as Fe2+, O2- react with H2O2 to generate a hydroxyl radical that changed to more reactive and cytotoxic than O2- or H2O2.10 Free radicals work as a scavenger to the several metalloenzymes such as, glutathione peroxidase, catalase, and superoxide dismutase as well as by the non-enzymatic antioxidant defense system such as vitamin C, lipoic Acid, ubiquinol, glutathione, metallothionein, Bilirubin, beta-carotene. That is the reason, much attention of nutritionists have focused to the possible role of enhancement of the defenses against ROS.11 Reactive nitrogen species (RNS) are generated because of the production of the ROS caused the loss of the antioxidant defense. ROS/RNS are usually too reactive and have a very short half-life period may be much shorter than second therefore allow direct measurement in body fluids, tissues and cell. It is also used in measuring stable metabolites e.g. nitrate and nitrite.12,13

Malondialdehyde (MDA) is a highly reactive three carbon compound which is also called di aldehyde. It produced as a
byproduct of polyunsaturated fatty acid peroxidation and during metabolism of arachidonic for synthesis of prostaglandin. MDA have ability to combine with several functional groups on molecules including protein, lipoprotein, RNA and DNA. MDA level are used for the important indicator of lipid peroxidation in vitro and vivo for various diseases. To check the cellular injury lipid peroxidation is a well established mechanism of cellular injury in both animals and plants and it is also used for indication the oxidative stress in tissues and cells. It is reported that MDA induced in various conditions such as hepatitis C infection, smoking, diabetes and HIV seropositive children.

Many reports indicated that glutathione plays an important role in defence against the reactive oxygen species. Cellular enzymes e.g; glutathione peroxidase and related enzymes glutathione s-transferase and glutathione reductase provides the principle mechanism against exogenous endogenous and exogenous free radicals and toxic substances damage the liver and other tissues. Glutathione is a naturally occurring antioxidant and it plays important role against the oxidative stress to reduce the disulphide linkages of the protein by scavenging hydro peroxides and lipid peroxide. Glutathione found both in reduced and oxidized form under the normal condition reduced form is 99.5% and oxidized form is 0.5% which may be cytotoxic. Glutathione is the detoxification enzyme which catalyzes the addition of the glutathione into various xenobiotic. Glutathione-α has high concentration in the liver cell hepatocytes crystal.

According to the experimental model it has been proposed that oxidative stress may lead to neoplastic transformation. Therefore presence of primary in the course of cirrhosis of HBV, HCV or other eithology may be associated with disturbances of organism antioxidant barrier.

The amount of SOD and GSH-Px reduced in liver which indicates decreased the capacity of liver and the antioxidant defense power of the patient with chronic hepatitis C. Researchers proved that lipid peroxidation increased due to viral inflammation and decreased the antioxidant level may be an early marker of the oxidative stress. Lipid peroxide can be chemotactic for the neutrophils causing increased inflammation which further increased the oxidative injury in the liver. Peripheral lymphocytes are also infected to the viral infection in the hepatitis C. lymphocytes which are infected to the hepatitis C produced the interferon to stimulate healthy cells against viruses. The mechanism through which HCV causes cell damage remain obscure, although it is claimed that oxidative stress may play pathogenic role in the infection. Hepatitis C patient exhibit increased production of TNF-α, a cytokine that can produced oxidative stress by stimulating the production of oxygen ROS. No vaccine against hepatitis C is available. IFNs belong to the large class of glycoproteins known as cytokines. Interferons are named after their ability to “interfere” with viral replication within host cells. IFNs have other functions: they activate immune cells, such as natural killer cells and macrophages; they increase recognition of infection or tumor cells by up-regulating antigen presentation to T lymphocytes; and they increase the ability of uninfected host cells to resist new infection by virus. Certain symptoms, such as aching muscles and fever, are related to the production of IFNs during infection. About ten distinct IFNs have been identified in mammals; seven of these have been described for humans. They are typically divided among three IFN classes: Type I IFN, Type II IFN, and Type III IFN. IFNs belonging to all IFN classes are very important for fighting viral infections. Alpha interferon has been used since 1980s in the chronic hepatitis C which are given as subcutaneous injection. Interferon and addition of oral ribiviron to them has brought their rate of long term viral clearance to them from <5% to now approximately who have never been received and 50% in those patients who never been received treatment. However side effects of the interferon are universal. This lead to the modifications of the dosage of the interferon or ribiviron in 35-45% patients of the patient treated with pegylated interferon in a large, randomized clinical trials and discontinuation of therapy in 14-19% of the patients. In each trial fatigue is the major side effect of the interferon and ribiviron. It is partly a direct side effect of interferon-alpha with stimulation of immune response to release cytokines and other factors that causes fatigue. This is partly related to anemia caused by ribiviron. Ribiviron is toxic to RBC because it directly breaks down the red blood cell, giving the median decline in hemoglobin approximately 3g/dl. Depression is also a major side effect which is 20-35% of patients treated with interferon and ribiviron. This can be one of the major morbidities related with treatment and practitioner screening for the development of suicidal and other ideas are essential when patient develops symptoms of depression. Patient prior history of depression or other mental illness by mental health professionals, with appropriate treatment before treatment is very important. Serotonin reuptake inhibitor (SSRIs) is found to be effective for the treatment of the depression associated with interferon therapy.

Ribiviron causes dosage dependent hemolytic anemia but interferon can stop red blood cell production in bone marrow, this result anemia. Likely >20% of patients are treated with ribiviron and pegylated interferon at 1000-1200mg/day. Anemia can be a most clinically significant side effect of the therapy and patients with condition that can be worse by anemia such as chronic obstructive pulmonary disease and coronary artery disease. Patient who is coinfected with HIV may be more susceptible to developing anemia due to treatment. Ribiviron dosage reduction should be the first step to manage the symptomatic anemia. The efficacy of ribiviron may be dosage related. Therefore doses should be maintained at 800mg/ day or more if possible. Manufacturer recommended reducing ribiviron dosage (to 600mg/day for Capegous and 200mg/day for Rebeto) with hemoglobin <10g/dl and discontinuing ribavirin with hemoglobin>8.5g/dl. Interferon affects the tongue and caused dysfunctioning of taste bud cells, killing taste bud entirely.

Materials and methods

Biochemical assays for liver function tests

One milliliter blood sample were taken and subjected to centrifuge at 3000-4000rpm for 10-15minutes for the separation of serum. The estimation of Aspartate Amino Transferase (AST), ALT Rietman and Frankle, ALP, Ochoa, was estimated by using commercially available Bio Merux and Randox kits and following their protocol.

Estimation of aspartate amino transferase (AST)

α - oxoglutarate reacts with L- aspartate in the presence of AST to form L glutamate plus oxaloacetate. The indicator reaction utilizes the oxaloacetate for a kinetic determination of NADH consumption

Anonymous

α -oxoglutarate + L- aspartate AST L - glutamate + oxaloacetate

Oxaloacetate + NADH + H+ MDL H - malate + NAD+

One milliliter reagent was taken in all the tubes by pipette and 100 µl of each serum sample was taken in tube. Contents of tube was mixed and then incubated at 37°C for 5min. after incubation 250µl of substrate from the kit was added to all the tubes. The contents were again mixed for 1.0 min and then absorbance of sample was measured after 1, 2 and 3 minutes at 404 nm Anonymous. The concentration of
AST was estimated by applying the following formula:
\[
\Delta A/\text{min} \times 1745 \text{ (at 37°C)} \text{ where } \Delta A = \text{ Change in absorbance}
\]

While Reference values for men and women were 15-40IU/L and 13-35IU/L respectively.

**Estimation of alanine aminotransferase (ALT)**

ALT converts \(\alpha\)-oxoglutarate and L-alanine to L-glutamate and pyruvate. This pyruvate is further converted to L-lactate and NADH to NAD\(^+\) Anonymous. The concentration of an unknown (sample) is determined by calculating from the linear equation generated from the rate of absorbance measured. The rate of absorbance measured represents the sum of reduced and oxidized glutathione in 412nm and oxidized glutathione GSSG. The amount of glutathione or DTNB) to produce a chromophore TNB with maximal absorbance at 412nm.

GSH reacts with Elman’s reagent (5, 5-dithio bis (Nitro benzoic acid) to produce a chromophore TNB with maximal absorbance 0.0167 at 37°C. The reaction mixture contained 0.01M phosphate buffer (pH 7.0) 2mM H\(_2\)O\(_2\), and the enzyme extract. The specific activity of Catalase was expressed in terms of units/gram of liver tissue. Absorbance values were compared with a standard curve generated from known Catalase.

\[
Y = -0.0065x + 0.1595
\]

\[
R^2 = 0.9934
\]

\[
X = \text{Concentration of GSH}
\]

\[
Y = \text{Absorbance of Sample}
\]

**Estimation of alkaline phosphate (ALP)**

In the presence of ALP p-nitrophenylphosphate is hydrated to p-nitrophenol and phosphate Anonymous. P-nitrophenylphosphate + H\(_2\)O ALP phosphate + p-nitrophenol

One milliliter reagent was taken in all the tubes by pipette and 100\(\mu\)l of each serum sample was taken in tube. Contents of tube was mixed and then incubated at 37°C for 5 min. after incubation 250\(\mu\)l of substrate from the kit was added to all the tubes. The contents were again mixed for 1.0 min and then absorbance of sample was measured after 1, 2 and 3 minutes at 404nm Anonymous. The concentration of ALP was estimated by applying the following formula:

\[
\Delta A/\text{min} \times 3433 \text{ (at 37°C)} \text{ where } \Delta A = \text{ Change in absorbance}
\]

While reference values for men and women were 10-40IU/L and 07-35IU/L respectively.

**Estimation of glutathione (GSH)**

Glutathione (GSH) was estimated according to the method of Moron et al. GSH reacts with Elman’s reagent (5, 5-dithio bis (Nitro benzoic acid) or DTNB) to produce a chromophore TNB with maximal absorbance at 412nm and oxidized glutathione GSSG. The amount of glutathione measured represents the sum of reduced and oxidized glutathione in the sample ([GSH] \(t = [GSH] + 2 \times [GSSG])\). The rate of absorbance change (\(\Delta A 412\text{nm/min})\) is made to be linear for the convenience and consistence of measurement, and is linearly proportional to the total concentration of GSH. The concentration of an unknown (sample) is determined by calculating from the linear equation generated from several standards of glutathione Tetze.

\[
Y = 0.0167 \times X - 0.0098
\]

**Estimation of catalase (CAT)**

Catalase was assayed according to the method of Aebi. The estimation was done using spectrophotometer by checking the decrease in absorbance at 230nm. The liver tissue was homogenized in M/150 phosphate buffer (pH 7.0) at 1-4°C and centrifuged at 5000rpm. The reaction mixture contained 0.01M phosphate buffer (pH 7.0) 2mM H\(_2\)O\(_2\), and the enzyme extract. The specific activity of Catalase was expressed in terms of units/gram of liver tissue. Absorbance values were compared with a standard curve generated from known Catalase.

\[
Y = -0.0065x + 0.1595
\]

\[
R^2 = 0.9934
\]

\[
X = \text{Concentration of CAT}
\]

\[
Y = \text{Absorbance of sample}
\]

**Estimation of thiobarbituric acid reactive substances (TBARS)**

Lipid peroxidation in liver tissues was estimated calorimetrically by measuring Thiobarbituric acid reactive substances (TBARS) by the method of Okawa. To 0.2ml of sample, 0.2ml of 8.1% Sodium dodecyl sulfate (SDS), 1.5ml of 20% acetic acid and 1.5ml of 0.8% TBA were added. After centrifugation at 3000rpm for 10 min the upper organic layer was taken and its OD was read at 532nm against an appropriate blank without the sample. The levels of lipid peroxides were expressed as millimoles of Thiobarbituric acid reactive substances (TBARS)/g of liver tissue using standard curve.

\[
Y = -0.129X + 0.592
\]

\[
X = 0.129 - Y
\]

\[
R^2 = 0.8952
\]

\[
X = \text{Concentration of CAT}
\]

\[
Y = \text{Absorbance of sample}
\]

**Estimation of Super Oxide Dismutase (SOD)**

Superoxide dismutase (SOD) activity was determined by the method of Kakkar. The assay mixture contained 0.1ml of sample, 1.2ml of sodium pyrophosphate buffer (pH 8.3, 0.052 M), 0.1ml of Phenazine methosulphate (186μm), 0.3ml of nitro blue tetrazolium (300μm), 0.2ml of NADH (750 mile). Reaction was started by addition of NADH. After incubation at 300C for 90sec, the reaction was stopped by the addition of 0.1ml of glacial acetic acid. The reaction mixture was stirred vigorously with 4.0ml of n-butanol. The mixture was allowed to stand for 10 min; centrifuged and n-butanol layer was separated. The color intensity of the chromogen in butanol

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layer was measured at 560nm against n-butanol and concentration of SOD was expressed as units/g of liver tissue. Absorbance values were compared with a standard curve generated from known SOD.

\[
\begin{align*}
Y &= 0.0162x + 0.0161 \\
X &= 0.0162 - Y \\
R^2 &= 0.889 \\
Y &= \text{Absorbance of the sample} \\
X &= \text{Concentration of SOD}
\end{align*}
\]

**Estimation of zinc**

Zinc present in the sample is chelated by 5-Br-PAPS 2-(5-bromo-2-pyridylazo)-5-(N-propyl-N-sulfopropylamino)-phenol in the reagent. The formation of this complex is measured at the wavelength of 560nm. Use all glassware were immersed in dilute HCl or dilute HNO₃ and then rinsed in DDH₂O. Special care was taken as rubber cap of commercial control serum or sample tube can be a cause of zinc contamination. Sample was properly mixed by vortex after the addition of deproteinizing reagent to sample. Reference value was taken as 10-17µmol/L.

\[
\begin{align*}
\text{Sample Conc} &= \text{Standard Conc} \\
A_{\text{sample blank}} &= A_{\text{standard blank}} \\
X &= \frac{A_{\text{sample}} - A_{\text{blank}}}{A_{\text{standard}} - A_{\text{blank}}} \\
Y &= \frac{X}{0.161}
\end{align*}
\]

**Statistical analysis**

Results have been expressed as Mean±SD (Standard Deviation). Statistical significance was determined by one way analysis of variance and Pearson correlation (Two Tailed) was used to correlate the different variables. The difference were considered significant at p<0.05.

**Results**

Hematological Profiles of Hepatitis C Patients Receiving Interferon Vs Interferon Intolerant HCV Subjects The results regarding hemoglobin and RBC profile in Figure 1 demonstrating statistically highly significant differences and consistent decreasing (hemoglobin) and increasing Vitamin E (Figure 1) in the interferon intolerant HCV subjects (P=0.04). The lowest value (10.42%) of hemoglobin was recorded in interferon subjects in group C and highest (15.37%) in group A (served as control). Vitamin E is a highly effective fat-soluble vitamin with a variety of cellular membrane stabilizing-antioxidant and non-antioxidant functions. Vitamin E has been suggested to prevent the oxidation of polyunsaturated fatty acids in red blood cell (RBC) membrane, thus inhibiting the premature erythrocyte lysis. Animal studies have shown that treatment with vitamin E results in increased number of colony forming units of erytroid precursors, enhanced erythropoiesis and improved blood hemoglobin levels in these animals.³²

Vitamins Profiles of Hepatitis C Patients Receiving Interferon Vs Interferon Intolerant HCV Subjects Vitamin A shows the highly significant difference (p=0.001) between the Vitamin B1. Lowest value of vitamin A is (0.63%) in interferon receiving patients and highest value is (1.02%) which is present in control served. HBV or HCV infection enhanced oxidative stress and lowered vitamin B in circulation. In order to avoid other healthy risk, nutrition status should be monitored and limitations or supplementation of certain nutrients might be helpful HBV or HCV infected patients.

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Lipid Profiles of Hepatitis C Patients Receiving Interferon Vs Interferon Intolerant HCV Subjects

The results depicted in Figure 3 reflecting that the lipid profile Total Cholesterol (TCh) and Triglyceride (Tg) shows differed significantly (P=0.002 and p=0.01 respectively) with the glutathione (GSH) Figure 4. The lowest value (4.26%) of Total Cholesterol was recorded in normal patients in group A and highest value (5.89%) was observed in interferon subjected patients in group C. In the triglyceride (TG) lowest value (1.17%) of was recorded in normal group A and highest value (1.79%) was observed in interferon intolerance HCV subjects in group B. The cumulative incidence rates of HCC were 24%, 51%, and 63% at 5 years, 10 years, and 15 years, respectively. Multivariate analysis identified hepatic steatosis, together with aging, cirrhosis, and no IFN treatment, as independent and significant risk factors for HCC (P = 0.0135, P = 0.0390, P = 0.0068, and P = 0.0142, respectively). In addition, hepatic steatosis was correlated with BMI, serum ALT levels, and triglyceride levels. Kazuyuki Ohata.

Vitamin E is a highly effective fat-soluble vitamin with a variety of cellular membrane stabilizing-antioxidant and non-antioxidant functions. When we look the hematological profile in which red blood cells and hemoglobin have lower values in interferon subjected HCV patients as compared to the interferon intolerance and control group patients? This may be due to the increased breakdown of the red blood cells and therefore decreased level of the RBC and Hb.

Vitamin E has been suggested to prevent the oxidation of polyunsaturated fatty acids in red blood cell (RBC) membrane, thus inhibiting the premature erythrocytolsis. Animal studies have shown that treatment with vitamin E results in increased number of colony forming units of elytroid precursors, enhanced erythropoiesis and improved blood hemoglobin levels in these animals.

AST is negatively correlated with the LDL, if AST level increases then LDL level decreases. Alkaline phosphates (ALP) exhibits highly significant (P=.03 respectively) differences between Vitamin E. The lowest value of aspartate aminotransferase (204.88%) was recorded in normal patients in group A and highest value (387.34%) was observed in interferon subjected patients in group C. The hepatic profile showed the highest values of AST, ALT, ALP and total bilirubin in the interferon subjected patients as compared to the interferon intolerance patients. We give interferon to the patients for cure purpose but it also have opposite results. Some research proved that interferon is itself Hepatotoxicity therefore it will damage the hepatocytes and...
the enzyme AST present in the cell membrane and the ALT and ALP present in cytoplasmic blood that indicates the hepatocyte damage.

An association of hepatitis C virus (HCV) with low-density lipoproteins (LDL) in serum of patients with chronic hepatitis C (CHC) has been suggested. We conducted a prospective study in CHC patients complicated with hyperlipidemia, to examine whether bezafibrate, which is commonly used for treatment of hyperlipidemia, reduces serum HCV-RNA titer and improves liver dysfunction. Fifteen patients received daily oral bezafibrate treatment (400 mg/day) for 8 weeks, and its effects on serum lipids, transaminases, HCV-RNA titers, and HCV-RNA titers bound to LDL were evaluated. Fifteen untreated patients with CHC and hyperlipidemia were used as controls.29 Triglyceride (TG) showed significantly with the glutathione (GSH). When we compare the lipid profile parameters the total cholesterol level is very high in interferon subjects as compared to the interferon intolerance subjects patients of HCV. Different research approved that cholesterol increased the proliferation of the hepatitis C virus. Therefore those individuals which have high level of lipids have more chances of HCV infection. Likewise low density lipoprotein which is harmful to the body was found at high level in the interferon subjects positive patients as compared to the interferon intolerance HCV patients. In contrast to its high density lipoprotein which is beneficial to the body, showed low level in interferon positive patients as compared to the interferon intolerance patients?

The reports from several studies have produced clear evidence that there exists a good correlation between type and severity of disease and antioxidant level in the blood.30 Although the main role of the immunological mechanisms in pathogenesis of the chronic viral hepatitis B and C was demonstrated, researchers also concentrate on the problem of oxidative stress in the pathology of the diseases. Oxidative stress develops when the disturbances between reactive oxygen forms are produced in excess and the factors preventing their harmful effects occur. Enzymatic antioxidant defense of the organism includes: SOD, CAT, and GSH-Px. Superoxide dismutase protects a cell from toxic effect of superoxide radicals as it catalyzes the dismutation reaction of the radicals.31 Glutathione peroxidase decomposes hydrogen peroxide but it also converts lipid peroxides to harmless molecules protecting the cells from the consequences of lipid peroxidation. GSH-Px removes \( \text{H}_2\text{O}_2 \) by the oxidation of reduced glutathione. Oxidized glutathione (GSSG) is produced and it is reduced again by glutathione reductase, and the NADPH (produced in pentose cycle).32 Oxidative stress has been detected in almost all clinical and experimental conditions of the chronic liver diseases.3334 There are many studies about the oxidant stress in chronic hepatitis C patients.35 It showed that MDA, a product of polyunsaturated fatty acid peroxidation, was elevated in the liver and the blood.36 Also demonstrated MDA-protein adducts immunohistochemically in infected liver tissues,9,36 showed that the peripheral blood mononuclear cells from patient of chronic hepatitis C had increased MDA concentrations, and enhanced SOD activity. MDA is reflection of lipid peroxidation and SOD is an important antioxidant defense enzyme that converts superoxide into hydrogen peroxide. Increased SOD activity appears to be an adaptive response to increased generation of the superoxide ions,40 showed higher serum malondialdehyde values in chronic hepatitis C patients than healthy subjects before the interferon treatment.

Current study showed that serum MDA has significant negative correlation to the glutathione in HCV patients receiving interferon. If the level of the MDA is increased then the level of the glutathione decreases and vice versa. The results presented confirm the involvement of the oxidative stress as a part of pathophysiology of HCV. Our findings support the existence of the oxidative stress in patients with chronic HCV infection. When we look the overall result of the stress biomarkers, malondialdehyde is produced as the byproduct of the lipid peroxidation. Therefore MDA level is increased in interferon positive patients as compared to the interferon intolerance patients. MDA is the cytotoxic metabolites which damage the DNA, protein and lipids leading to the cell death. Other stress biomarkers which are catalase, glutathione, and superoxide dismutase are antioxidants which are used to stop the lipid peroxidation, have low level in the interferon receiving HCV patients as compared to the interferon intolerance and control patients. This may be due to the disturbance of the oxidants and antioxidants in the infected cells.

Glutathione (GSH), the most abundant non-enzymatic antioxidant present in the cell, plays an important role in the defense against oxidative-stress-induced cell injury. In the cells glutathione is present mainly in its reduced form. Reduced GSH can be converted to oxidized glutathione (GSSG) with the glutathione reductase (GR). Cells are also equipped with the enzymatic antioxidant mechanisms that play an important role in the elimination of free radicals.38 Suarez studied the group of 100 individuals with chronic hepatitis C. They demonstrated GSH decrease in the blood serum of patients who were not treated with interferon. In contrast, GSH concentration was higher in the blood of patients responding satisfactorily to the treatment. The author set forth a theory explaining the cytopathic effect of a virus causing glutathione deficiency. Kramer studied antioxidant enzymes and they found diminished GSH-Px level in the blood serum and the erythrocytes, in patients with abnormal liver function.39,40 Pak also demonstrated low activities of GSH-Px and MDA in adults with acute hepatitis B. Results of our study are consistent with the reports by Suarez and associates. In the current study we demonstrated that glutathione peroxidase level and superoxide dismutase level are decreased in erythrocytes of patients with chronic hepatitis C. It probably decreased the antioxidant barrier efficiency in studied CHC patients. When the activity of the enzymes mentioned above is insufficient, an organism is not capable to neutralize free oxygen radicals that are produced in excess. In consequence, it leads to hepatocyte lesion.45 They demonstrated that GSH and catalase levels decreased both in group of children with chronic hepatitis C, and B. Kramer and associates et al, studied the small group of children with abnormal function of liver cell, and demonstrated decreased activity of SOD in comparison to the healthy children. Loginov et al. studied antioxidant system in adults with chronic active hepatitis, and demonstrated SOD decrease correlating with severity of inflammatory process. Study by Yasuyama showed decrease of SOD levels in liver tissues of patients with acute10 and chronic hepatitis accompanied by fatty degeneration while comparing with patients with the liver inflammatory diseases of different etiology. In another study Irshad et al.,42 found SOD activity significantly low in the chronic active hepatitis C patients.

Conclusion

My great finding in these results is that zinc (Zn) shows the lower level in interferon intolerance patients as compared to the interferon positive patients and control groups. And this might be reason to the intolerance of the interferon. According to my finding if a person who is intolerant to the interferon should be submitted to zinc along with the interferon therapy.

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Conflict of interest

The authors declare that there is no conflict of interest.

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