Metformin Inhibits Propofol-Induced Apoptosis of Mouse Hippocampal Neurons HT-22 Through Downregulating Cav-1

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Objective: To elucidate the neuroprotective function of metformin in suppressing propofol-induced apoptosis of HT-22 cells.

Methods: HT-22 cells were treated with 0, 10 or 100 $\mu\text{mol/L}$ propofol, followed by determination of their proliferative ability. Subsequently, changes in proliferation and apoptosis of propofol-treated HT-22 cells induced with metformin were assessed. Apoptosis-associated genes in HT-22 cells were detected by Western blot. At last, regulatory effects of Cav-1 on propofol and metformin-treated HT-22 cells were examined.

Results: Propofol treatment dose-dependently decreased proliferative ability and increased apoptosis ability in HT-22 cells, which were partially blocked by metformin administration. Upregulated Bcl-2 and downregulated Bax were observed in propofol-treated HT-22 cells following metformin administration. In addition, Cav-1 level in HT-22 cells was regulated by metformin treatment. Notably, metformin reversed propofol-induced apoptosis stimulation and proliferation decline in HT-22 cells via downregulating Cav-1.

Conclusion: In our study, we found that propofol could induce apoptosis of HT-22 cells and metformin could rescue the apoptosis effect regulated by propofol. Then, we found that metformin protects propofol-induced neuronal apoptosis via downregulating Cav-1.

Keywords: metformin, propofol, Cav-1, apoptosis

Introduction

Propofol (2,6-diisopropylphenol) belongs to a type of short-acting intravenous anesthetics. Propofol is commonly used to induce and maintain anesthesia in adults owing to its rapid induction and recovery time.^{1,2} Since it has a good antiemetic effect, propofol is also widely applied in the induction and maintenance of pediatric and obstetric anesthesia.³ Nevertheless, a certain dose of propofol may induce neurotoxicity as the nervous system is extremely sensitive to changes in internal and external environment.^{4,5} Recent studies reported that propofol administration would induce neuronal apoptosis^{6,7} and may impair learning, memory and cognitive function in children during brain development.⁸ Therefore, it is necessary to clarify the mechanism underlying the neurotoxicity of propofol. So far, several mechanisms have been identified for explaining propofol-induced neurotoxicity, including mitochondrial dysfunction, translocation of apoptosis-inducing factors and the mTOR pathway.^{9,10} It is worth noting that propofol can regulate a variety of cellular pathways.¹⁰

Apoptosis results from changes in plasma membrane phospholipids, cell shrinkage and nuclear DNA condensation and cleavage.¹¹ Apoptosis is regulated by

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abundant genes, especially the Bcl-2 family.¹² Among them, Bax, Bad and Bak exert pro-apoptotic effects, while Bcl-2 and Bcl-xL are anti-apoptotic genes.¹³ The balance between expression levels of pro-apoptotic proteins (Bax, Bak, and Bad) and anti-apoptotic proteins (Bcl-2 and Bcl-xL) is critical for neuronal survival.¹⁴ Apoptosis may be a crucial target during the process of propofol-induced neurotoxicity. Suppression of neuronal apoptosis could be a novel strategy for the treatment of propofol-induced neurotoxicity.

Metformin is a first-line antidiabetic drug. Metformin exerts hypoglycemic outcomes through reducing hepatic gluconeogenesis, increasing glucose utilization and improving insulin sensitivity.¹⁵ Abundant evidence have suggested the potential function of metformin in anti-inflammation, anti-apoptosis and anti-oxidation in central nervous system diseases (ie, ischemic stroke, intracerebral hemorrhage and multiple sclerosis).¹⁶ As a result, metformin is believed to exert the function of neuronal protection.

Caveolins is a membrane protein family composed of three members with 21–24 kDa. Caveolin-1 (Cav-1) is abundantly expressed in endothelial cells, and caveolin-3 (CAV-3) is mainly expressed in muscle cells.¹⁷ Cav-1 is a structural protein responsible for maintaining vesicle morphology. It is involved in the disease progression of ischemia/reperfusion (I/R).^{18–20} A recent study demonstrated the extensive functions of Cav-1 in regulating multiple cellular behaviors.²¹ A recent study reported that Cav-1 knockdown increases the therapeutic sensitivity of lung cancer to cisplatin-induced apoptosis.²² Besides, β -Carotene induces apoptosis in esophageal squamous cell Carcinoma via the Cav-1/AKT/NF- κ B signaling pathway.²³ The previous study showed that Cav-1 was strongly associated with apoptosis. However, the role of Cav-1 in propofol-induced apoptosis remains unclear.

In this paper, we mainly explored the potential regulatory effects of metformin and Cav-1 on propofol-induced neurotoxicity.

Methods

Cell Culture of HT-22

Immortalized mouse hippocampal neuron HT-22 (CL-0595) cells were provided by Cell Bank, Shanghai. Cells were cultured in DMEM containing 10% FBS, 100 μ M penicillin and 100 ng/mL streptomycin (Invitrogen, Carlsbad, CA, USA). They were maintained at 37°C, 5%

CO₂. Cells were inoculated in 6-well plates with 1×10^5 cells per well, and incubated with 0, 1, 10 or 100 μ M propofol for 24 hrs, respectively.

Transfection

HT-22 cells were treated with 100 μ M propofol and 10 μ M metformin for 24 h. Then, si-NC, si-Cav-1, pcDNA-NC and pcDNA-Cav-1 purchased from Invitrogen (Invitrogen, CA, USA) were transfected into HT-22 cells using Lipofectamine 3000. After transfection for 4–6 h, medium containing 10% FBS was replaced.

Cell Counting Kit-8 (CCK-8) Assay

Cells were seeded in the 96-well plate and cultured for 80% confluence. Twenty microliters of CCK-8 (Houston TX, USA) solution was added in each well and reacted at 37°C in dark for 2 h. Absorbance (A) at 450 nm was recorded at the appointed time points using the CCK-8 kit (Dojindo Laboratories, Kumamoto, Japan) for depicting the viability curves.

5-Ethynyl-2'-Deoxyuridine (EdU) Assay

Cells were labeled with 100 μ L of EdU (Ribobio, Guangzhou, China) reagent (50 μ M/L) per well for 2 h. After PBS washing, cells were fixed in 50 μ L of fixation buffer, decolorized with 2 mg/mL glycine and permeated with 100 μ L of 0.5% Triton X-100 (Solarbio, Beijing, China). Cells were dyed with Apollo for 30 min, followed by Hoechst in dark for 10 min. EdU-positive cells, Hoechst-labeled cells and their merged images were determined under a fluorescent microscope.

Flow Cytometry

Cells were prepared into suspension with 1×10^6 /mL. One hundred microliters of suspension was added in each tube, incubated with 10 μ L of AnnexinV, 380 μ L of buffer and 10 μ L of PI (propidium iodide). After 15-min incubation in dark, apoptosis was analyzed by a BD FACSCalibur flow cytometer (BD Bioscience, USA).

TdT-Mediated dUTP Nick-End Labeling (TUNEL)

For TUNEL assay (Roche, Shanghai, China), cells were subjected to 30-min fixation in 4% paraformaldehyde, followed by 30-min incubation in H₂O₂ to inactivate the endogenous enzyme. Cells were immersed in 0.2% Triton X-100 solution for 5 min to enhance cell membrane permeability, and further incubated with deoxynucleotide

terminal transferase (rTdT) at 37°C for 1 h. The cell nucleus was stained brown. Each section was randomly selected for 5 fields. Apoptotic rate (TUNEL-positive ratio) was finally calculated (magnification 400×).

Western Blot

Cellular protein was extracted and quantified by BCA method. After electrophoresis, proteins were transferred on a PVDF membrane. Membranes were immersed in 5% skim milk for blocking the non-specific sites. Two hours later, membranes were reacted with primary and secondary antibodies. Band exposure was achieved by ECL and analyzed by Image Software. The antibodies purchased from Proteintech used in our research were as follows: Bax (50599-2-Ig, 1:1000), Bcl-2 (26593-1-AP, 1:1000), Cav-1 (16447-1-AP, 1:1000), Gapdh (60004-1-Ig, 1:1000).

Statistical Processing

SPSS 22.0 was used for data analyses. Data were expressed as mean ± standard deviation. Differences were analyzed by one-way ANOVA. $P < 0.05$ was considered as statistically significant.

Results

Propofol Administration Inhibited Proliferation and Induced Apoptosis in HT-22 Cells

CCK-8 assay revealed a dose-dependent decline in the viability of HT-22 cells after propofol administration (Figure 1A). EdU assay further depicted the dose-dependently declined EdU-positive ratio in propofol-treated HT-22 cells (Figure 1B and C). After treatment of increased doses of propofol, the apoptotic rate gradually increased (Figure 1D and E). TUNEL-positive ratio was dose-dependently elevated by propofol treatment in HT-22 cells (Figure 1F and G). Apoptosis-associated genes were determined by Western blot. As data revealed, Bcl-2 was downregulated and Bax was upregulated in propofol-treated hippocampal neurons in a dose-dependent way (Figure 1H).

Metformin Treatment Reversed Propofol-Induced Apoptosis in HT-22 Cells

To elucidate the influence of metformin on HT-22 cells, they were administrated with metformin and propofol.

Interestingly, the declined viability owing to propofol treatment was reversed following metformin administration (Figure 2A). Similarly, decreased EdU-positive ratio in propofol-treated HT-22 cells was partially blocked by metformin (Figure 2B and C). Decreased apoptotic rate was observed after metformin administration in propofol-treated HT-22 cells (Figure 2D and E). Compared with those treated with propofol, TUNEL-positive ratio decreased in HT-22 cells treated with both propofol and metformin (Figure 2F and G). As data revealed, Bcl-2 was downregulated and Bax was upregulated in propofol-treated hippocampal neurons which were reversed by metformin (Figure 2H). As a result, metformin effectively reversed propofol-induced proliferation inhibition and apoptosis stimulation in hippocampal neurons.

Metformin Regulated Cav-1 Level

Western blot analysis uncovered that the protein level of Cav-1 dose-dependently upregulated in propofol-treated HT-22 cells (Figure 3A and B). Furthermore, metformin treatment downregulated Cav-1 level in propofol-treated HT-22 cells (Figure 3C and D). Hence, metformin was able to regulate propofol-induced Cav-1 upregulation.

Propofol-Induced Apoptosis in HT-22 Cells by Upregulating Cav-1

To further explore the function of propofol in inducing neuronal apoptosis, we constructed si-Cav-1 and pcDNA-Cav-1. Transfection efficacy of si-Cav-1 was determined in HT-22 cells by Western blot (Figure 4A). The protein level of Cav-1 was markedly upregulated after propofol treatment, which was then downregulated by transfection of si-Cav-1 (Figure 4B). CCK-8 assay showed that knock-down of Cav-1 partially reversed the declined viability in HT-22 cells treated with propofol (Figure 4C). EdU assay obtained a similar result (Figure 4D and E). The increased apoptotic rate in propofol-treated HT-22 cells was slightly reduced after transfection of si-Cav-1 (Figure 4F and G).

Metformin Reversed Propofol-Induced Neuronal Apoptosis Through Downregulating Cav-1

Next, transfection efficacy of pcDNA-Cav-1 was verified in HT-22 cells (Figure 5A). Transfection of pcDNA-Cav-1 inhibited the upregulated protein level of Cav-1 in propofol-treated HT-22 cells (Figure 5B). Metformin reversed the declined viability and EdU-positive ratio in

Figure 1 Propofol-induced apoptosis in HT-22 cells. **(A)** CCK-8 assay results showed viability in HT-22 cells treated with 0, 1, 10 and 100 μ M propofol, respectively. **(B and C)** EdU assay results showed EdU-positive HT-22 cells treated with 0, 1, 10 and 100 μ M propofol, respectively **(B)**. Quantitative analysis of EdU-positive ratio **(C)**. **(D and E)** Flow cytometry results showed distribution of apoptotic cells, necrotic cells and survival cells following the treatment of 0, 1, 10 and 100 μ M propofol in HT-22 cells, respectively **(D)**. Quantitative analysis of apoptosis rate **(E)**. **(F and G)** TUNEL results showed TUNEL-positive cells following the treatment of 0, 1, 10 and 100 μ M propofol in HT-22 cells, respectively **(F)**. Quantitative analysis of TUNEL-positive rate **(G)**. **(H)** Protein levels of Bcl-2 and Bax in HT-22 cells treated with 0, 1, 10 and 100 μ M propofol, respectively (* p <0.05 compared to control group).

propofol-treated HT-22 cells, which were further reduced after overexpression of Cav-1 (Figure 5C–E). In the meantime, overexpression of Cav-1 enhanced apoptosis in propofol-treated neurons even after metformin treatment (Figure 5F and G). The above data proved that Cav-1 was responsible for protective effects of metformin on propofol-induced neurotoxicity.

Discussion

Increasing evidence have proved that propofol administration is able to induce developmental neurotoxicity, thereafter leading to long-term cognitive and learning abnormalities.

Hence, the safety usage of propofol in pediatric anesthesia is well concerned.²⁴ Fredriksson et al²⁵ proposed that propofol administration dose-dependently aggravates the apoptotic degree in neurons of young rats, which could impair the long-term learning and memory ability of rats. In 7-day-old neonatal SD rats, propofol treatment markedly induces apoptosis in hippocampal neurons.²⁶ Our results revealed that propofol treatment dose-dependently decreased proliferative ability and increased apoptosis in HT-22 cells.

Apoptosis is of significance in maintaining homeostasis and normal development. Nevertheless, excessive apoptosis leads to adverse biological consequences. For instance,

Figure 2 Metformin reversed propofol-induced apoptosis in HT-22 cells (**A**) CCK-8 assay results showed viability in propofol-induced HT-22 cells either treated with 10 μ M metformin or not. (**B** and **C**) EdU assay results showed EdU-positive HT-22 cells with propofol induction, followed by 10 μ M metformin treatment or not (**B**). Quantitative analysis of EdU-positive ratio (**C**). (**D** and **E**) Flow cytometry results showed distribution of apoptotic cells, necrotic cells and survival cells in propofol-induced HT-22 cells either treated with 10 μ M metformin or not (**D**). Quantitative analysis of apoptosis rate (**E**). (**F** and **G**) TUNEL results showed TUNEL-positive cells in propofol-induced HT-22 cells either treated with 10 μ M metformin or not (**F**). Quantitative analysis of TUNEL-positive rate (**G**). (**H**) Protein levels of Bcl-2 and Bax in propofol-induced HT-22 cells either treated with 10 μ M metformin or not (* p <0.05 compared to control group; & p <0.05, compared to propofol (100 μ M) group).

uncontrolled apoptosis is closely linked to ischemic heart disease, AIDS and Alzheimer's disease, Parkinson's disease and other neurodegenerative diseases.²⁷ Cell apoptosis has been well concerned in biology and medical researches.

Metformin has been verified as a safe and effective oral hypoglycemic agent extensively used in the treatment of diabetes.^{28,29} In nervous system, metformin stimulates the growth of newly formed hippocampal neurons and improves spatial learning and memory in mice.^{30,31} Through activating the S1P1-dependent ERK1/2 pathway, metformin protects rabbits from sevoflurane-induced neuronal apoptosis.³²

Besides, metformin another AMPK activator AICAR activates AKT S473 which provides survival signal for cells by inhibiting apoptotic signaling pathways.³³ In this analysis, metformin treatment protected propofol-induced proliferation inhibition and apoptosis stimulation in HT-22 cells. Therefore, we have proved the protective effect of metformin on propofol-induced neurotoxicity.

Cav-1 is a 22 kDa plasma membrane scaffold protein.³⁴ A previous study reported that Cav-1 regulates Caspase 3-mediated apoptosis pathway by downregulating Survivin.³⁵ Cav-1 deficiency has protective effects on both intracellular and extracellular apoptosis.³⁶ In addition, through interacting

Figure 3 Metformin treatment regulated Cav-1 level. **(A and B)** Protein level of Cav-1 in HT-22 cells treated with 0, 1, 10 and 100 μ M propofol, respectively **(A)**. Grey value analysis of Cav-1 **(B)**. **(C and D)** Protein level of Cav-1 in propofol-induced HT-22 cells either treated with 10 μ M metformin or not **(C)**. Grey value analysis of Cav-1 **(D)** (* p <0.05 compared to control group; & p <0.05, compared to propofol (100 μ M) group).

Figure 4 Propofol-induced apoptosis in HT-22 cells through upregulating Cav-1. **(A)** Transfection efficacy of si-Cav-1 in HT-22 cells. **(B)** Protein level of Cav-1 in propofol-induced HT-22 cells transfected with si-NC or si-Cav-1. **(C)** CCK-8 assay results showed viability in propofol-induced HT-22 cells transfected with si-NC or si-Cav-1. **(D and E)** EdU assay results showed EdU-positive cells in propofol-induced HT-22 cells transfected with si-NC or si-Cav-1 **(D)**. Quantitative analysis of EdU-positive ratio **(E)**. **(F and G)** Flow cytometry results showed distribution of apoptotic cells, necrotic cells and survival cells in propofol-induced HT-22 cells transfected with si-NC or si-Cav-1 **(F)**. Quantitative analysis of apoptosis rate **(G)** (* p <0.05 compared to control group; & p <0.05, compared to propofol (100 μ M) group).

with Ras, Cav-1 promotes the activation of the Ras/Raf/ERK pathway, an important pathway participating in cell proliferation and apoptosis.^{37–40} Here, Cav-1 level was upregulated in propofol-treated HT-22 cells, which was downregulated

following metformin treatment. In particular, knockdown of Cav-1 suppressed propofol-induced apoptosis in HT-22 cells. Rescue experiments further confirmed that overexpression of Cav-1 abolished the anti-apoptosis function of metformin in

Figure 5 Metformin reversed propofol-induced apoptosis in HT-22 cells through downregulating Cav-1. **(A)** Transfection efficacy of pcDNA-Cav-1 in HT-22 cells. **(B)** Protein level of Cav-1 in propofol-induced HT-22 cells transfected with pcDNA-NC or pcDNA-Cav-1 with either metformin treatment or not. **(C)** CCK-8 assay results showed viability in propofol-induced HT-22 cells transfected with pcDNA-NC or pcDNA-Cav-1 with either metformin treatment or not. **(D and E)** EdU assay results showed EdU-positive cells in propofol-induced HT-22 cells transfected with pcDNA-NC or pcDNA-Cav-1 with either metformin treatment or not **(D)**. Quantitative analysis of EdU-positive ratio **(E)**. **(F and G)** Flow cytometry results showed distribution of apoptotic cells, necrotic cells and survival cells in propofol-induced HT-22 cells transfected with pcDNA-NC or pcDNA-Cav-1 with either metformin treatment or not **(F)**. Quantitative analysis of apoptosis rate **(G)** (* $p < 0.05$ compared to control group; $^{\#}p < 0.05$, compared to propofol (100μM) group).

propofol-treated HT-22 cells. Collectively, metformin protected propofol-induced neurotoxicity through Cav-1.

Conclusions

In our study, we found that propofol treatment dose-dependently decreased proliferative ability and increased apoptosis in HT-22 cells. Besides, metformin could rescue the apoptosis effect of propofol induced in HT-22. In addition, we found that propofol induced the apoptosis of HT-22 by up-regulating Cav-1 and metformin protects

propofol-induced neuronal apoptosis via downregulating Cav-1.

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Disclosure

The authors report no conflicts of interest in this work.

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