Study of mechanism and condition for ischemic preconditioning induced by volatile anesthetics pretreatment in the liver

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Research Project Project/Area Number 12470314 **Research Category** Grant-in-Aid for Scientific Research (B) **Allocation Type** Single-year Grants Section 一般 Research Field Anesthesiology/Resuscitation studies **Research Institution** Kanazawa University **Principal Investigator INABA Hideo** Kanazawa University, Graduate School of medicine, Professor, 大学院·医学系研究科, 教授 (60159952) Co-Investigator(Kenkyū-buntansha) INAGAKI Nobuya Akita University, School of medicine, Professor, 医学部, 教授 (30241954) TANIGUCHI Takumi Kanazawa University, Graduate School of Medicine, Associate Professor, 大学院·医学系研究科, 助教授 (30301196) GOTO Yoshikazu Kanazawa University, School of Medicine Hospital, Lecturer, 医学部附属病院, 講師 (60282167) NAKAE Hajime Akita University, School of Medicine. Assistant Professor, 医学部, 助手 (10254781) Project Period (FY) 2000 - 2002 **Keywords**

Research Abstract

Halothane, isoflurane and sevoflurane attenuated ischemia-reperfusion injury in the perfused liver of fasted rats, suggesting a possible involvement of ischemic preconditioning.

In the rat liver slice incubation, isoflurane attenuated apoptosis induced by hypoxic challenge. Suppression of apoptosis is predominant in the hepatocyte.

volatile anesthetics / ischemic preconditioning / isolated perfused liver / liver slice incubation / apoptosis / patch clamp / K-ATP channel

Direct effect of isoflurane on K-ATP channel was studied in isolated pancreatic β cells using whole cell patch clamp method. Isoflurane suppressed K-ATP channel in a dosedependent manner. Repeated treatment of isoflurane enhanced the suppression of K-ATP channel.

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