

The device is used mainly for the 140 KeV γ -rays from ^{99m}Tc . A multi-channel tungsten collimator with 4681 holes are employed to project an image of the subjects on the crystal.

Radioisotope angiography of the cases with control and various cardiopulmonary disorders were studied. After i.v. inj. of 10mCi of $^{99m}\text{TcO}_4$ saline solution the process of ^{99m}Tc bolus filling and escape from the heart chambers was recorded, processed later as the cumulative images of dilution process in one second intervals. Also with specially designed

collimated light sensor, dilution processes of each heart chamber was obtained by replaying the tape. This method is very useful because the numerical data from a specific picture area recorded on the tape can be read out any time.

This camera has the advantages of simplicity, good sensitivity for both low and medium-energy γ -rays, and no dead time problems for large dose of radioisotope administered. Radioisotope angiography is one of the most promising application of this γ -camera.

A Method of Lung Profile Scanning

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The purpose of this report is to describe our method of lung profile scanning using ^{131}I -MAA and ^{133}Xe and our equipment with driving unit and speed-changing gears.

Two collimated counters with 2 inches crystal detector were placed behind the lungs of a subject, whose back was attached to the lucent plastic plate. The linear profile scanning was performed from the apex to the bottom of the lungs. Various speeds of scanning were obtained by combination of an induction motor and gears. Pre-setting markers set in the equipment determined the position of the detector to the lungs.

The apparent shift and decrease of peak values of the pulmonary blood distribution curves were examined, which resulted from combination of various scanning speeds and time constants of the rate meter using ^{131}I -MAA. When apparent shifts of curves were estimated by the variation of the upper to lower blood distribution ratio (U/L ratio), a real distribution curve was obtained in the following conditions, i.e., when time constant was 0.5 sec., the scanning speed was 20 mm/sec. or less, and the speed for time constant of 1 sec. was 5 mm/sec. or less. The decrease of peak value was not found in the following conditions, i.e., when time constant was 0.5

sec., the scanning speed was 30 mm/sec. or less, and for time constant of 1 sec. the speed was 10 mm/sec. or less, respectively. So that, optimal conditions of scanning speeds and time constants were determined to be a combination of 0.5 sec of time constant with 20 mm/sec. or slower of speed, and that of 1.0 sec. with 5 mm/sec. or slower.

The ratio of U/L in ^{133}Xe -pulmogram was slightly larger than that in ^{131}I -MAA pulmogram under the same conditions. This seems to be resulted from the alteration of the lung volume by breath-holding follows ^{133}Xe injection and the energetic difference between two isotopes. That is ^{133}Xe has low energy level (80 KeV) in comparison with that of ^{131}I -MAA (364 KeV). Since gamma-emitter (^{133}Xe) distributed in the deep and front part of the lungs was absorbed to considerable extent by the tissues, the rear detector could not detect sufficiently its emission. In this respect, two detectors were confronted, one in the front and the other in the rear of the lung so that an efficient and even detection of the gamma-emission was accomplished.

As the conclusion, our method presently described is useful in establishing clinical diagnosis and elucidation of pulmonary blood flow distribution.