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Lungaggregate™Reagent [Aggregated Albumin (Human)] has eight important advantages for pulmonary scintigraphy.

The first one is obvious:

1. Particles Presuspended in Solution.

Lungaggregate Reagent is the only Tc 99m-labeled MAA agent containing albumin aggregate particles that are already suspended in an aqueous solution. There is less chance for radiation exposure to the user since no visual inspection is required after radioactive labeling.

2. Soft Particles for Rapid Lung Clearance.

The uniform-size particles in Lungaggregate Reagent have a biological half-time of 4.77 hours.

3. Quick, Easy Preparation.

No thawing, reconstitution of lyophilized particles, or ultrasonic agitation are required.

4. Conveniently Stable.

Lungaggregate Reagent, labeled with Tc 99m, may be used up to 24 hours after preparation when stored as directed. A supply of Tc 99m-Lungaggregate Reagent is therefore available when emergency studies are required.

5. Multi-Dose Economy.

Each vial can be used to give several patient doses since Lungaggregate Reagent contains a preservative.

6. Imaging Excellence.

Tc 99m is the radionuclide of choice for scintigraphy. With a 4 mCi dose of Tc 99m-Lungaggregate Reagent, up to 500,000 counts can be

obtained in two to three minutes on a gamma camera.

7. High Lung/Liver Activity Ratio.

The ratio of lung to liver-andspleen activity is over 10/1.

8. Patient Safety.

No adverse reactions have been reported. See the brief summary section below.

For a monograph summarizing clinical experience with Lungaggregate Reagent, or for additional information, call Medi-Physics toll free: (800) 772-2446 in California or (800) 227-0483 outside California.

Brief Summary

(For full product information including method of preparation and administration procedure, see package insert.)

Description: Lungaggregate[™] Reagent is a sterile, apyrogenic, buffered, preserved, aqueous preparation of aggregated albumin from human plasma.

Indications: For imaging regional pulmonary perfusion in the presence of clinically suspected regional ischemia.

Contraindications: This agent is contraindicated (1) in the presence of large right-toleft cardiovascular shunts which could allow direct entry of macroaggregates into systemic circulation; (2) in patients with cyanosis or evidence of severely restricted pulmonary blood flow, as in pulmonary hypertension; (3) in pregnant or lactating women and in patients under 18 years, unless expected benefits outweigh risks involved.

Warnings: Whenever protein-containing materials such as Tc 99m-labeled Lungaggregate Reagent are used in man, hypersensitivity reactions are possible. Have epinephrine, antihistamines, and corticosteroid agents available.

Precautions: Note—Follow aseptic techniques in preparing this agent to minimize the possibility of contamination with microorganisms. Take steps to minimize exposure to patient and attending personnel, including use of minimum dosage to achieve useful diagnostic data. Make injection slowly. Use an 18-21 gauge needle. After withdrawal from the vial the material should be administered promptly; also avoid aspirating blood and tissue fluids into the syringe.

Adverse reactions: None reported in over 4,000 patient studies.





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- 99m Tc Polyphosphate-Tin
 100mg Polyphosphate and 2mg Stannous Chloride
- 99m Tc DTPA-Tin

5mg DTPA and 0.25mg Stannous Chloride

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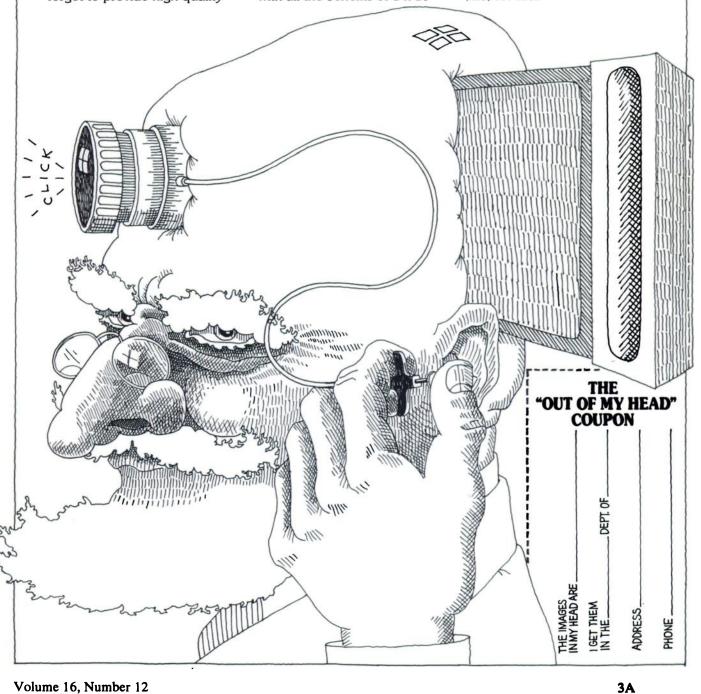
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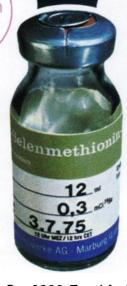
Volume 16, Number 12



L-Selenomethionine (Se-75)

new

For pancreas scintigraphy as a simple detection method for space occupying lesions like tumors or cysts and alterations of parenchyme.



Already after 10 min maximum count rate At least 75% of the initial activity after 60 min

Low radiation dose for 100 µCe in liver, pancreas and kidneys Whole body dose: 0.8 rd High radiochemical purity (98 %) at calibration date Recommended dose: 300 µCi

Specification

L-Selenomethionine-(Se-75) Less than 5% D-Selenomethionine. Concentration of activity: 0.2 mCi Se-75/mI Specific activity: 5-10 mCi Se-75/mg Selenomethionine

Pack

L-Selenomethionine-(Se-75) in physiological saline for injection (12ml beaded rim vial)

Order No.: SE-515

Calibration day: 1st of the month

Dispatch: daily from the 1st of

the previous month on

Shelf life: 3 months from the day of first dispatch

Contraindications

Radioactive material should be handled with special care to insure minimum radiation exposure to personnel and patients.

Unless strictly indicated, radiopharmaceuticals should not be administered to pregnant or nursing women or to juvenile patients.

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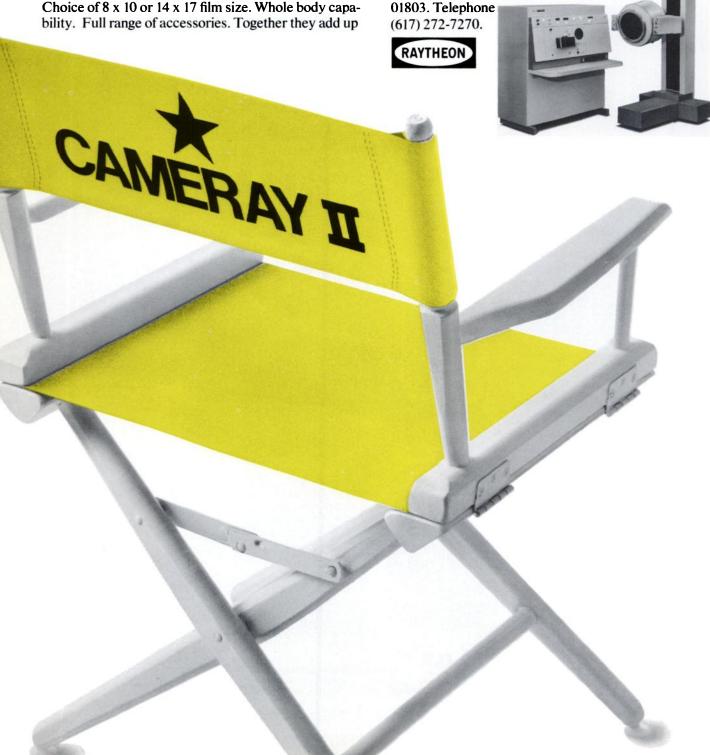
Any scintillation camera that's a top performer has to put a lot of good operating characteristics together. System and energy resolution. Uniformity. Linearity. Count rate. Price. Consider all these together and you'll find Cameray II at the top. There are other reasons too. Choice of 8 x 10 or 14 x 17 film size. Whole body capability. Full range of accessories. Together they add up

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On the current nuclear medicine scene



MINITEC® (Technetium 99m) Generator

The Technetium 99m Generator using fission product molybdenum to produce technetium 99m. MINITEC is unlike any generator you've ever used—made small to make sense.

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Designed for easy elution

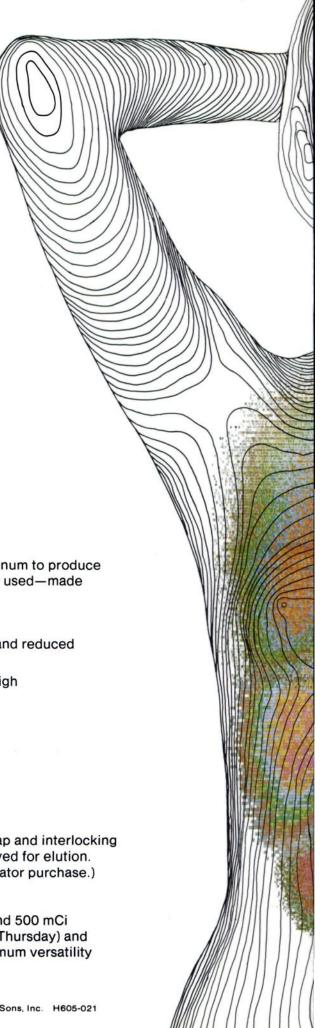
- Sets up in seconds
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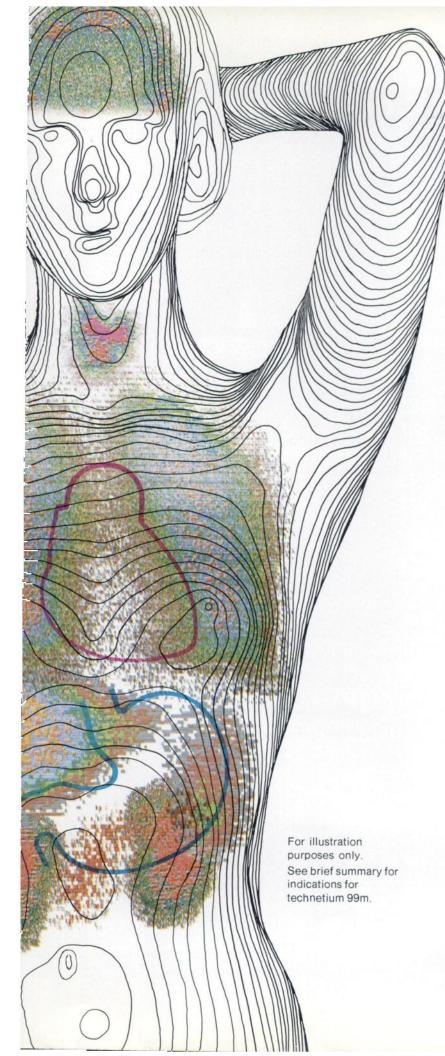
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- No exposed tubing when eluting
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Designed for convenience

MINITEC Generator is available in 50,100, 200, 300, 400 and 500 mCi
potencies. Delivery on Monday AM (precalibrated through Thursday) and
Wednesday (precalibrated through Monday) provides maximum versatility
to satisfy technetium requirements of your lab's work load.





Minitec[®] (Technetium 99m) Generator

Minitec® (Technetium 99m) Generator provides a means of obtaining a sterile, non-pyrogenic supply of technetium 99m (**Tc) as sodium pertechnetate

Indications: Sodium pertechnetate *** Tc is indicated for brain imaging, thyroid imaging, salivary gland imaging, blood pool imaging, and placenta local-

Contraindications: At present, there are no known contraindications to the use of sodium pertechnetate ***Tc.

Warnings: Radiopharmaceuticals should be used only by physicians who are qualified by specific training in the safe use and safe handling of radionuclides, produced by nuclear reactor or cyclotron, and whose experience and training have been approved by the appropriate federal or state agency authorized to license the use of radionuclides.

This radiopharmaceutical should not be administered to women who are pregnant or who may become pregnant or during lactation unless the information to be obtained outweighs the possible potential risks from the radiation exposure involved. 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.

Since radioactive pertechnetate is secreted in milk during lactation, formula-feedings should be substituted for breast-feedings.

Important: Since material obtained from the generator may be intended for intravenous administration, aseptic technique must be strictly observed in all handling. Only the eluent provided should be used to elute the generator. Do not administer material eluted from the generator if there is any evidence of foreign matter.

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

At the time of administration, the solution should be crystal clear.

Adverse Reactions: At present, adverse reactions have not been reported following the use of sodium pertechnetate ***Tc.

For complete prescribing information, consult package insert.

How Supplied: Minitec (Technetium 99m) Generator is available in potencies of 50, 100, 200, 300, 400, and 500 mCi. Supplied with the generator are vials of eluent containing 5 ml. of a sterile, non-pyrogenic solution of $0.9\bar{\%}$ sodium chloride in water for injection. Also supplied is suitable equipment for eluting, collecting, and assaying the technetium 99m.

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linear or log scale.

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PICTURES BY PEOPLE

Easy use: full plain text dialogue separated from display enables sophisticated use under people control without the usual secret code of computers.

Protocols: routine procedures may be chained into a protocol, with comment, for full automatic machine control.

Identification: it is impossible to have unidentified displays or to mix patient records in these systems.

PICTURES FOR PEOPLE

Dynamic: flexible visualization and quantification of physiological processes promotes positive diagnoses.

Static: finally available, static images significantly better than the raw camera output promote earlier more effective clinical diagnoses.

PICTURES FOR MORE PEOPLE

Dual Cameras: systems for two cameras with simultaneous dynamic capability without interference or record confusion.

Multi-tasking: the BETA executive automates the computer functions for clinical use, or permits the computer-orientated to access FORTRAN or ASSEMBLER and to multi-task up to 7 functions (memory size option permitting) simultaneously.

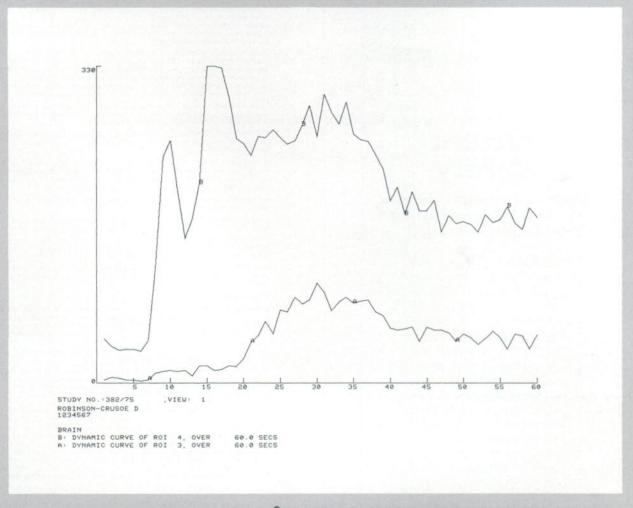
Multi-accessing: background tasks may be run such as radio immunoassay, E.K.G., radiotherapy planning, etc., simultaneously with gamma camera use (which has, of course, priority).

System Growth: a start may be made with a low-cost budget system. Large comprehensive systems may be built from standard modules.

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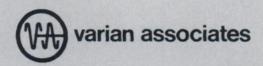
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Aggregated Albumin (Human) Kit

DESCRIPTION - The kit contains 6 sterile vials containing 9-11 mg, of pyrogen-free aggregated albumin (numan), 0.67 - 0.83 mg, stamous chloride, and 18 mg, sodium chloride. When sterile, pyrogen-free sodium particinetate Tc99m is added to the vial, technetium-tabelled macroaggregated human serum albumin (Technetium MAA Tc 98m Technetium Macroaggregates) is formed. The particles of aggregated albumin in the kit are formed by the denaturation of Normal Serum Albumin (Human) USP through heat and pri adjustment. Sodium hydroxide of hydrochloric acid may be present in variable amounts. Al least 95% of the macroaggregated particles are between 10 and 100 micrors in size, the great bulk (as seen on a microscope side) being an average of 10 to 70 micrors. Vial counts indicate that each vial contains 6.8 ± 0.8 million particles per mg. The labelling efficiency is essentially quantitative and the bound Tc-MAA remains stable in vitro throughout the useful period after preparation.

Application has been filed with the U. S. Nuclear Regulatory Commission for distribution of this reagent kit to persons licensed pursuant to §35.14 and §35.100, Group III of CFR Part 35, or under equivalent licenses of agreement states, and is still pending.

ACTIONS - Following intravenous injection, Technetium MAA Tc 99m is rapidly transported by the blood stream to the lungs. The apprepaies do not enter the tissues of the lungs, but remain in the pulmonary vasculature. When pulmonary blood flow is normal, the material is carried throughout the entire lung field; when pulmonary blood flow is diminished or obstructed by a disease process, the particles are correspondingly prevented in part of in whole from passage through the affected portion of the pulmonary vasculature.

Technetium Macroaggregates remain in the lungs for variable amounts of time depending on particle size. The particles disappear from the lungs in exponential tashion with the larger-sized aggregates having the longer half-life; particles ranging from 10 to 90 microns in diameter usually have a half-life of 2 to 8 hours. Apparently, the aggregates are temporarily trapped by the narrow pulmonary cigilitaries where the particles are broken down until they are small enough to pass. In rats 4.3% of the Tc 99m remains in the lungs after 24 hours.

Although the particles of macroaggregates remain for a time in the pulmonary capillaries, they do not appear to interfere even temporarily with pulmonary blood flow or ventilation in the dosage required for lung scanning. This is evidenced by the fact that these doses do not produce any respiratory distress nor any tachycardia, even in patients severely ill with pulmonary and/or cardiac disorders.

Once the albumin particles leave the lungs, they are carried to the liver, where they are removed from the blood stream primarily by the Kupfter cells. There, the particles are phagocytized and rapidly metabolized.

INDICATIONS - Scintilation scanning of the lungs with Technetium Macroaggregates is indicated as an adjunct to other diagnostic procedures whenever information about pulmonary vascutature is desired. The most useful clinical applications of lung scanning have been outlined by one investigator: 1) The diagnosts of pulmonary embolism: 2) differentiation of local conditions such as butlae or cysts from diffuse pulmonary disorders; 3) determination of the degree of pulmonary vascular obliteration in parenchymal disease; and 4) evaluation of the patient's ability to withstand pulmonary surgery.

Perhaps the most frequently useful indication for the lung scan has been the early detection of pulmonary emboli. The lung scan is uniquely able to demonstrate the existence of an embolism shelver addological signs become apparent. Although an area of increased radiolucency on the chest film may suggest an embolism. X-ray findings do not usually become apparent until the embolism has produced signs of ischemia or infarction. Once an embolism has been diagnosed, information obtained from the scan is of value in determining the desirability of surgical embolectomy, while subsequent scans provide information on the effectiveness of surgical or articoappulant therapy.

Lung scanning is similarly helpful in the diagnosis of various types of malignancies affecting the lungs. Again, scanning is of value in locating the affected areas, in determining the need for and probable effectiveness of surgery or of radiation therapy, and in following up the benefits of treatment.

Useful information is also provided by the scan in the diagnosis or evaluation of other pulmonary problems, such as pneumonia, arelectasis pleural effusion, pulmonary suberculosis, parenchymal disease, emphysema and chronic asthmatic bronchitis.

CONTRAINDICATIONS - The presence of right to left shurts which would allow Technetium MAA Tc 99m injected in a systemic veri to reach a systemic artery is contraindication to the use of this material. Particulate material such as Technetium MAA Tc99m should not be administered to patients with evidence of severe restriction to pulmonary blood flow such as may be present in pulmonary hypertension.

WARPHINGS - Technetium MAA Tc99m should not be administered to patients who are pregnant, or during lactation unless the benefits to be gained outweigh the potential hazards.

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

Radiopharmaceuticats should be used only by physicians who are qualified by specific training in the sale use and handling of radionuclides produced by nuclear reactor or particle accelerator and whose experience and training have been approved by the appropriate government agency authorized to license the use of radionuclides.

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

To insure the integrity of this product use needles in gauge sizes 18 to 21.

ADVERSE REACTIONS - No adverse reactions have been observed with this product. However Vincent et al.

(3) have recorded the only immediate and tetal reaction following infusion of Tc 99m macroaggregates (technetium labelled macroaggregates). This was in a seven-year-old child who had severe pulmonary secular disease. The exact size of the particles used was not disclosed, and in the summary of the publication "it is suggested that this type of reaction will continue to be rare and that it will probably be somewhat predictable on the basis of clinical and laboratory evidence of severe pulmonary hypertension. Such a patient might be scanned safely by strict control of macroaggregates dose, size range and mean particle size".

The literature has recorded two adverse reactions to lung scanning with I-131 labelled macroaggregates. Wagner et al (4) observed that urticaria developed in a young girl several hours after lung-scanning procedure with lodins-131 macroaggregates where Lugol's solution was administered to block the thyroid gland. The subject had a history of anglo-edema. The reaction may have been caused by either material. Dworkin et al (5, 6) reported "I-131-labelled macroaggregated albumin highly suspect as the causative agent" in the death of a woman who was scanned for the possibility of demonstrating pulmonary embolism. With a 249-year history or admonscrationar of the breast she had severe and rapidly progressive edema. Prior to scanning, the nasal administration of oxygen was interrupted. "Within or 2 minutes after injection of 300 uCi of I-131 labelled macroaggregates abumin (11 mg, of albumin or 0.219 mg, per kilogram of body weight) she complained of laintness and became cyanotic, daphonetic, and aglated with distended neck veins. The initial pulse rate of 50 rose to 140 with a tall in blood pressure to 100/30. Oxygen thrappy releved the prolound dyspone and cyanosis. An electrocardiogram 40 minutes later was compatible with acute cor pulmonale. Within several hours she had returned to her pre-scan status, but on the next day the temperature rose, dyspone increased and she died 26 hours after the lung scan. We have continued lung scanning but limit the albumin to 0.020 mg, per kilogram, reject lots with more than 15 percent of particles over 40 micrors and require two minutes for injection."

More recently, Williams (7) has reported a severe reaction immediately after injection of macroaggregated albumin (MAA) particles followed by death six hours later (while the patient was undergoing right-heart catheterization). Like those previously reported, it occurred in a patient with severe chronic pulmonary hypertension due to disease of the pulmonary vascular bed. The patient ded in right heart failure. Post-mortem examination reveated "severe atheroma and thickening of all the pulmonary arteries but no macroscopic evidence of emboti. The right heart was hypertrouphile and dilated".

Transient neurological complications following intra-arterial injection of I-131 labelled macroaggregates have been reported (3).

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 Dworkin, J. J., Smith, J. R. and Bull, F. E., N. Engl. J. Med. 275, 376 (1966).
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- 7. Williams, J. O., Brit. J. Radiol. 47, 61-63 (1974).

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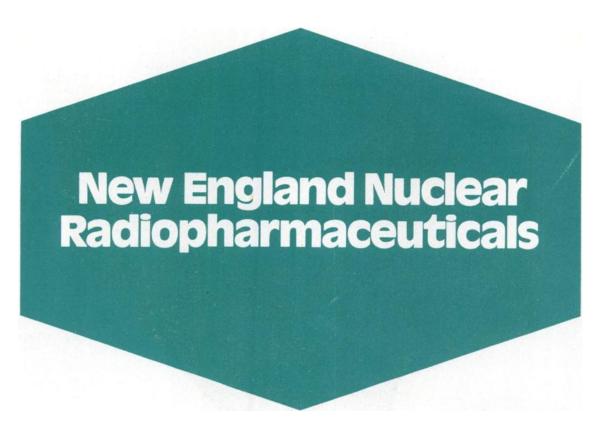
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INDICATIONS: Pertechnetate Sodium Tc 99m is used for brain imaging, thyroid imaging, salivary gland imaging, placental localization and blood pool imaging.

CONTRAINDICATIONS: To date, there are no contraindications to the use of Pertechnetate Sodium Tc 99m.

WARNINGS: This radiopharmaceutical should not be administered to pregnant or lactating women unless the information to be gained outweighs the potential hazards.

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 the menses.

Radiopharmaceuticals should be used only by physicians who are qualified by specific training in the safe use and handling of radionuclides produced by nuclear reactor or particle accelerator, and whose experience and training have been approved by the appropriate governmental agency authorized to license the use of radionuclides.

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

At the time of administration the solution should be crystal clear.

IMPORTANT: Refer to Operating Instructions on the proper use of the New England Nuclear Generator. These instructions are enclosed with each generator.

ADVERSE REACTIONS: To date, no adverse reactions based on the use of this agent have been reported.

DOSAGE AND ADMINISTRATION: Pertechnetate Sodium Tc 99m is usually administered by intravenous injection but can be given orally. The dosage employed varies with each diagnostic procedure.

The suggested dose range employed for various diagnostic indications in the average patient (70 kg) is:

Brain Imaging: 10-20mCi Thyroid Imaging: 1-10mCi Salivary Gland Imaging: 1-5mCi Placental Localization: 1-3mCi Blood Pool Imaging: 10-20mCi

Note: Up to 1 gram of reagent grade potassium perchlorate in a suitable base or capsule may be given orally prior to administration of Pertechnetate Sodium Tc 99m injection for brain imaging, placental localization and blood pool imaging.

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



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Volume 16, Number 12



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The kit is a simple radioassay – a matter of routine for any clinical laboratory with a gamma counter. Please write or 'phone for further information.



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The Radiochemical Centre Amersham The Radiochemical Centre Limited, Amersham, England. Tel: 024-04 444.
In the Americas: Amersham/Searle Corp. Illinois 60005. Tel: 312-593-6300.
In W.Germany: Amersham Buchler GmbH & Co., KG, Braunschweig. Tel: 05307-4693-97.

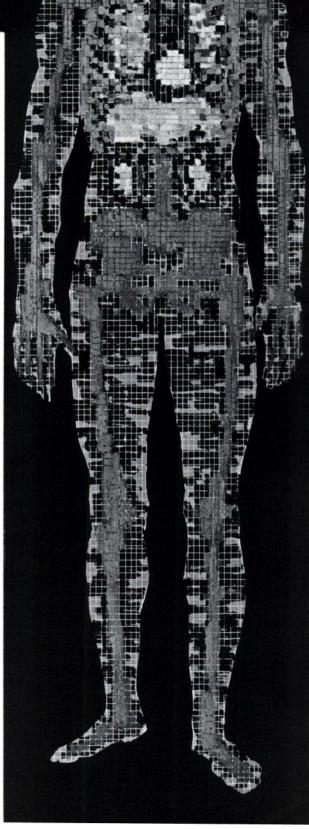
Volume 16, Number 12





Automatic RIA Gamma Counter

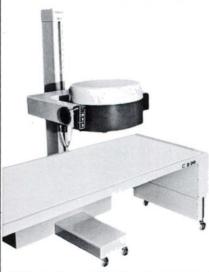
180-1260 sample capacity. Accepts most standard sized test tubes. Rack/tray based to save handling time. Built-in calculator-printer. Optional teletype printout/punchout.





NEW Gamma Cameras

Two gamma cameras to choose from: one designed to give you exceptional image resolution — the second, an economy model. The short dead time of both give you high information density even with short frame times. Maximum count rates up to 200,000 cps.



Whole Body Imaging Accessory

Space saving imaging accessory greatly extends the camera's usefulness.

in-vitro.. Elscint gives maximum performance at minimum price.

IMAGING ACCESSORIES



NEW LOW COST Videodisplay Processor

Full 16 color or 32 gray shades, and on-line or off-line display of camera or scanner images facilitates accurate patient diagnosis. Includes memory for data study or manipulation, photography and printout on Elscint color printer. Easy to use. Interfaces to rectilinear scanners and gamma cameras.



Color Dot Printer

Provides full color hard copies of images, printed on regular paper, either minified or at actual body size.

DATA PROCESSING



Dynamic Image/Function Processor

Advanced version of Elscint's Videodisplay Processor. Adds large dual disc memory for extensive non-destructive data processing. Displays time functions in real time; provides up to 8 regions of interest; eliminates artifacts and non-uniformities. Upgradable to Analyzing Image Processor.

Analyzing Image Processor

Most sophisticated image processing system available today.
Offers all capabilities of our Dynamic Image Processor plus many other features needed for in-depth study of renal functions, regional cerebral blood flow, cardiac and many other studies. Simultaneous acquisition and processing is a standard feature.

Whole Body Scanner

Scan the whole body or any part of the body with pushbutton ease. Every desirable performance feature is built into these scanners. You'll get long dependable service at an unbelievably low price. A full range of options meets every need.



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FEATURING

- No serum blanks
- Range 0.1 32 ng/ml
- Will measure Folate concentration in whole blood
- ½ hour incubation

Diagnostic Products Corporation, the prime producer of the ³H Folate Assay Kit, has developed an ¹²⁵I Folate Kit with all the characteristics that have enabled us to maintain our leadership in the Folate RIA market. The unexcelled simplicity, accuracy and reproducibility of our tritiated products is characteristic of our ¹²⁵I Folate, T-3 RIA, T-4 RIA.

T-4 RIA

- No extraction
- Range $0.3 32 \,\mu g/100 ml$
- 30 minute incubation
- Cross-reactivity with T-3 0.015%

T-3 RIA

- No extraction
- Range 10 800 ng/100ml
- 30 minute incubation
- Cross-reactivity with T-4 0.002%

Also Available:

[3H] Aldosterone

· (No Chromatography)

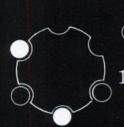
[3H] Cortisol RIA

[57CO] Vitamin B-12

[3H] Digoxin

[3H] Cyclic AMP

[3H] Cyclic GMP



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Cardiac Gate



Opti Imager

Cardiac Gate is designed to synchronize the cardiac image exposure with predetermined phases of the cardiac cycle.

The Cardiac Gate has two modes of operation: manual and automatic. In the manual mode, delay and exposure time parameters are set manually, using the R wave of the electrocardiogram as a reference. In the automatic mode, microprocessor circuitry automatically tracks the cardiac cycle and computes the position of end-systole and end-diastole. In the automatic mode, end-systole and end-diastole exposures are made without any calibration settings.

The dual gating operation mode allows recording of both end-systole and end-diastole simultaneously in a split screen two image format.

The cardiac cycle can even be divided into nine equal time segments and the image corresponding to each displayed simultaneously in a nine image format.

The Cardiac Gate includes a complete electrocardiograph module. The built in heated stylus strip chart recorder records both the ECG trace and the gating intervals.

The Cardiac Gate provides both ECG and gating outputs for computer interface.



Opti-Imager is designed to provide an organ image with effects due to respiratory motion minimized. Opti-Imager has two distinct modes of operation: continuous motion correction and respiratory gating. In the continuous motion correction mode, the motion of the organ is tracked and corrected electronically without the need to attach any sensors to the patient. The distribution of counts within the organ image is monitored and corrections are applied to continuously shift the image before it is displayed to compensate for organ motion. Correction is made for motion in both the X and Y direction. Thus, the gamma camera is not gated and all the counts provided by the detector are recorded. The time required to attain a statistically satisfactory image is the same for both a motion corrected and an uncorrected image. In the gating mode,

The Cardiac Gate and Opti-Imager can be synchronized to yield a combination of both cardiac and respiratory gating. Mail coupon to receive detailed information and sample clinical studies.

inspiration plateau and expiration plateau

split screen two frame format. Dual scalers

record the number of counts in each image.

images are recorded. The dual gating operation

mode allows recording of both inspiration and

expiration plateau images simultaneously in a

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The Quiet Evolution of a Unique Gamma Camera



The New SYSTEM SEVENTY SEVEN from Baird-Atomic

For the past forty years, Baird-Atomic has set the pace in high-technology instrumentation for a wide variety of applications, and especially for nuclear medicine. The accent has always been on innovation—taking a fresh look at each problem and devising an original way to solve it. Our original scanning gamma camera, System Seventy, was an ideal example.

In the earliest stages of the system's design we realized that existing mono-crystal systems had inherent disadvantages which would inhibit their use as clinical studies became more sophisticated and higher count rates became necessary for statistical accuracy and integrity. The answer was a multi-crystal detector. The decision to design and build it — a long, difficult, and expensive process — became the critical step in the evolution of a unique gamma camera system versatile enough to accommodate every future change in clinical procedures.

That was only the first step, however. So many refinements and improvements have since been made that we've given it a new name. System Seventy Seven.

Briefly noted, some of the new features: A fully comprehensive program of nuclear medicine software, eliminating the time consuming work of converting data to clinically useful formats. An image processing console that analyzes 200,000 observed counts per second, at any energy level, with a minicomputer as its storage and data manipulation base. Computer controlled bed motion, virtually eliminating collimator dead space and optimizing resolution for uniform, always reproducible imaging. A computer console with pushbutton simplicity, one that backlights only legitimate subsequent operations.

There are more. And more details about these. They're all described in our new brochure about the new System Seventy Seven. You can join the evolution simply by sending or calling for it.



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Photo insert: Wall motion of the left ventricle, a typical example of the kind of selective imaging possible with System Seventy Seven's unique data processing capabilities. Zones of interest and histograms of selectively specific target areas can be routinely obtained, and as many as four can be simultaneously manipulated. The operator has total control in determining the shape and size of the region examined, as well as the time/count scale of the histogram. From 10 to 20 cycles of systole and diastole, recorded during the first passage of the radionuclide, may be reformatted into a single representative cardiac cycle of maximum retrievable depth, detail, and accuracy. Study courtesy of Dr. Robert H. Jones, Duke University.

GAMMA CAMERA CALIBRATION KIT

The radioactive sources and phantom of the AECL Gamma Camera Calibration Kit provide an effective means of routinely checking the vital characteristics of your camera system.

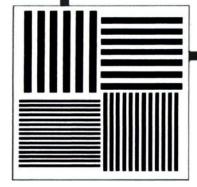
Sources are safe, light and easy to carry in the attractive carrying case provided.

Sources are approved for licensing in U.S.A. and Canada.

FLOOD FIELD SOURCE

A rapid and convenient way of making the daily check of your camera response. It is a flat plastic disc 12 inches in diameter containing 3 mCi of Gadolinium-153 (100 KeV photopeak, 242 day half life) dispersed uniformly to give an output better than $\pm 5\%$ over the whole surface.





BAR PHANTOM Used with a Flood Field Source to provide an efficient check of the inherent and system resolution of your camera system. It can also be used to check image size and linearity.

The Bar Phantom consists of four groups of lead bars embedded in a plastic holder 13.5 inches square and 0.37 inches thick. The bars are 0.125 inches thick and 0.500, 0.375, 0.250 and 0.187 inches wide respectively. The spacing between the bars is equal to the width of the bars for each group.

RESOLUTION REFERENCE

SOURCE A convenient way of checking the resolution of your gamma camera and scanner. The source contains a grid of radioactive lines which vary in spacing. Most cameras should be able to resolve the finest part of the grid. By adjusting the distance of the source from the collimator, the depth resolution of your camera can also be measured. Total activity of the source is 3 mCi of Gadolinium-153.

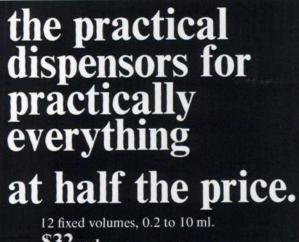


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27A Volume 16, Number 12

CARDIONUCLEAR Heart Attack... is the nation's number ANALYSIS:

... is the nation's number one killer. In 1972, it claimed 683,100 lives. An estimated 3,940,000 Americans have a history of heart attack and/or angina pectoris.¹

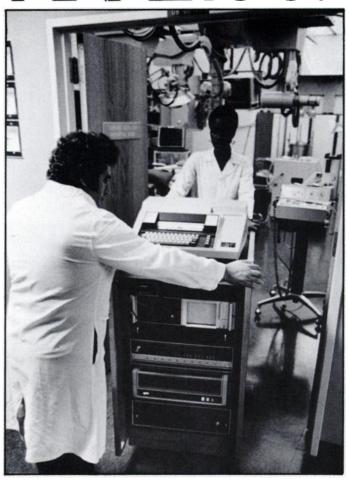
This year, approximately 350,000 myocardial infarct patients will die before they can reach medical help.

Cardiac catheterization, an invasive procedure with attendant morbidity and occasional mortality², does not fulfill all the diagnostic requirements of patients with heart disease. The cardiologist therefore requires aid in determining the status of patients both upon admission and during the course of therapy.

Cardionuclear Analysis serves as a cardiac catheterization screen to determine the probability of cardiac disease non-invasively. Additionally, it can provide indications of pre and post surgical cardiac function, and enhance hot and cold spot scanning of myocardial ischemia. Cardionuclear Analysis allows convenient review of study data as many times as desired.

10 analyses of cardiovascular functions can be determined from only one bolus injection. Because of sophisticated programming, the total time involved can be as little as ten minutes using simple keyboard commands.

And because of the mobility of the MODUMED PADtm, image processing can now take place in the ICU, CCU, Recovery Room, Cath Lab, ER or anywhere you take your scintillation camera.



The mobile MODUMED PADtm provides immediate diagnostic support in the ICU, CCU, Recovery Room, ER, Cath Lab, and other special care facilities.

Medical Data Systems Corporation, a subsidiary of Warner-Lambert, has been developing diagnostic imaging products for several years. As the leading company in Digital Nuclear Medicine, our commitment is to support the growth of Nuclear Medicine as both an art and a science.

- 1. Source: American Heart Association. Heart Facts 1975
- Hamilton, Glen, M.D.: Kennedy, J. Ward, M.D.: Assessment of Left Ventricular Function: Current Methods and Clinical Significance. Presented at the Symposium on Cardiovascular Investigation with Radionuclides at the University of Miami School of Medicine, Mt. Sinai Medical Center, Miami, March 12-16, 1975.
- 3. Hamilton, Kennedy, As Above.



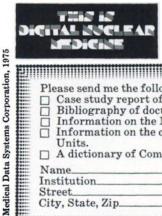
TEN DETERMINATIONS

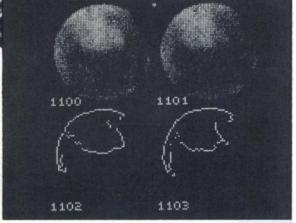
ECG GATED STUDIES

- 1. EJECTION FRACTION*
- 2. BLOOD POOL
- 3. WALL MOTION
- ANEURYSM

Cine Display:

The complete flow of a radio-tracer can be dynamically reproduced on the system display scope in a cine display without flicker. Even the dynamics of an individual heartbeat can be analyzed in patients with dyskinesis, aneurysm, and shunt.





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Please send me the following:

Case study report of Cardionuclear Analysis

Bibliography of documented use of Cardionuclear Analysis techniques.

Information on the Mobile MODUMED PADTM Image Processing Unit. Information on the complete series of MODUMEDTM Image Processing

☐ A dictionary of Computer Terms used in Digital Nuclear Medicine.

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DYNAMIC STUDIES

- 5. CARDIAC OUTPUT
- 6. STROKE VOLUME
- 7. CARDIAC INDEX
- 8. VENTRICLE VOLUME
- 9. TRANSIT TIMES

10. LEFT TO RIGHT SHUNTS *EJECTION FRACTION can also be determined in dynamic studies.

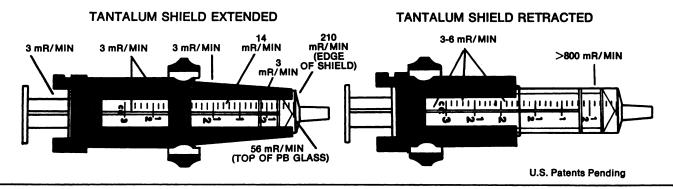
Additionally, static studies of Myocardial Infarct can be evaluated through hot spot imaging with phosphates and cold spot imaging with thallium. The MODUMEDtm System is used for image enhancement, area quantification, background suppression and wall definition.

Because radionuclide angiography will very likely become one of the most common Nuclear Medicine procedures3, it would be to the practitioner's advantage to have data that is quantified in a variety of parameters, repeatable, and readily available.

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FILM EXPOSURE AT THE SURFACE 29.2 mCi for 99mTc in 1cc of a 3cc PLASTIC SYRINGE



*CONCEIVED and DESIGNED BY: J. Howley, H. Tipton, A. Jones, M. Dickinson, M. Green, and G. Johnston. National Institutes of Health, Bethesda, Md.

SHIELDING PERFORMANCE CHART COMPLIMENTS OF: J. Howley, Radiation Safety Services, National Institutes of Health, Bethesda, Md.

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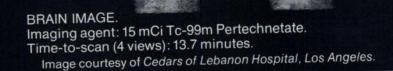
a quiet revolution in WHOLE BODY and ORGAN imaging.

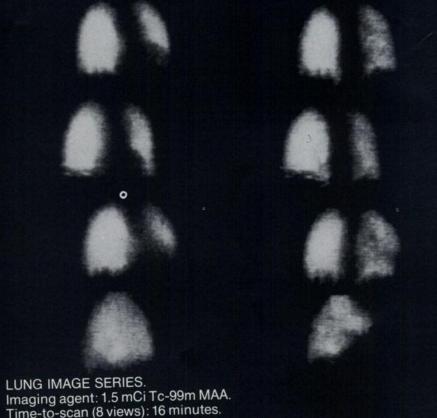
The Cleon Imager fills basic needs in the busy nuclear medicine department. In "WHOLE BODY MODE," it handles patient caseloads three to five times as rapidly as a conventional rectilinear scanner, providing dual anterior and posterior skeletal images of such clarity and sharpness that repeat small-area scans to confirm diagnoses rarely are needed. Yet it can provide, in "ORGAN MODE," small-area organ images with speed comparable to (and in-depth resolution better than) a gamma camera.

Large crystal area (109 square inches in each detector head) gives high information density with reproducible results for given scan times. Interchangeable focused collimators permit use with various nuclides for skeletal and organ imaging, as well as tumor-screening. (The Imager has proved successful in detecting abnormalities in soft tissue when used with Ga-labelled agents.)

The Imager's display and recording options, enhancement of photo-images, and the capability to playback stored data greatly increase its clinical usefulness. Reliability, rapidity of operation, and high patient turnover mean increased utilization and economy, along with improved diagnostic efficiency.

BONE IMAGE OF 58-YEAR-OLD MALE. Imaging agent: 15 mCi Tc-99m Pyrophosphate. Time-to-scan (2 views) 24.8 minutes. Image courtesy of Cedars of Lebanon Hospital, Los Angeles.





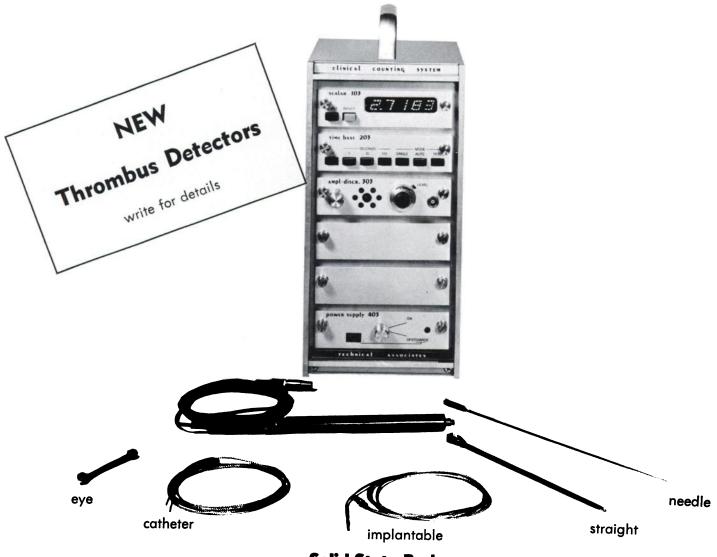
Time-to-scan (8 views): 16 minutes. Image courtesy of Leonard Morse Hospital, Natick, MA.



LIVER AND SPLEEN IMAGE OF PATIENT SHOWING SPLENOMEGALY AND CIRRHOTIC LIVER. Imaging agent: 1.5 mCi TC-99m Sulfur Colloid. Time-to-scan (4 views) 14 minutes. Image courtesy of Cedars of Lebanon Hospital, Los Angeles.



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Ian Falvey found that doing our new Ab-TRAC*digoxin test was easy.

Getting his mom's permission was tough.

lan is ten years old. His mom is a senior medical technologist who knows that tests like our new Ab-TRAC RIA [125] kit for digoxin should only be performed by trained professional technologists.

We agreed. But we also wanted to illustrate how easy the new Ab-TRAC solid phase digoxin [125] kit was to use. Its solid phase design combines 3 steps into 1 (the color coded anti-body and tracer are contained within the Ab-TRAC tubes). New "wet" serum standards require no reconstitution and there is only one incubation. All designed for accuracy and reproducibility, resulting in time saving for the technologist.

Shelley Falvey MT (ASCP) finally consented. Providing she could supervise ("but no coaching, Mom") How did lan do? Just great. His chart was on a par with his mom's when she did the test. Here are lan's comments: "It's easy. It turns colors so you know where to put the stuff."

And here's what Mrs. Falvey said: "This new kit is so easy, even my ten year old boy can do it."

The Falveys' conclusion? The new Schwarz/Mann RIA Ab-TRAC digoxin kit is going to make things a lot

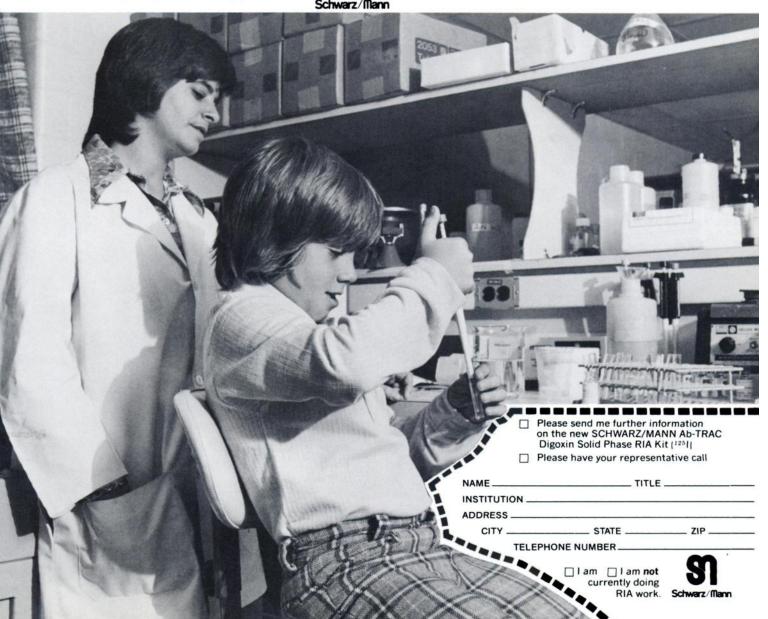
easier for technologists all over the country.

And they don't even have to get their mom's permission.

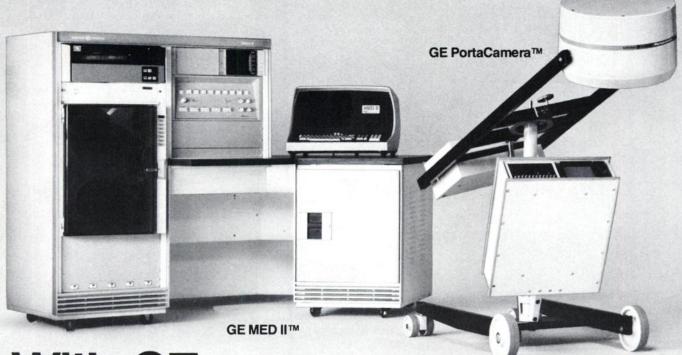
*Ab-TRAC stands for anti-body and tracer contained in tubes. This saves technologists time and eliminates a source of potential pipetting error.

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permits precise, motorless positioning by hand. Performs all Technetium 99m studies with high performance results.

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GE MED STOR™ is a modestly-priced image storage and processing system which can be used with any scintillation camera. Provides computer controlled acquisition of static and dynamic function data, selection of up to 4 regions of interest and simultaneous generation of up to 4 time/activity histograms.



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Volume 16, Number 12

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- 5. Aspirate and wash.
- 6. Count the coated tubes.

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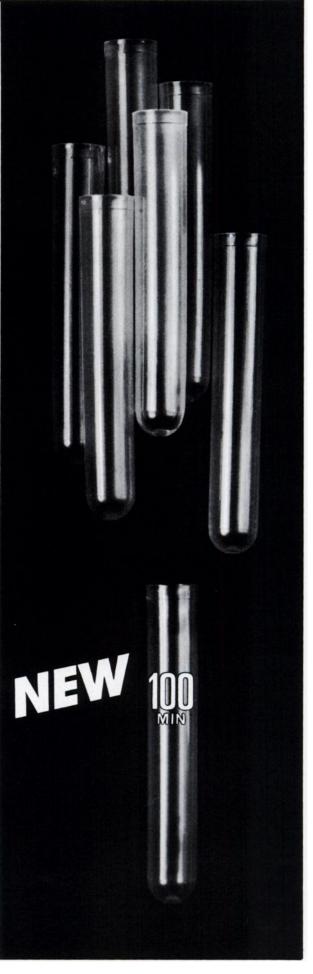
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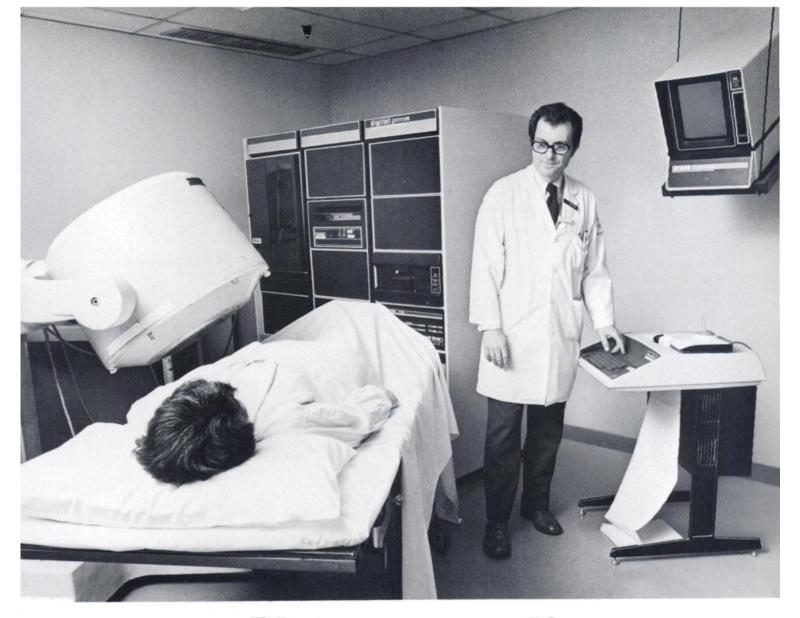
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SPECIFICATIONS

RANGE: 30, 100, 300, 1000, 3000 cps

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TIME CONSTANT: Fast 2 sec., slow 14 sec.

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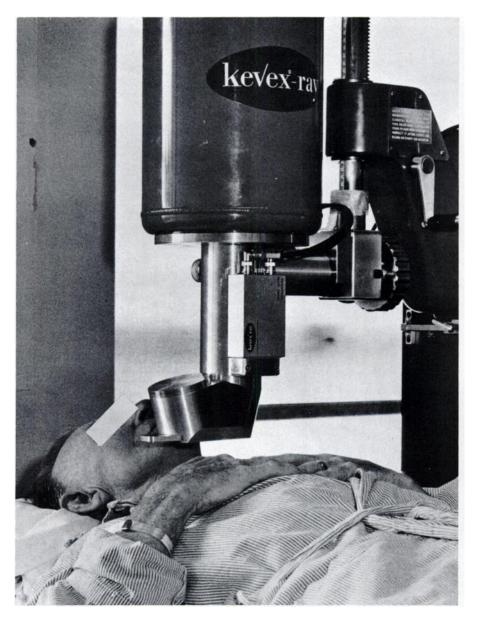
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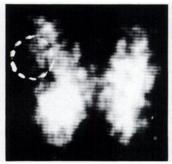
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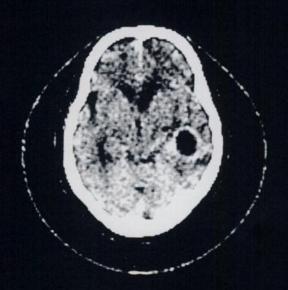
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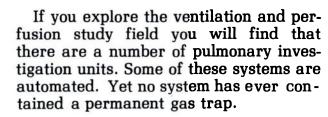
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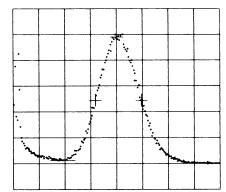
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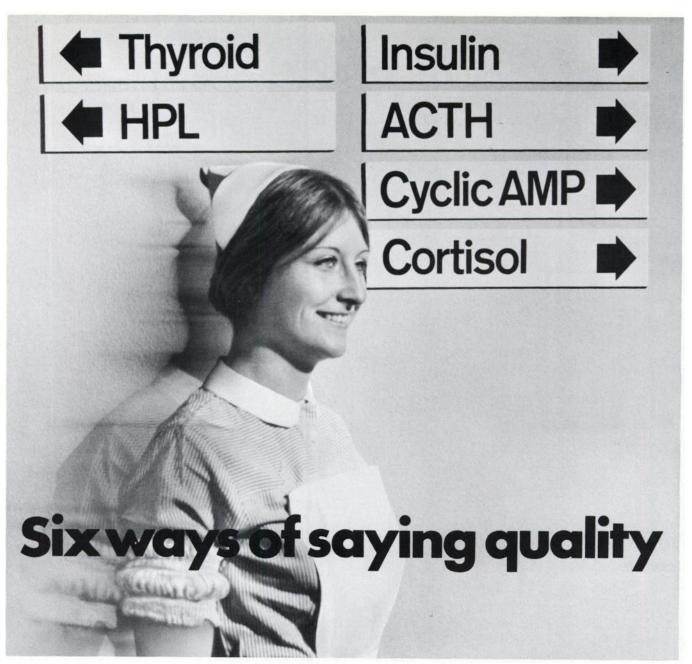
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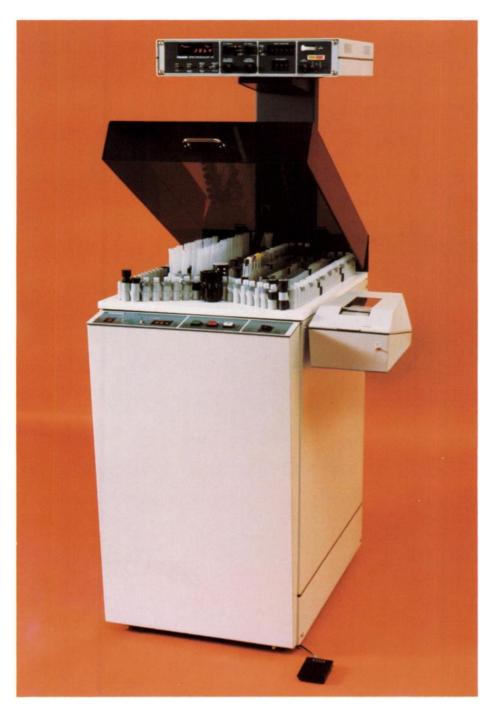
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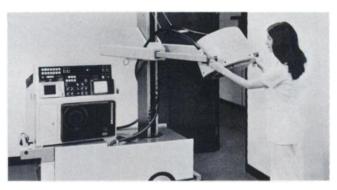
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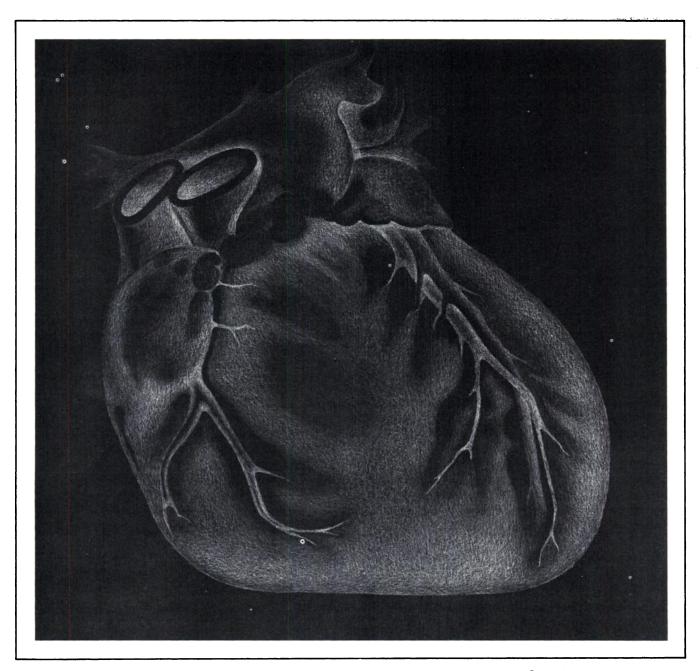


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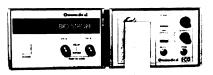
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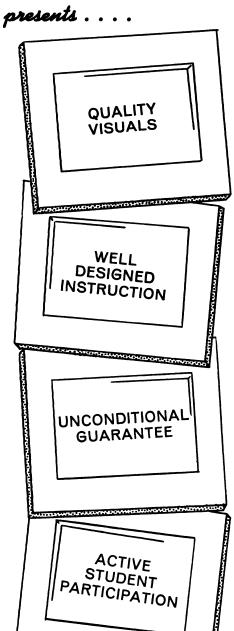
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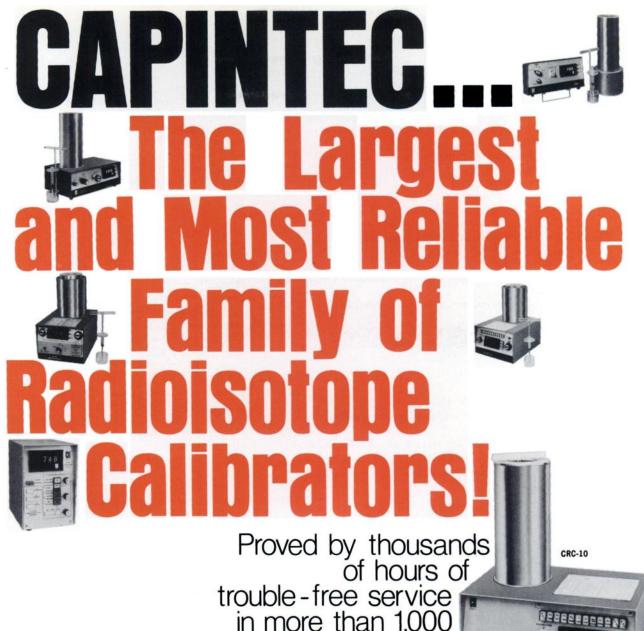
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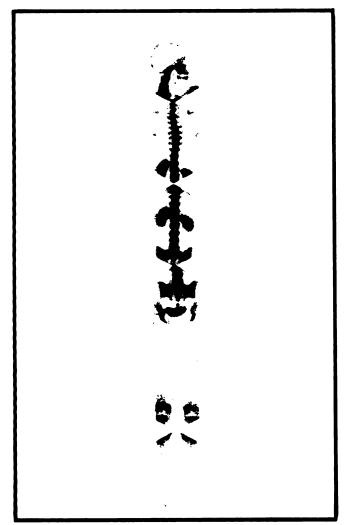
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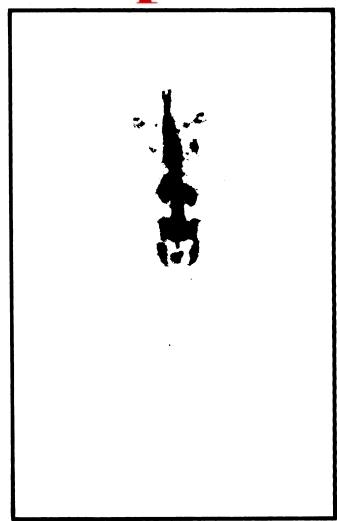
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Study performed with Ohio-Nuclear Series 110 Wide Field Radioisotope Camera.

Study performed with Ohio-Nuclear Series 110 Wide Field Radioisotope Camera equipped with Series 110-8 AreaScan.

35 year old female: normal scan

Study was performed in supine position with posterior

view taken from beneath the table

Collimator: medium resolution (Model 14W11013)

Centerline: 140 keV Window: 20%

Isotope: 20mCi 99mTc Pyrophosphate

Time Begun: 4 hours post dose

Composite View AreaScan

700,000 counts per view except legs were 100,000

counts per view

Total Scan Time: 30 minutes (included positioning)

Total Scan Time: 12.2 minutes



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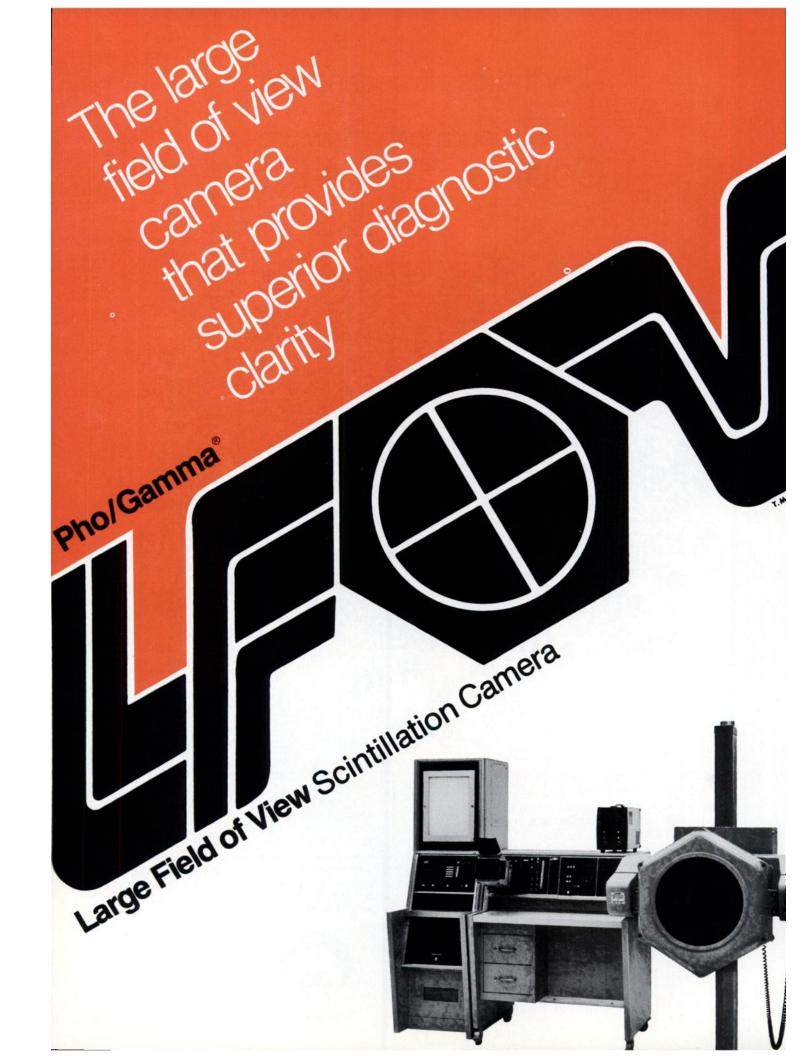
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0-20 second frame . Approx., 300,000 counts 20mCi in Spirometer • 133 Xe

300,000 counts in this posterior ventilation image provide enough photon sufficiency to visualize small ventilatory defects. The patient, a 21 yr. old female, has cystic fibrosis.

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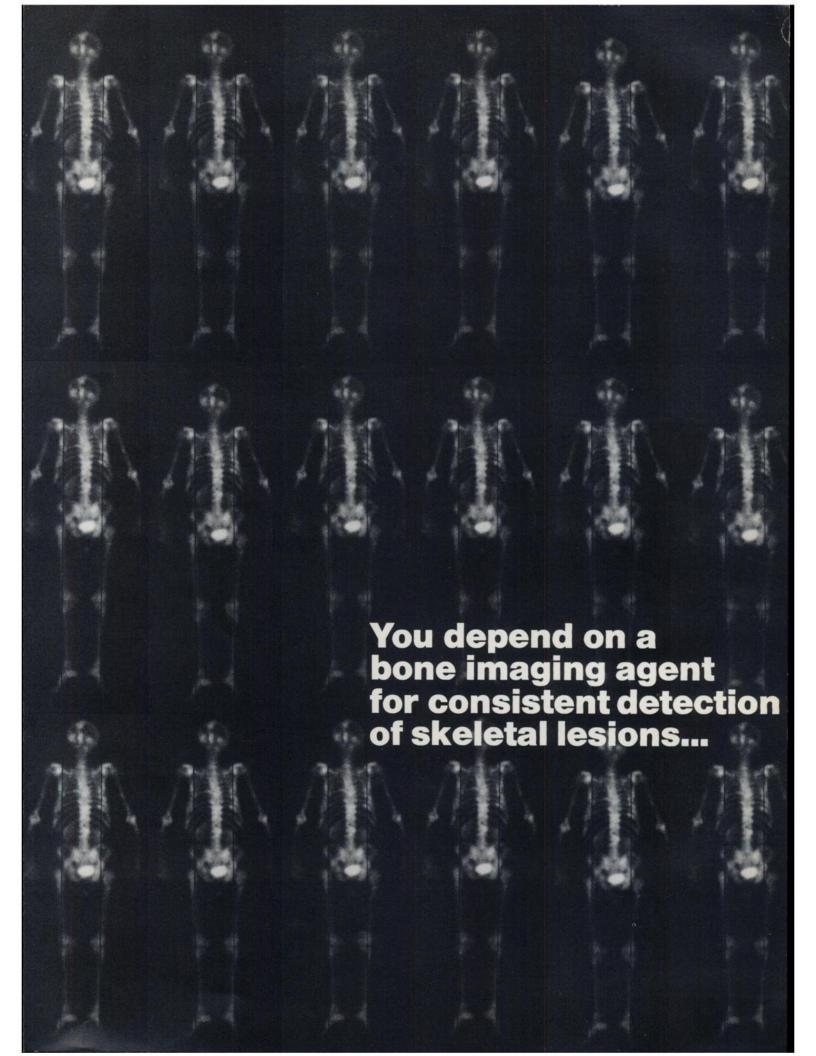


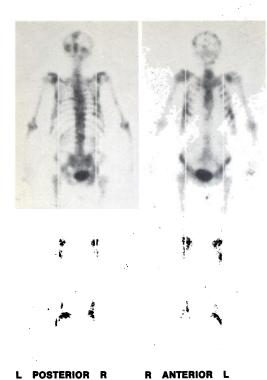
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A 65-vear-old patient with known carcinoma of the prostate. Note pelvic, skull, rib, ster-num and vertebral le-

Imaging Agent: 15 mCi

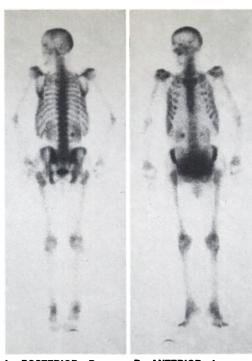
99mTc-OSTEOSCAN Anterior Count per Time:

1,000,000/30 min Posterior Count per Time:

1,000,000/30 min Instrument:

Searle Pho/Gamma® HP camera with whole body table, Microdot Imager® and high-sensitivity collimator Scanned:

3 hours postiniection



66-year-old male with prostatic carci noma and no conclu sive evidence of me tastasis to bone.

Imaging Agent: 15 mCi 99mTc-OSTEOSCAN Posterior Count pe Time: 636,690/35 min Anterior Count pe Time: 613,007/35 min Instrument: Picker Dynacamera 2C with Omniview table and ultrafine collimator

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POSTERIOR R





An 82-year-old patient with extensive meta static bone disease secondary to known carcinoma of the prostate.

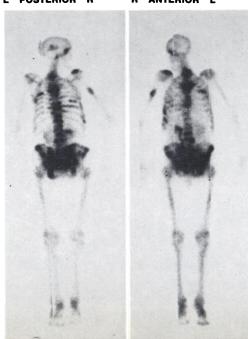
Imaging Agent: 99mTc-OSTEOSCAN Anterior Count per

561,220/30 min Posterior Count per

631,388/30 min Instrument: Picker Dynacamera®

2C with Omniview® table and ultrafine collimator

4 hours postiniection



A 79-year-old mal with known prostati carcinoma metastati to bone. Multiple le sions are seen through out skeletal system.

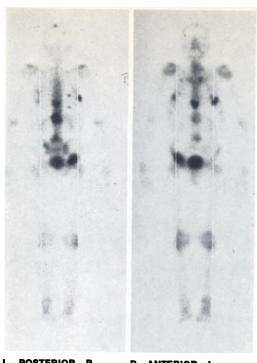
Imaging Agent: 99mTc-OSTEOSCAN Posterior Count pe

621.153/26 min Anterior Count De

649,702/31 min Instrument:

Picker Dynacamera 2C with Omniview table and ultrafin collimator

4 hours postinjectio



A 58-year-old male with a 41-year history of smoking displays extensive metastatic disease in ribs, vertebrai bodies, pelvis, sternum and skull, secondary to known carcinoma of the lung.

Imaging Agent:

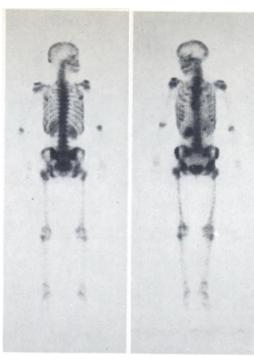
99mTc-OSTEOSCAN Anterior Count per

> 1,000,000/30 min Posterior Count per

> 1.000.000/30 min Instrument:

Searle Pho/Gamma® HP camera with whole body table. Microdot Imager® and high-sensitivity collimator Scanned:

3 hours postinjection



49-year-old female with previous right radical mastectomy for malignancy, having rib pain. Increased uptake in ribs suggests metastatic disease.

Imaging Agent:

99mTc-OSTEOSCAN Posterior Count per

500,381/28 min Anterior Count per

508,462/27 min Instrument:

Picker Dynacamera® 2C with Omniview® table and ultrafine collimator

4 hours postinjection

L POSTERIOR R

R ANTERIOR L

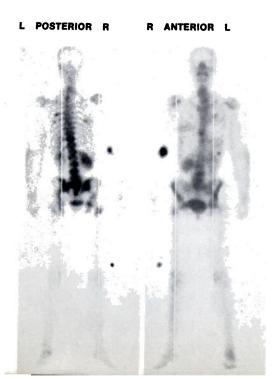
L POSTERIOR R

R ANTERIOR L

OSTEOSCAN consistently delivers:

- Clear, sharp images
- High-quality lesion detection

See following page for brief summary of package insert.



A 43-year-old female with known metastatic disease secondary to carcinoma of the left breast. Swollen left arm is secondary to lymphedema, a result of radical mastectomy. (Note negative defect in region of left breast as a result of prosthesis.) Metastatic disease clearly visualized in vertebral bodies and ribs. Uptake at elbow is extravasation at injection site.

Imaging Agent: 15 mCi

99mTc-OSTEOSCAN Anterior Count per Time:

> 1,000,000/30 min Posterior Count per Time:

> 1,000,000/30 min Instrument:

Searle Pho/Gamma® camera with whole body table, Microdot imager® and high-sensitivity

Scanned: 3 hours postiniection POSTERIOR R R ANTERIOR L ANTERIOR

A 61-year-old male following thoracotomy for carcinoma of the left lung.Two rib fractures (anterior view) of unknown etiology. thumbuptake (posterior view) secondary to arthritic changes.

Imaging Agent:

99mTc-OSTEOSCAN Anterior Count per

Time: 1,000,000/30 min Posterior Count per Time:

1,000,000/30 min Instrument:

Searle Pho/Gamma® camera with whole body table, Microdot imager® and high-sensitivity

Scanned: 5 hours postinjection

DSTEOSCAN...Clear, sharp

rief summary of Package Insert. Before using, please consult e full Package Insert included in each kit.

ESCRIPTION

ach vial of OSTEOSCAN contains 5.9 mg disodium etidronate nd 0.16 mg stannous chloride as active ingredients. Upon addion of ADDITIVE-FREE 99mTc-pertechnetate, these ingredients pmbine with 99mTc to form a stable soluble complex.

CTIONS (CLINICAL PHARMACOLOGY)

When injected intravenously, 99mTc-labeled OSTEOSCAN has a pecific affinity for areas of altered osteogenesis. Areas of bone hich are undergoing neoplastic invasion often have an unusully high turnover rate which may be imaged with 99mTc-labeled STEOSCAN.

hree hours after intravenous injection of 1 ml 99mTc-labeled DSTEOSCAN, an estimated 40-50% of the injected dose has been taken up by the skeleton. At this time approximately 50% has been excreted in the urine and 6% remains in the blood. A small amount is retained by the soft tissue. The level of 99mTc-abeled OSTEOSCAN excreted in the feces is below the level letectable by routine laboratory techniques.

NDICATIONS

DSTEOSCAN is a skeletal imaging agent used to demonstrate reas of altered osteogenesis.

CONTRAINDICATIONS

None.

WARNINGS

This radiopharmaceutical should not be administered to patients who are pregnant or lactating unless the information to be gained outweighs the potential hazards.

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.

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

The 99mTc-generator should be tested routinely for molybdenum preakthrough and aluminum. If either is detected, the eluate should not be used.

PRECAUTIONS

Both prior to and following 99mTc-labeled OSTEOSCAN adminstration, patients should be encouraged to drink fluids. Patients should void as often as possible after the 99mTc-labeled OSTEO-SCAN injection to minimize background interference from accunulation in the bladder and unnecessary exposure to radiation.

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

ADVERSE REACTIONS

lone.

OSAGE AND ADMINISTRATION

The recommended adult dose of 99mTc-labeled OSTEOSCAN is ml with a total activity range of 10-15 mCi. 99mTc-labeled OSTEOSCAN should be given intravenously by slow injection over a period of 30 seconds within three (3) hours after its preparation. Optimum scanning time is 3-4 hours postinjection. The patient dose should be measured by a suitable radioactivity alibration system immediately prior to administration.

..Clear, sharp images for high quality lesion detection... consistently





(5.9 mg disodium etidronate 0.16 mg stannous chloride)

SKELETAL IMAGING AGENT

jnm/placement

POSITIONS OPEN

THE SEATTLE VA HOSPITAL AND the Division of Nuclear Medicine, University of Washington are seeking qualified applicants for a full-time academic position at the Assistant or Associate Professor level. Board eligibility or certification in Nuclear Medicine is required. Candidates with demonstrated clinical, teaching, and research ability are requested to C. V. and reprints to Glen W. Hamilton, M.D., Veterans Administration Hospital, 4435 Beacon Avenue South, Seattle, Washington 98108. Phone: (206) 762-1010, Ext. 459. Non-discrimination in employment.

STANFORD UNIVERSITY—FULLTIME position in Nuclear Medicine now available at Assistant or Associate Professor level. University teaching hospital, Nuclear Medicine facilities expanding. Desired qualifications: Proven excellence in teaching and in all aspects of clinical Nuclear Medicine, plus research experience Excellent opportunity for advancement. Stanford University is an equal opportunity through affirmative action employer. Interested persons please send complete curriculum vitae and names and addresses of references to: Joseph P. Kriss, M.D., Division of Nuclear Medicine, Stanford University Medical Center, Stanford, Calif. 94305.

STAFF NUCLEAR MEDICINE TECHnologist needed for expanding imaging section in large multispecialty clinic and hospital in Big Ten University town. Must have registry now or in near future and have excellent references. Contact: R. A. Baylor, M.D., Department of Radiology, Carle Clinic, Urbana, Ill. 61801. Phone 217-387-3270.

RESEARCH MEDICAL OFFICER. CAreer Federal service. Conduct studies Human Nutrition Research Laboratory, Agricultural Research Service, Grand Forks, North Dakota. Including a whole body gamma scintillation counter located in a steel room. This organization has neutron activation facilities and other support, and well-equipped nuclear medicine laboratory. Applicants must have completed training in nuclear medicine and metabolism. Salary range \$31,552 to \$36,000. Equal opportunity employer. Mary Dickson, USDA, ARS, 2000 W. Pioneer Parkway, Peoria, Ill. 61614, AC 309-673-9061, Ext. 532.

CLINICAL COORDINATOR, SANTA FE Community College, Gainesville, Florida. Position available immediately coordinating clinical involvement in a 2-year nuclear medicine technology program with 4 teaching hospitals in a university setting. Must be registered and experienced in all aspects of nuclear medicine. Good salary proportional to qualifications and excellent fringe benefit package is included. Contact: Robert A. Short, Radiologic Technology Programs, Santa Fe Community College, P.O. Box 1530, Gainesville, Fla. 32601, Phone: (904) 377-5161.

NUCLEAR MEDICINE TECHNOLOgist. Immediate opening in our 1200 plus

bed, centrally located, Indiana teaching hospital. Registry required. Good benefits, salary commensurate with experience. Contact: Personnel Department, Methodist Hospital of Indiana, Inc., 1604 North Capitol Avenue, Indianapolis, Ind. 46202 or call collect 317-924-8931.

NUCLEAR MEDICINE TECHNICIAN. Registered ARRT or ASCP or eligible to registry. Will be cross trained in EMI Scanner use. Competitive salary and benefits. Interested candidates should reply to Director of Personnel, St. Bernardine Hospital, 2101 N. Waterman, San Bernardino, Calif. 92404. Phone 714-883-8711.

NUCLEAR MEDICINE RESIDENCY.
Two-year program affiliated with University of CA/Irvine. Opening available July '76. Active clinical and research program in 1691 bed GM&S teaching hospital. Contact K. P. Lyons, M.D., Chief, Nuclear Medicine Service, V.A. Hospital, Long Beach, Calif. 90801. An equal opportunity employer.

EDUCATIONAL COORDINATOR POSItion available. Methodist Hospital of Indiana, at Indianapolis, a 1200-bed progressive medical center in the heart of
the mid-west, has a recently approved Nuclear Medicine Program. We are looking
for an Educational Coordinator with teaching experience to coordinate the program.
The salary is commensurate with education (degree preferred) and experience,
and we offer excellent benefits. If you
are interested, we invite you to call collect
(317) 924-8931 or send your resume to
Mrs. Mary M. Shaw, Assistant Director of
Employee Relations, Methodist Hospital
of Indiana, 1604 North Capitol Avenue,
Indianapolis, Ind. 46202.

REQUIRED: RADIOCHEMIST FOR ACtive Division of Isotopes in University of Alberta Hospital, involved in service research development and teaching. Ph.D. in related field preferred. Salary dependent on experience. Apply to: Dr. H. E. Bell, Chairman, Dept. of Laboratory Medicine, University of Alberta Hospital, Edmonton, Alberta, T6G 2B7, Canada.

POSITIONS FOR RESEARCH PHYSIcians with training in nuclear medicine, endocrinology, or hematology, at Donner Laboratory, Lawrence Berkeley Laboratory, University of California. California medical license needed. Clinical responsibility and medical school or hospital affiliations optional. Send resume to James L. Born, M.D. or Thomas F. Budinger, M.D., Ph.D., 467 Donner Laboratory, Lawrence Berkeley Laboratory, University of California, Berkeley, Calif. 94720. An equal opportunity employer.

NUCLEAR MEDICINE TECHNOLOgist. Staff Technologist for new 363-bed hospital located in community of 35,000, unlimited recreation, educational and cultural opportunities, in Southeastern Ohio. Excellent fringe benefits. Salary open; call collect: William B. Montgomery, Assistant Administrator (614) 454-4601. Bethesda Hospital, 2951 North Maple Ave., Zanesville, Ohio.

POSITION WANTED

NUCLEAR AND DIAGNOSTIC RADIologist; 33, certified ABNM and ABR. University trained including fellowship in nuclear medicine, seeks position in radiology and/or nuclear medicine, near large metropolitan area. Full capabilities incuding angiography and ultrasound. Reply to: Box 1200, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

NUCLEAR MEDICINE TECHNOLOgist desires to relocate. Graduate of prestige university with many years field experience. Versed in opening and managing Nuclear Deaprtments. Reply: Box 1201, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

NUCLEAR MEDICINE PHYSICIAN, ABNM (Radiology background), well trained and experienced in all aspects of Nuclear Medicine, administrative experience, desires full-time position in clinical Nuclear Medicine, prefers Southwest or Pacific Coast, reply with job description and potential, available 7/1/76. Box 1202, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

NUCLEAR MEDICINE DEPARTMENtal administrator and technical director of education, university affiliated medical center technologists course. Sixteen years experience, B.S. degree, certification N.M.T. (ASCP-ARRT). Desires position in comprehensive nuclear medicine department. West coast area or south west but will consider relocating elsewhere. Curriculum vitae upon request. Box 1208, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

NUCLEAR MEDICINE PHYSICIAN, completed two-year fellowship with university affiliated hospital, with medical background, seeks full-time position with clinical and research responsibilities. Reply to Box 1204, Society of Nuclear Medicine, 475 Park Avenue South, New York, N.Y. 10016.

M.D., NUCLEAR MEDICINE BOARD certified, experienced in university teaching, desires full-time position in clinical and academic nuclear medicine. Box 1206, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

M.S., ASCP/AART NUCLEAR MEDIcine Technologist desires a position as Chief Technologist. Four years field experience. Reply Box 1206, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

CHIEF NUCLEAR MEDICINE TECHnologist, ARRT registered. Eight years
experience. Capabilities include in vivo and
in vitro applications. Expert with most
equipment and procedures. Interested in
planning, organizing and managing established or new facilities. Prefer to relocate
north west or north east U.S. Reply to
Box 1207, Society of Nuclear Medicine, 475
Park Avenue South, New York, N.Y. 10016.

NUCLEAR MEDICINE RESIDENCY

Two-year approved program offering broad clinical and basic science experience including pediatrics, oncology, radiation safety, radiopharmacy management and research. An integrated program at State University of New York at Buffalo School of Medicine.

Contact: Merrill A. Bender, M.D. Program Director

r Monte Blau, Ph.D. Chairman Dept. of Nuclear Medicine 666 Elm Street Buffalo, New York 14263

RESIDENCY AND FELLOWSHIPS IN NUCLEAR MEDICINE NOW AVAILABLE

For information contact:

John A. Burdine, M.D.
Chief, Nuclear Medicine Section
Department of Radiology
Baylor College of Medicine
Texas Medical Center
Houston, Texas 77025
Phone (713) 521-2272

VETERANS ADMINISTRATION HOSPITAL UNIVERSITY OF MINNESOTA NUCLEAR MEDICINE RESIDENCY PROGRAM

Resident positions are available in an AMA-approved two-year training program beginning January 1, 1976. A combined University of Minnesota VA Hospital program with active clinical and research opportunities. Minimum stipend \$13,100.

AN EQUAL OPPORTUNITY EMPLOYER

For further information, contact:

Merle K. Loken, M.D., Ph.D.
Director, Division of Nuclear Medicine
University of Minnesota Hospitals
Box 382, Mayo Memorial Building
Minneapolis, Minn. 55455

OR

Rex B. Shafer, M.D.
Chief, Nuclear Medicine Service (172)
Veterans Administration Hospital
54th Street & 48th Avenue South
Minneapolis, Minn. 55417

EXPERIENCED NUCLEAR PHYSICIAN

Massachusetts General Hospital Harvard Medical School Nuclear Medicine Division Department of Radiology

ABNM Certification Required

Clinical and Research Competency
Emphasized

CONTACT

Juan M. Taveras, M.D. Radiologist-in-Chief

OI

Majic S. Potsaid, M.D.

Director, Nuclear Medicine Division

Department of Radiology

Massachusetts General Hospital

Boston, Massachusetts 02114

Equal Opportunity Employer

INTRODUCTORY ONE WEEK PHYSICIAN COURSE IN NUCLEAR MEDICINE Cleveland, Ohio

Contact:

D. Bruce Sodee, M.D., DirectorNuclear Medicine Institute6780 Mayfield RoadCleveland, Ohio 44124

1976

January 12–16, 1976 March 8–12, 1976 May 10–14, 1976 July 12–16, 1976 November 8–12, 1976

ASSOCIATE DIRECTOR NUCLEAR MEDICINE

450 Bed Teaching Hospital Northern New Jersey

Internal Medicine Background Preferred
Must be Board Eligible or Certified

Send resume and salary requirements to Box 1209

Society of Nuclear Medicine 475 Park Ave. South New York, N.Y. 10016

JNM CLASSIFIED PLACEMENT SERVICE SECTION

This section in the Journal of Nuclear Medicine contains "Positions Open", "Positions Wanted", and "For Sale" listings. Nondisplay "Positions Wanted" ads by members of the Society are billed at 30¢ per word for each insertion with no minimum rate. Nondisplay "Positions Wanted" ads by nonmembers and all nondisplay "Positions Open" and "For Sale" ads by members and nonmembers are charged at 65¢ per word, with a minimum of \$15. Display advertisements are accepted at \$50 for ½ page, \$90 for ¼ page, \$165 for ½ page, and \$295 for a full page. Closing date for each issue is the 15th of the second month preceding publication. Agency commissions and cash discounts are allowed on display ads only. Box numbers are available for those who wish them. All ads must be prepaid.

Please note our new address.

JOURNAL OF NUCLEAR MEDICINE 475 Park Ave. South, New York, N.Y. 10016

Immediate opening, Assistant Chief Technologist, ARRT or ASCP registered to aid in supervising an expanding imaging laboratory of a 300-bed teaching hospital. Overtime and standby pay, with paging device provided. Emergency call once every six weeks. Excellent growth opportunity. Outstanding recreational area. Competitive salary and benefits. An equal opportunity employer. Contact

> PAUL CHRISTIAN, Division of Nuclear Medicine, University of Utah Medical Center, Salt Lake City, Utah 84132. Telephone: (801) 581-2716.

NUCLEAR MEDICINE CHIEF TECHNOLOGIST

ASRT or ARRT registered, B.A. or B.S. in Nuclear Medicine, 3 years experience minimum; must have working knowledge of Rectilinear and camera scanning, and complete "wet lab" procedures, including RIA. Managerial experience helpful. Must perform not only test procedures, but handle ordering, budgeting, radiation safety program and personnel items.

Send resumes to Box 1208, Society of Nuclear Medicine, 475 Park Ave. South, New York, N.Y. 10016.

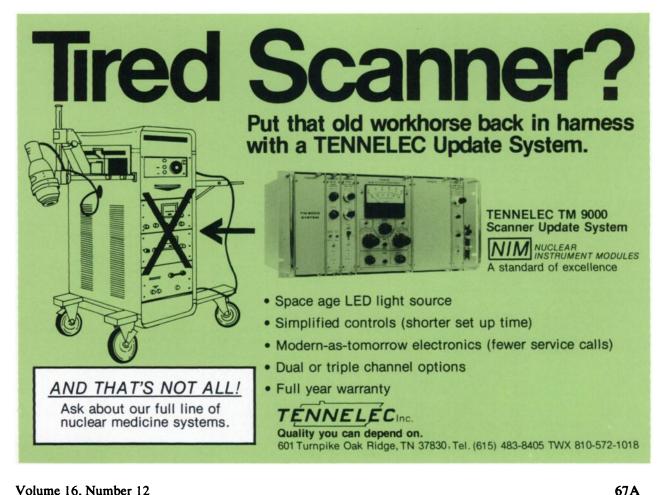
Western Regional Registry Review and Continuing Education Meeting

Sponsored by

The Technologist Section of the Northern California Chapter, Society of Nuclear Medicine February 19-21, 1976 Harrah's Convention Center Reno, Nevada

The Western Regional Registry Review and Continuing Education Meeting will offer points of interest for those students who will be taking the registry examination, for the physician who will be taking the ABNM exam and a well rounded continuing education program for the technologist. Topics will include the Quality Assurance of Scintillation Camera Programs developed by William R. Hendee (Bureau of Radiological Health, Food and Drug Administration, DHEW,) RIA Program, Radiopharmacology, Radiation Safety, Physiology and Anatomy, and Administration. Registration will open on Thursday evening Feb. 19th; exhibits will be open at that time and there will be a no-host cocktail party.

Contact: Jean Lynch, Exec. Secy., P.O. Box 40279, San Francisco, Calif. 94140



Volume 16, Number 12



INDICATIONS. Technetium Tc 99m MAA is indicated as a lung imaging agent to be used as an adjunct in the evaluation of pulmonary perfusion.

Specifically, the distribution of the agent reflects regional pulmonary perfusion and may be helpful in the evaluation of such clinical conditions as pulmonary embolus, chronic obstructive lung disease, congenital anatomic abnormalities, and pulmonary abscess. It can also be used in conjunction with a suitable liver imaging agent for the performance of lung-liver scans to detect subphrenic abscesses.

CONTRAINDICATIONS. The safety of aggregated albumin in patients with right-to-left cardiac shunts has not been demonstrated, and its use in such patients is contraindicated. The use of Tc 99m macroaggregated albumin is contraindicated in persons with a history of hypersensitivity reactions to products containing human serum albumin.

WARNINGS. Although not reported to date, the possibility of allergic reactions should be considered in patients who receive multiple doses. This radiopharmaceutical preparation should not be administered to pregnant or lactating women, or persons under 18 years of age unless the benefits to be gained outweigh the potential hazards.

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.

Theoretically, the intravenous administration of any colloid material such as aggregated albumin imposes a temporary small mechanical impediment to blood flow. While this effect is probably physiologically insignificant in most patients, the administration of aggregated albumin is possibly hazardous in acute cor pulmonale and other states of severely impaired pulmonary blood flow. Although not reported with NEN's Tc 99m Aggregated Albumin, the literature contains four reports of deaths occurring after the administration of aggregated albumin to patients with pre-existing severe pulmonary hypertension.

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

Radiopharmaceuticals should be used only by physicians who are qualified by specific training in the safe use and handling of radionuclides produced by a nuclear reactor or particle accelerator and whose experience and training have been approved by the appropriate governmental agency authorized to license the use of radionuclides.

The labeling reactions involved in preparing the agent depend on maintaining the tin in reduced state. Any oxidant present in the Pertechnetate Sodium Tc 99m supply may thus adversely affect the quality of the prepared agent. Hence, Pertechnetate Sodium Tc 99m containing oxidants, or other additives, should not be employed without first demonstrating that it is without adverse effect on the properties of the resulting agent.

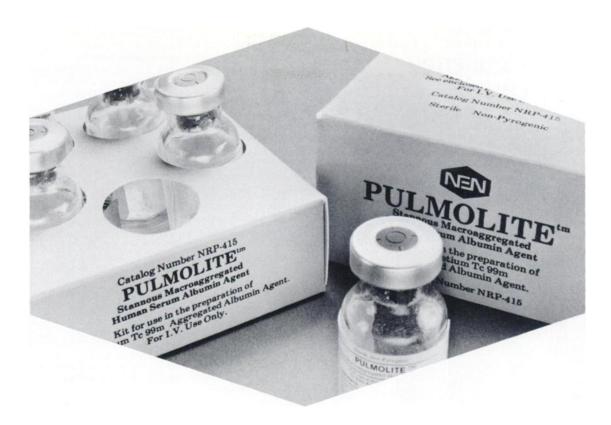
PRECAUTIONS. The contents of the vial are sterile and non-pyrogenic. It is essential that the user follows the directions carefully and adheres to strict aseptic procedures during preparation of the product.

PULMOLITE Agent should be used within 8 hours after reconstitution with Pertechnetate Sodium Tc 99m. Refrigerate after reconstitution.

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

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

ADVERSE REACTIONS. Although no adverse reactions have been reported using NEN Technetium Tc 99m Aggregated Albumin, rare instances of hemodynamic or idiosyncratic reactions to other preparations of Tc 99m labeled macroaggregated albumin have been recorded.



Stat lung scan

Just add Tc 99m, shake, inject, and do your scan.

Convenient – No special storage conditions or equipment required

Flexible – You can reconstitute with 2-8ml containing 15-80mCi of TcO₄

Economical – Lyophilized preparation may be stored at room temperature for up to one year, allowing large quantity purchases at a savings **Labeling efficiency** – Typical efficiency is over 95% to give you high quality imaging

Uniform particle size – Typically 90% of the aggregates are well within the range of 5-75μm, and none larger than 150μm

Contact your NEN Representative or Customer Service for further details



Atomlight Place, North Billerica, Mass. 01862 Telephone 617-667-9531

Los Angeles: 213-321-3311 Miami: 305-592-0702

Canada: NEN Canada Ltd, Lachine, Quebec. Tel: 514-636-4971 Europe: NEN Chemicals GmbH, D6072 Dreieichenhain, W. Germany, Siemensstrasse 1. Tel: Langen 06103-85035

A kid with leukemia can die from a cold.



Leukemia is a disease of the blood-forming tissues. It keeps the body from producing the necessary amounts of normal white blood cells to fight infection.

An infection that means a day in bed for a normal child is a threat to the life of a child with leukemia.

Today research has made enormous progress. At one time, leukemia victims lived only a few months.

Now, in some cases, we can prolong their lives a few years. That's good. But not good enough.

Even though we're closer to a cure, leukemia is still the major cause of disease and death in kids between the ages of 3 and 14.

We want to save the life of every leukemia victim.

We can't do it without a healthy contribution from you.

We want to wipe out cancer in your lifetime. Give to the American Cancer Society.

This space contributed by the published

PHOSPHOTEC®

Technetium 99m-Stannous Pyrophosphate Kit

Phosphotec provides all the nonradioactive components required to prepare ^{99m}Tc-stannous pyrophosphate complex. Each vial contains a sterile, nonpyrogenic lyophilized powder prepared from 40 mg. tetrasodium pyrophosphate decahydrate (equivalent to 23.9 mg. tetrasodium pyrophosphate) and 1.0 mg. stannous fluoride; pH is adjusted with sodium hydroxide or hydrochloric acid. The product does not contain a preservative. At the time of manufacture, the air in the vials is replaced by nitrogen.

Reconstitution of Phosphotec with sterile sodium pertechnetate-^{99m}Tc results in an aqueous solution of Technetium 99m-Stannous Pyrophosphate Complex.

INDICATIONS: Technetium 99m-Stannous Pyrophosphate Complex is indicated for use as a bone imaging agent to define areas of altered blood flow in osseous tissues.

CONTRAINDICATIONS: At present, there are no known contraindications to the use of ^{99m}Tc-stannous pyrophosphate complex.

WARNINGS: The contents of the Phosphotec (Technetium 99m-Stannous Pyrophosphate Kit) vial are intended only for use in the preparation of ^{99m}Tc-stannous pyrophosphate complex and **are NOT** to be directly injected into a patient prior to labeling.

Phosphotec (Technetium 99m-Stannous Pyrophosphate Kit) is not radioactive. However, after 99mTc-sodium pertechnetate is added, adequate shielding of the resulting preparation must be main-

Radiopharmaceuticals should be used only by physicians who are qualified by specific training in the safe use and safe handling of radionuclides, produced by nuclear reactor of cyclotron, and whose experience and training have been approved by the appropriate federal or state agency authorized to license the use of radionuclides.

This radiopharmaceutical should not be administered to patients who are pregnant or during lactation unless the information to be gained outweighs the possible potential risks from the radiation exposure involved.

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.

PRECAUTIONS: It is essential that the user follow the directions carefully and adhere to strict aseptic procedures during preparation of the product.

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

To minimize visualization of the bladder, the patient should be encouraged to void immediately prior to the examination; prior hydration of the patient may be useful.

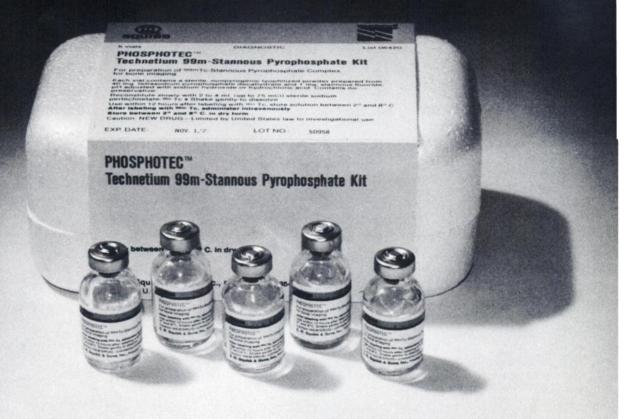
Use the preparation within 12 hours after labeling with ^{99m}Tc.

ADVERSE REACTIONS: At present, adverse reactions have not been reported following the administration of ^{99m}Tc-stannous pyrophosphate complex.

HOW SUPPLIED: Phosphotec (Technetium 99m-Stannous Pyrophosphate Kit) is supplied in a kit containing five vials.

SQUIBB® 'The Priceless Ingredient of every product is the honor and integrity of its maker.'TM

Now available for skeletal imaging



PHOSPHOTEC®

Technetium 99m-Stannous Pyrophosphate Kit

20.5

(ratio of Pyrophosphate to Stannous Tin)

SQUIBB QUALITY—THE PRICELESS INGREDIENT

Unlike many companies involved in nuclear medicine, Squibb is also a broad line pharmaceutical house ... and has been for over a century. So when it comes to formulation and quality control procedures, we wrote the book. Consider that before you purchase any radiopharmaceutical. At Squibb, quality is a way of life.

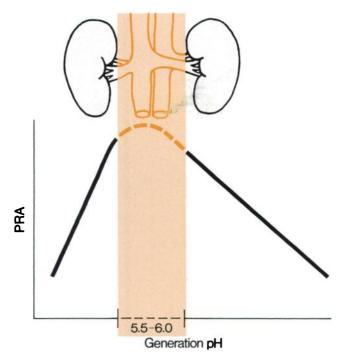
SQUIBB HOSPITAL DIVISION

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The Angiotensin I [1251] Kit For Plasma Renin Activity



The original Angiotensin I RIA kit with a buffered generation system

The original Angiotensin I RIA kit utilizing an optimized pH for generation of Angiotensin I

The original Angiotensin I RIA kit employing a mono-iodinated, highly immunoreactive, stabilized tracer

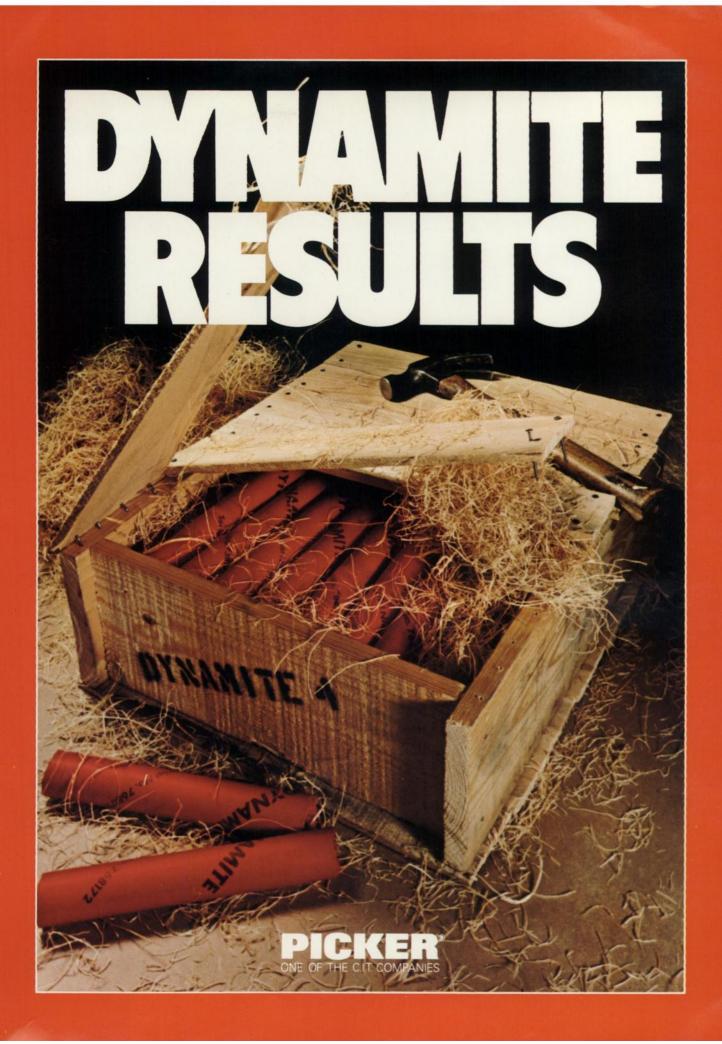
Now the first commercial Angiotensin I RIA kit with time-saving, individual, pre-calibrated, lyophilized standards

Send for clinical validation and technical data.



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Brain Study Bone Study Lung Study

For dynamite clinical results, rely on the DynaCamera 4 System, presently the only system with choice of three detector sizes: 10", 12" or 15". Contact your Picker representative or write: Picker Corporation, 12 Clintonville Road, Northford, CT 06472.

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We've added...

PTH!!~PARATHYROID HORMONE and PROLACTIN!!

I. ENDOCRINOLOGY	Meximum Turn-Aroun
THYROID	Time (Days)
1. Y ₃	- i -
2. T.	<u>i</u> _
3. T, or FTI	<u>_</u>
4. PBI	1
5. TBI	1
6. Free Thyrexine * * *	2
7. E.T.R. (Effective Thyrexine Ratio)	3
8. T ₃ Radioimmunoessay	4
9. T ₄ Radioimmunoessay	7
ANTERIOR PITUITARY	
10. T.S.H. (Thyroid Stimulating Hormone)	5
11. F.S.H. (Follicle Stimulating Hormone)	7
12. L.H. (Luteinizing Hermone)	7
13. H.G.H. (Human Grawth Hormone)	7
14. A.C.T.H.	15
15. Prolactin	10
PLACENTA	
16. H.C.G. (Human Chorionic Gonadotropin) B/Sub Unit	
17. H.C.S. (Human Chorionic Sematemammetropin)	7
GONADS	
18. Estragons, Total	
19. Estradiol	7
20. Estriol	
21. Estrone	7
22. Progesterone	7.
23. Testosterone	
ADRENAL CORTEX	
24. Cortisol	3
25. Aldosterene	10
PARATHYROID	
26. Parathyroid Hermone	15
27. Calcitonin	15
PANCREAS	
28. Insulin	5

II. CARDIOVASCULAR
Toro-Around Time (Board
29. Digestin J Hrs.
30. Digitekin 2 Days
31. Angiotensin I (Plasma Renin Activity) 4
III. HEMATOPOIETIC
32. Vitemin B ₁₂ 5
33. Felic Acid 5
34. Serum Iron (Total & Unceturated binding exceeded) 5
34. Serum Iron (Total & Uncotorated binding expectly) 5
IV. IMMUNE
35. H.A.A. (Hopatitis Associated Antigon) 1 36. IgE 7
37. Cercinoembryonic Antigen 4
V. MISCELLANEOUS
38. Morphine 3
39. Cyclic AMP 4
40. Gestrin 4
41. T.B.G. (Thyroid Binding Globulin) 3
VI. SPECIAL PACKAGES
42. Allergy Assays (RAST)
Over Fifty (50) Different Allergens — 43. OB and GYN Complete Package
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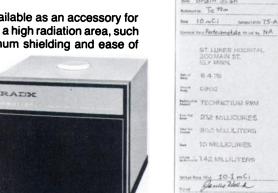
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operation. When the remote chamber is connected, the Melétron's internal chamber is deactivated.



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The Melefile permanent record storage system — instant NRC (AEC) accountability. RADE

SENAL (II) mes Smith, Mary in 1275127 Tuber No. 0287653 tem No. 220A more Marcus Welby no 6/4/75 me Brain Scan Reference 10 mCi temperature 75-A(23

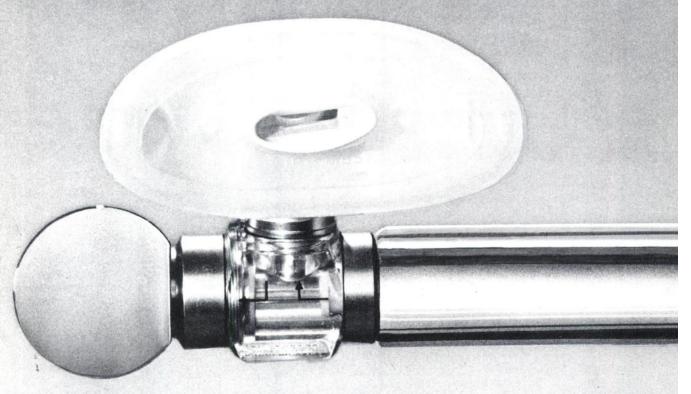
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Volume 16, Number 12 77A

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Multi-Imager 1

Multi-Imager 1 employs the CRT of the gamma camera to record static, dynamic, and whole body imaging procedures on transparency format. The highly versatile Multi-Imager 1 offers film size formats of 5x7 and 8x10, yielding superior quality transparency scintiphotos recorded on a wide range of x-ray film processor compatible films. Up to 30 images can be recorded on a single sheet of film in ten different formats. In addition to the usual 1, 4, and 16 image formats, Multi-Imager 1 offers seven further choices to yield the exact diagnostic format required. For example, Multi-Imager 1 offers a 6 image format to allow recording of static studies that require a fifth and sixth view, and a 30 image format for dynamic studies that require more than sixteen frames. For whole body imaging, the 2 image format records side by side AP and PA views on the same sheet of film. Static, dynamic, and different size images can be mixed on the same sheet of film.



Multi-Imager 4

Multi-Imager 4 yields unmatched performance in gamma camera hard copy recording. A built in high resolution CRT, state of the art microprocessor technology, and electronically synchronized multiple lens optics provide a very small dot size on 8x10 format without increasing the pulse pair resolution dead time of the gamma camera system. The fast lens system of Multi-Imager 4 is compatible with both conventional x-ray film and the slower single emulsion radiographic films that provide the best image quality. Up to 64 images can be recorded in ten different formats. The dual intensity recording mode allows simultaneous acquisition of whole body or static views at two different intensity levels. Positive patient indentification is achieved through a nine digit keyboard LED system.

Both Multi-Imager 1 and Multi-Imager 4 can provide thousands of dollars in annual film cost savings and are compatible with all gamma cameras. Mail coupon to receive detailed information and sample clinical studies.

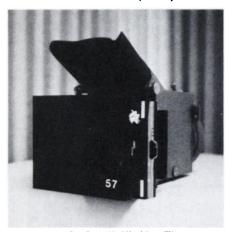
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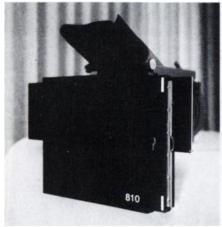
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- Model No. 1114 For your "special" requirements (3 "Y" positions)
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- Works with triple or single lens cameras
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^{*}As shown at the 22nd Annual Meeting of the S.N.M. in Philadelphia, PA.

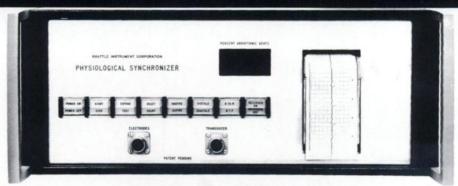
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Help your cardiologist study heart kinetics non-invasively with Brattle-gated scintiphotos.



The RAO view shows akinesis of the lower antero-lateral wall and apex; and contraction of the inferior wall and high up the antero-lateral wall. The LAO view shows good contrac-

tion posteriorly and akinesis of the septal aspect of the chamber. Patient was injected IV with 20mCi of ^{99m}Tclabelled Human Serum Albumin. The agent was prepared using the New England Nuclear Electrolysis Kit for labelling HSA. Write or call for a portfolio of Brattle-gated lung, liver and heart studies.



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The spartan panel above tells the second-best part of our story. If you want to photograph peak systole, press the SYSTOLE button. If, say, you want systole only at full expiration, press the EXPIRATION button as well. If only breathing is relevant, don't press the heart button.

The Brattle is connected to the patient and to your gamma (or x-ray or ultrasonic) camera. Whenever the patient is in the selected phase, both the scope and the scaler on your gamma camera are gated ON, and film is exposed. Otherwise, they are OFF.

Brattles lock onto patients – and stay locked on

It doesn't matter if the patient's heart rate and breathing depth change while he's under the collimator because we stay right with him. Brattles contain an ECG to track heart, a plethysmograph to track respiration, and a tiny computer to deduce systole and diastole times from the heart signal. And because it's all built in, your operator need not be a physiologist.

We don't cover our tracks—we print them

The panel lights flash whenever the patient reaches the selected phases; and pushing the RECORDER-ON button gets you an ECG tracing marked with breathing and cameraon times. You can verify function before, during and after exposure.

A single pair of axillary electrodes captures both heart and breath It's easy. And we supply disposable, pre-filled electrodes.

Some Brattles have been in clinical use for over three years—in community and major hospitals

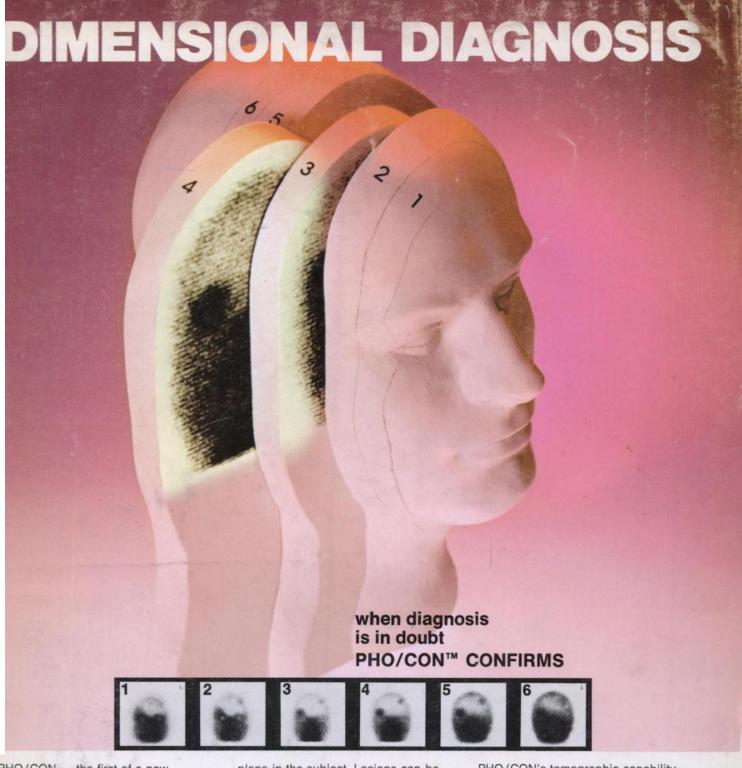
More than half of our instruments are in community hospitals and the list is growing rapidly. Upon request, we'll supply names of happy users in your area.

What's the next step? Get in touch

Ask your NEN man about Brattles and HSA Kits. He can show you a portfolio of clinical pictures and arrange to have one of our people give you a demo. Or write or call us direct. We'll send you brochures on this and other models, and will give you your own set of clinical pictures and a bibliography on gated scintigraphy. If you wish, we'll even make you a Brattle owner. (This is the best part of our story.)

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