

Protective Effects of N-acetylcysteine Against the Statins Cytotoxicity in Freshly Isolated Rat Hepatocytes

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ABSTRACT

Purpose: Hepatotoxicity is one of the most important side effects of the statins therapy as lipid-lowering agents. However, the mechanism(s) of hepatotoxicity induced by these drugs is not clearly understood yet, and no hepatoprotective agent has been developed against this complication.

Methods: The protective effect of N-acetylcysteine (NAC) against statins-induced cytotoxicity was evaluated by using freshly isolated rat hepatocytes. Hepatocytes were prepared by the method of collagenase enzyme perfusion via portal vein. This technique is based on liver perfusion with collagenase after removal of calcium ion (Ca²⁺) with a chelator (ethylene glycol tetra acetic acid (EGTA) 0.5 mM). The level of parameters such as cell death, ROS formation, lipid peroxidation, mitochondrial membrane potential (MMP) in the statins-treated hepatocytes were determined. Additionally, the mentioned markers were assessed in presence of NAC.

Results: Incubation of hepatocytes by the statins resulted in cytotoxicity characterized by an elevation in cell death, increasing ROS generation and consequently lipid peroxidation and impairment of mitochondrial function. Administration of NAC caused reduction in amount of ROS formation, lipid peroxidation and finally, cell viability and mitochondrial membrane potential (MMP) were improved.

Conclusion: This study confirms that oxidative stress and consequently mitochondrial dysfunction is one of the mechanisms underlying the statins-induced liver injury and treating hepatocytes by NAC (200 μM) attenuates cytotoxicity.

Introduction

Statins (3-hydroxy-3-methylglutaryl co-enzyme A reductase inhibitors) are the most prescribed agents for cholesterol lowering and consequently prevention of obstructive cardiovascular events in the world.^{1,2} These pharmaceuticals block the mevalonic acid pathway by inhibition of the rate limiting step in the hepatic *de novo* cholesterol biosynthesis.³ Severe adverse effects of the statins involving myopathy and hepatotoxicity sometimes limit their usage as lipid lowering agents. Hepatotoxicity characterized by elevation of plasma transaminases is most common and the frequency of elevation is 0.5-5% and dose dependent.^{4,5} However the precise molecular mechanisms underlying statins cause hepatotoxicity are not understood.² Statins are metabolized by cytochrome P450 (3A4) enzymes.¹ The dose- and time-dependent impairment of mitochondrial function caused by statins has been observed in different *in vitro* models.^{6,7}

Different antioxidants administration has been shown to be useful in reducing oxidative stress and cell death in hepatocytes. Efficacy of N-acetylcysteine (NAC) administration as a free radical scavenger in multiple models of oxidative stress induction has been studied.⁸ In the present study, isolated rat hepatocytes were exploited to gain insight into statins hepatotoxicity and to investigate the role of supplementation with NAC in reducing toxicity. Markers like reactive oxygen species (ROS), lipid peroxidation, mitochondrial membrane potential and cell death in the statins-treated freshly isolated rat hepatocytes were assessed and the results were compared to those in NAC-treated hepatocytes. The hepatoprotective properties of NAC(200 μM) supplementation against the statins-induced hepatotoxicity were related to the reduction of ROS generation, lipid peroxidation and maintenance of mitochondrial membrane potential.

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Material and Methods

Chemicals

Atorvastatin, Simvastatin and Lovastatin were purchased from sigma-aldrich. NAC was provided from Acros (New Jersey, USA). 2-vinyl pyridine, Triethanolamine and (4-(2-hydroxyethyl) 1-piperazine-ethanesulfonic acid (HEPES) were obtained from Acros (New Jersey, USA). Albumine bovine type was purchased from Roche diagnostic corporation (Indianapolis USA). Rhodamine 123, 5,5'-dithio-bis(2-nitro-benzoic acid)(DTNB), 2,7-Dichlorofluorescein diacetate (DCFDA) and clostridium histolyticum extracted Collagenase (Type II), were obtained from Sigma Aldrich (St. Louis, USA). Trichloroacetic acid (TCA), Ethyleneglycol-bis (ρ -aminoethylether)-N,N,N',N'-tetra acetic acid (EGTA), and Trypan blue were obtained from Merck (Darmstadt, Germany). Thiobarbituric acid (TBA) was obtained from SERVA (Heidenberg, New York). All salts used for preparing buffer solutions were of analytical grade and obtained from Merck (Darmstadt, Germany).

Animals

Male Sprague–Dawley rats (weight range: 250-300g) were provided from Tabriz University of Medical Sciences, Tabriz, Iran. The animals were housed in an animal house unit (temperature 21-23°C and 50-60% relative humidity, fed a standard chow diet and water *ad libitum* were used. The animals were handled and used according to the animal handling protocol of Tabriz University of Medical Sciences, Tabriz, Iran which was approved by a local ethic committee.

Hepatocytes isolation

Hepatocytes were isolated from male Sprague–Dawley rats by a two-step collagenase perfusion as described previously.⁹ Approximately 85–90% of hepatocytes excluded Trypan blue (0.1%, w/v) at the time of isolation. The cells were suspended (1×10^6 cell/ml) in Krebs–Henseleit buffer containing 12.5 mM HEPES and incubated under a stream of 95% O₂ and 5% CO₂ in continuously rotating round-bottomed 50 ml flasks at 37°C water bath. Hepatocytes were kept under the relevant atmosphere for 30 minute to achieve equilibrium between gas and liquid phases before the addition of chemicals.

Cell viability

After hepatocyte isolation process, cell viability was assessed by the extent of plasma membrane intactness as determined by Trypan blue exclusion test.¹⁰ Viability was determined immediately after isolation and after one, two and three hours after statins incubation. In experiments NAC (200 μ M) and statins were added contemporarily (There were no significant difference between before and co-adding of chemicals).

Reactive oxygen species (ROS) formation

To assess the rate of hepatocytes ROS formation during statins metabolism, 2,7-dichlorofluorescein diacetate (DCF-DA) 1.6 μ M was added to the hepatocyte incubate. DCFH-DA became hydrolyzed to non-fluorescent dichlorofluorescein (DCFH) in hepatocytes. Dichlorofluorescein then reacted with reactive oxygen species to form the highly fluorescent dichlorofluorescein. One ml (10^6 cells) of hepatocytes was taken and the fluorescence intensity was measured using a Jasco® FP-750 spectrofluorometer with excitation and emission wavelengths of 500 and 520 nm, respectively.¹¹

Lipid peroxidation

Hepatocyte lipid peroxidation was determined by measuring the amount of thiobarbituric acid-reactive substances (TBARS) such as malondialdehyde (MDA), formed during the decomposition of lipid hydroperoxides by following the absorbance at 532 nm in a Pharmacia Biotech Ultrospec 2000 spectrophotometer after treating a 1.0 ml aliquot of hepatocyte suspension (10^6 cells/ml) with trichloroacetic acid (70%, w/v) and boiling the supernatant with thiobarbituric acid (0.8%, w/v) for 20 min.¹²

Mitochondrial membrane potential assay

Mitochondrial membrane potential (MMP) in hepatocytes was assessed by monitoring the uptake of rhodamine 123, a cationic dye, as described.^{13,14} Isolated cells were extracted, and then resuspended in original media containing 1.5 μ M rhodamine 123. After 10 min of incubation, the cells were centrifuged and the supernatant was measured with a Jasco FP-750 spectrofluorimeter. The amount of dye remaining in the supernatant was inversely proportional to the membrane potential of the cells. The results are reported as the difference in fluorescence intensity between control and treated cells and expressed as percentage of control.

Statistical analysis

The results are shown as mean \pm SD, measured at least for 3 different batches of hepatocytes. The differences among the control and experimental groups were determined by one way ANOVA followed by Tukey's post hoc analysis and a $p < 0.05$ was considered significant.

Results

By adding different concentrations of atorvastatin, simvastatin and lovastatin to isolated rat hepatocytes and by use of trypan blue exclusion test, the LC₅₀s of statins (the statin concentration that caused 50% death after 120 minutes) were found as 450 μ M for atorvastatin, simvastatin 200 μ M and lovastatin 200 μ M (data not shown). As shown in Table 1, simvastatin 200 μ M is the most cytotoxic statin, because the amount of dead cell in third hour is more noticeable. Additionally, an optimum effective dose for NAC that provided an appropriate protection was found as 200 μ M. Hepatocytes were treated

with NAC at the same time that statins were added. It was found that NAC (200 μ M) effectively prevented cell death induced by the statins (Table 1). Markers such as ROS formation, lipid peroxidation, and mitochondrial membrane potential were assessed to investigate the mechanism by which NAC protects hepatocytes against statins-induced toxicity and elucidate the cause of cell

death induced by statins. A significant amount of reactive oxygen species were formed when hepatocytes were treated with atorvastatin (Figure 1), simvastatin (Figure 2) and/or lovastatin (Figure 3) and supplementation with NAC (200 μ M) significantly reduced the statins-induced ROS generation (Figure 1-3).

Table 1. Prevention of statins cytotoxicity by NAC.

Cytotoxicity (% Trypan blue uptake)			
Addition Incubation time (min)	60	120	180
Control (only hepatocytes)	18 \pm 1	22 \pm 2	24 \pm 1
+NAC(N-acetylcysteine) 200 μ M	15 \pm 1	17 \pm 1	21 \pm 1
+Atorvastatin 450 μ M	36 \pm 1 ^a	52 \pm 2 ^a	68 \pm 1 ^a
+NAC200 μ M	23 \pm 1 ^b	29 \pm 1 ^b	38 \pm 2 ^b
+Simvastatin 200 μ M	38 \pm 2 ^a	54 \pm 2 ^a	88 \pm 2 ^a
+ NAC200 μ M	25 \pm 1 ^b	33 \pm 2 ^b	47 \pm 2 ^b
+Lovastatin 200 μ M	39 \pm 2 ^a	52 \pm 2 ^a	74 \pm 3 ^a
+ NAC200 μ M	28 \pm 2 ^b	38 \pm 2 ^b	45 \pm 4 ^b

Statin doses are LD₅₀ (The statins concentration caused 50% death after 120 minutes).
 NAC (200 μ M) was added at the time of statins addition.
 Data represent Mean \pm SE for three independent experiments.
^a: Significantly different from control group (P<0.05).
^b: Significantly different from statins-treated group (P<0.05).

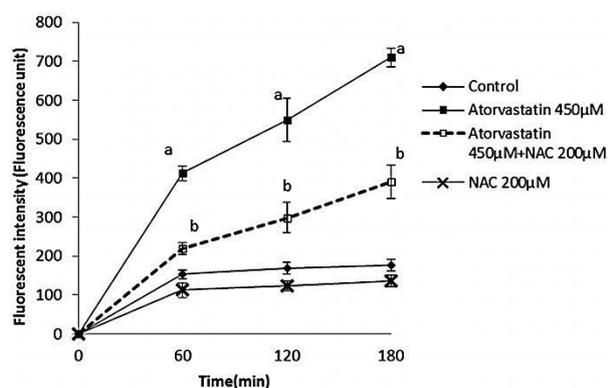


Figure 1. Atorvastatin- induced ROS formation in isolated rat hepatocytes and the protective effect of NAC.

NAC (200 μ M) was added at the time of Atorvastatin(450 μ M) addition.

Data are shown as Mean \pm SE for three independent experiments.

^a: significant VS control group(p<0.05).

^b: significant VS Atorvastatin treated group(p<0.05).

It has been shown that ROS formation is usually followed by lipid peroxidation.¹⁵ Lipid peroxidation was determined after 120 and 180 minutes after the statins treatment and it was found that NAC (200 μ M) significantly prevented lipid peroxidation in the statins-treated hepatocytes (Figure 4-6).

The effect of statins on mitochondria as an essential organelle in energy production was assessed. It was found that statins caused mitochondrial membrane potential (MMP)depression (Figure 7-9). NAC

supplementation showed a significant elevation of mitochondrial membrane potential.

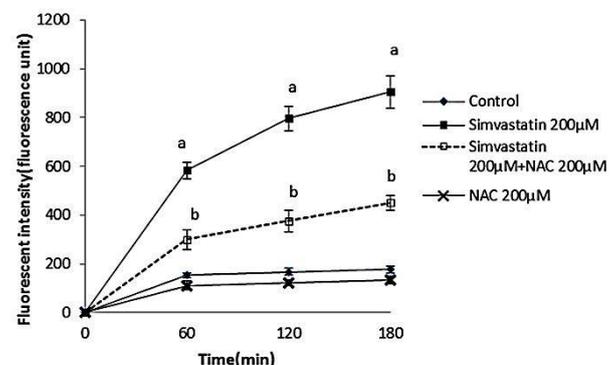


Figure 2. Simvastatin- induced ROS formation in isolated rat hepatocytes and the protective effect of NAC.

NAC (200 μ M) was added at the time of Simvastatin (200 μ M) addition.

Data are shown as Mean \pm SE for three independent experiments.

^a: significant VS control group(p<0.05).

^b: significant VS Simvastatin - treated group(p<0.05).

Discussion

The results of our study showed the beneficial protective effects of NAC against statin-induced hepatotoxicity. Inhibition of the respiratory chain (complex I and III), depolarization of mitochondrial membrane⁷ and releasing of Ca²⁺, are known as the results of statins exposure.¹⁶ The statins are metabolized by liver cytochrome P450 (CYP 3A4)

enzymes.¹ The cytochrome P450 include a superfamily of monooxygenases whose members function critically in xenobiotics catabolism.¹⁷ The major metabolic reactions include oxidation of aliphatic carbons, alkenyl groups and aromatic rings. After oxidation, subsequent reactions involve N-dealkylations, dehydration to form C=C double bonds, dehydrogenation of primary alcohols to an aldehyde and then a carboxylic acid, and finally glutathione conjugation of epoxides.¹⁸ Mitochondria or cytochrome P450-dependent metabolism act as ROS production systems and participate in cell death processes.¹⁹ Our data showed that treating hepatocytes with the statins produces a significant amount of ROS, induces lipid peroxidation, reduces mitochondrial membrane potential and promotes cytotoxicity as compared to the control group and these effect are dose- and time-dependent (different doses of the statins were assessed but data are not shown). Our results are in accordance with the previous study.⁷ The amount of ROS formation by Simvastatin was more than that with the others. The intracellular ROS generation is accompanied with cell death in cultured hepatocytes.²⁰ The efficacy of N-acetylcysteine (NAC) administration as a free radical scavenger in multiple models of oxidative stress induction has been studied.^{8,12} The administration of N-acetylcysteine (NAC), as an excellent source of intracellular cysteine and free radical scavenger, has been shown to have clinical applications in conditions such as HIV infection, cancer, heart disease, as well as in smoking, kidney and liver diseases.²¹ Adding NAC to the cells incubated with the statins resulted in prevention of ROS formation, lipid peroxidation, mitochondrial dysfunction, and finally improving the cells survival. Therefore, reactive oxygen species formed during the statins metabolism are scavenged by NAC and this may have a role in its protective effects in statins cytotoxicity.

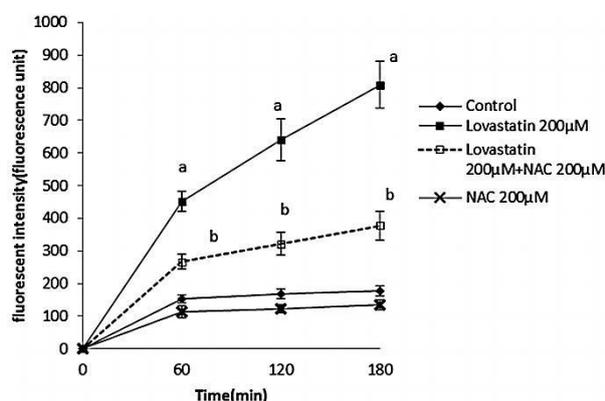


Figure 3. Lovastatin- induced ROS formation in isolated rat hepatocytes and the protective effect of NAC. NAC(200 µM) was added at the time of Lovastatin(200 µM) addition. Data are shown as Mean±SE for three independent experiments. a : significant VS control group($p<0.05$). b: significant VS Lovastatin - treated group($p<0.05$).

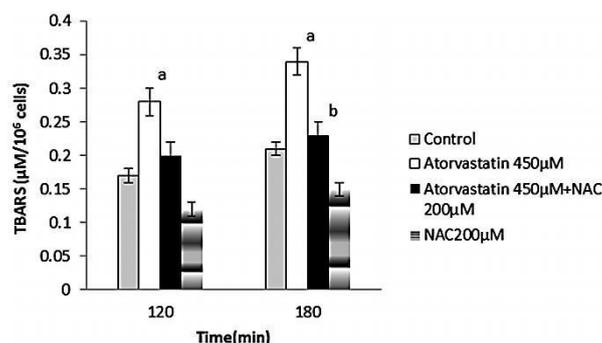


Figure 4. Atorvastatin-induced lipid peroxidation and the protective effect of NAC. NAC (200 µM) was added at the time of Atorvastatin(450 µM) addition. Data are shown as Mean±SE for three independent experiments. a : significant VS control group($p<0.05$). b : significant VS Atorvastatin treated group($p<0.05$).

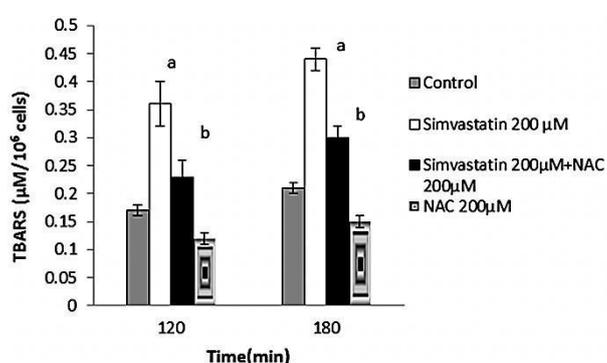


Figure 5. Simvastatin- induced lipid peroxidation and the protective effect of NAC. NAC (200 µM) was added at the time of Simvastatin(200 µM) addition. Data are shown as Mean±SE for three independent experiments. a : significant VS control group($p<0.05$). b : significant VS Simvastatin - treated group($p<0.05$).

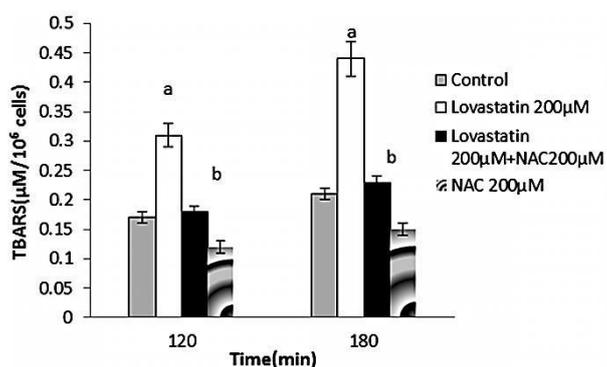


Figure 6. Lovastatin- induced lipid peroxidation and the protective effect of NAC. NAC (200 µM) was added at the time of Lovastatin (200 µM) addition. Data are shown as Mean±SE for three independent experiments. a : significant VS control group($p<0.05$). b : significant VS Lovastatin - treated group($p<0.05$).

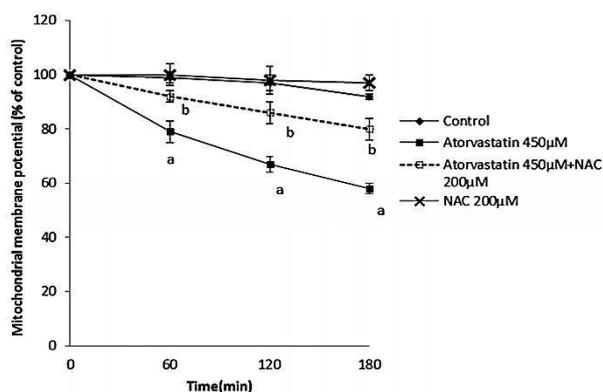


Figure 7. Effect of Atorvastatin on mitochondrial membrane potential and the protective role of NAC. NAC (200 µM) was added at the time of Atorvastatin (450 µM) addition.

Data are shown as Mean±SE for three independent experiments.

a : significant VS control group($p<0.05$).

b: significant VS Atorvastatin -treated group($p<0.05$).

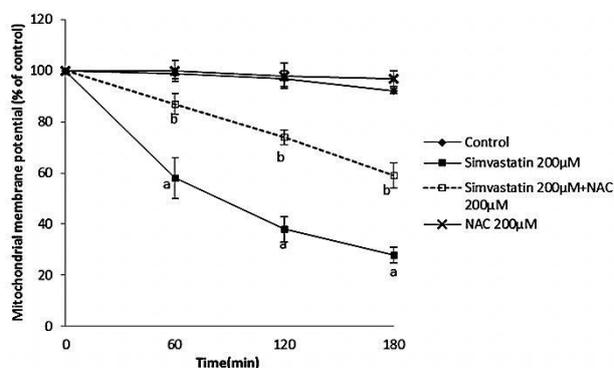


Figure 8. Effect of simvastatin on mitochondrial membrane potential and the protective role of NAC. NAC (200 µM) was added at the time of Simvastatin(200 µM) addition.

Data are shown as Mean±SE for three independent experiments.

a : significant VS control group($p<0.05$).

b: significant VS Simvastatin -treated group($p<0.05$).

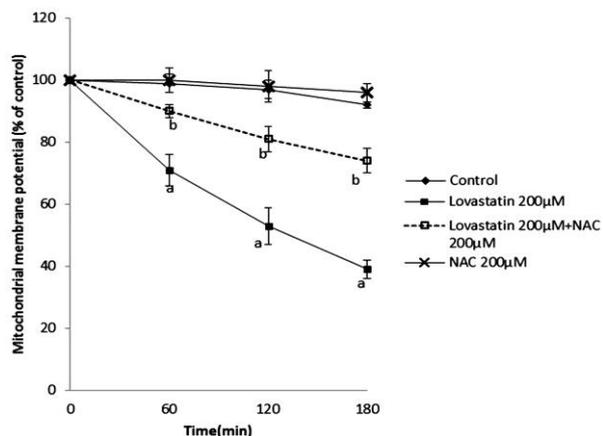


Figure 9. Effect of lovastatin on mitochondrial membrane potential and the protective role of NAC. NAC (200 µM) was added at the time of lovastatin (200 µM) addition.

Data are shown as Mean±SE for three independent experiments.

a : significant VS control group($p<0.05$).

b: significant VS lovastatin -treated group($p<0.05$).

Conclusion

In summary, in accordance with a previous study⁷, we have shown that, statins act as the inducers of oxidative stress that severely inhibits mitochondria respiration and decreases mitochondrial membrane potential in rat hepatocytes. They also promote peroxidation of lipids and subsequently induce cytotoxicity and cell death. The treatment of hepatocytes with NAC protected cells from the statins-induced ROS production, lipid peroxidation, mitochondrial impairment and cell death. Hence, supplementation with NAC may be an effective therapeutic strategy for the prevention and treatment of clinical conditions caused by statins.

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Conflict of Interest

The authors declare that they have no conflict of interest.

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