

IMSUT Hospital

Department of Radiology

放射線科

Associate Professor	Hiroyuki Akai, M.D., D.M.Sc.
Senior Assistant Professor	Toshihiro Furuta, M.D., D.M.Sc.
Assistant Professor	Shimpei Kato, M.D., D.M.Sc.
Project Assistant Professor	Naomasa Okimoto, M.D., D.M.Sc.

准教授	博士(医学)	赤井宏行
講師	博士(医学)	古田寿宏
助教	博士(医学)	加藤伸平
特任助教	博士(医学)	沖元斉正

Department of Radiological Technology

放射線部

Associate Professor	Hiroyuki Akai, M.D., D.M.Sc.
Head Radiologic Technologist	Kenji Ino, RT

准教授	博士(医学)	赤井宏行
放射線技師長		井野賢二

The Department of Radiology undertakes radiology service at IMSUT hospital. Our expertise includes general diagnostic radiology, neuroradiology, clinical nuclear medicine, and radiation therapy. Board-certified radiologists at the Department of Radiology conduct all examinations of CT, MRI, and nuclear medicine. Radiological reports are made by the radiologists. In addition, several clinical studies are being conducted in collaboration with other departments or institutions. We also investigate the technical aspects of molecular imaging with intact small animals for its application to preclinical studies using an optical imaging system and MRI.

The Department of Radiological Technology constitutes the hospital radiology service together with the Department of Radiology. Plain radiography, dual-energy X-ray absorptiometry, and barium studies are also available at the Department of Radiological Technology, other than CT, MRI, and radioisotope examinations. More than 10,000 patients visit our department every year. Radiologic technologists at the department make an effort to provide high-quality medical images in daily practice as well as to reasonably reduce radiation exposure of a patient during the examination.

CT lymphangiography of the thoracic duct in mice: direct mesenteric versus popliteal lymph node puncture.

Kato S, Sugawara H¹, Furuta T, Abe O², Akai H.

¹ Department of Diagnostic Radiology, McGill University, ² Department of Radiology, Graduate School

of Medicine, University of Tokyo

In the present study, we aimed to evaluate the efficacy of computed tomography (CT) lymphangiography after direct mesenteric lymph node injection for thoracic duct (TD) visualization in mice. Twelve female BALB/c mice were injected with 35 μ L of iodi-

nated contrast medium (iomeprol 350 mgI/mL) into the mesenteric (mesenteric group) or popliteal (popliteal group) lymph nodes. CT images were acquired before injection and 1 min, 3 min, 5 min, 10 min, and 15 min after injection using a micro-CT scanner. Contrast ratios (CRs) were measured at the cisterna chyli and three levels of the TD (diaphragm, carina, and venous angle). Two experienced radiologists qualitatively assessed images as good, fair, or poor. As a result, the mesenteric group had significantly higher mean ($\pm$ standard deviation) CRs than the popliteal group for all examined regions at 1 min after injection: cisterna chyli (14.01 ± 4.77 versus 1.47 ± 1.21 , $p < 0.001$), diaphragm (7.28 ± 2.50 versus 0.85 ± 0.61 , $p = 0.0011$), carina (10.33 ± 3.42 versus 0.44 ± 0.40 , $p < 0.001$), and venous angle (6.26 ± 2.02 versus 0.79 ± 0.75 , $p < 0.001$). For the TD between the cisterna chyli and the diaphragm, 6/6 mice in the mesenteric group showed strong enhancement, whereas 5/6 mice in the popliteal group showed minimal or no enhancement. The visual scores of the mesenteric group were significantly higher than those of the popliteal group for all the evaluated regions ($p = 0.002$). To sum up, CT lymphangiography via mesenteric lymph node injection provides better imaging of the TD in mice than popliteal lymph node injection.

Pulmonary Toxicity of Ethiodized Oil in the Venous Circulation in Mice: Radiological Findings and Pathological Correlation.

Kato S, Sugawara H¹, Okimoto N, Furuta T, Abe O², Ota Y³, Akai H.

³ Department of Pathology, Institute of medical sci-

ence, University of Tokyo

This study investigated the acute and chronic pulmonary toxicity and determine the median lethal dose (lethal dose 50% [LD50]) of intravenously administered ethiodized oil given the limited data on its systemic toxicity despite its known risk of inadvertent venous entry during lymphangiography. Twenty female BALB/c mice received tail vein injections of saline ($n = 4$) or ethiodized oil (Lipiodol; Guerbet, Villepinte, France) at 10 μ L ($n = 4$), 30 μ L ($n = 8$), or 50 μ L ($n = 4$). Survival outcomes, computed tomography (CT) imaging findings, and histopathological features including hemosiderin deposition were evaluated. Chronic effects were assessed at 10 weeks through histopathology and hemosiderin deposition quantification. Human equivalent doses (HEDs) were calculated using body weight ratio conversion. As a result, The LD50 was determined to be 30 μ L in mice, corresponding to an HED of 94.7 mL for a 60-kg human. All mice receiving 50 μ L died immediately, whereas the 30- μ L group showed 50% survival. CT imaging revealed dose-dependent ethiodized oil accumulation predominantly in pulmonary vasculature, with regional heterogeneity in distribution patterns. Surviving mice from the 30- μ L group exhibited significantly higher hemosiderin deposition compared with the 10- μ L group ($P = .0054$), indicating chronic microvascular damage. In conclusion, intravenous ethiodized oil induces dose-dependent pulmonary embolism with an LD50 of 30 μ L in mice, and even sublethal doses cause chronic pulmonary injury, highlighting the need for clinical monitoring in cases of suspected venous exposure.

Publications

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