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# A new cell model of glutaric acidemia type I and the toxicity research

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Glutaric acidemia (GA-1) is an autosomal recessive disorder of lysine, hydroxylysine, and tryptophan metabolism caused by deficiency of glutaryl-CoA dehydrogenase. It results in the accumulation of 3-hydroxyglutaric and glutaric acid. Affected patients can present with brain atrophy and macrocephaly and with acute dystonia secondary to striatal degeneration in most cases triggered by an intercurrent childhood infection with fever between 6 and 18 months of age. The most common animal model of GA-1 is GCDH deficient mouse. We are trying to establish a new cell model of GA-1 to study the mechanism of neurotoxicity and for the future establishment of the new animal model.

Three *Gcdh* shRNA sequences were recombined to the lentivirus vector carrying the GFP reporter gene. And the *Gcdh*-shRNA vector with the highest knock-down efficiency was transfected into the primary striatal neuron cells. Real-time PCR and Western Bot showed the knock-down efficiency was about 65%. And compared with the control group, the cells transfected with *Gcdh* shRNA vector shranked obviously and gathered together, the neurites disappeared, and cell debris increased significantly. Hoechst staining of the nucleus showed the increased apoptotic rate in the knock-down group. MTT indicated the decreased mitochondrial function after *Gcdh* shRNA interference. These results suggest that *Gcdh* shRNA interference in rat primary striatal neuron cells could increase the apoptotic rate, which could be due to the activation of mitochondria mediated apoptosis, triggered by the dysfunction of the mitochondria.

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