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franime download videos . and promote the establishment of an effective treatment system. Fetal/neonatal brain injury There is evidence to support hypoxic damage to the fetal and neonatal brain after an acute hypoxic-ischemic event. This brain damage leads to distinct long-term health consequences in later life. Although it is challenging to model this brain damage in a laboratory setting due to the lack of available animal models that mimic the disease process, there are a number of animal and in vitro models that can be used to study hypoxic brain injury. For example, middle cerebral artery occlusion, which is a common model of stroke, causes a hypoxic-ischemic event in an animal. This model can be used to study the acute phase of hypoxic-ischemic brain injury, including the immediate cellular response in the brain such as increased extracellular glutamate levels and the calcium toxicity that occurs after ischemia. Another animal model of hypoxic-ischemic injury, preterm brain injury, has been widely used to understand the long-term consequences of hypoxic-ischemic injury and is a model that can be used to study the effects of therapeutic interventions. In this model, animals are exposed to a hypoxic event in the first week after birth. An important aspect of this model is that it allows a greater window for interventions than those done in the middle cerebral artery occlusion and term brain injury models. Other models Other models of hypoxia-ischemia injury include brain trauma, which occurs when there is a blunt impact to the head. Brain trauma can occur due to a high-speed car crash or as a result of an assault: however, the degree of injury depends on the severity of the impact. Brain injury is characterized by cell death, inflammation, edema, and disruption of the blood-brain barrier. The resulting injury causes an elevation in extracellular glutamate levels, which in turn leads to calcium toxicity. This model can also be used to study the long-term consequences of the injury. In this model, the animal is exposed to a hypoxic-ischemic event in the first 2 weeks after birth, and then a mild brain injury is induced at P0-P21 with an impact that can be graded by the severity of the impact. Barbiturates and hypoxia Other models of hypoxia-ischemia injury include exposure to systemic,

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