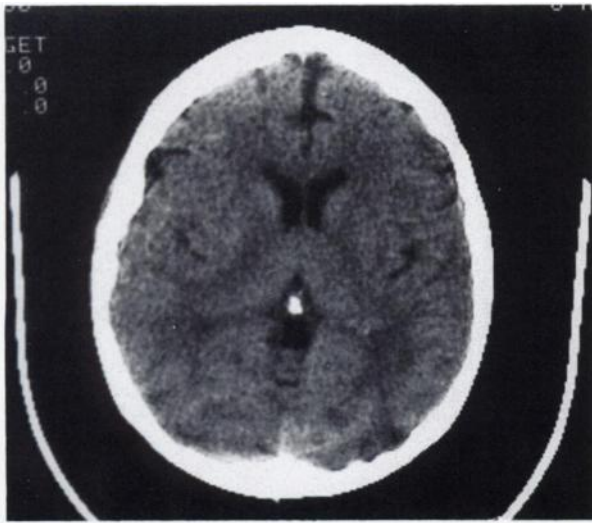


SPECTamine[®]

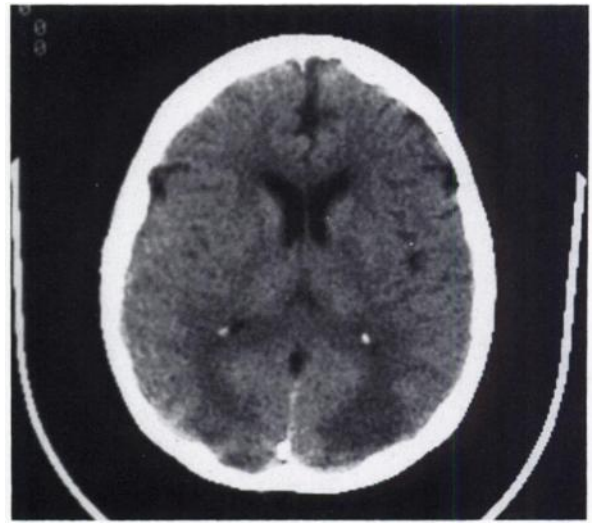
Iofetamine HCl I 123 Injection

STUDY: Confirms Diagnosis



Initial CT (5/6/88)

Showed multiple low-density regions involving white and gray matter in the parietal and occipital areas. Thought to be related to an inflammatory process, less likely an embolic insult. A confirmatory diagnosis was not possible.



Repeat CT (5/16/88)

Showed some change to the low attenuation areas—appearing larger and more confluent than previously noted. The pattern was atypical for infarction. Again, infectious etiology for the abnormality was entertained and diagnosis nonconfirmatory.

*Images Courtesy of
Deaconess Hospital
Boston, MA*

For additional information on the use of SPECTamine in stroke diagnosis, contact your local Medi-Physics Territory Manager, or call the SPECTamine[®] Hotline 1-800-451-7732.

A Case Study:

61-year-old female postop aorta bifemoral graft with a complicated 7-month postoperative course including renal failure, diverticulitis, Candida sepsis, multiple enteric cutaneous fistulous with multiple surgical procedures.

On 5/6/88, patient was noted to have two generalized seizures.

On 5/16/88, patient began to deteriorate neurologically. Complained of blindness.



SPECT Study (5/19/88)

Demonstrated bilaterally posterior cerebral artery infarction. Subsequent neurologic exams and clinical course confirmed diagnosis.

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Circle Reader Service No. 30

SPECTamine[®]

Iofetamine HCl I 123 Injection

For complete product information, consult package insert, a brief summary of which follows:

DIAGNOSTIC—FOR INTRAVENOUS USE

DESCRIPTION: SPECTAMINE[®] Iofetamine HCl I 123 Injection, is supplied as a sterile, apyrogenic, aqueous, isotonic sodium chloride solution for intravenous administration. Each milliliter of the solution contains 37 megabecquerels (1 millicurie) of Iofetamine HCl I 123 at calibration time, 0.15 milligram Iofetamine HCl, 0.017 millimole sodium phosphate, and 8.0 milligrams sodium chloride for isotonicity. The pH is adjusted to 4.5-6.0 with sodium hydroxide or hydrochloric acid. SPECTAMINE contains no bacteriostatic preservative. The radionuclidic composition at calibration time is not less than 94.7 percent I 123, not more than 4.8 percent I 124, and not more than 0.5 percent all others (I 125, I 126, I 130 and Te 121). The radionuclidic composition at the 6-hour expiration time is not less than 93.1 percent I 123, not more than 6.2 percent I 124, and not more than 0.7 percent all others.

INDICATIONS AND USAGE: SPECTAMINE (Iofetamine HCl I 123 Injection) is recommended for use as a lipid-soluble brain-imaging agent. It has been shown to be useful in the evaluation of nonlacunar stroke especially when used within 96 hours of onset of focal neurological deficit. The rates of agreement between abnormal images and the neurological examination suggestive of ischemic cerebrovascular insufficiency, appear to increase with the severity of symptoms. Its usefulness for the measurement of cerebral blood flow has not been established.

CONTRAINDICATIONS: None known.

WARNINGS: SPECTAMINE (Iofetamine HCl I 123 Injection) should not be administered to individuals with known hypersensitivity to sympathomimetic amines or to those individuals taking monoamine oxidase inhibitors.

PRECAUTIONS:

General

Some primate (Macaca fascicularis) studies have shown marked eye uptake of Iofetamine HCl I 123. Localization has not been studied in the isolated human eye although in vivo images suggest the concentration of Iofetamine HCl I 123 is below the limit of detection. Individual human variations in pharmacokinetics of this drug and the long-term effect on the eye have not been elucidated. The contents of the vial are radioactive. Adequate shielding of the preparation must be maintained at all times.

Do not use after the expiration time and date (6 hours after calibration time) stated on the label. Potassium Iodide Oral Solution should be administered before the examination to minimize thyroid uptake of Iodine 123.

The prescribed Iofetamine HCl I 123 dose should be administered as soon as practical from the time of receipt of the product (i.e., as close to calibration time or before, if possible), in order to minimize the fraction of radiation exposure due to relative increase of radionuclidic contaminants with time.

To minimize radiation dose to the bladder, the patient should be encouraged to drink fluids and void frequently.

SPECTAMINE, as well as other radioactive drugs, must be handled with care. Appropriate safety measures should be used to minimize radiation exposure to clinical personnel. Care should also be taken to minimize radiation exposure to the patient consistent with proper patient management. Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe use and handling of radionuclides, and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

Drug Interactions

There has been a single report of elevated diastolic hypertension (about 30 mm Hg) occurring 18 hours after administration of SPECTAMINE in a patient maintained on therapeutic doses of valproic acid.

Concurrent use of monoamine oxidase (MAO) inhibitors and compounds containing the amphetamine structure has been known to result in hypertensive crisis. Caution, therefore, should be exercised when administering SPECTAMINE (Iofetamine HCl I 123 Injection) to individuals taking medications known to potentiate the effects of sympathomimetic amines. It is recommended that SPECTAMINE not be administered during or within 14 days following administration of MAO inhibitors.

Sympathomimetic amines may affect the biodistribution of SPECTAMINE and, thus, may influence the image quality and diagnostic utility of the image.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term animal studies have been performed to evaluate carcinogenic potential, mutagenic potential or effects on fertility in male or female animals. The Ames test was negative for mutagenic effects.

Pregnancy Category C

Animal reproduction studies have not been conducted with SPECTAMINE. It is also not known whether SPECTAMINE can cause fetal harm when administered to a man or a pregnant woman or can affect reproduction capacity. SPECTAMINE should be given to a pregnant woman only if clearly needed.

Ideally, examinations using radiopharmaceuticals, especially those elective in nature, in women of childbearing capability, should be performed during the first few (approximately ten) days following the onset of menses.

Nursing Mothers

Since Iodine I 123 is excreted in human milk, formula feeding should be substituted for breast feeding if the agent must be administered to the mother during lactation.

Pediatric Use

Safety and effectiveness in children have not been established.

ADVERSE REACTIONS: In a clinical study in 93 patients with sudden onset of focal neurological deficit, e.g., cerebral infarction, 7 patients died within 2 to 55 days after administration. The deaths were considered to be a result of the disease state. Although there was no concurrent control group, statistics from historical controls support this evaluation.

There is evidence suggesting that the administration of 1 to 2 milligrams of Iofetamine HCl, the carrier in SPECTAMINE, may increase systolic blood pressure by about 10 mm Hg. In a patient with a history of hypertension, there has been a single report of sudden onset of hypertension and dizziness with transient chest tightness which occurred 5-10 minutes after administration of SPECTAMINE. One case of transient unilateral hearing loss also was reported several hours after the use of SPECTAMINE in a patient with a coincidental upper respiratory infection.

As with all organic-iodine-containing compounds, the possibility of allergic reactions must be considered.

HOW SUPPLIED: SPECTAMINE is supplied in nominal 3.5 ml vials as a sterile, apyrogenic, aqueous, isotonic sodium chloride solution for intravenous injection. Each milliliter contains 37 megabecquerels (1 mCi) of Iofetamine HCl I 123 at calibration time.

It is available in individual vials containing 111 megabecquerels (3 mCi) of Iofetamine HCl I 123 at calibration time in a volume of 3 ml.

Vials are packaged in individual lead shields with plastic outer container.

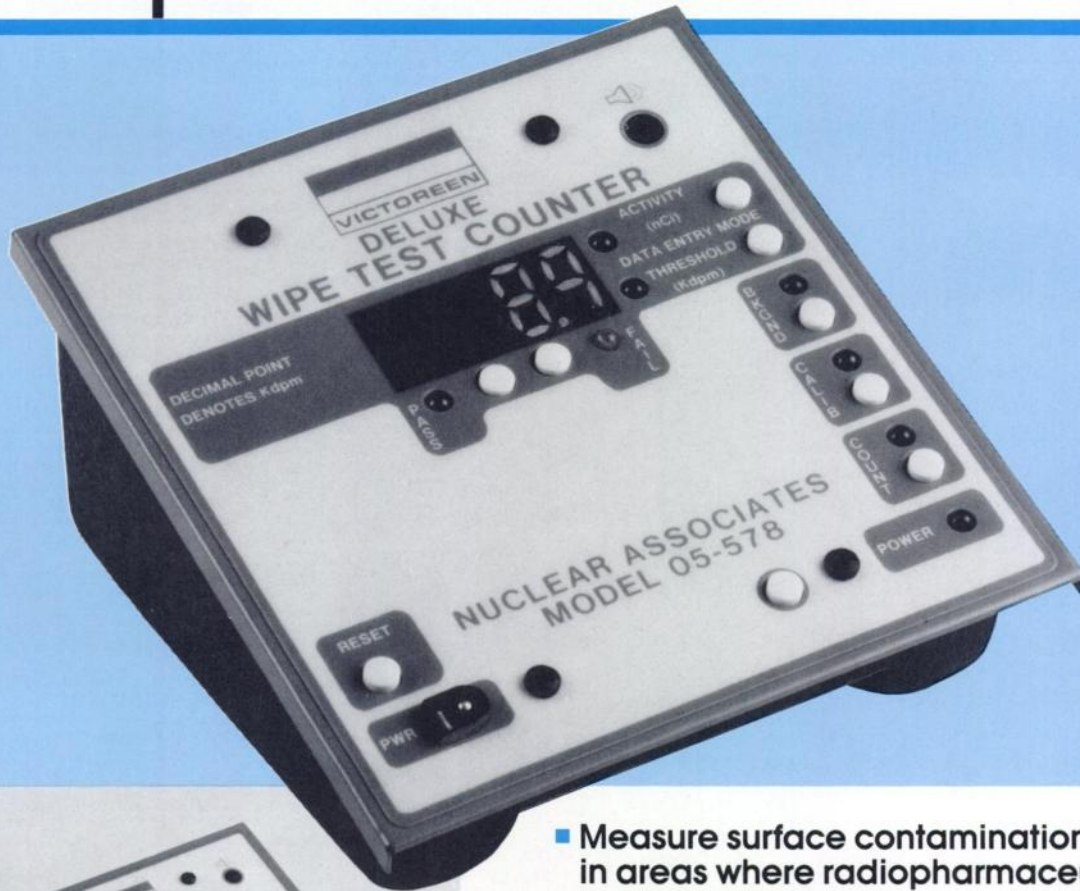
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The nerve injury illustration (Fig. 1) was shot on PolaBlue 35mm Instant Slide Film to provide a bright, high quality, white-on-blue slide in minutes. PolaBlue is the most cost-effective way to get white-on-blue slides of text and illustrations. And like all Polaroid 35mm instant films, PolaBlue can be exposed in virtually any 35mm camera and developed in minutes using the Polaroid 35mm PowerProcessor.

This fundus image (Fig. 2) was made with a fundus camera on Polaroid Colorgraph Type 691 Transparency film. This full-color film creates small format overhead transparencies in just minutes.

The X-ray copy of the pelvic area (Fig. 3) was made on PolaPan 35mm Instant Slide Film. PolaPan produces a black and white continuous tone image.

This slide of an ear illustration (Fig. 4) was made on PolaGraph 35mm Instant Slide Film. It produces a high contrast black and white slide in minutes.

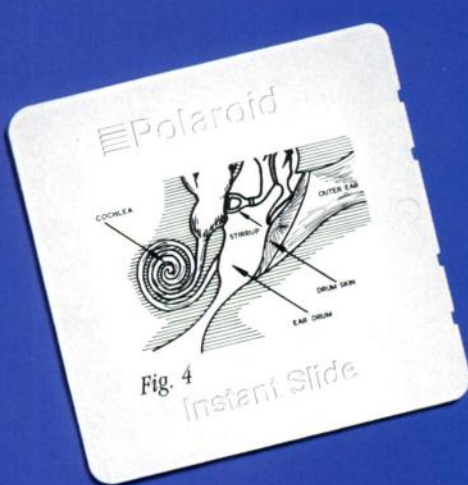
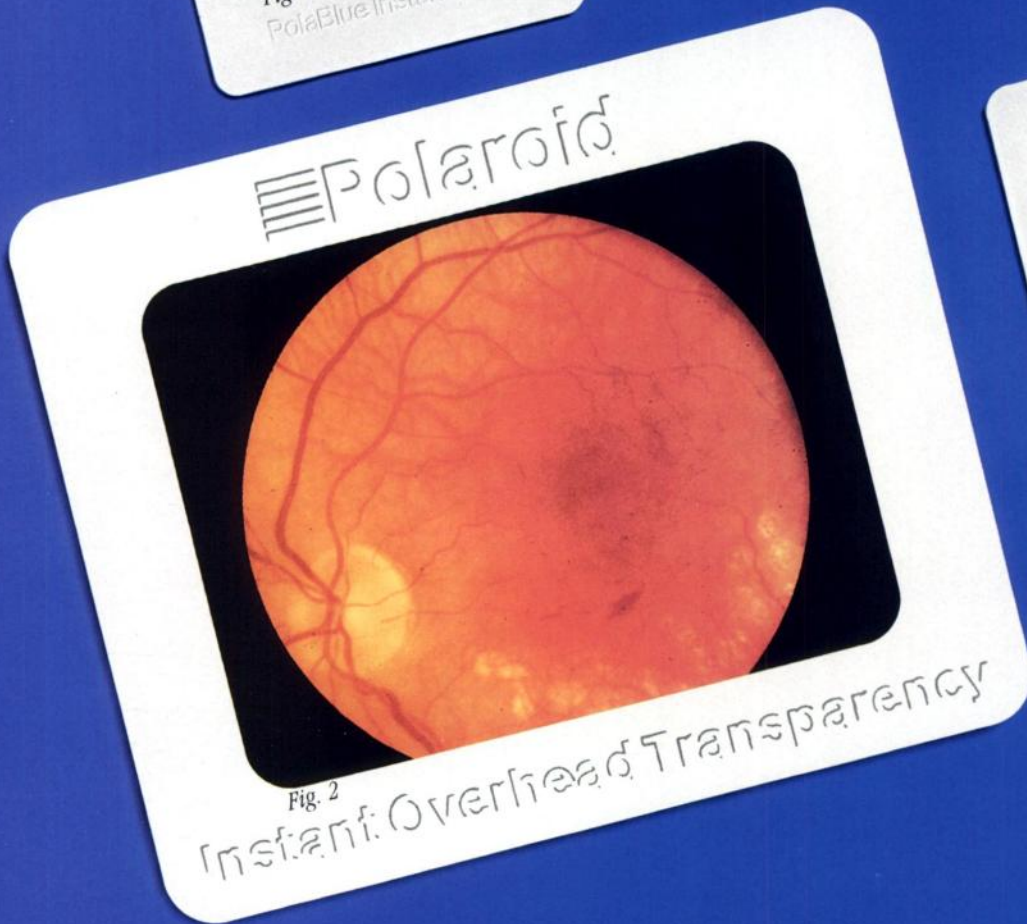
This close-up slide of a surgical procedure (Fig. 5) was made on High Contrast PolaChrome 35mm Instant Slide Film. This film is specifically balanced to give visuals rich, saturated color and pure whites.

This Doppler Color Flow Mapping image of a normal heart during systole (Fig. 6) was also captured on High Contrast PolaChrome 35mm Instant Slide Film using the Polaroid FreezeFrame Medical Video Image Recorder.

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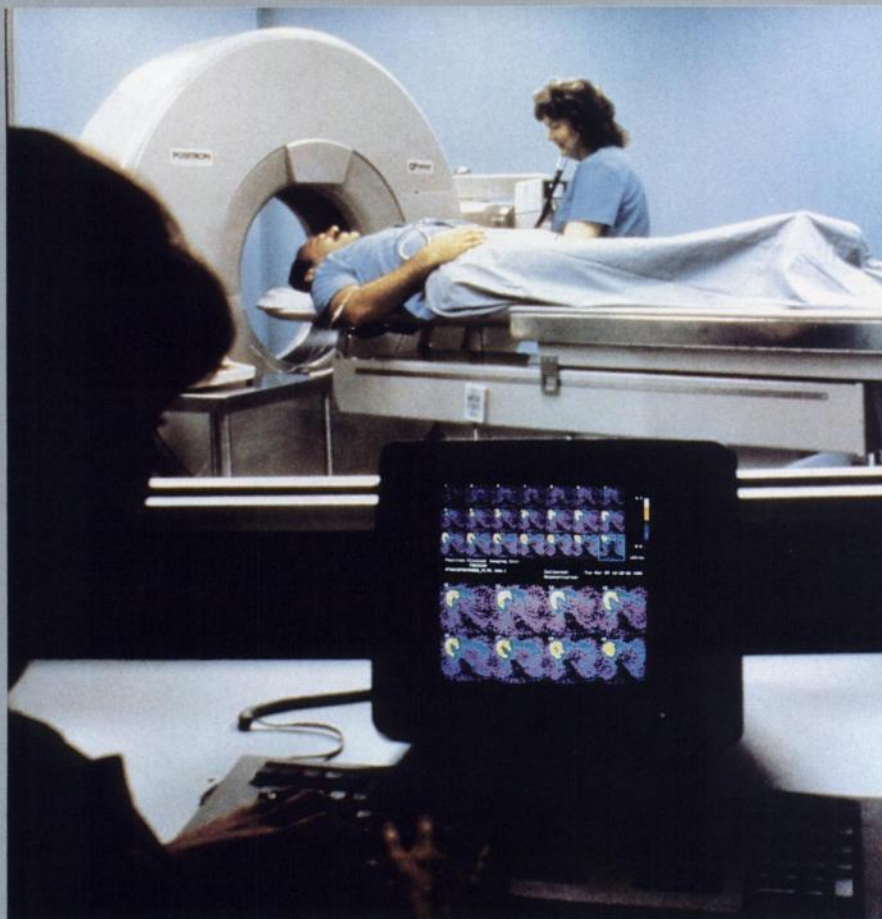


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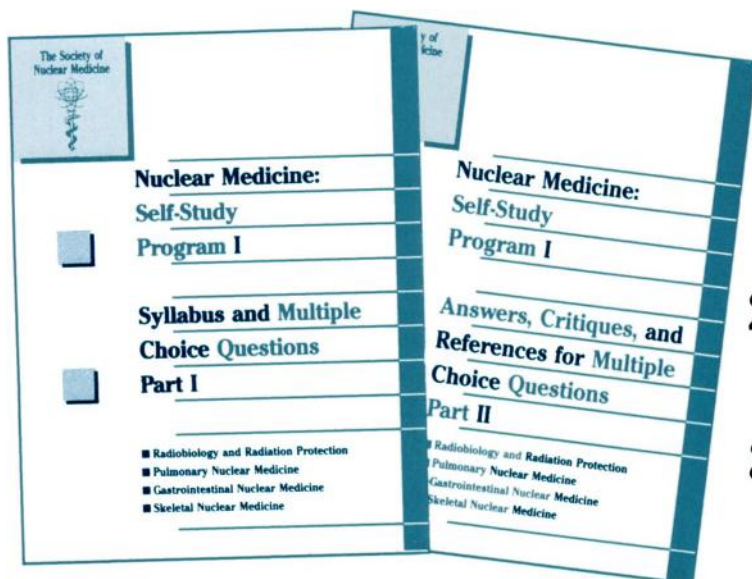
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- 3.** *Personal Psychometric Evaluation*—Available to those who return the answer form before **December 15, 1988**. Included here is a norms table that indicates your percentile ranking in each subject area by comparison with your peers.

The Society of Nuclear Medicine presents *Nuclear Medicine: Self-Study Program I*, the first volume of a comprehensive series that will cover all areas of nuclear medicine. It has been designed to help physicians, scientists, pharmacists, and technologists expand their knowledge of the clinical, basic science and technical aspects of nuclear medicine.

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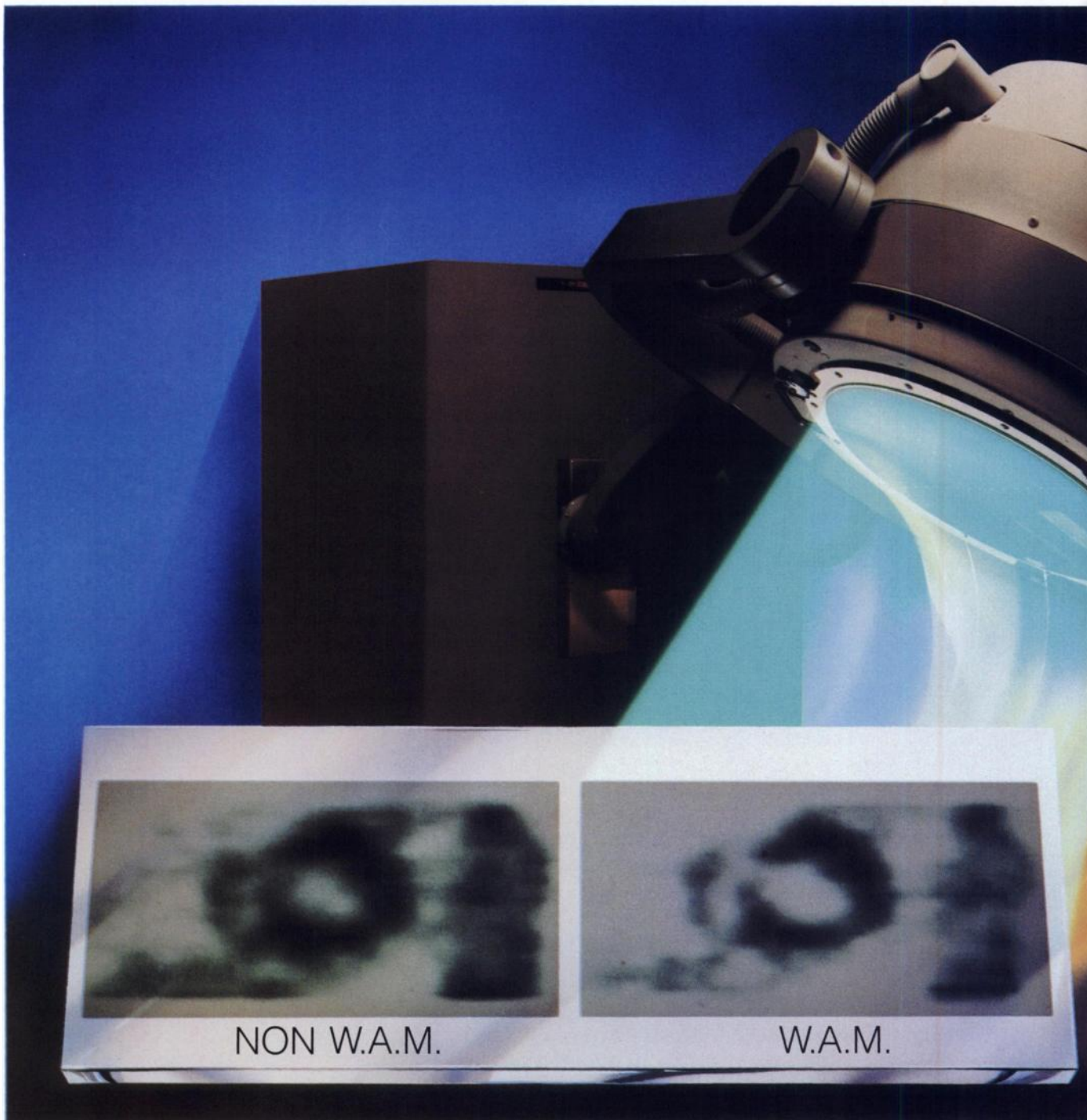
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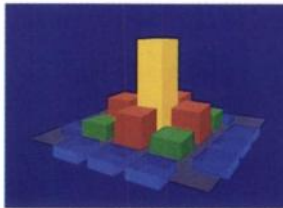
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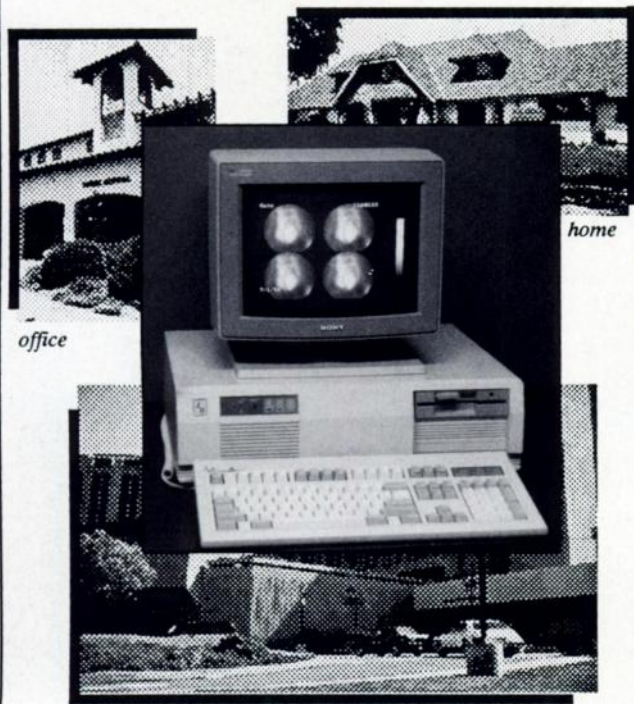
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BRIEF SUMMARY

THALLOUS CHLORIDE TI 201 INJECTION

Diagnostic - For Intravenous Use

DESCRIPTION—Thallous Chloride TI 201 Injection is supplied in an isotonic solution as a sterile, non-pyrogenic diagnostic radiopharmaceutical for intravenous administration. Each milliliter contains 37 megabecquerels (1 millicurie) Thallous Chloride TI 201 at calibration time, made isotonic with 9 milligrams sodium chloride and preserved with 0.9% (v/v) benzyl alcohol. The pH is adjusted to between 4.5 to 7.0 with hydrochloric acid and/or sodium hydroxide. Thallium TI 201 is cyclotron produced. At the time of calibration it contains no more than 1.0% Thallium TI 200, no more than 1.0% Thallium TI 202, no more than 0.25% radionuclides Leads and no less than 98% Thallium TI 201 as a percentage of total activity. No carrier has been added.

INDICATIONS AND USAGE—Thallous Chloride TI 201 may be useful in myocardial perfusion imaging and for the diagnosis and localization of myocardial infarction.

It may also be useful in conjunction with exercise stress testing as an adjunct in the diagnosis of ischemic heart disease (atherosclerotic coronary artery disease).

It is usually not possible to differentiate recent from old myocardial infarction, or to differentiate exactly between recent myocardial infarction and ischemia.

CONTRAINDICATIONS—None known.

WARNINGS—When studying patients suspected or known to have myocardial infarction or ischemia, care should be taken to assure continuous clinical monitoring and treatment in accordance with safe, accepted procedures. Exercise stress testing should be performed only under the supervision of a qualified physician and in a laboratory equipped with appropriate resuscitation and support apparatus.

PRECAUTIONS—Data are not available concerning the effect on the quality of Thallous Chloride TI 201 images of marked alterations in blood glucose, insulin or pH (such as is found in diabetes mellitus). Attention is directed to the fact that thallium is a potassium analog, and since the transport of potassium is affected by these factors, the possibility exists that the Thallous Chloride TI 201 may likewise be affected.

General—This drug should not be used after six (6) days from the calibration date, or nine (9) days from date of manufacture, whichever comes first.

As in the use of any radioactive material, care should be taken to minimize radiation exposure to the patient consistent with proper management and to insure minimum radiation exposure to occupational workers.

Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe handling of radionuclides and whose experience and training have been approved by the appropriate governmental agency authorized to license the use of radionuclides.

Carcinogenesis, Mutagenesis, Impairment of Fertility—No long-term animal studies have been performed to evaluate carcinogenic potential, mutagenic potential or whether this drug affects fertility in males or females.

Pregnancy Category C—Animal reproductive studies have not been conducted with Thallous Chloride TI 201. It is also not known whether Thallous Chloride TI 201 can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. Thallous Chloride TI 201 should be given to a pregnant woman only if clearly needed.

Ideally, examination using radiopharmaceutical drug products — especially those elective in nature — of women of childbearing capability should be performed during the first ten days following the onset of menses.

Nursing Mothers—It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, as a general rule nursing should not be undertaken when a patient is administered radioactive material.

Pediatric Use—Safety and effectiveness in children below age 18 have not been established.

ADVERSE REACTIONS—A single adverse reaction to the administration of Thallous Chloride TI 201 has been reported consisting of hypotension accompanied by pruritus and a diffuse rash which responded to antihistamines and steroids within one hour.

DOSAGE AND ADMINISTRATION—The recommended adult (70 kg) dose of Thallous Chloride TI 201 is 37 to 74 MBq (1 to 2 mCi). Thallous Chloride TI 201 is intended for intravenous administration.

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration, whenever solution and container permit. Do not use if contents are turbid.

Waterproof gloves should be worn during the handling procedures.

The patient dose should be measured by a suitable radioactivity calibration system immediately prior to administration.

With a shielded sterile syringe, aseptically withdraw the material for use.

For resting Thallous Chloride TI 201 studies, imaging should begin 10 to 20 minutes after injection. Myocardial-to-background ratios are improved when patients are injected upright and in the fasting state; the upright position reduces the hepatic and gastric Thallium TI 201 concentration.

When utilized in conjunction with exercise stress testing, Thallous Chloride TI 201 should be administered at the inception of a period of maximum stress which is sustained for approximately 30 seconds after injection. Imaging should begin within ten minutes after administration to obtain maximum target-to-background ratios. Several investigators have reported that within two hours after the completion of stress testing the target-to-background ratios may decrease significantly in lesions that are attributable to transient ischemia.

HOW SUPPLIED—Thallous Chloride TI 201 is supplied in a sterile, non-pyrogenic solution for intravenous administration. Each ml contains 37 MBq (1 mCi) Thallous Chloride TI 201 at calibration time, 9 mg sodium chloride and 0.9 percent (v/v) benzyl alcohol. The pH is adjusted to between 4.5 to 7.0 with hydrochloric acid and/or sodium hydroxide solution. Vials are available in the following quantities of radioactivity: 74, 111, 148, 296, and 333 megabecquerels (2, 3, 4, 8, and 9 millicuries) of Thallium TI 201.



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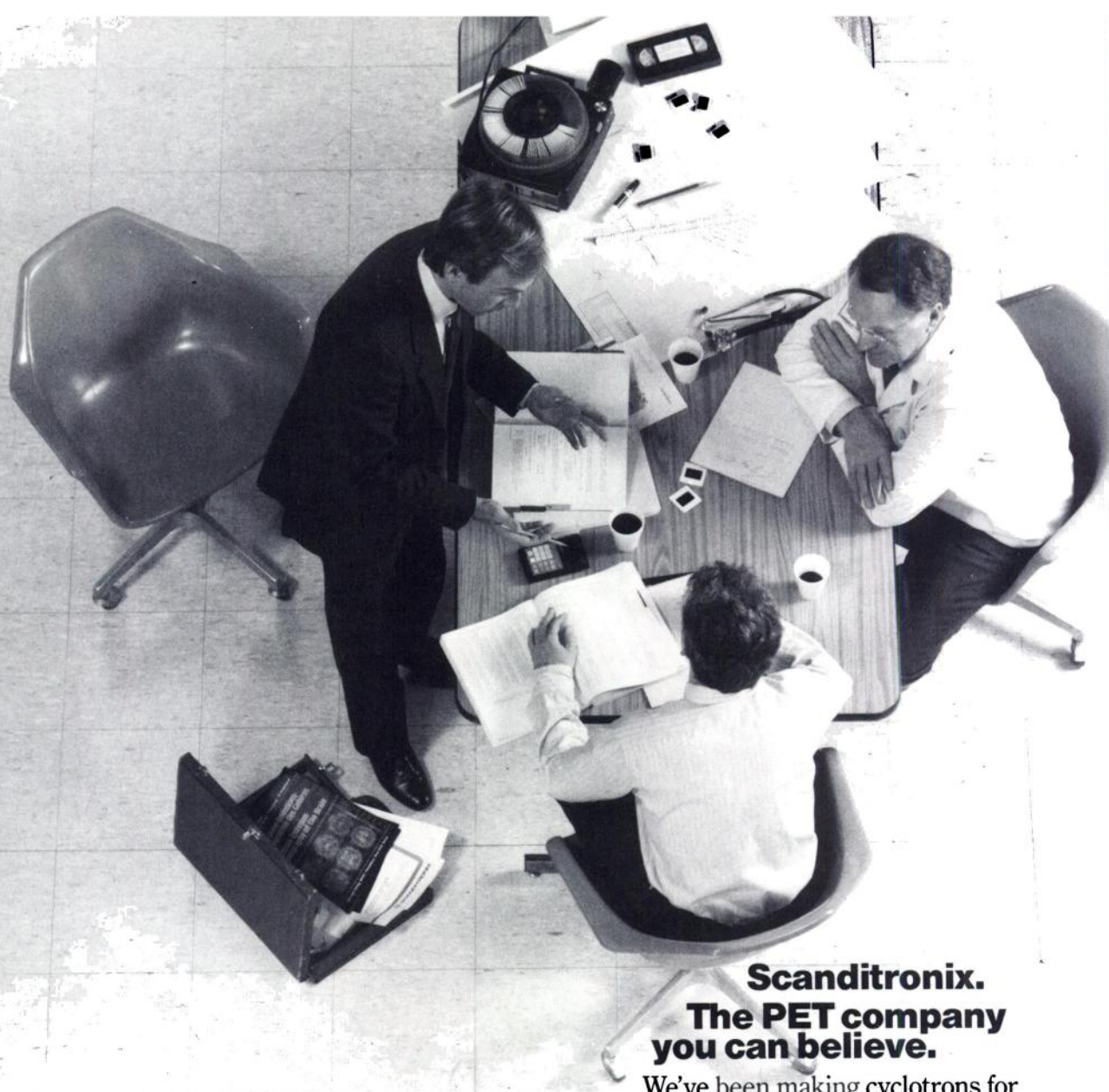
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*Includes 10% overfill. Please see next page for brief summary.



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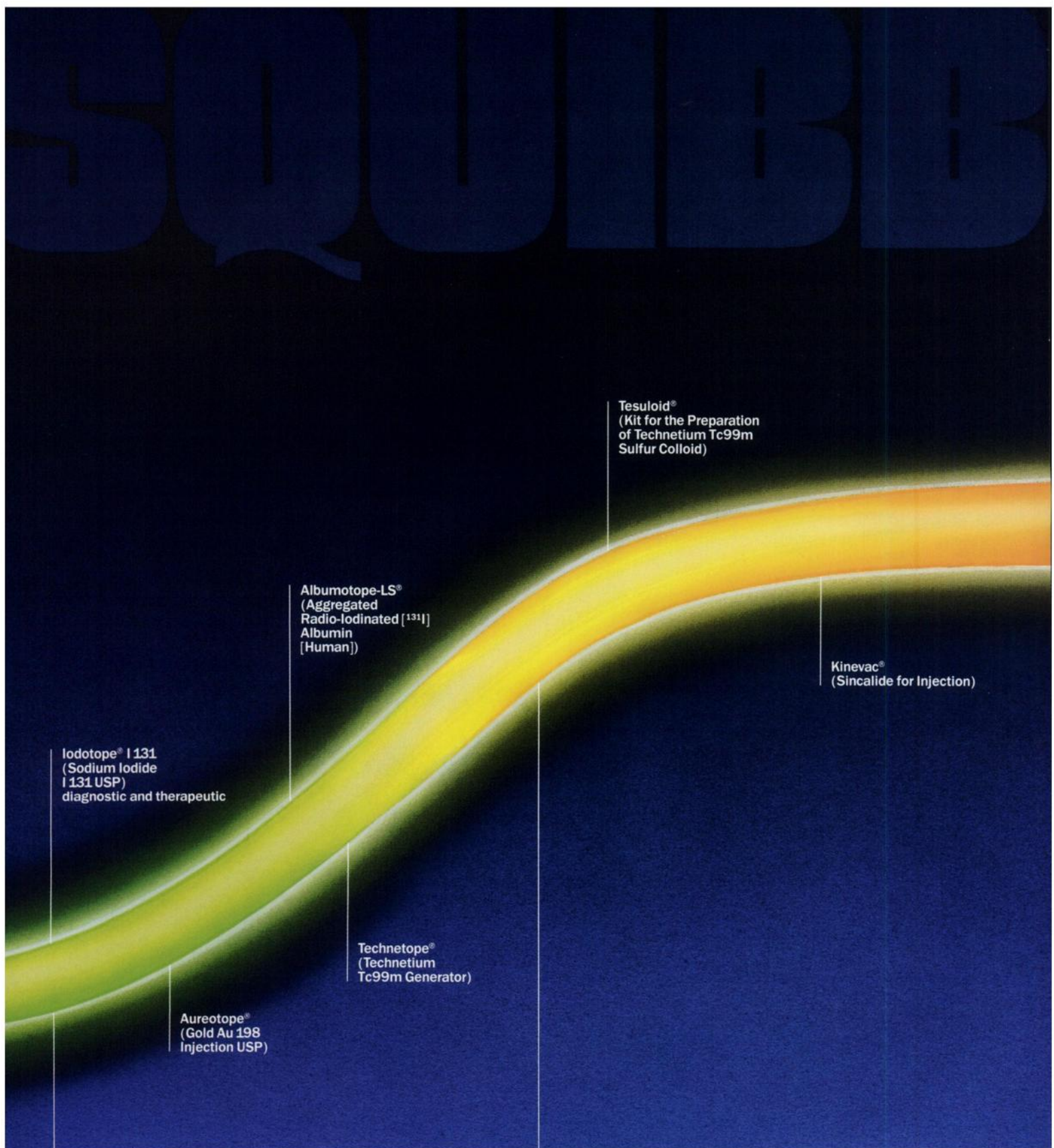
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SQUIBB



A Profile of Progress in Nuclear Medicine



The Years of Growth

Nuclear medicine emerged from the experimental stage into a phase of rapid clinical growth. The number of procedures performed rose rapidly during the 1960s. During this same period, Squibb Diagnostics developed

and introduced important products and services for nuclear medicine, including the first sterile technetium generator, nuclear medicine training seminars and technical support through the Technical Associates Program.

The Years of Refinement

The '70s saw the development of other imaging modalities which drew procedures away from nuclear medicine and slowed its growth. Developments and advances continued, however, and Squibb

introduced a variety of radiopharmaceutical products, including Macrotec. Squibb's Choletec* was introduced in 1987, and quickly became the premier hepatobiliary imaging agent.

Nuclear Medicine: A Distinguished Past, A Promising Future

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Preparation of
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New heart imaging agents

New brain imaging agent

The Years of Promise
The future of nuclear medicine is bright, and Squibb's contributions to it continue. New Squibb brain and heart agents are now in clinical development. In addition to extensive research and development, the Squibb

contribution to nuclear medicine continues with technical support and professional education programs.

*See brief summary on following page.

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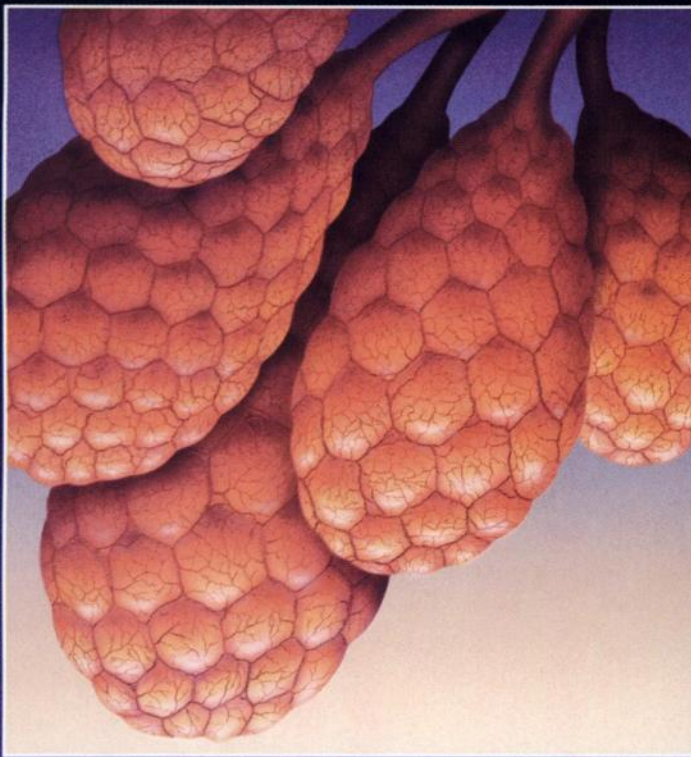


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*Callahan RJ, Swanson DP et al. A multi-institutional in vitro evaluation of commercial Tc 99m Macroaggregated Albumin Kits. *J Nucl Med Tech*;14:(No. 4)206-209, 1986

MACROTEC demonstrated the highest radiochemical purity (RCP) as measured by supernatant activity

LOW SUPERNATANT ACTIVITY—for high target-to-background ratios

OPTIMAL PARTICLE SIZE—more than 90% of particles in the 10-90 micron range for diagnostic efficacy and reduced activity in the liver and nontarget areas

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A UNIQUE INDICATION—the only MAA product indicated for use in isotopic venography

MACROTEC[®] Kit for the Preparation of Technetium Tc 99m Albumin Aggregated Diagnostic—For Intravenous Use

DESCRIPTION

Macrotec is a sterile, nonpyrogenic, lyophilized preparation of albumin aggregated. Each 5 mL vial of Macrotec contains 1.5 mg of Albumin Aggregated, 10.0 mg Albumin Human, 0.07 mg (minimum) stannous chloride ($\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$) and 0.19 mg total tin, maximum (as stannous chloride, $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$), 1.8 mg of sodium chloride with trace amounts of sodium acetate, acetic acid and hydrochloric acid. Macrotec contains no preservatives. The pH of the reconstituted product is between 3.8 and 8.0.

The aggregated particles are formed by denaturation of Albumin Human in a heating and precipitation process. Each vial contains 2-7 million particles, 90% of which are between 10 and 90 microns in size. The average size is 20 to 40 microns; no particles are greater than 150 microns.

Reconstitution of Macrotec with sterile sodium pertechnetate Tc 99m forms an aqueous suspension of Technetium Tc 99m Albumin Aggregated for diagnostic use by intravenous injection. No less than 90% of the pertechnetate Tc 99m added to the reaction vial is bound to the aggregates at preparation time and remains bound throughout the 6-hour lifetime of the suspension.

INDICATIONS AND USAGE

Lung Imaging

Macrotec (Kit for the Preparation of Technetium Tc 99m Albumin Aggregated) is a lung imaging agent which may be used as an adjunct in the evaluation of pulmonary perfusion in adults and children. It is useful in the early detection of pulmonary emboli and in the evaluation of the status of the pulmonary circulation in such conditions as pulmonary neoplasm, pulmonary tuberculosis and emphysema.

Isotopic Venography

Macrotec is also indicated for use in isotopic venography as an adjunct in the screening, diagnosis and management of deep vein thrombosis in the lower extremities.

Combined isotopic venography of the lower extremities and the pulmonary vasculature may be performed.

CONTRAINDICATIONS

Technetium Tc 99m Albumin Aggregated Injection should not be administered to patients with severe pulmonary hypertension.

The use of Technetium Tc 99m Albumin Aggregated Injection is contraindicated in persons with a history of hypersensitivity reactions to products containing human serum albumin.

WARNINGS

The literature contains reports of deaths occurring after the administration of Albumin Aggregated to patients with pre-existing severe pulmonary hypertension. Instances of hemodynamic or idiosyncratic reactions to preparations of Technetium Tc 99m Albumin Aggregated have been reported.

PRECAUTIONS

General

In patients with right to left heart shunts, additional risk may exist due to the rapid entry of Albumin Aggregated into the systemic circulation. The safety of this agent in such patients has not been established.

Hypersensitivity reactions are possible whenever protein-containing materials such as pertechnetate labeled Albumin Aggregated are used in man. Epinephrine, antihistamines and corticosteroids should be kept available for immediate use.

The intravenous administration of any particulate material such as Albumin Aggregated imposes a temporary, small mechanical impediment to blood flow. While this effect is probably physiologically insignificant in most patients, the administration of Albumin Aggregated is possibly hazardous in acute cor pulmonale and other states of severely impaired pulmonary blood flow.

The components of the Macrotec (Kit for the Preparation of Technetium Tc 99m Albumin Aggregated) are sterile and nonpyrogenic. It is essential to follow directions carefully and adhere to strict aseptic procedures during preparation.

Contents of the vial are intended only for use in the preparation of Technetium Tc 99m Albumin Aggregated Injection and are NOT to be administered directly to the patient.

The contents of the kit before preparation are not radioactive. However, after the sodium pertechnetate Tc 99m is added, adequate shielding of the final preparation must be maintained.

The technetium Tc 99m labeling reactions involved depend on maintaining the stannous ion in the reduced state. Hence, sodium pertechnetate Tc 99m containing oxidants should not be employed.

The preparation contains no bacteriostatic preservative. Technetium Tc 99m Albumin Aggregated Injection should be stored at 2-8°C and discarded 6 hours after formulation.

Technetium Tc 99m Albumin Aggregated Injection is a physically unstable suspension and consequently the particles settle with time. Failure to agitate the vial adequately before use may result in non-uniform distribution of radioactive particles.

If blood is drawn into the syringe, unnecessary delay prior to injection may result in clot formation.

Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe use and handling of radionuclides and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

As in the use of any other radioactive material, care should be taken to minimize radiation exposure to patients consistent with proper patient management, and to minimize radiation exposure to clinical personnel.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term animal studies have been performed to evaluate carcinogenic potential or whether Technetium Tc 99m Albumin Aggregated Injection affects fertility in males or females.

Pregnancy Category C

Animal reproduction and teratogenicity studies have not been conducted with Technetium Tc 99m Albumin Aggregated Injection. It is also not known whether Technetium Tc 99m Albumin Aggregated Injection can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. There have been no studies in pregnant women. Technetium Tc 99m Albumin Aggregated Injection should be given to a pregnant woman only if clearly needed.

Ideally, examinations using radiopharmaceuticals, especially those elective in nature, in women of childbearing capability, should be performed during the first few (approximately 10) days following the onset of menses.

Nursing Mothers

Technetium Tc 99m is excreted in human milk during lactation. Therefore, formula-feedings should be substituted for breast-feedings.

Pediatric Use

The lowest possible number of particles should be used in the right-to-left shunting, in neonates and in severe pulmonary disease.

ADVERSE REACTIONS

Although adverse reactions specifically attributable to the Technetium Tc 99m Albumin Aggregated Injection have not been noted, the literature contains reports of deaths occurring after the administration of Albumin Aggregated to patients with pre-existing severe pulmonary hypertension. Instances of hemodynamic or idiosyncratic reactions to preparations of Technetium Tc 99m Albumin Aggregated have been reported.

HOW SUPPLIED

Macrotec (Kit for the Preparation of Technetium Tc 99m Albumin Aggregated) is supplied as a kit containing 10 reaction vials (5 mL size), 10 pressure sensitive labels and 1 package insert. (J3-436L)

CIRCLE 67 ON READER SERVICE CARD

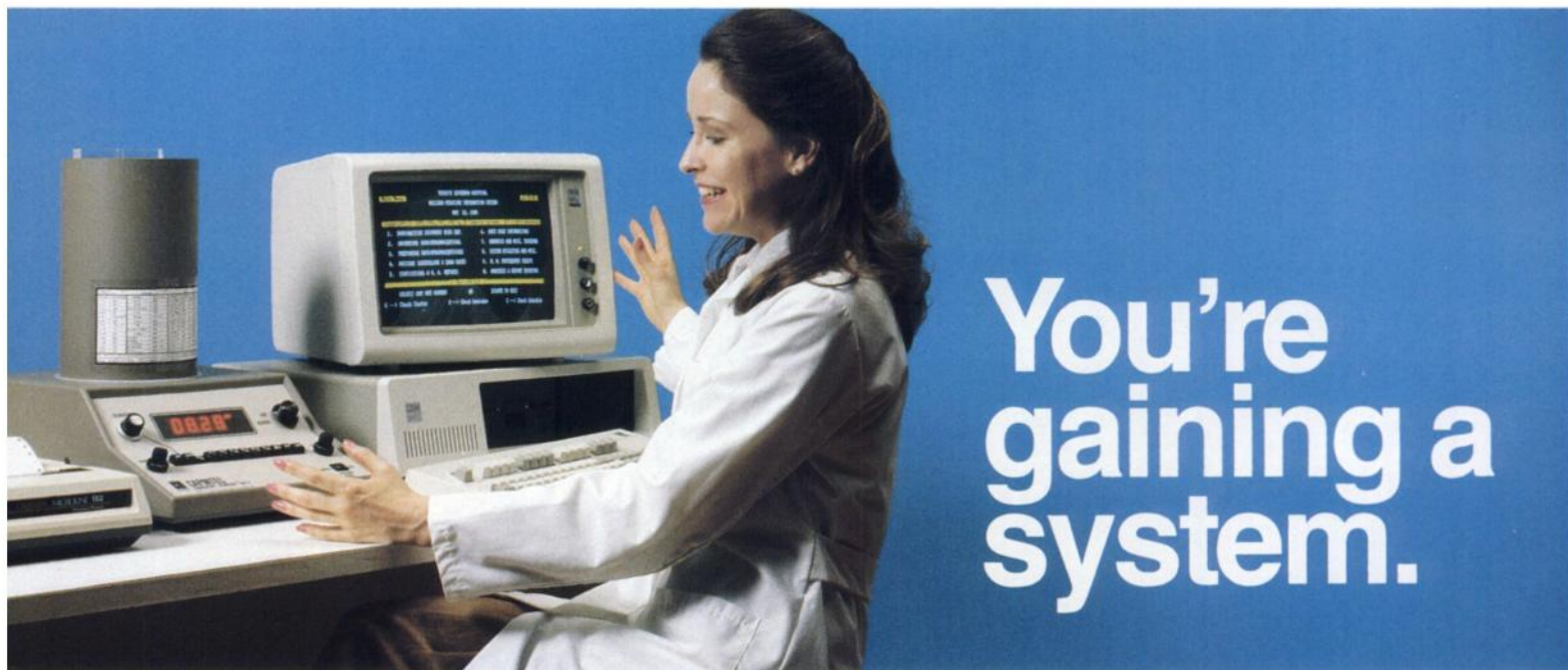
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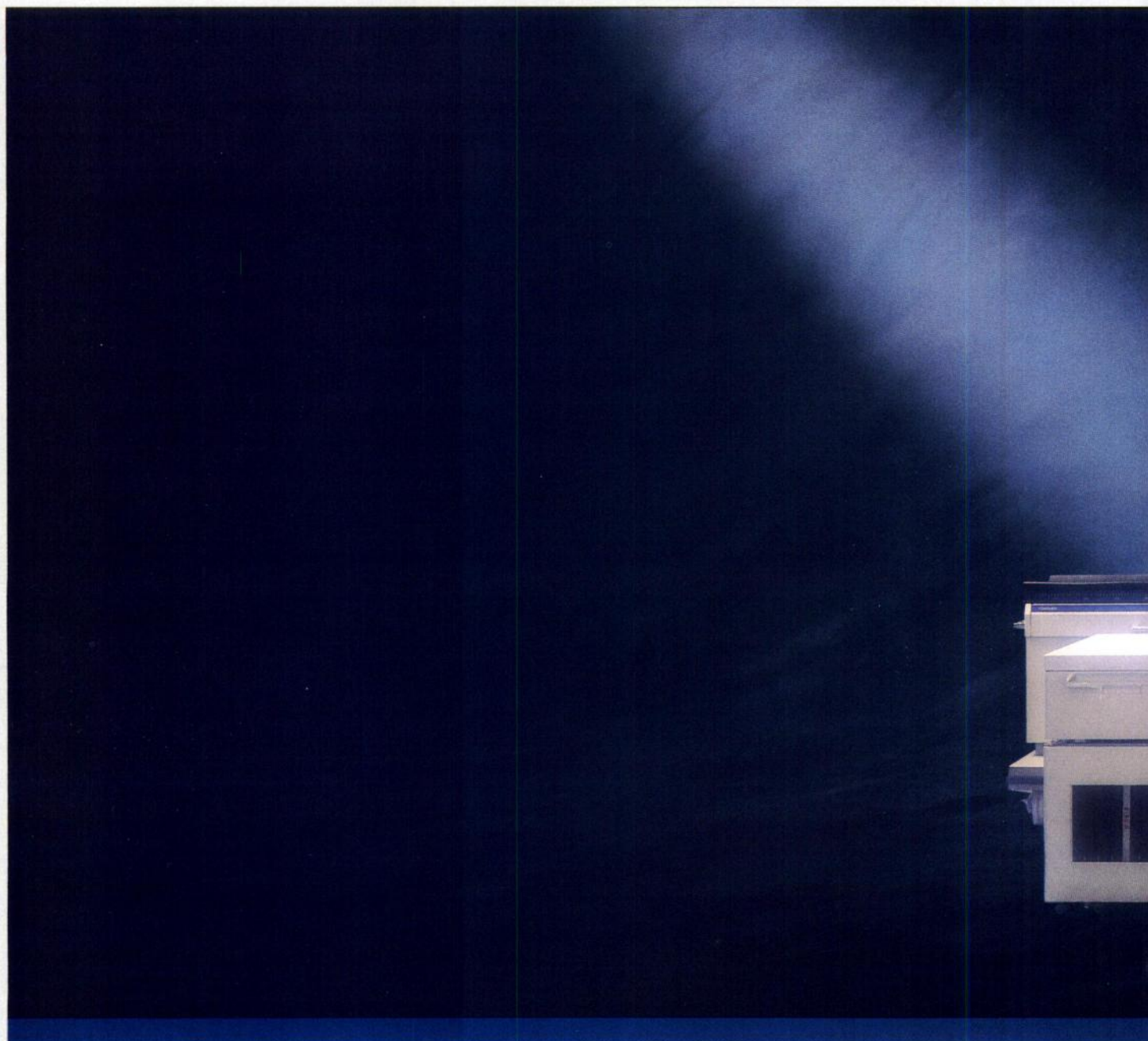


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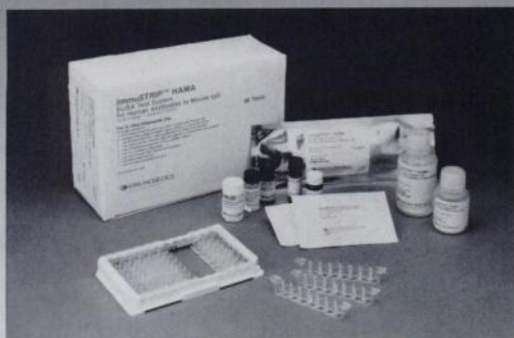
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Chromatography of Technetium-99m Radiopharmaceuticals

—A Practical Guide

By Philip J. Robbins

To provide up-to-date information about the most accurate procedures for ensuring quality control of radiopharmaceuticals, The Society of Nuclear Medicine has published *Chromatography of Technetium-99m Radiopharmaceuticals—A Practical Guide*.

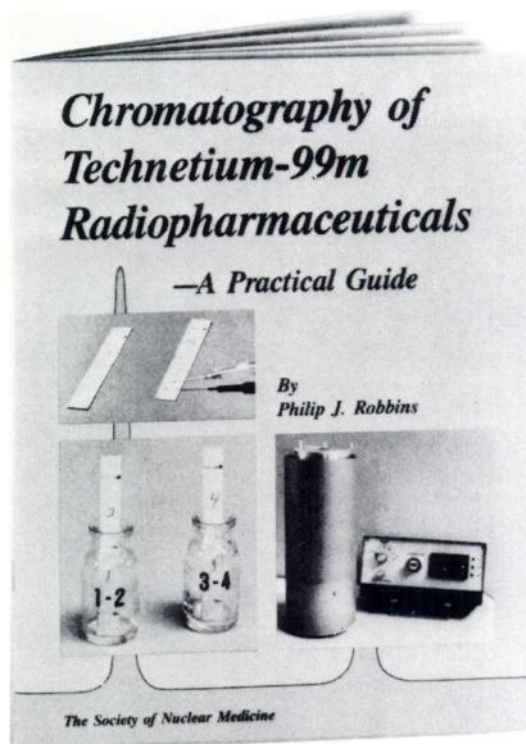
This important manual offers readers a collection of miniaturized chromatographic methods for the rapid and precise determination of the radiochemical purity of commonly used Tc-99m radiopharmaceuticals.

Topics covered include the nature and source of impurities, principles and classic techniques of chromatography, methods for counting miniature chromatographic strips, and pitfalls of miniature methods and how to avoid them. Also contained herein is a listing of each radiopharmaceutical with the USP criteria for radiochemical purity, typical scans of impure products, and standards and inter-laboratory comparisons for miniaturized systems.

Prepared to aid nuclear medicine personnel in implementing voluntary quality-assurance programs, the material may also be used as a training resource for individuals preparing for professional licensure and certification.

Ordering Information:

Add \$2.50 postage and handling for each book ordered. Prepayment required in U.S. funds drawn on U.S. banks only. For payments made in U.S. dollars, but drawn on a foreign bank, add a bank processing fee of \$4.50 for Canadian bank drafts or \$40.00 for all other foreign bank drafts. Check or purchase order must accompany all orders. Make checks payable to: The Society of Nuclear Medicine. *Prices are in U.S. dollars and are subject to change without notice.*



8½ × 11" softcover, 48 pages

\$ 8.00 SNM members;

\$10.00 non-members

Publication Date: January 1984

**The Society of Nuclear Medicine, Book Order Dept. 1287J,
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The Society of
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SNM
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Tuesday, June 13–
Friday, June 16, 1989

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Call for Abstracts for Scientific Papers Call for Abstracts for Scientific Exhibits

The 1989 Scientific Program Committee and Scientific Exhibits Subcommittee solicits the submission of abstracts from members and nonmembers of The Society of Nuclear Medicine for the 36th Annual Meeting in St. Louis. Abstracts accepted for the program will be published in a special supplement to the May issue of *The Journal of Nuclear Medicine*. Original contributions on a variety of topics related to nuclear medicine will be considered, including:

- INSTRUMENTATION
- COMPUTERS AND DATA ANALYSIS
- RADIOASSAY
- RADIOPHARMACEUTICAL CHEMISTRY
- DOSIMETRY/RADIOBIOLOGY
- NUCLEAR MAGNETIC RESONANCE
- CLINICAL SCIENCE APPLICATIONS
- Bone/Joint
- Cardiovascular
- Endocrine
- Gastroenterology
- Infectious Disease and Immunology
- Neurology
- Oncology/Hematology
- Pediatrics
- Pulmonary
- Renal/Hypertension

Authors seeking publication for the full text of their papers are strongly encouraged to submit their work to the *JNM* for immediate review.

The official abstract form may be obtained from the September, 1988 issue of the *JNM* or by calling or writing:

The Society of Nuclear Medicine
Att: Abstracts
136 Madison Avenue, New York, NY 10016-6760
Tel: (212)889-0717
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Deadline for receipt of abstracts for Scientific Exhibits is Thursday, January 19, 1989.*

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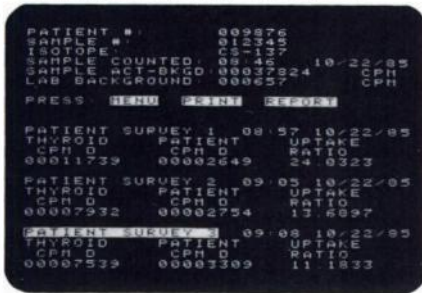
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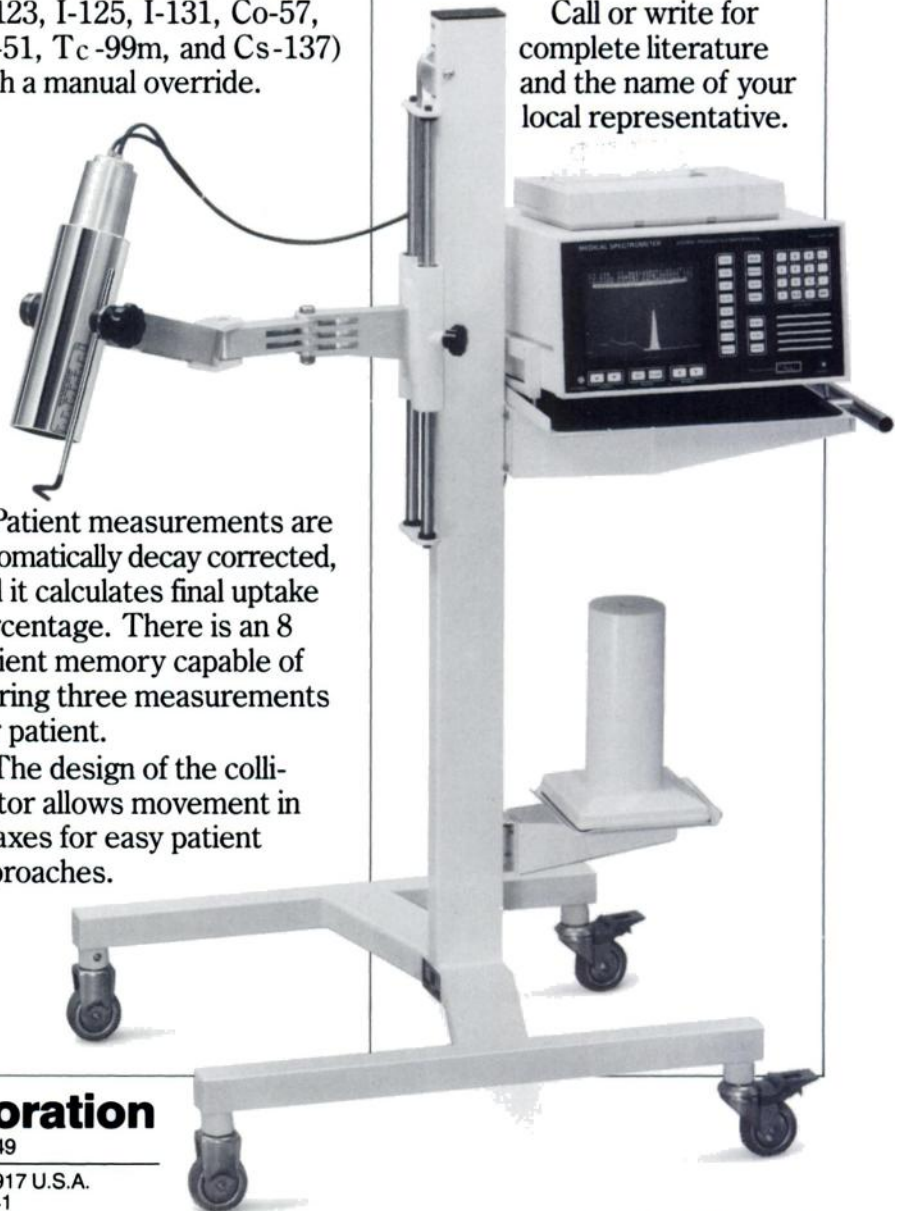
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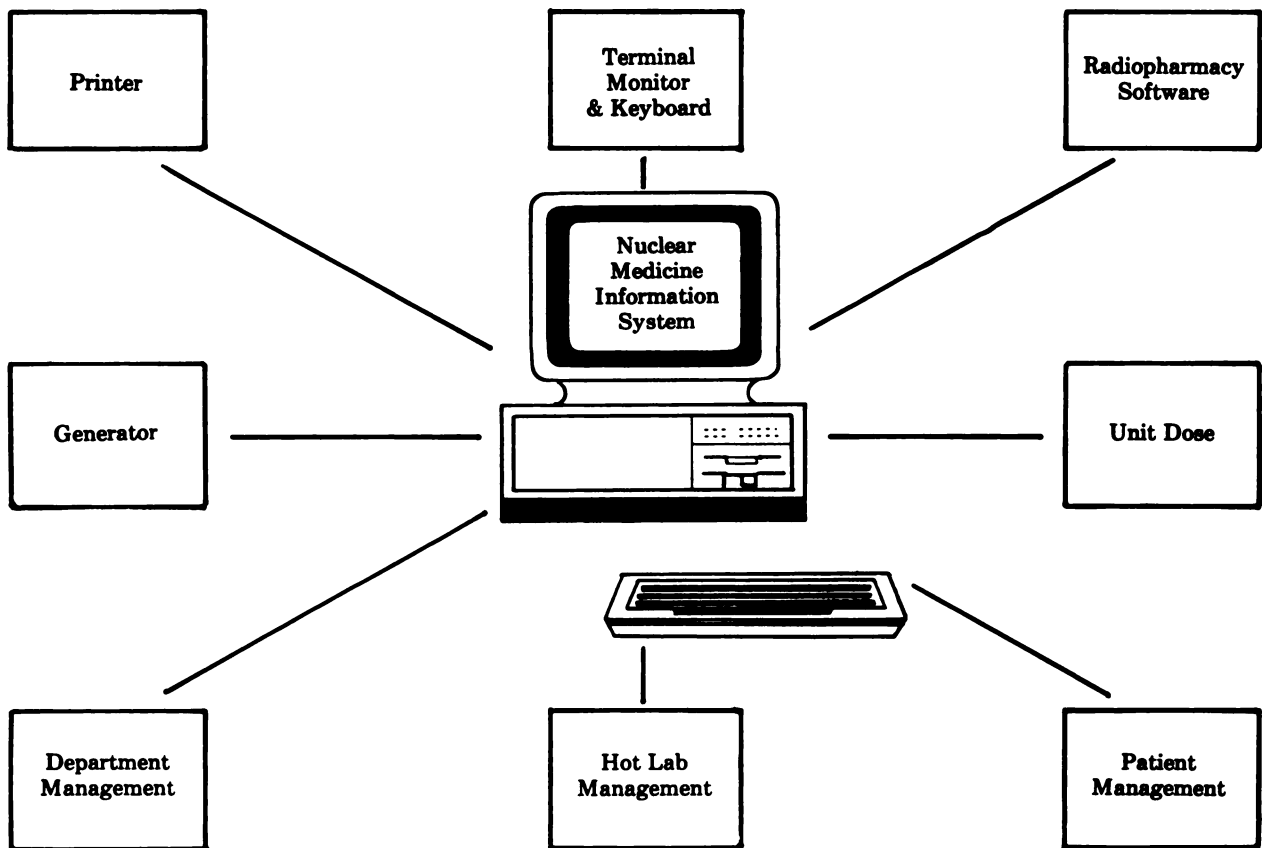
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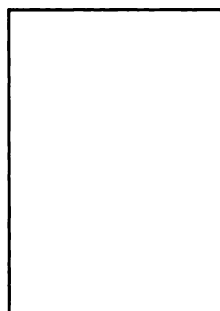
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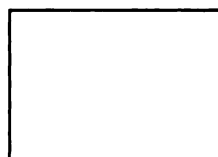
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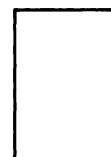
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RADIOLOGIST with additional training in nuclear medicine needed to join seven-member fee-for-service radiology group practicing in medical school-affiliated hospital. Applicant will help direct a laboratory performing 8,500 examinations per year on five gamma cameras (two SPECT cameras). Clinical research and teaching are encouraged in this division that trains both diagnostic radiology residents and nuclear medicine fellows. Position which leads to partnership in the group is available immediately. Contact: Keith C. Fischer, MD, Chief, Division of Nuclear Medicine, The Jewish Hospital of St. Louis, Washington Univ. Medical Center, 216 S. Kingshighway, St. Louis, MO 63110. EOE.

Washington, DC area. Board certified RADIOLOGIST with interest and/or experience in nuclear medicine to join eight radiologists practicing all aspects of imaging, including MRI, in a 400-bed hospital and office setting. Nuclear cardiology expertise desirable. Submit CV to: Michael H. Friedman, MD, Chairman, Dept. of Radiology, The Alexandria Hospital, 4320 Seminary Rd., Alexandria, VA 22304. EOE.

Technologist

NUCLEAR MEDICINE TECHNOLOGISTS. STAT Tech Services, Inc. has full-time positions available for Registered (ARRT) or Certified (NMTCB) NMT's in our Greater Los Angeles/Orange County region. We offer full benefits including medical and dental coverage with the opportunity for professional growth and career advancement as you work with state-of-the-art (SPECT) equipments. Since we are an in-house, full service nuclear medicine, and ultrasound imaging company, there are no portables to push. Contact: STAT Tech Services, Inc., 5315 Laurel Canyon Blvd. #102, North Hollywood, CA 91607 (818)766-0840. EOE.

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Albany Memorial Hospital is seeking a Chief **NUCLEAR MEDICINE TECHNOLOGIST** to oversee operations of its busy nuclear medicine department. Equipment includes three gamma cameras with full computerization and SPECT. Extensive experience in computer work as well as baccalaureate education in a health science field is preferred. This is an opportunity to lead a dynamic growing service in a department that is known for excellence. Salary negotiable based on experience and education. Send resume in confidence to: David J. Fanning, Manager, Medical Imaging, Albany Memorial Hospital, 600 Northern Blvd., Albany, NY 12204; (518)471-3280. EOE.

NUCLEAR MEDICINE TECHNOLOGIST. Immediate full-time positions available for registered nuclear medicine technologists at the Cox Medical Center South, which is a new, 510-bed general hospital in Springfield, MO, located in the vacation land of the Ozark Mountains. Excellent salary and benefits are offered. Please contact: Personnel Office, Cox Medical Center South, 3801 S. National Ave., Springfield, MO 65807; (417)885-6125. EOE.

NUCLEAR MEDICINE TECHNOLOGISTS. The Queen's Medical Center, a 506-bed acute care teaching facility located in downtown Honolulu has immediate full-time positions available for nuclear medicine technologists. Qualified candidates must be registered (ARRT, NMTCB) or registry eligible nuclear medicine technologists. Our large, newly constructed progressive department offers state-of-the-art equipment including multiple SPECT camera/computer systems. Enjoy all your outdoor activities year-round with our warm and temperate climate. Interested applicants may call collect, Jerrie Balsai, Employment Specialist, (808)547-4355, or send resume to: The Queen's Medical Center, Human Resources Division, 1381 Punchbowl St., Honolulu, HI 96813. EOE.

NUCLEAR MEDICINE TECHNOLOGIST. Huntington Memorial Hospital, a 606-bed acute care facility, has a part-time opening. The successful candidate will have ARRT or NMTCB. We offer an excel-

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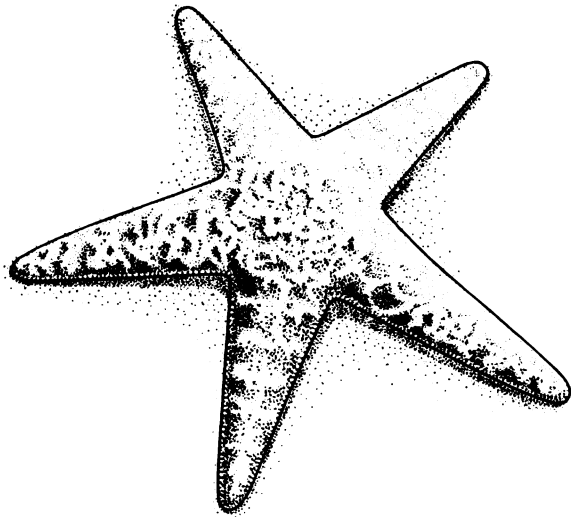
NUCLEAR MEDICINE TECHNOLOGIST. SwedishAmerican Hospital is seeking a staff nuclear medicine technologist to be the third member of our busy nuclear medicine department. We are a 427-bed general acute care hospital designated as the Radiation Oncology Center for this region of Illinois. As part of our department you will be involved with all imaging, including nuclear cardiology and related duties. This position requires a registered or registry eligible nuclear medicine tech. We offer a starting salary commensurate with experience, additional pay for on-call and excellent benefits including flexible paid time off. Rockford, Illinois, a city of 140,000, is located in north-central Illinois. For further information, contact: Faith Reese, Personnel Dept., SwedishAmerican Hospital, 1400 Charles St., Rockford, IL 61104; (815) 966-2080. Equal Opportunity Employer m/f.

NUCLEAR MEDICINE TECHNOLOGIST. Position available for technologist at the VA Medical Center, Syracuse, NY. Located in Central New York adjacent to the Syracuse University Campus and affiliated with the SUNY Health Science Center. We offer competitive benefits and salary. Call our Personnel Dept. at (315)476-3950 or send resume to: VA Medical Center, Personnel (05), 800 Irving Ave., Syracuse, NY 13210. An equal opportunity employer.

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Northwestern Memorial Hospital
250 East Superior Street
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The Liverpool Hospital is situated in The South Western Region of Sydney, 30 km from the centre of Sydney. The Department caters for the needs of a 460 bed Hospital and offers a wide range of Nuclear Medicine procedures, including computerised studies, real-time ultrasound and radio-immunoassays.

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Applicants should be accredited or be eligible for accreditation by the Australian and New Zealand Society of Nuclear Medicine. Reciprocity exists with Canadian Association of Medical Radiation Technologists.

Applications in writing giving full details of qualifications and experience together with names and addresses of two referees should be forwarded to:

The Personnel Manager
South Western Sydney Area Health Service
Private Mailbag 17,
Liverpool, NSW 2170 Australia.

NUCLEAR CARDIOLOGY TECHNOLOGIST

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Director, Dept. of Physics—Nuclear Medicine
Mt. Sinai Medical Center
Box 1141
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The Journal of Nuclear Medicine

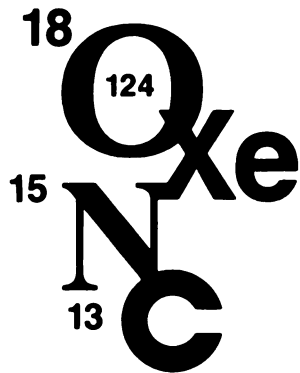
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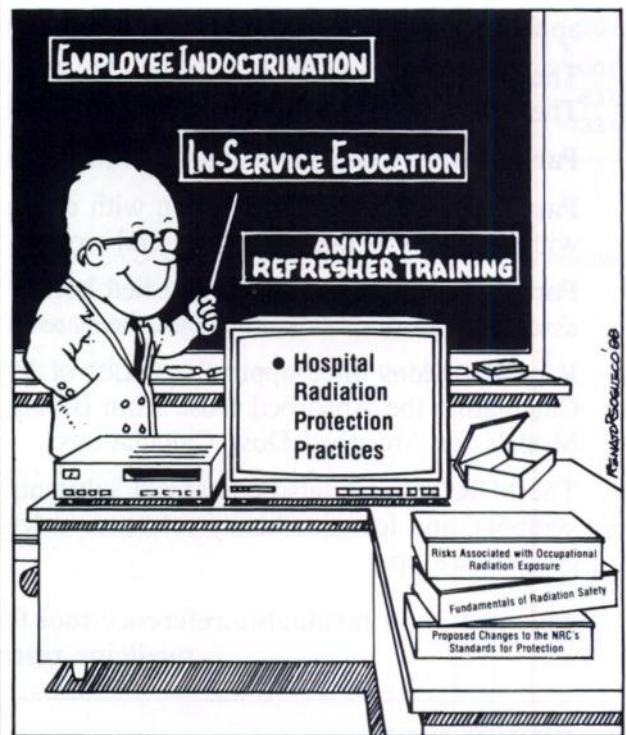
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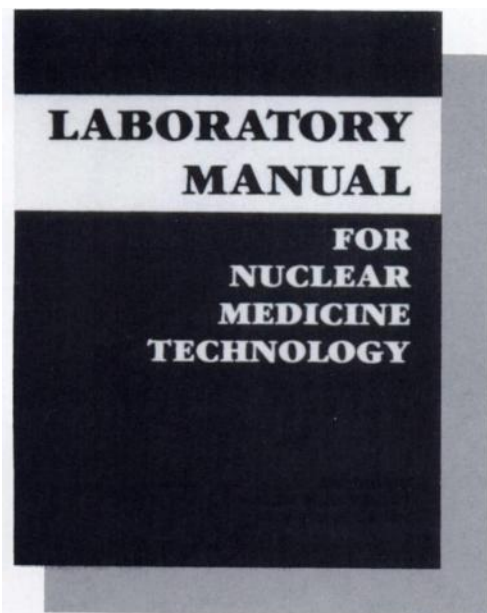
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Softcover format, 8½ x 11", 163 pp. Publication date: July 1984

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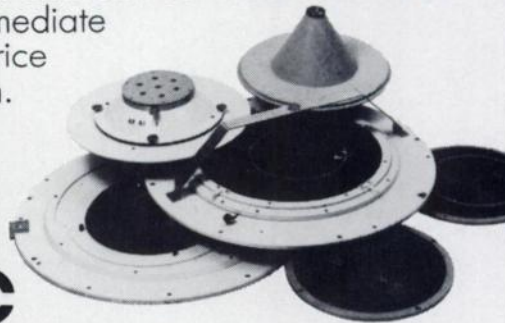
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The Technologist Section of The Society of Nuclear Medicine is a scientific organization formed with, but operating autonomously from, the Society to promote the continued development and improvement of the art and science of nuclear medicine technology. Membership in the Section is open to any member of the Society regardless of category, who can provide evidence of training and/or experience in nuclear medicine technology that is satisfactory to the Membership Committee of the Section.

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- Awards for outstanding achievements, and contributions to the technologist meetings, publications, and exhibits.
- Enrollment in the computerized continuing education accounting system (VOICE).

For more information, contact the Membership Department at:

The Society of Nuclear Medicine

**136 Madison Avenue
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THE SOCIETY OF NUCLEAR MEDICINE

Application for Membership

(see reverse side for instructions)



Last Name Dr. Mr. Mrs. Ms. Miss (CIRCLE ONE) _____	First Name _____	Middle Initial _____
---	-------------------------	-----------------------------

Check Degree(s) Earned:

MD ___ PhD ___ MA ___ MS ___ BA ___ BS ___ AA ___ AS ___ Other _____

Indicate Board Certification(s): ABNM ABR ABP ABIM ABSNM ABHP NMTCB

ASCP ARRT(N) ARRT(T) ARRT(R) Other _____

Please check ONE box for preferred mailing address, but complete both columns for our files:

<input type="checkbox"/> Institutional	<input type="checkbox"/> Home Address
--	---------------------------------------

DIVISION	STREET ADDRESS	APT. NO.
DEPARTMENT	CITY	STATE/PROVINCE/COUNTRY
INSTITUTION OR COMPANY	AREA CODE	TELEPHONE NO.
STREET ADDRESS	PRESENT POSITION (TITLE)	
CITY	STATE/PROVINCE/COUNTRY	ZIP CODE
AREA CODE	BUSINESS TELEPHONE NO.	EXT.

IN-TRAINING STATUS
 YES NO

Program Director _____

Projected Completion Date: _____ month/year

PROGRAM DIRECTOR'S TELEPHONE NO. _____

Would you like to join the TECHNOLOGIST SECTION? Yes No

COUNCIL MEMBERSHIP (OPTIONAL)

<input type="checkbox"/> Academic Council	<input type="checkbox"/> Correlative Imaging Council	<input type="checkbox"/> Radioassay Council
<input type="checkbox"/> Cardiovascular Council	<input type="checkbox"/> Instrumentation Council	<input type="checkbox"/> Radiopharmaceutical Council
<input type="checkbox"/> Computer Council		

NAME OF SNM MEMBER WHO SUGGESTED THAT YOU JOIN _____ (optional)

APPLICANT'S SIGNATURE _____ DATE _____



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<input type="checkbox"/> Full	<input type="checkbox"/> TS _____	CHAIRMAN, MEMBERSHIP COMMITTEE (sign)
APPLICATION FEE _____	<input type="checkbox"/> AM <input type="checkbox"/> R _____	TECHNOLOGIST SECTION DESIGNEE (sign)
CHAPTER _____	<input type="checkbox"/> TM <input type="checkbox"/> IT _____	
ACCOUNT # _____	<input type="checkbox"/> AF	

THE SOCIETY OF NUCLEAR MEDICINE

Instructions to Application for Membership

1. Please complete and sign the enclosed application form, either printing or typing the information. Make sure you have completed all information requested in order to avoid unnecessary delays in processing.
2. A membership category will be assigned to you in accordance with the Society's Bylaws based on the information supplied on your application form.
3. To be eligible for "In-Training" status, at least 90 days must be remaining in your formal training program. No application processing fee is required.
4. Upon acceptance by the Society, you will automatically become a member of the regional chapter that covers your area of residence. If you wish membership in some other chapter, you should submit your request with your application. If no regional chapter exists for the area of your residence, you will be assigned "Membership-at-Large."
5. **A \$10.00 non-refundable processing fee must accompany the completed application form. Otherwise applications will not be processed.**
6. Receipt of your application will be acknowledged. Allow 4-6 weeks for processing and for receipt of the appropriate journals. DO NOT prepay your dues. An invoice will be sent to you upon approval of your application.

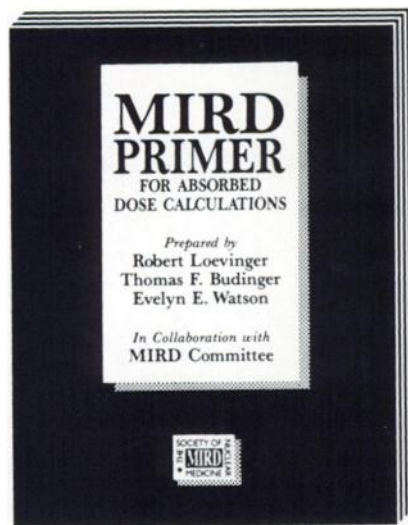
Guide to Membership Dues—1988

Membership Categories	Society	Technologist Section	Total
Full	\$100.00	—	\$100.00
Full-in-training	50.00	—	50.00
With Tech Section membership			
Doctoral degrees (MD, DO, PhD)	80.00	\$33.00	113.00
Doctoral degrees-in-training	40.00	16.50	56.50
All other degrees	75.00	33.00	108.00
All other degrees-in-training	37.50	16.50	54.00
Associate	75.00	—	75.00
Associate-in-training	37.50	—	37.50
With Tech Section membership			
Doctoral degrees	50.00	33.00	83.00
Doctoral degrees-in-training	25.00	16.50	41.50
All other degrees	50.00	33.00	83.00
All other degrees-in-training	25.00	16.50	41.50
Technologist (must be Tech Section member)	35.00	33.00	68.00
Technologist-in-training	17.50	16.50	34.00
Doctoral degrees	80.00	33.00	113.00
Doctoral degrees-in-training	40.00	16.50	56.50
Affiliate	100.00	—	100.00
With Tech Section membership	50.00	33.00	83.00
Doctoral degrees	100.00	33.00	133.00

- Society and Technologist Section chapter dues are additional and vary by chapter. A chapter dues table is available upon request.
- Council dues are an additional \$5.00 per Council.
- Dues for those applicants joining during the year are prorated to January 1st.

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MIRD PRIMER

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Prepared by
**Robert Loevinger
Thomas F. Budinger
Evelyn E. Watson**

In Collaboration with the MIRD Committee

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Part 4 contains three appendices: List of MIRD Pamphlets, A Revised Schema for Calculating the Absorbed Dose from Biologically Distributed Radionuclides, and Kinetic Models for Absorbed Dose Calculations.

The MIRD Primer also contains a substantive index, a detailed glossary and list of symbols, and for your handy reference calculation tables on the inside front and back covers; 128 pp.

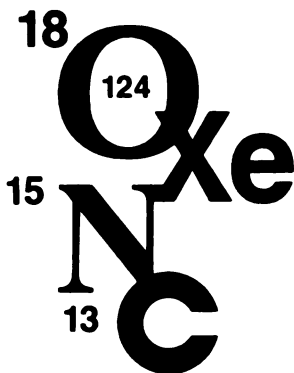
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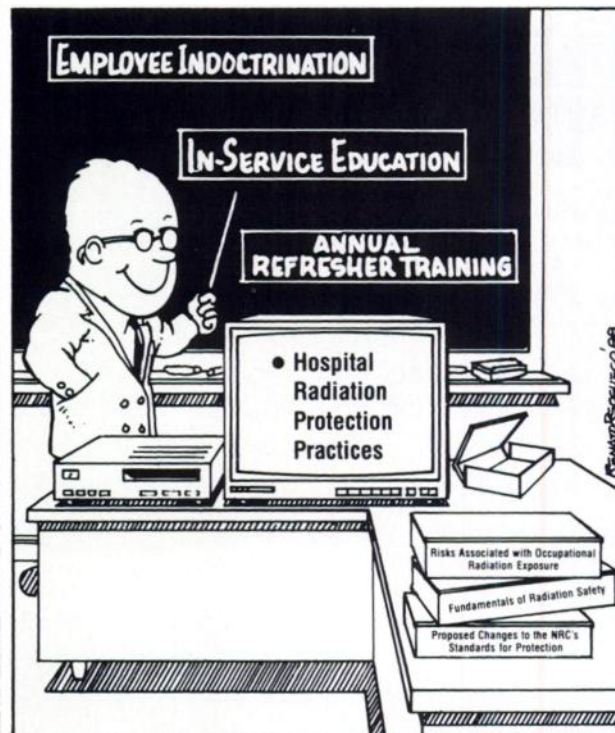
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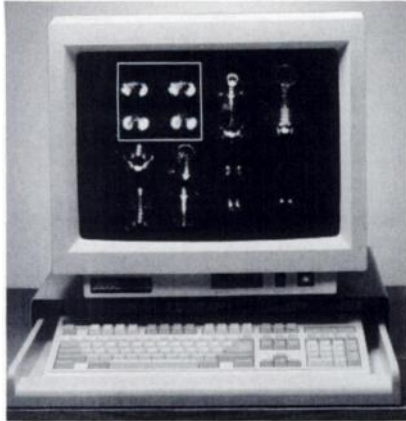
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Each description of the products below was condensed from information supplied by the manufacturer. The reviews are published as a service to the professionals working in the field of nuclear medicine and their inclusion herein does not in any way imply an endorsement by the Editorial Board of The Journal of Nuclear Medicine or by The Society of Nuclear Medicine.

Display Workstation



ADAC Laboratories announces the high performance VIEWPOINT nuclear medicine display workstation. Typically used in conference rooms or physicians' offices, the VIEWPOINT provides the physician with immediate access to patient studies for reviewing and reporting. Independent and economical, VIEWPOINT display systems provide powerful image manipulation and come in 512-resolution and 1024-resolution configurations. VIEWPOINT interactive digital viewing sessions replace conventional film reading without impacting the clinical imaging systems. Digital viewing, which is displayed on a high resolution 13" color monitor, enables the physician to: compare and contrast multiple studies in a simultaneous high resolution 1280 x 1024 display; improve diagnostic image quality by normalizing images, resetting grey scale, and manipulating color windows; and reference special cases or research studies by quickly recalling them from any acquisition or processing station. The VIEWPOINT system is not hard to operate with a mouse-driven, pull down menus screen environment. Grey scale windowing is available from every menu with simple mouse interaction and each menu contains an on-line HELP feature to assist the operator. Other display functions include: brightness, contrast, and grey map manipulation controls for image enhancement; cut and paste features that allow for on-screen editing; real time magnifier al-

lows inspection of critical detail; and motion picture display for visualization of dynamic studies. **ADAC Laboratories, 540 Alder Dr., Milpitas, CA 95035 Attn: Bruce Quill. (408)945-2990.**

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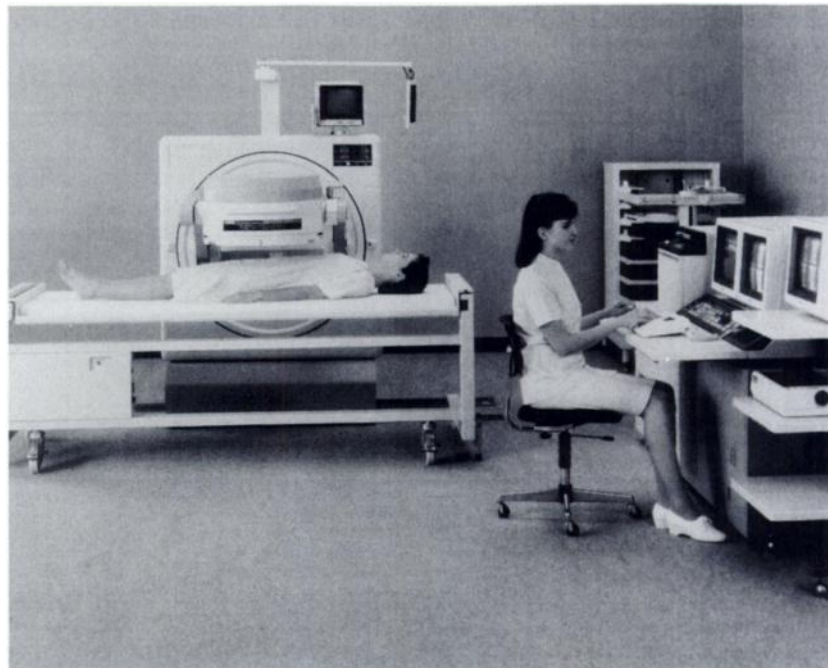
Radiation Shielding Material

Now handling, disposing, and storing low-level radioactive waste has been made easy with the development of a new radiation shielding material called maXmRAY manufactured by The Branford Companies. maXmRAY is currently being used by the company to manufacture a radioactive waste containment system called RAD BAG. Dielectrically sealed, RAD BAG is fabricated to specific thickness in

order to capture energy levels from Alpha, Beta, and Gamma emitters. Produced in sizes accommodating 55 and 30 gallon drums or 5 gallon pails, each RAD BAG comes with an easily positioned cover. In tests conducted by Teledyne Isotopes, average net count of isotope gamma emitted radiation when RAD BAG was not used was 249 ± 10 , however, when RAD BAG was used this dropped dramatically to 13 ± 7 . The attenuation factor was 0.9478 for energy levels of 0.136 to 0.170 MeV. Branford engineers state they are able to offer RAD BAG in a variety of sizes and shapes from tool kit bags to overbags for drums and tanks. Various closures are offered. **The Branford Companies, Box 713, Shelton, CT 06484 (203)735-6415, FAX (203)736-9102.**

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Five-Function Nuclear Medicine System



Toshiba Medical Systems introduces a new digital gammacamera which allows operators to perform five major system functions at once. The GCA-602A digital gammacamera permits simultaneous digital image data acquisition, analog data acquisition, image processing, data transfer, and computer communication. The benefits of five-function simultaneity to both clinicians and patients promise to be noteworthy. The use of the GCA-602A permits the total exam and processing time to be significantly reduced. The system allows operators to scan and process more effi-

ciently, patients spend less time on the scan couch, and throughput is improved. The GCA-602A is designed to complement Toshiba's GCA-901 whole-body gammacamera with hardware optimized for more efficient scanning of cardiac and neurological analyses. The system also features 1024 x 1024 acquisition and display. **Toshiba Medical Systems, 2441 Michelle Dr., Tustin, CA 92680 Attn: Steve Petras. (714)730-5000 or (800)421-1968.**

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For complete prescribing information consult package insert, a brief summary of which follows:

DIAGNOSTIC - FOR INTRAVENOUS USE

DESCRIPTION: The kit consists of 10 multidose reaction vials which contain the sterile, non-pyrogenic, non-radioactive ingredients necessary to produce Technetium Tc 99m Albumin Aggregated Injection for diagnostic use by intravenous injection.

Each 10 mL reaction vial contains 2.5 mg of Albumin Aggregated, 5.0 mg of Albumin Human, 0.06 mg (minimum) stannous chloride (maximum stannous and stannic chloride 0.11 mg) and 1.2 mg of sodium chloride; the contents are in a lyophilized form under an atmosphere of nitrogen. Sodium hydroxide or hydrochloric acid has been used for pH adjustment. No bacteriostatic preservative is present.

The Albumin Human was non-reactive when tested for hepatitis B surface antigen (HB_sAg) by radioimmunoassay. The aggregated particles are formed by denaturation of Albumin Human in a heating and aggregation process. Each vial contains 4 to 8 million particles. By light microscopy, more than 90% of the particles are between 10 and 70 micrometers, while the typical average size is 20 to 40 micrometers; none is greater than 150 micrometers.

Technetium Tc 99m Albumin Aggregated Injection for intravenous use is in its final dosage form when sterile isotonic sodium pertechnetate solution is added to each vial. No less than 90% of the pertechnetate Tc 99m added to a reaction vial is bound to aggregate at preparation time and remains bound throughout the 6 hour lifetime of the preparation.

INDICATIONS AND USAGE: Technetium Tc 99m Albumin Aggregated Injection is a lung imaging agent which may be used as an adjunct in the evaluation of pulmonary perfusion in adults and children.

CONTRAINDICATIONS: Technetium Tc 99m Albumin Aggregated Injection should not be administered to patients with severe pulmonary hypertension. The use of Technetium Tc 99m Albumin Aggregated Injection is contraindicated in persons with a history of hypersensitivity reactions to products containing human serum albumin.

WARNINGS: Although adverse reactions specifically attributable to Technetium Tc 99m Albumin Aggregated have not been noted, the literature contains reports of deaths occurring after the administration of Albumin Aggregated to patients with pre-existing severe pulmonary hypertension. Instances of hemodynamic or idiosyncratic reactions to preparations of Technetium Tc 99m Albumin Aggregated have been reported.

PRECAUTIONS: General

The contents of the kit before preparation are not radioactive. However, after the sodium pertechnetate Tc 99m is added, adequate shielding of the final preparation must be maintained.

In patients with right to left heart shunts, additional risk may exist due to the rapid entry of Albumin Aggregated into the systemic circulation. The safety of this agent in such patients has not been established.

Hypersensitivity reactions are possible whenever protein-containing materials such as pertechnetate labeled Albumin Aggregated are used in man. Epinephrine, antihistamines, and corticosteroids should be available for immediate use.

The intravenous administration of any particulate materials such as Albumin Aggregated imposes a temporary small mechanical impediment to blood flow while this effect is probably physiologically insignificant in most patients; the administration of Albumin Aggregated is possibly hazardous in acute cor pulmonale and other states of severely impaired pulmonary blood flow.

The components of the kit are sterile and non-pyrogenic. It is essential to follow directions carefully and to adhere to strict aseptic procedures during preparation.

Contents of the vials are intended only for use in the preparation of Technetium Tc 99m Albumin Aggregated Injection and are NOT to be administered directly to the patient.

The Technetium Tc 99m labeling reactions involved depend on maintaining the stannous ion in the reduced state. Hence, sodium pertechnetate Tc 99m containing oxidants should not be employed.

The preparation contains no bacteriostatic preservative. Technetium Tc 99m Albumin Aggregated Injection should be stored at 2-8°C and discarded 6 hours after reconstitution.

Technetium Tc 99m Albumin Aggregated Injection is physically unstable and consequently the particles settle with time. Failure to agitate the vial adequately before use may result in non-uniform distribution of radioactive particles.

If blood is drawn into the syringe, unnecessary delay prior to injection may result in clot formation *in situ*.

Do not use if clumping of the contents is observed.

Technetium Tc 99m Albumin Aggregated, as well as other radioactive drugs must be handled with care. Once sodium pertechnetate Tc 99m is added to the vial, appropriate safety measures must be used to minimize radiation exposure to clinical personnel. Care must also be taken to minimize the radiation exposure to patients in a manner consistent with proper patient management.

Radiopharmaceuticals should be used only by physicians who are qualified by training and experience in the safe use and handling of radionuclides and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

Carcinogenesis, Mutagenesis, Impairment of Fertility

No long term animal studies have been performed to evaluate carcinogenic potential or whether Technetium Tc 99m Albumin Aggregated Injection affects fertility in males or females.

Pregnancy Category C

Animal reproduction and teratogenicity studies have not been conducted with Technetium Tc 99m Albumin Aggregated Injection. It is also not known whether Technetium Tc 99m Albumin Aggregated Injection can cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. There have been no studies in pregnant women. Technetium Tc 99m Albumin Aggregated Injection should be given to a pregnant woman only if clearly needed.

Ideally, examinations using radiopharmaceuticals, especially those elective in nature, of a woman of childbearing capability, should be performed during the first few (approximately 10) days following the onset of menses.

Nursing Mothers

Technetium Tc 99m is excreted in human milk during lactation. Therefore, formula feedings should be substituted for breast feedings.

Pediatric Use

The lowest possible number of particles should be used in right-to-left shunting, in neonates, and in severe pulmonary disease.

ADVERSE REACTIONS: The literature contains reports of deaths occurring after the administration of Albumin Aggregated to patients with pre-existing severe pulmonary hypertension. Instances of hemodynamic or idiosyncratic reactions to preparations of Technetium Tc 99m Albumin Aggregated have been reported (see Warnings).

HOW SUPPLIED:

MPI MAA Kit
Kit for the Preparation of Technetium Tc 99m Albumin Aggregated Injection
Product No. 4432

Each kit contains 10 reaction vials, each vial containing in lyophilized form, sterile and non-pyrogenic:

Albumin Aggregated	2.5 mg
Albumin Human	5.0 mg
Stannous Chloride (minimum)	0.06 mg
(Maximum stannous and stannic chloride)	0.11 mg
Sodium chloride	1.2 mg

HCl or NaOH has been used for pH adjustment. The vials are sealed under an atmosphere of nitrogen.

Twenty labels with radiation warning symbols and a package insert are supplied in each carton.

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