

You can see the difference.

2 ml Ampul

**AGGREGATIN**  
**LUNGACIN**

intravenous injection  
Indicated Adult Dose

See Insert for  
Federal (U.S.)  
before use



# 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-and-spleen 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-to-left 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.



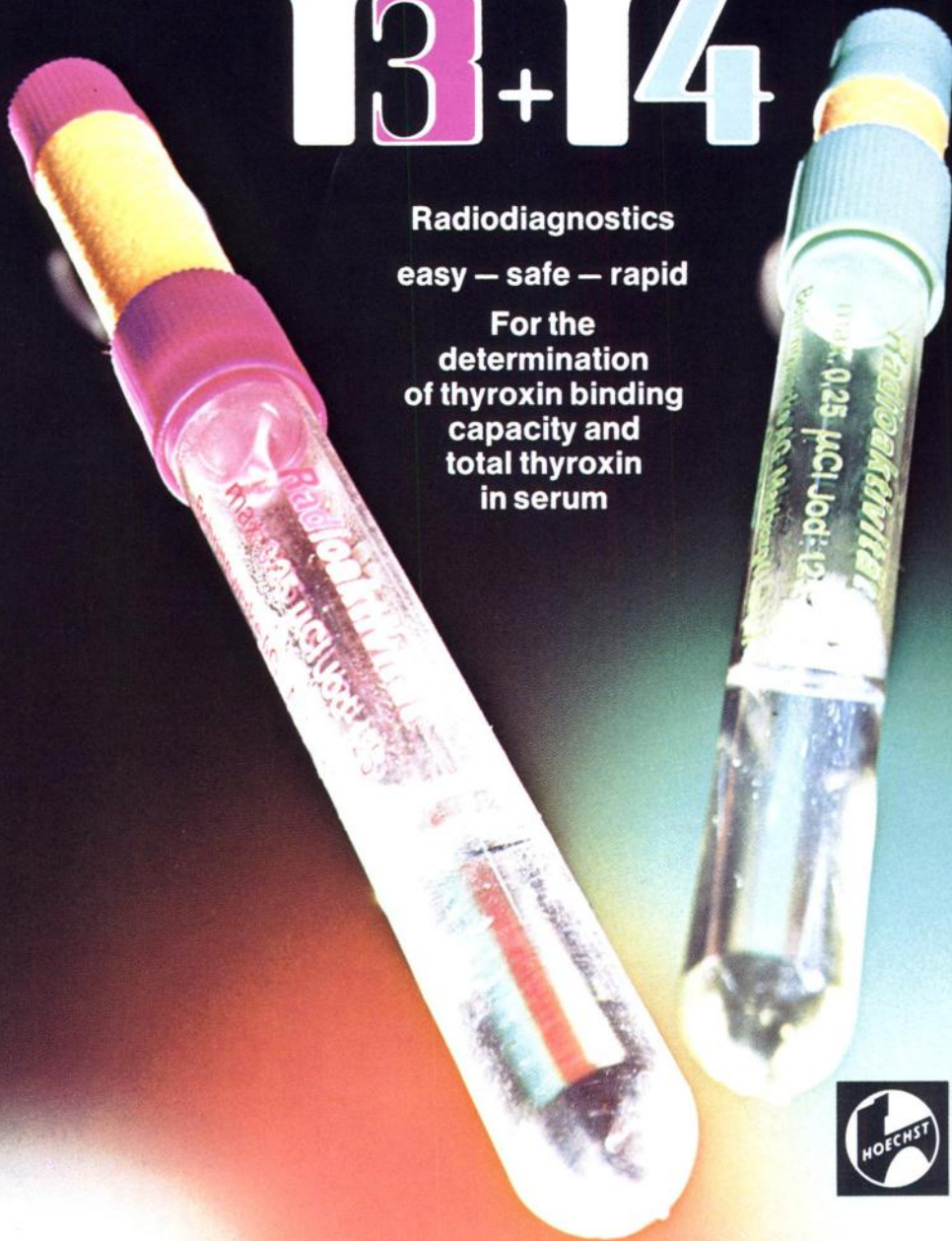
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**Storage:** store protected from light in the refrigerator at +4° to +6° C  
**Stability:** 8 weeks at proper storage. The expiry date is indicated on the package.

Order No.: J 5113  
for T3      1 package 12 tests

**Contents T 4 kit:** 12 calibrating tubes with 3.3 ml TBG-T 4- (J-125)- solution each • total activity: 1  $\mu$ Ci J-125 • preservative: 0,02% sodium azide • 12 adsorption tubes • 1 standard serum of defined T 4-concentration •

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**RAYTHEON**





A large blue diamond-shaped graphic with a white border, centered on a white background. Inside the diamond, the text "New England Nuclear Radiopharmaceuticals" is written in white, bold, sans-serif font.

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Patient Name: Smith, Mary		
Room No. 02B7653	Lot No. 25127	
Physician: Marcus Welby Date: 6/4/75		
Title: Brain Scan		
Radionuclide: Tc 99m		
Dose: 10 mCi		
Injection Site: 75-A123		
Contact: Perfectionplate No. 10 NA		
ST. LUKE'S HOSPITAL		
205 MAIN ST.		
ELY MINN.		
Date of Birth: 6-4-75		
Time of Day: 0900		
Radionuclide: TECHNETIUM 99M		
Dose: 212 MILLICURIES		
Vol of Dose: 30.0 MILLILITERS		
Dose: 10 MILLICURIES		
Vol Admin: 1.42 MILLILITERS		
Spray Area: 10.1 mCi		
Signed: <i>Dennis Black</i>		

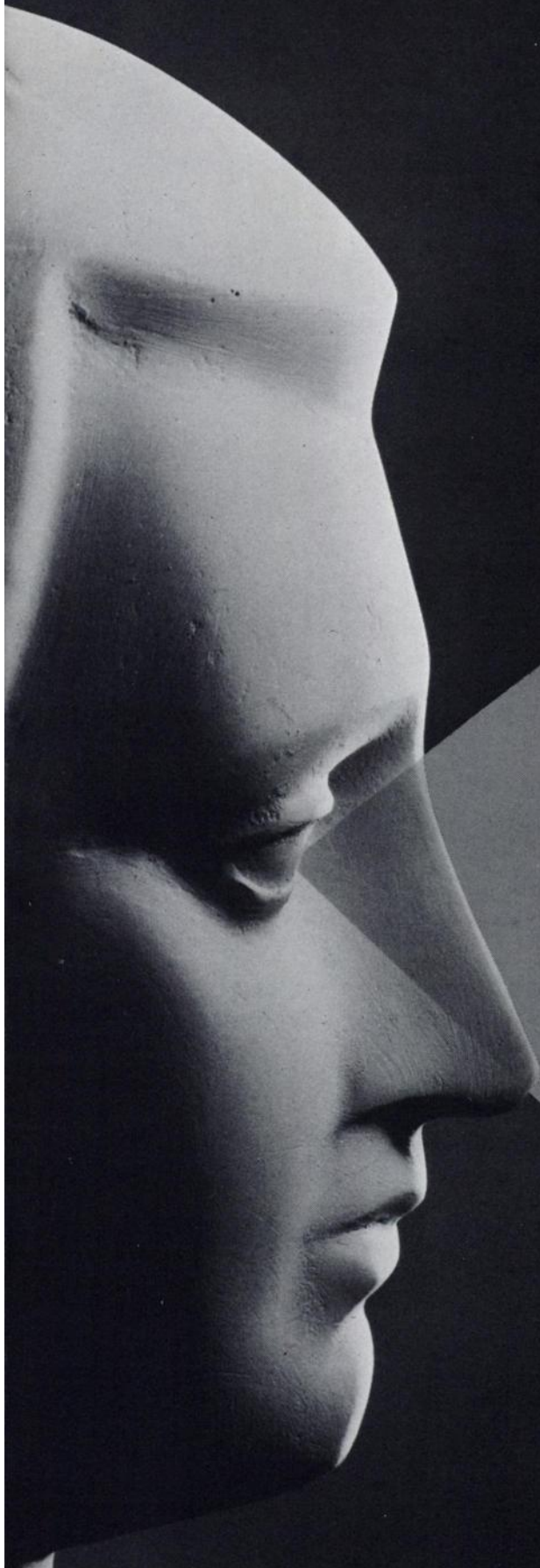
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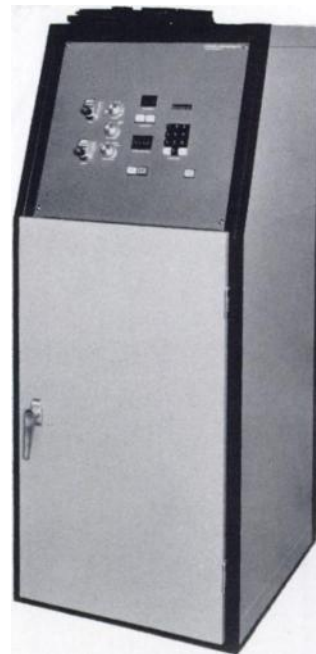


# State of the art in gamma camera hard copy recording.



## 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 identification 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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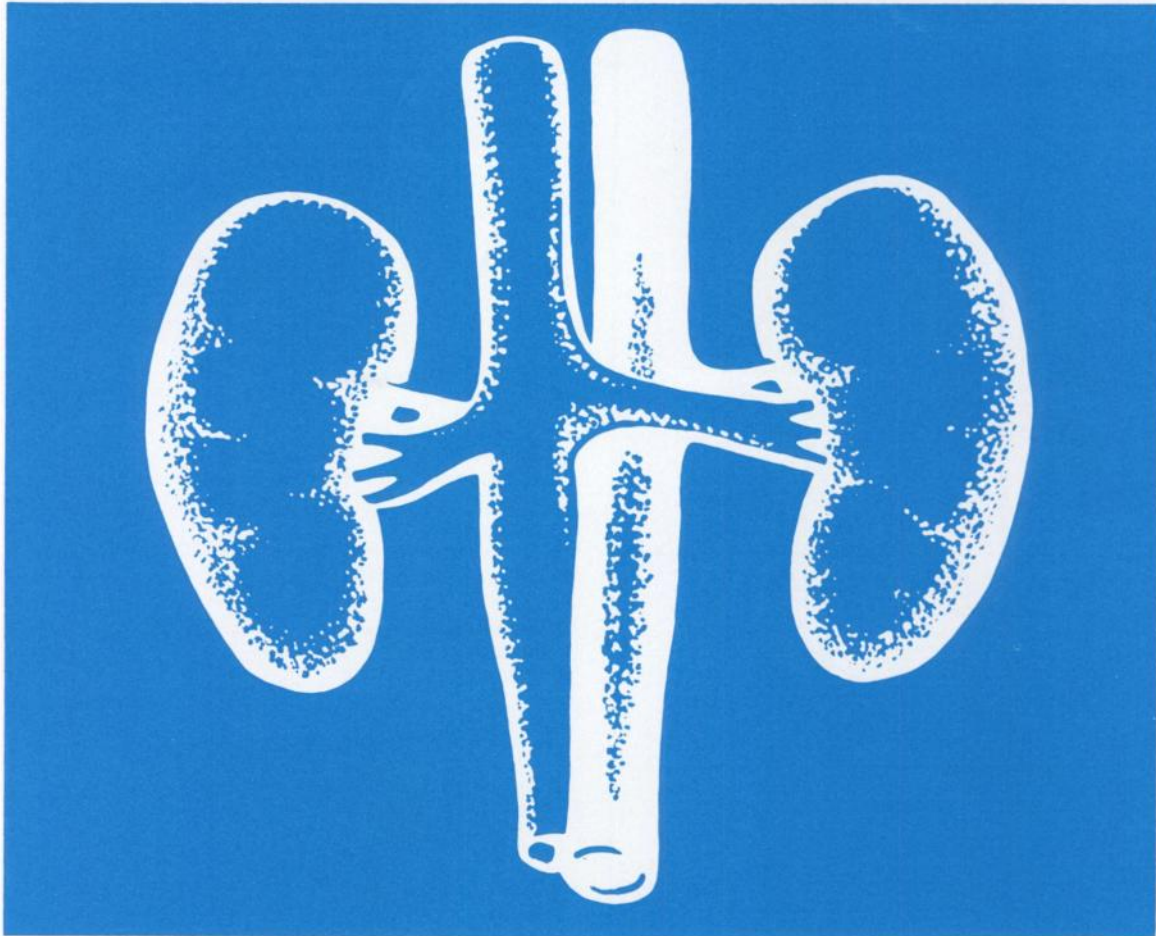
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A complete explanation and description of the procedure is yours for the asking. Or call direct for RIA Technical Service: 617-667-2743.

References: 1. Osmond, D.H., Ross, L.J. and Scaiff, K.D., *Can. J. Physiol. Pharmacol.* 51, 705 (1973).  
2. Sealey, J.E. and Laragh, J.H., *Circ. Res.* (Supplement 1 to Vol. 36 and 37), 10-16, June 1975.



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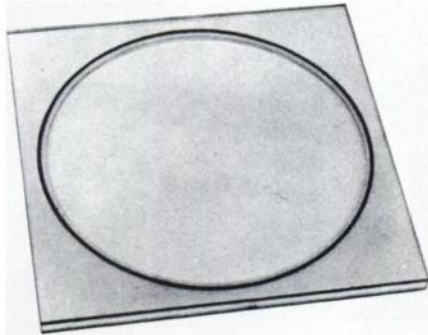
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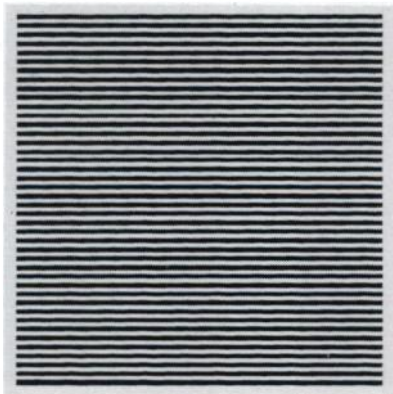
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## STRAIGHT BAR PHANTOM

Model CP-708, 716



- Available with 1/8" or 3/16" wide straight bars.
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CP-708 1/8" Straight Bar Phantom ..... \$265.00  
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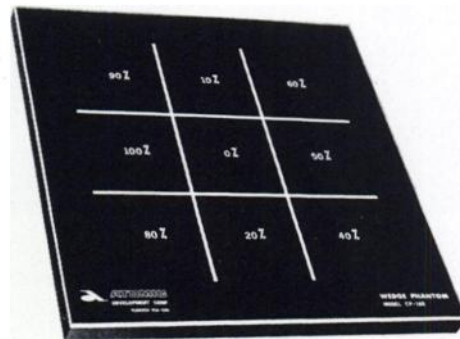
## LARGE AREA HIGH RESOLUTION BAR PHANTOM

Model CP-812



- Check field size and linearity.
- Bar pattern covers 16" x 16" or 14 1/4" x 14 1/4" area.
- Evaluate intrinsic resolution of scintillation cameras.
- Check collimator spatial resolution of cameras and scanners.
- Bar widths available in 3/8", 1/4", 3/16" and 3/32" or 1/2", 3/8", 1/4" and 3/16".

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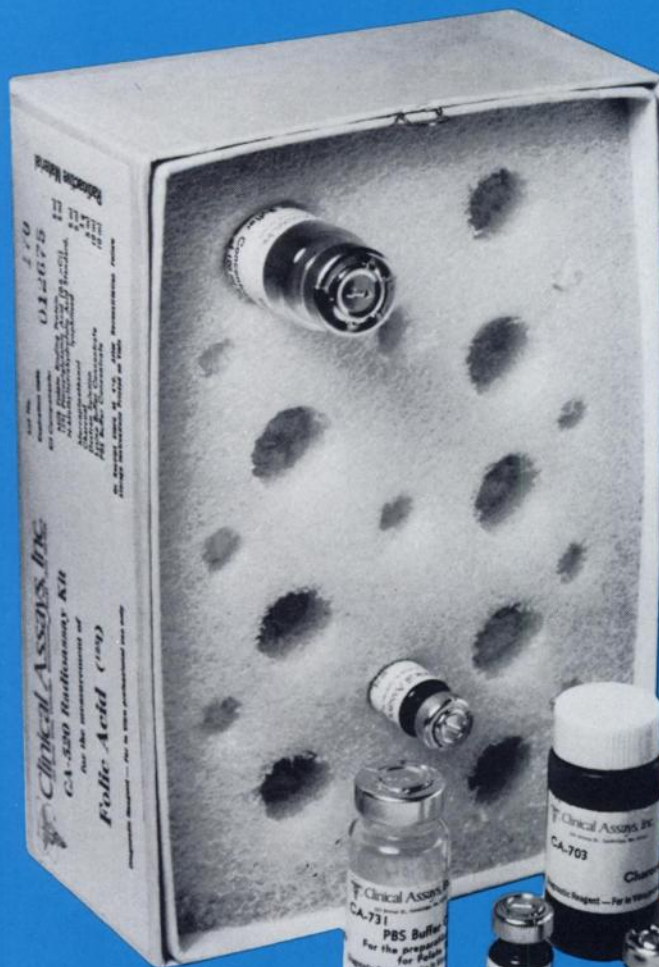
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#### Other kits available:

GammaCoat Digoxin ( <sup>125</sup> I)	Digoxin ( <sup>3</sup> H)
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For Full Details Contact:

## Clinical Assays, Inc.

237 Binney Street • Cambridge, Mass. 02142  
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References: 1) Dunn, R. T.; Foster, L. B.;  
Clin. Chem. 19, No. 10,1101, 1973.





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(5.9MG DISODIUM ETIDRONATE, 0.16MG STANNOUS CHLORIDE)

SKELETAL IMAGING AGENT

In Europe, contact: Philips-Duphar B.V.,  
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See following page for a brief summary of package insert.



PROCTER & GAMBLE

# OSTEOSCAN<sup>®</sup>

(5.9MG DISODIUM ETIDRONATE, 0.16MG STANNOUS CHLORIDE)  
SKELETAL IMAGING AGENT



Brief summary of Package Insert. Before using, please consult the full Package Insert included in each kit.

## DESCRIPTION

Each vial of OSTEOSCAN contains 5.9 mg disodium etidronate and 0.16 mg stannous chloride as active ingredients. Upon addition of ADDITIVE-FREE <sup>99m</sup>Tc-pertechnetate, these ingredients combine with <sup>99m</sup>Tc to form a stable soluble complex.

## ACTIONS (CLINICAL PHARMACOLOGY)

When injected intravenously, <sup>99m</sup>Tc-labeled OSTEOSCAN has a specific affinity for areas of altered osteogenesis. Areas of bone which are undergoing neoplastic invasion often have an unusually high turnover rate which may be imaged with <sup>99m</sup>Tc-labeled OSTEOSCAN.

Three hours after intravenous injection of 1 ml <sup>99m</sup>Tc-labeled OSTEOSCAN, 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 <sup>99m</sup>Tc-labeled OSTEOSCAN excreted in the feces is below the level detectable by routine laboratory techniques.

## INDICATIONS

OSTEOSCAN is a skeletal imaging agent used to demonstrate areas 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 <sup>99m</sup>Tc-generator should be tested routinely for molybdenum breakthrough and aluminum. If either is detected, the eluate should not be used.

## PRECAUTIONS

Both prior to and following <sup>99m</sup>Tc-labeled OSTEOSCAN administration, patients should be encouraged to drink fluids. Patients should void as often as possible after the <sup>99m</sup>Tc-labeled OSTEOSCAN injection to minimize background interference from accumulation in the bladder and unnecessary exposure to radiation.

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.

## ADVERSE REACTIONS

None.

## DOSAGE AND ADMINISTRATION

The recommended adult dose of <sup>99m</sup>Tc-labeled OSTEOSCAN is 1 ml with a total activity range of 10-15 mCi. <sup>99m</sup>Tc-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 calibration system immediately prior to administration.

## Aggregated Albumin (Human) Kit

**DESCRIPTION** - The kit contains 6 sterile vials containing 9-11 mg. of pyrogen-free aggregated albumin (human), 0.67 - 0.83 mg. stannous chloride, and 18 mg. sodium chloride. When sterile, pyrogen-free sodium pertechnetate Tc99m is added to the vial, technetium-labelled macroaggregated human serum albumin (Technetium MAA Tc 99m 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 pH adjustment. Sodium hydroxide of hydrochloric acid may be present in variable amounts. At least 95% of the macroaggregated particles are between 10 and 100 microns in size, the great bulk, (as seen on a microscope slide) being an average of 10 to 70 microns. None are larger than 150 microns. 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 aggregates 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 or 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 fashion 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 capillaries 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 Kupffer cells. There, the particles are phagocytized and rapidly metabolized.

**INDICATIONS** - Scintillation scanning of the lungs with Technetium Macroaggregates is indicated as an adjunct to other diagnostic procedures whenever information about pulmonary vasculature is desired. The most useful clinical applications of lung scanning have been outlined by one investigator: 1) The diagnosis of pulmonary embolism; 2) differentiation of focal conditions such as bullae 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 before radiological 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 anticoagulant 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, atelectasis pleural effusion, pulmonary tuberculosis, parenchymal disease, emphysema and chronic asthmatic bronchitis.

**CONTRAINDICATIONS** - The presence of right to left shunts which would allow Technetium MAA Tc 99m injected in a systemic vein 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.

**WARNINGS** - 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 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.

**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 fatal reaction following infusion of Tc 99m macroaggregates (technetium labelled macroaggregates). This was in a seven-year-old child who had severe pulmonary vascular 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 Iodine-131 macroaggregates where Lugol's solution was administered to block the thyroid gland. The subject had a history of angio-edema. The reaction may have been caused by either material. Dworin 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 2½-year history of adenocarcinoma of the breast she had severe and rapidly progressive edema. Prior to scanning, the nasal administration of oxygen was interrupted. "Within 1 or 2 minutes after injection of 300 uCi of I-131 labelled macroaggregates albumin (11 mg. of albumin or 0.219 mg. per kilogram of body weight) she complained of lightheadedness and became cyanotic, diaphoretic, and agitated with distended neck veins. The initial pulse rate of 50 rose to 140 with a fall in blood pressure to 100/30. Oxygen therapy relieved the profound dyspnea 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, dyspnea 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 microns 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 died in right heart failure. Post-mortem examination revealed "severe atheroma and thickening of all the pulmonary arteries but no macroscopic evidence of emboli. The right heart was hypertrophied and dilated".

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

## REFERENCES

1. Suprenant E. L., Webber M.M., Bennett L. R., *International Journal of Applied Radiation and Isotopes*, 20, 77-79 (1969).
2. De Paoli T., Hager A., Nicolini J., *International Journal of Applied Radiation and Isotopes*, 17, 551-556 (1966).
3. Vincent W. R., Goldberg, S. J. and Disikes, D., *Radiology* 91, 1181-1184 (1968).
4. Wagner, H. N., Jr., et al., *N. Engl. J. Med.* 271, 377-384 (1964).
5. Dworin, J. J., Smith, J. R. and Bull, F. E., *N. Engl. J. Med.* 275, 376 (1966).
6. Dworin, J. J., Smith, J. R. and Bull, F. E., *Am. J. Roentgenol Ther. Nucl. Med.* 98, 427-433 (1966).
7. Williams, J. O., *Brit. J. Radiol.* 47, 61-63 (1974).

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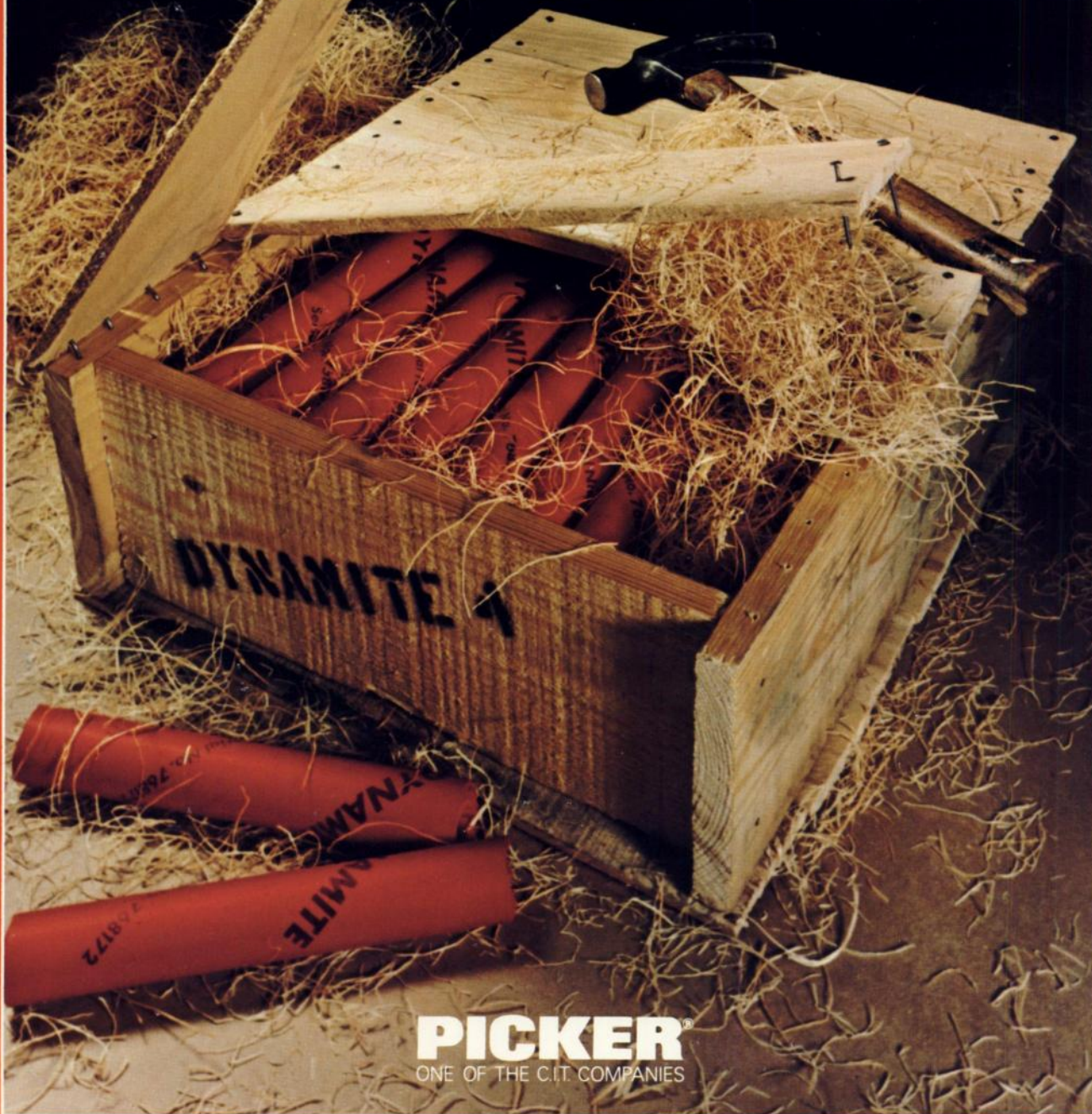
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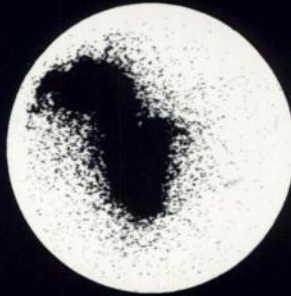
Cardiac Flow Study  
Picker Large Field (15" diameter) Detector  
10mCi  $^{99m}\text{Tc}$  Sodium Pertechnetate



1



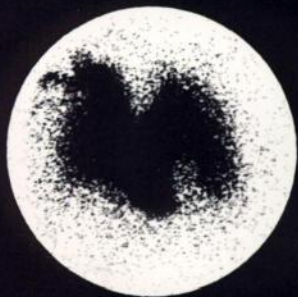
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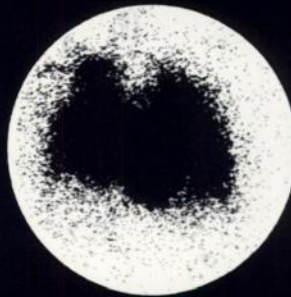
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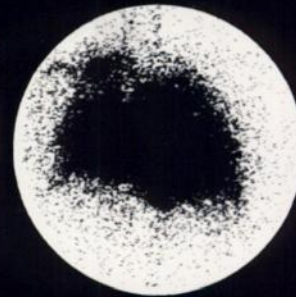
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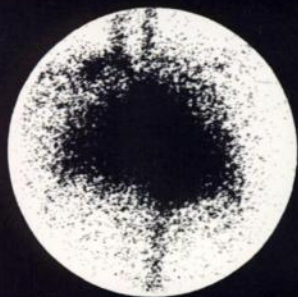
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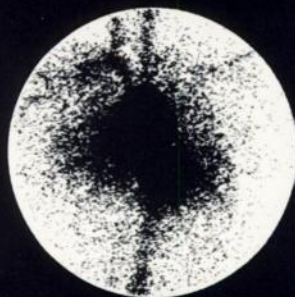
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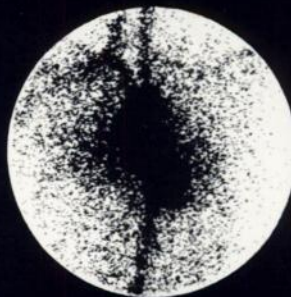
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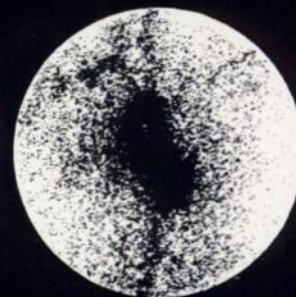
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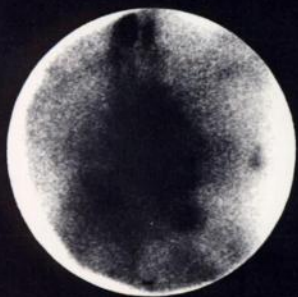
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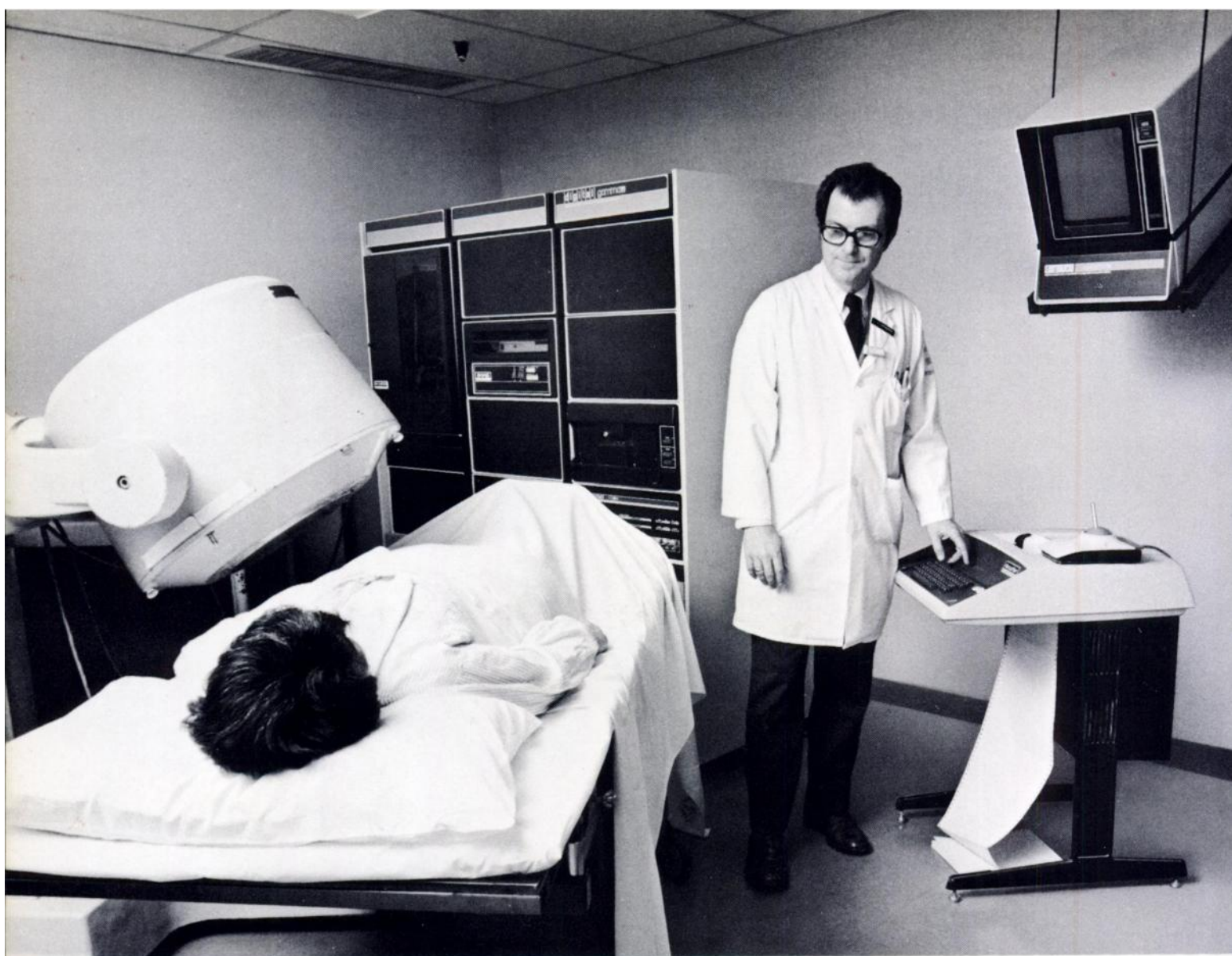
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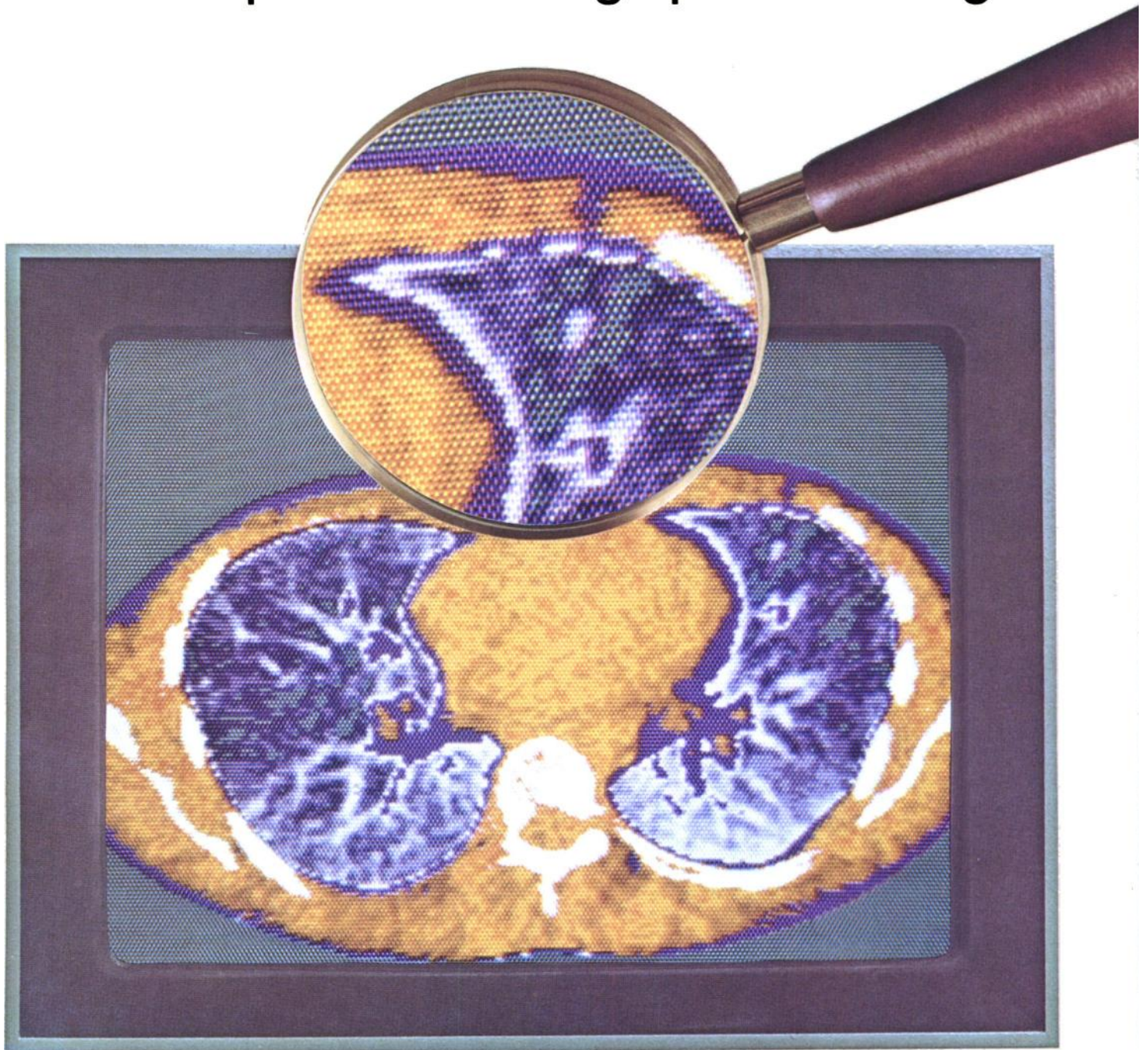
Photo of Gamma-11 installation at  
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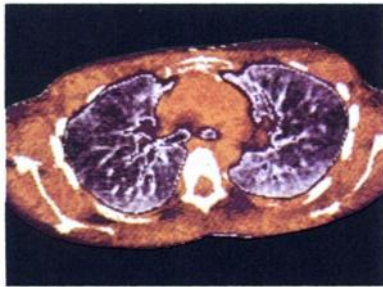


WHOLE BODY COMPUTERIZED TOMOGRAPHIC SCANNER

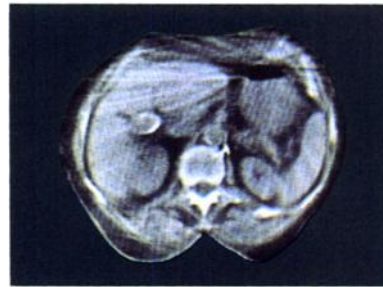
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Up to 57,600 absorption values are now actually measured for translation into the finished ACTA-scan with the recently developed 320 matrix.

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*Thoracic 320 Scan.  
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*Abdominal 320 Scan.  
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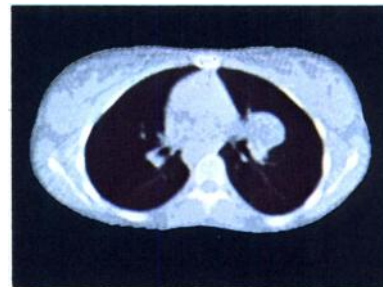
**Multiple windows let you see more**

With the Pfizer ACTA-Scanner, multiple windows can be imposed upon the image, allowing tissues with great density differences to be viewed at the same time in a single ACTA-scan.

This capability greatly facilitates interpretation of scans in the thoracic and abdominal areas.



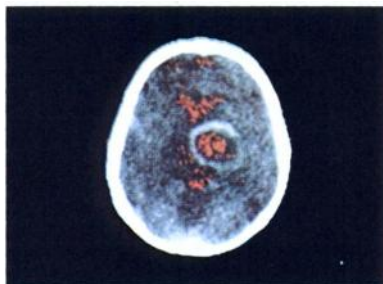
*Thoracic scan with multiple windows. Mass in right lung.*



*Same area as scanned at left, without imposition of the multiple window capability.*

**And you can enlarge selected areas**

A special cursor – or movable dot – allows the operator of the ACTA-Scanner to enlarge selected areas of interest by a factor of 2 in diameter (4 in area).



*Pituitary Adenoma*



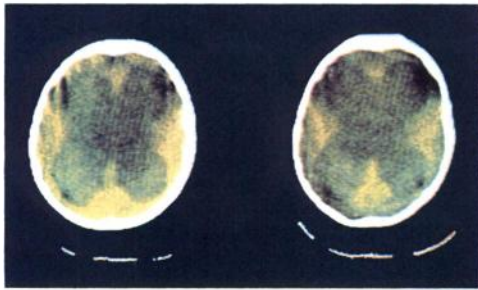
*Pituitary Adenoma.  
Instantaneous enlargement of pathologic area.*



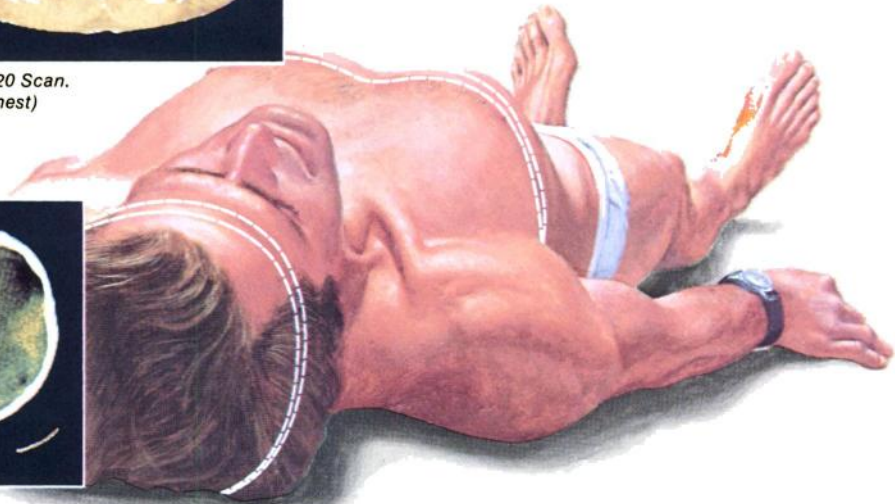
Pathology in virtually any part of the body can be visualized and evaluated.



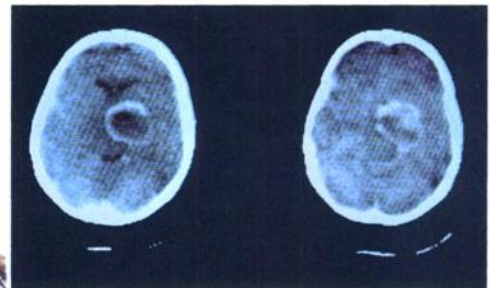
Thoracic 320 Scan.  
(Normal Chest)



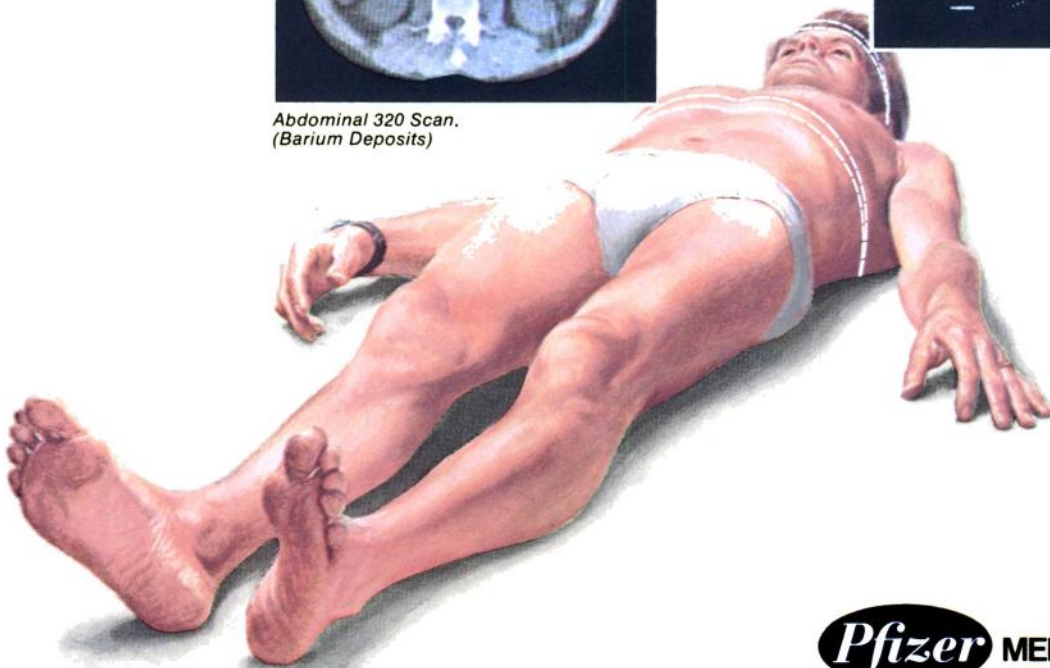
Marked Hydrocephalus



Abdominal 320 Scan.  
(Barium Deposits)



Pituitary Adenoma



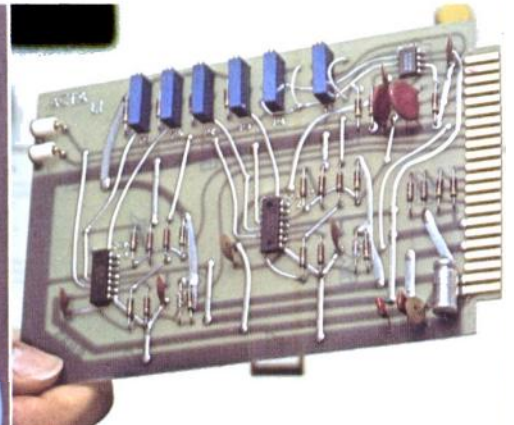
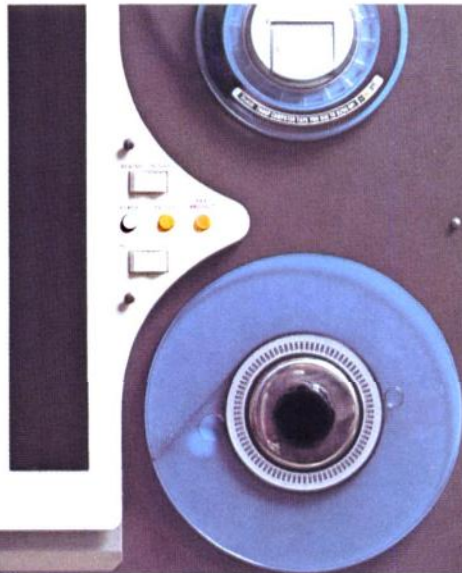
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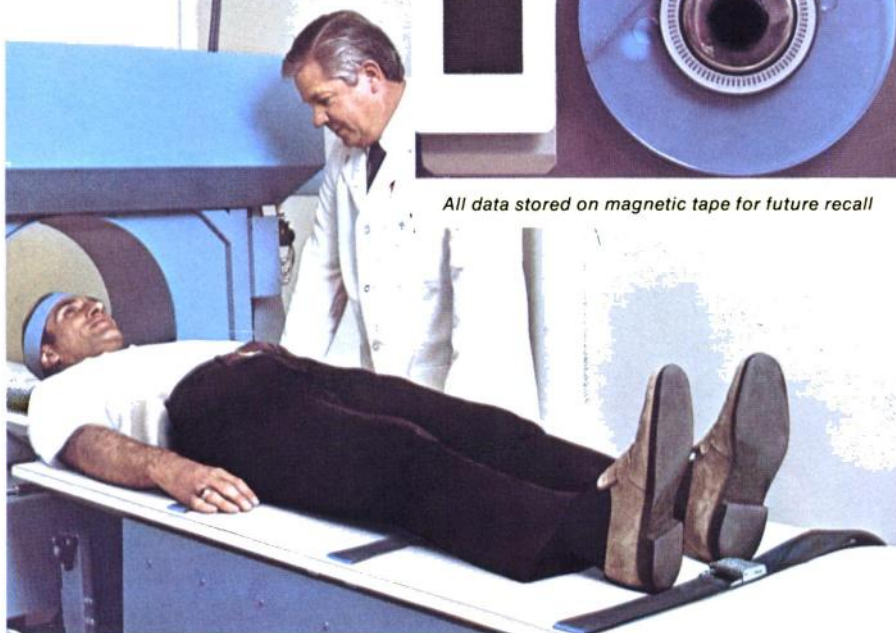


*Field engineer readily available*

The Pfizer Medical Systems Field Engineer is an integral component of the ACTA-Scanner Service Program.

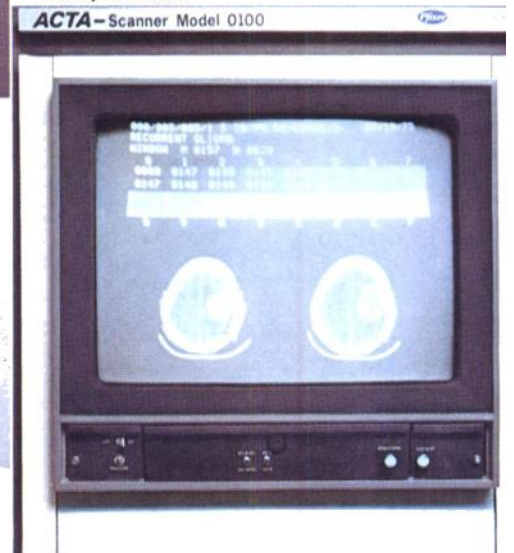


*Replacement parts, if needed, are readily available*



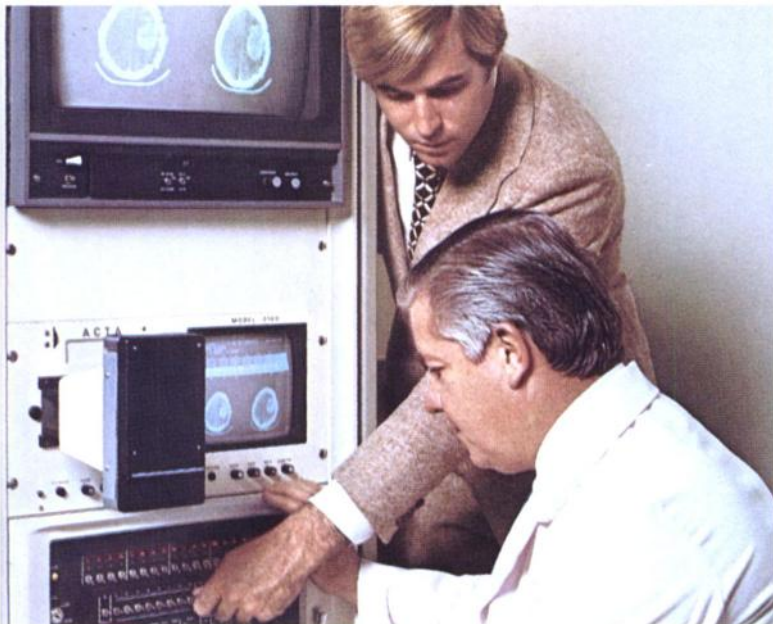
*All data stored on magnetic tape for future recall*

*Minimal patient preparation before scan*

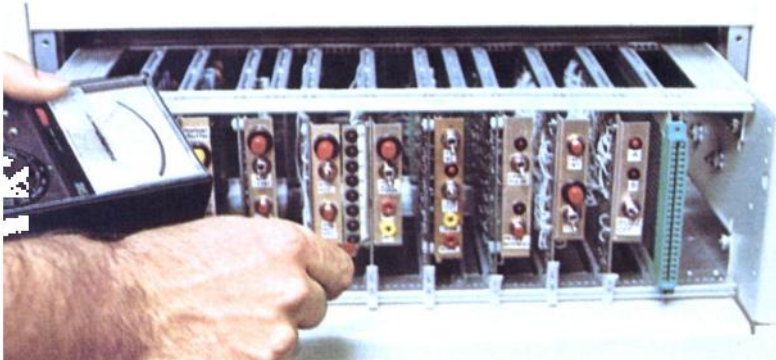


*Instant display following completion of scan*





*Field engineer provides continued updating on capabilities*



*Frequent visits by field engineer keep ACTA-Scanner at peak performance*

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211	-3	112	13
164	0	100	14
128	1	89.1	15
100	2	79.4	16
78.0	3	70.7	17
60.9	4	63.0	18
47.5	5	56.2	19
37.0	6	50.0	20
28.9	7	44.5	21
22.5	8	39.7	22
17.6	9	35.4	23
13.7	10	31.5	24
10.7			24

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Before prescribing please consult the complete product information, a summary of which follows:

**CONTRAINDICATIONS**—The safety of *TechneScan MAA Tc 99m* in patients with a known right-to-left cardiac shunt has not been established and its use in such patients is contraindicated.

**WARNINGS**—In acute cor pulmonale the administration of aggregated albumin is theoretically hazardous due to the temporary small additional mechanical impediment to pulmonary blood flow. Although not reported with *TechneScan MAA Tc 99m* there are three reports in the literature of deaths occurring after the administration of radioiodinated aggregated albumin as a result of pre-existing primary pulmonary hypertension.<sup>1,2,3</sup>

The contents of the *TechneScan MAA* reaction vial are intended only for use in the preparation of *TechneScan MAA Tc 99m* and are not to be directly administered to the patient.

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

This radiopharmaceutical preparation should not be administered to patients with severe kidney disease unless the benefits to be gained outweigh the potential hazards. Similar care should be observed with patients who are pregnant or who are lactating.

Ideally, examinations using radiopharmaceuticals, especially those elective in nature, of a woman of childbearing capacity 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.

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

**ADVERSE REACTIONS**—Although no anaphylactoid reactions have been reported in patients following the administration of *TechneScan MAA Tc 99m*, the possibility should be considered that hypersensitivity reactions may occur rarely in patients who, after the initial administration, receive additional doses a number of weeks after the initial dose.

<sup>1</sup>Dworkin, H. J., Smith, J. R. and Bull, F. E.: Reaction after Administration of Macroaggregated Albumin for a Lung Scan. *New England J. Med.*, 275:376, August 18, 1966.

<sup>2</sup>Roberts, H. J.: Fatal hemoptysis in pulmonary embolism probably precipitated by pulmonary scanning—Report of a case and suggested precautions. *Angiology*, 21:270, 1970.

<sup>3</sup>William, J. O.: Death following injection of lung scanning agent in a case of pulmonary hypertension. *Br. J. Radiol.* 47:61, 1974.



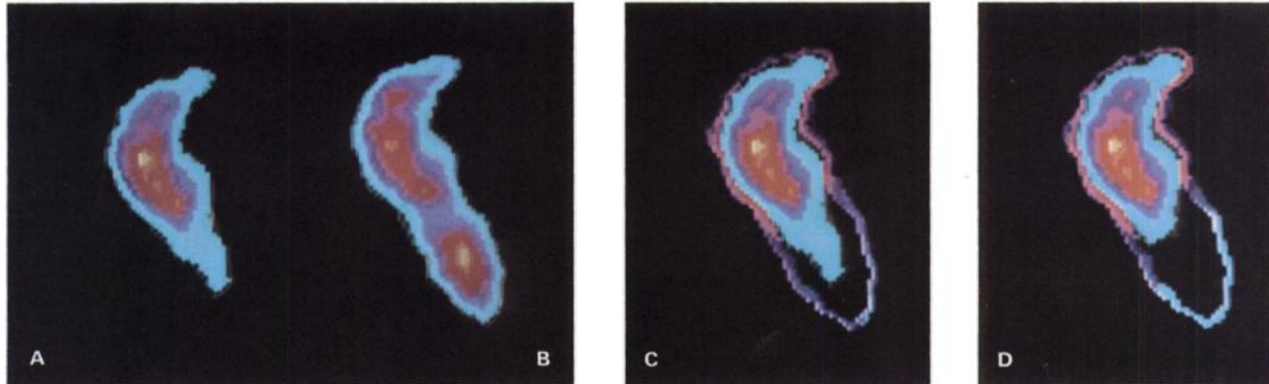
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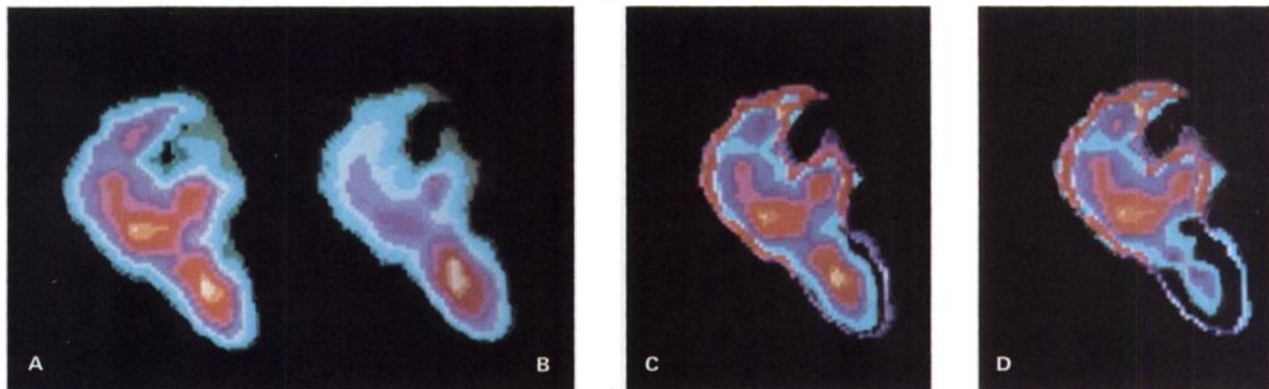


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## 1. Comprehensive, first-pass dynamics of cardiac wall motion



**NORMAL PATIENT. Anterior View. Ejection Fraction 63%.** (A) Image at End Systole shows volume displacement flow is maximum in the aorta and volume is minimum in the ventricle. (B) Image shows that volume displacement flow is minimum in the aorta and volume is maximum in the ventricle at End Diastole. (C) ES, with perimeter at ED superimposed, shows normal volume displacements and symmetric wall motion band due to motion of the septal and lateral walls. (D) Subtraction of stroke volume from ES, with ED perimeter superimposed, shows that all volume displacements in the stroke volume exceed volume components in residual distribution at ES.



**ABNORMAL PATIENT. Anterior View. Ejection Fraction 34%.** (A) ES, showing spatial distribution of volume components. Abnormally high residual volume at ES in the ventricle compared to volume flow components in the aorta. (B) ED, showing distribution of left heart volume components. Comparison with ES suggests relative lack of ventricular volume displacement during systole. (C) Lack of wall motion is indicated by very narrow wall motion band between ED perimeter and the ES distribution along the septal wall to the apex. Wall motion of the lateral wall is closer to normal. (D) Volume component in ES distribution exceeds stroke volume displacement because of reduced anterior or posterior wall motion proximal to the septal wall.

Shown here are stop-action data extracted from the representative cycle of first-pass images showing hemodynamics of the left heart, including volume distribution of end systole, end diastole, end systole with the end diastolic perimeter superimposed, stroke volume subtracted from end systole with end diastolic perimeter superimposed. These images provide the basis for the clinical diagnosis of ventricular wall motion, in addition to providing data for a closer examination of specific areas for evidence of hypokinesia, akinesia, or dyskinesia.

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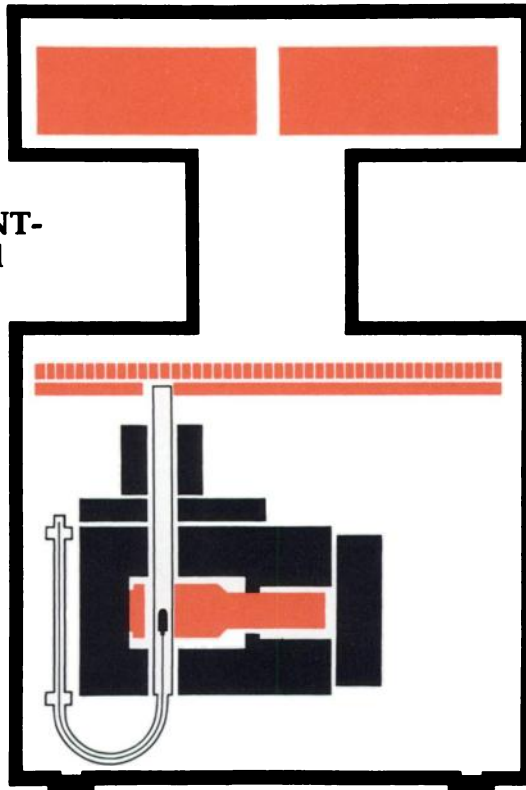
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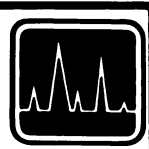
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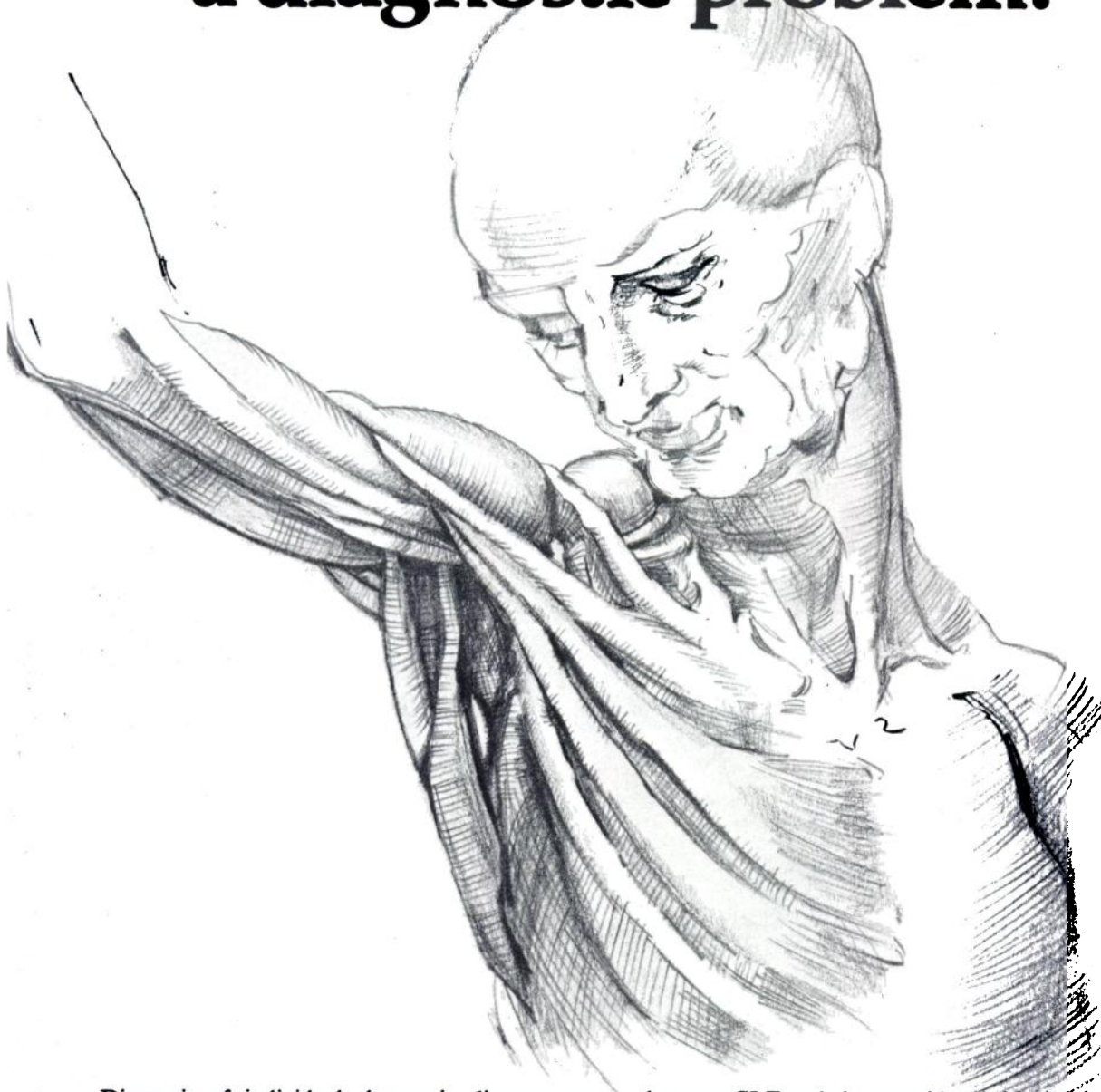
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Diagnosis of individual rheumatic diseases can present problems. Our simple test, the anti-DNA Kit, can give vital information to aid that diagnosis.

The kit provides the first standardized assay to consistently and reliably measure anti-DNA antibodies. High circulating levels of these antibodies are closely linked with systemic lupus erythematosus (SLE). In doubtful cases, the kit offers excellent discrimination

between SLE and rheumatoid arthritis and is particularly valuable as a follow-up to ANF tests. Results show that the kit is also useful as a means of monitoring disease activity, providing the physician with guidance on drug therapy.

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.



**The Radiochemical Centre  
Amersham**

## Anti-DNA kit

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.

0395

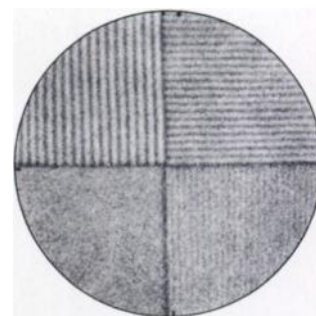
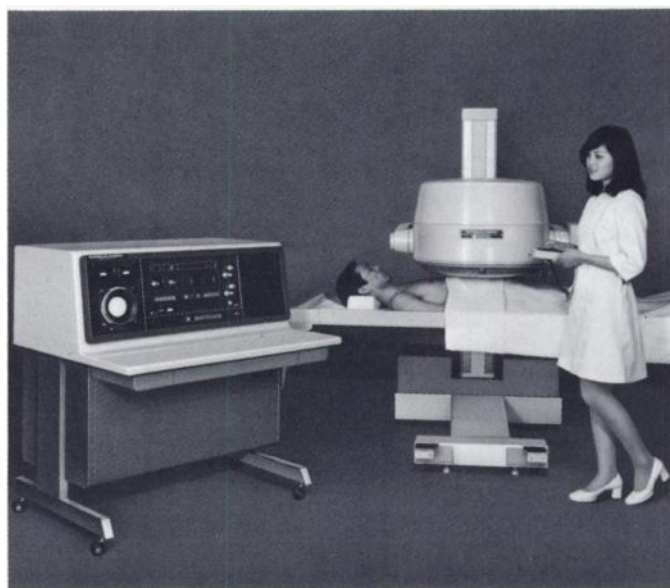


# Why All the Interest in Toshiba's Newest Jumbo Gammacamera?

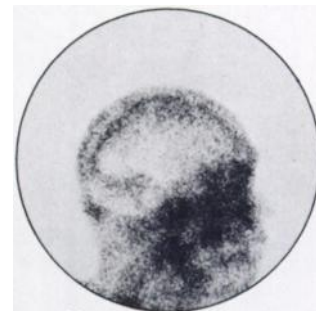
Since its introduction at the First World Congress of Nuclear Medicine, our newest high performance delay line Gammacamera, GCA-401, has been generating world-wide interest. In fact, several sets have been, or soon will be installed in Europe, Australia, and Japan. The features that make this unit so attractive include:

- High intrinsic resolving capability (3.2mm lead pattern using  $^{99m}\text{Tc}$ .)
- 35cm usable field of view, large enough to image both lungs or a large organ.
- Programable setting of measuring conditions
- Compact, easy-to-operate control console
- Adaptable for whole-body imaging
- Compatible with any data processing system
- Reliability assured through utilization of Toshiba's world renowned IC electronics

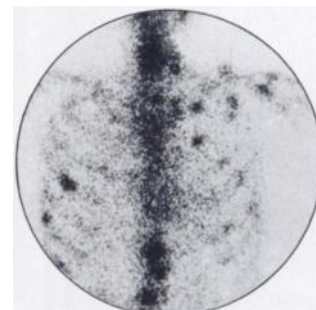
If we've caught your interest too, please write. We'll be pleased to send you all the information you need on the GCA-401.



**Intrinsic Resolution**  
 $^{57}\text{Co}$  999 K-counts,  
Window; 20%  
Pb-Bar pattern; 2.4, 3.2, 4.0,  
4.8 mm



$^{99m}\text{Tc}$ -DTPA, 24m Ci,  
300 K-counts, Window; 20%  
Collimator; High resolution.



$^{99m}\text{Tc}$ -pyrophosphate, 13m Ci,  
200 K-counts, Window; 20%  
Collimator; High resolution.

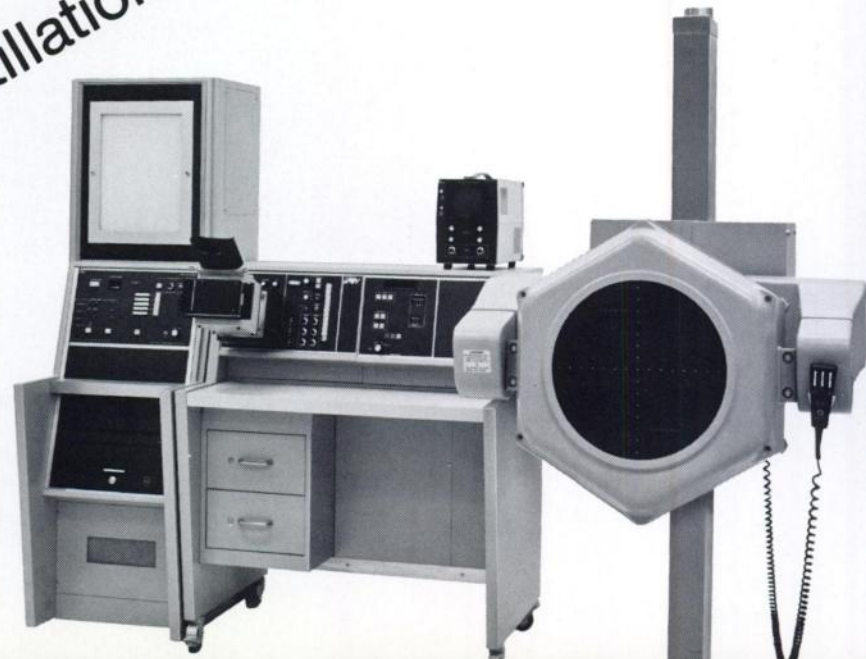
 **TOSHIBA**  
TOKYO SHIBAURA ELECTRIC CO., LTD.

Producer Goods Export Division  
1-6, Uchisaiwaicho 1-chome, Chiyoda-ku, Tokyo, 100 Japan  
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The large  
field of view  
camera  
that provides  
superior diagnostic  
clarity

Pho/Gamma<sup>®</sup>

Large Field of View Scintillation Camera



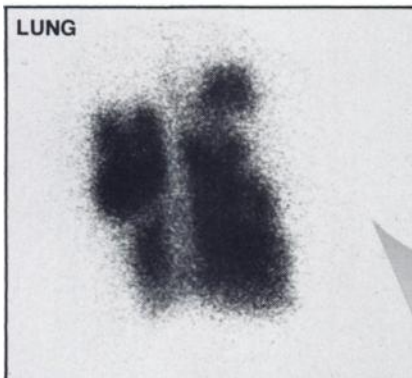


- 15¼ inch (390 mm) field of view
- Hexagonal detector head design for ease of positioning
- Superior resolution at depth
- Totally new electronic design
- Large selection of parallel hole and converging collimators
- Backed by the world's largest and most experienced nuclear service facilities

#### LUNG VENTILATION STUDY

Important LFOV applications include functional and anatomical studies of the lungs using Xenon 133. The images are of such quality that the physician is able to define more anatomical detail than previously possible. Images obtained during breath holding intervals eliminate motion artifact and still can contain over 300K counts because of the unique design of the LFOV and its parallel hole collimators.

Pulmonary studies demonstrate quality in the ventilatory image which is near that obtained in perfusion images. In



many cases, segmental and even sub-segmental defects have been observed.

#### Posterior Ventilation

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

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.

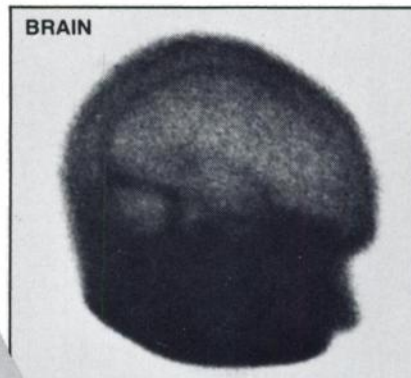
#### BRAIN STUDY

Large diameter crystal cameras require well designed converging collimation in order to image the brain adequately... studies of the brain obtained with the LFOV provide a remarkable level of image quality as observed in over 2000 clinical comparative studies. The vascular structures are clearly seen, both spatially and temporally. Even in the lateral view, the deep veins leading to the jugular systems are readily seen.

#### Lateral Brain Study

400,000 counts • 19mCi • 99mTc • 185 seconds

The improved resolution with depth allows clear separation of the sagittal sinus from the lower activity in the skull, and the scalp activity which is somewhat greater than that within the skull. In addition, the sinus is defined in its entirety as it proceeds downward from the torcular through the sigmoid sinus into the jugular bulb. Other midline structures including the floor of the anterior and middle cranial fossa, as well as the region of the pituitary, are clearly demarcated. There is a zone of activity extending superiorly and posteriorly from the region of the pituitary in line with the sphenoid ridge which is more likely the inferior sagittal sinus.



*For additional sample studies performed on the LFOV and associated collimator information, ask your Searle representative for our latest Large Field of View Scintillation Camera brochure; or write:*

**SEARLE**

#### Searle Radiographics Inc.

Subsidiary of G. D. Searle & Co.  
2000 Nuclear Drive  
Des Plaines, Illinois 60018 U.S.A.  
*Offices in Principal Cities  
throughout the World*

SR511





**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.)

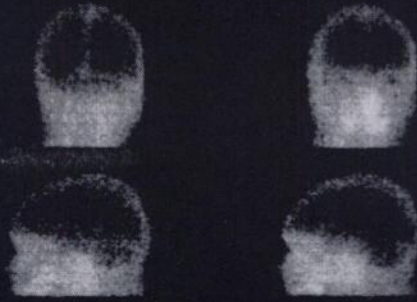
The Imager's display and recording options, enhancement of photo-images, and the capability to playback stored data greatly increase its clinical usefulness. Reliability, rapidity of operation, and high patient turnover mean increased utilization and economy, along with improved diagnostic efficiency.



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.





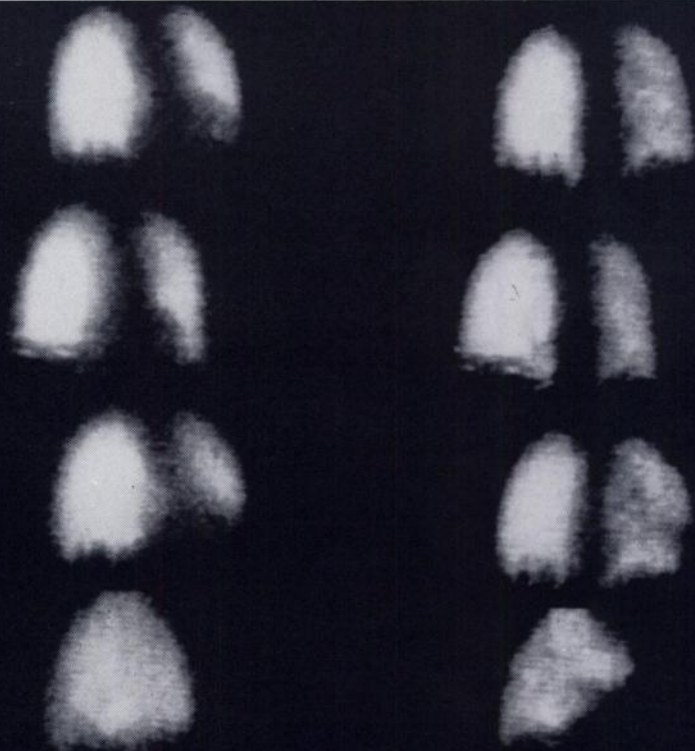
**BRAIN IMAGE.**

Imaging agent: 15 mCi Tc-99m Pertechnetate.

Time-to-scan (4 views): 13.7 minutes.

Image courtesy of Cedars of Lebanon Hospital, Los Angeles.

---



**LUNG IMAGE SERIES.**

Imaging agent: 1.5 mCi Tc-99m MAA.

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.

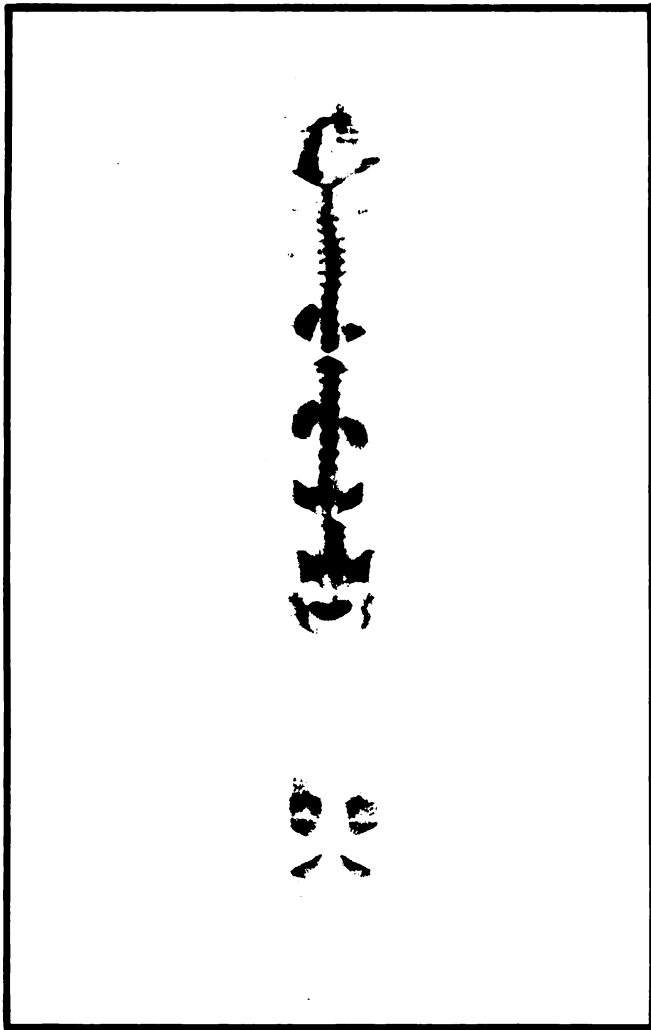
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**cleon**  
CORPORATION

150 Gould Street, Needham, Massachusetts 02192/Telephone 617-444-2494

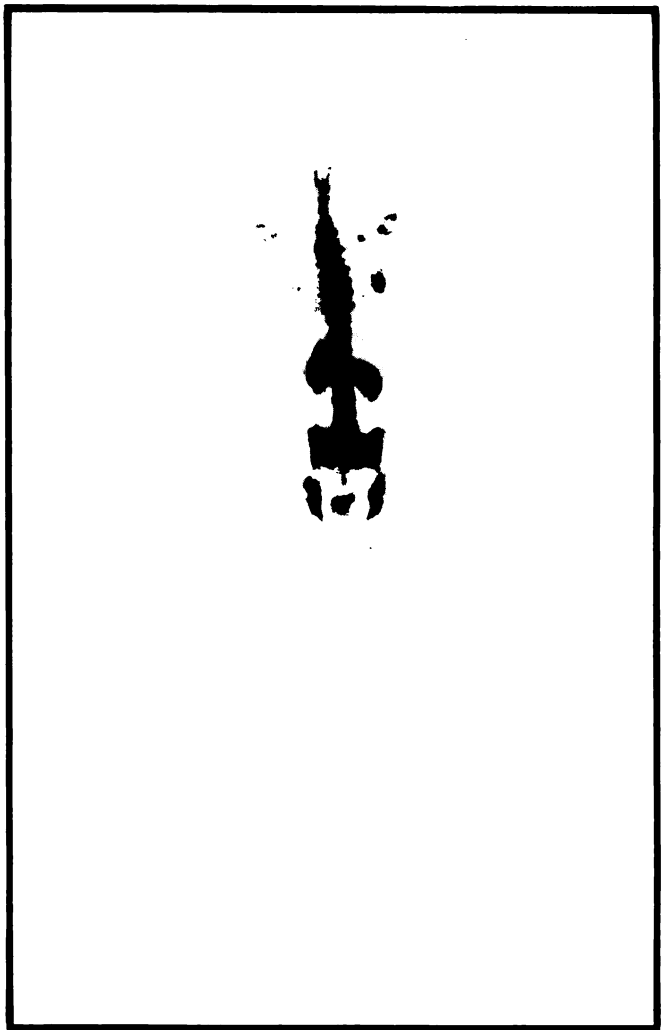


# An Unbiased Comparison



***Our Wide Field***

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



***Our Wide Field***

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 <sup>99m</sup>Tc Pyrophosphate  
Time Begun: 4 hours post dose

Composite View  
700,000 counts per view except legs were 100,000 counts per view  
Total Scan Time: 30 minutes (included positioning)

AreaScan

Total Scan Time: 12.2 minutes



**ohio-nuclear, inc.**  
A subsidiary of Technicare Corporation

U.S. — 6000 Cochran Road • Solon Ohio 44139

Telephone (216) 248-8500

TWx. 810-627-2688

Telephone Staines 51444

U.K. — Ohio Nuclear U.K. • Radix House • Central

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# From the makers of Thyopac...\*

... a T3RIA kit, with the performance, simplicity and reproducibility of the Thyopac range.

A good T3RIA is the most sensitive assay for hyperthyroidism and the only specific *in vitro* test for T3 thyrotoxicosis. It is also a valuable follow-up for treated patients, when T4 values may mislead.

Our T3RIA Kit has been acclaimed by some very critical users, and we're sure you'll appreciate it too.

Contact us for complete information on T3RIA, Thyopac-3, -4 and -5, and our latest monograph on thyroid function testing. With something as good as this, you don't want to miss it.

**Simple** direct serum assay on 50 $\mu$ l sample, pre-dispensed serum standards, no centrifugation step.

**Flexible** incubate for 1 hour at 37°C, or overnight at room temperature.

**Reproducible** 5-7% coefficient of variation.

**Specific** minimal T4 cross-reactivity.

**Versatile** use T3RIA with Thyopac-3 (binding capacity test), with Thyopac-4 (T4 CPB), or with Thyopac-5 (T4 CPB+NTR).

**Reliable** every batch of kits is tested to the highest standards of quality control before despatch.

# T3RIA

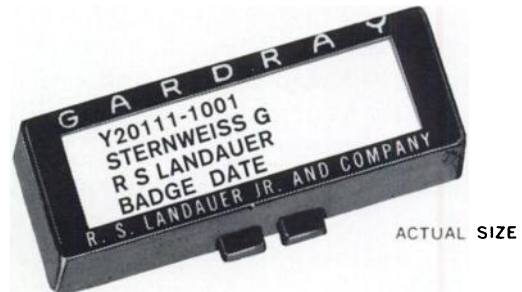


The Radiochemical Centre  
Amersham

The Radiochemical Centre Limited, Amersham, England. Telephone: 024-04-4444  
In the Americas: Amersham/Searle Corp., Illinois 60005. Telephone: 312-593-6300  
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\*Trade Mark. 0243

“Make  
the  
best  
available  
better!”



“Work on the ultimate, but in the meantime, make the best available better.”


Our people have always accepted the challenge and it's what makes us the leader.

We agree that all things considered the Landauer Gardray 8 film badge system is the best available personnel dosimeter. And, although we are always looking for the ultimate, we have continued to work hard and invest money and time to make it better.

Greatly simplified ordering procedures – permanently encoded unique numbering of film, which is independent of film darkening – new improved techniques for analyzing the film for anomalies that may affect the “meaning” of the exposure and new N.R.C. annual statistical summary reports available now, are just some of the ways our people are working hard to make it better for you.

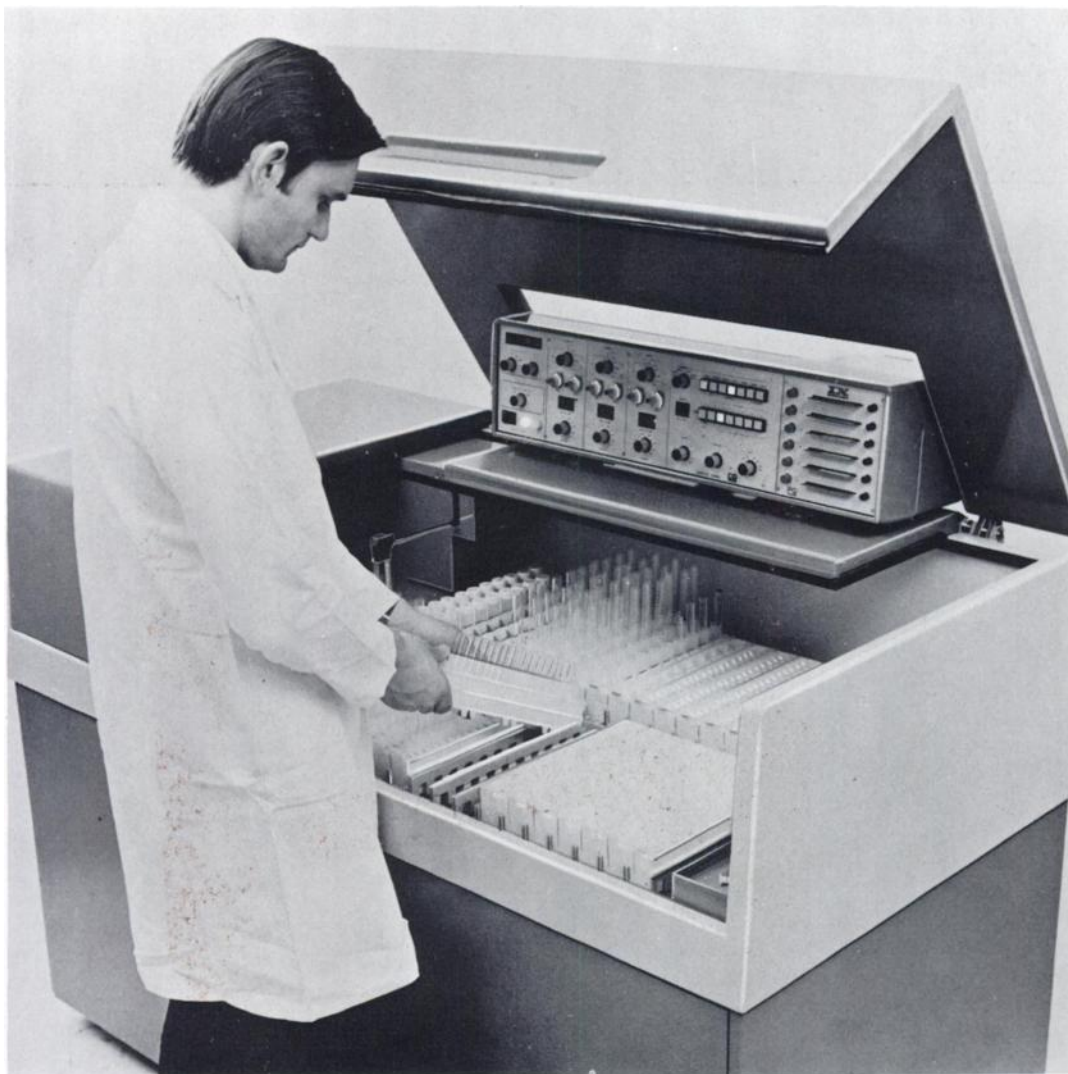
Write or call for more details.

*Landauer*

R.S. LANDAUER JR. & CO. A  COMPANY  
Glenwood Science Park  
Glenwood, Illinois 60425 . (312) 755-7000



# INTERTECHNIQUE CG-4000 AUTOMATIC GAMMA COUNTERS INCREASE LABORATORY OUTPUT.



**400-760 TUBE RACK-TYPE TRANSPORT** for fast, flexible sample handling. Accepts intermixed tube diameters up to 28 mm. Insures fast, reliable tube transfer to detector.

**WELL-TYPE DETECTORS AND CASCADED LINEAR AMPLIFIER** for maximum resolution, efficiency and long term stability with low and high energy isotopes.

**BUILT-IN RIA CALCULATOR** computes average NSB and  $B_0$  or Total activity from replicates, then

calculates %  $\left[ \frac{\text{CPM} - \text{NSB}}{B_0 - \text{NSB}} \right]$

or %  $\left[ \frac{\text{net CPM}}{\text{Total}} \right]$

**ADVANCED RIA COMPUTATION** with the 16-32 K Multi-Mat<sup>®</sup> 4000 computer option. Programmable in LEM, delivered with general purpose and kit-oriented RIA programs.

**MULTI-USER PROGRAMMER** selects windows, presets, background subtract parameters and specific RIA programs by individual Control Cards<sup>®</sup>.



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cable address INTERTEC Telex: IN 692 642 F  
U.S.A 50 Van Buren Avenue WESTWOOD,  
New Jersey 07675 Tel : (201) 664-7070

# Buy the capabilities of a \$70,000 computer...

## ADAC Clinical Data System



### Standard Features

- 40K x 16 Memory
- Dual 7" x 9" Video Displays
- 64 Gray Shades
- 512 x 512 Image Display Matrix
- Light Pen (8 regions-of-interest)
- Dual Disc Drives
- High Speed (110 characters/sec) Printer
- Clinical Software Package
- In Vivo and In Vitro Data Analysis
- Free Software Upgrades
- Installation and Training
- One-Year Warranty

### Optional Features

- Third Disc Drive (allows simultaneous processing while acquiring data)
- 64K x 16 Memory
- 8 x 10 X-ray Film Camera
- Polaroid Camera
- FORTRAN IV Compiler
- Paper Tape Reader (for input of RIA data)
- High Speed Disc Drive (list mode, mass storage)
- Remote Dual Displays
- Color Display
- Image Transmission via Telephone
- Microdot (Searle TM) Interface



# ... for the price of a \$25,000 tape system!

The new ADAC Clinical Data System may be just what you're looking for . . . it's more than a video tape recording system, more than a hard-wired, tape-based data system (and easier to use), and comparable in performance and flexibility to the minicomputer systems that cost \$50,000 to \$75,000.

Quantitative organ function analysis will play a significant role in the future of nuclear medicine. Effective data acquisition and analysis is becoming more of a necessity than ever before. Until now, your choice was between expensive programmable computers and less costly (and less effective) tape-oriented systems. Recent advancements in computer technology give you a third choice that closes the gap!

For less than \$30,000 you can have a programmable computer that incorporates the very latest technology, giving you a system that is low priced, easy to use, versatile, and expandable to meet the most demanding needs of the future.

Call or write today for a sample clinical study and the locations of our current installations.

**ADAC**

10300 Bubb Road  
Cupertino, California 95014  
Telephone (408) 255-6353



# New England Nuclear Radiopharmaceuticals

**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 train-

ing 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.

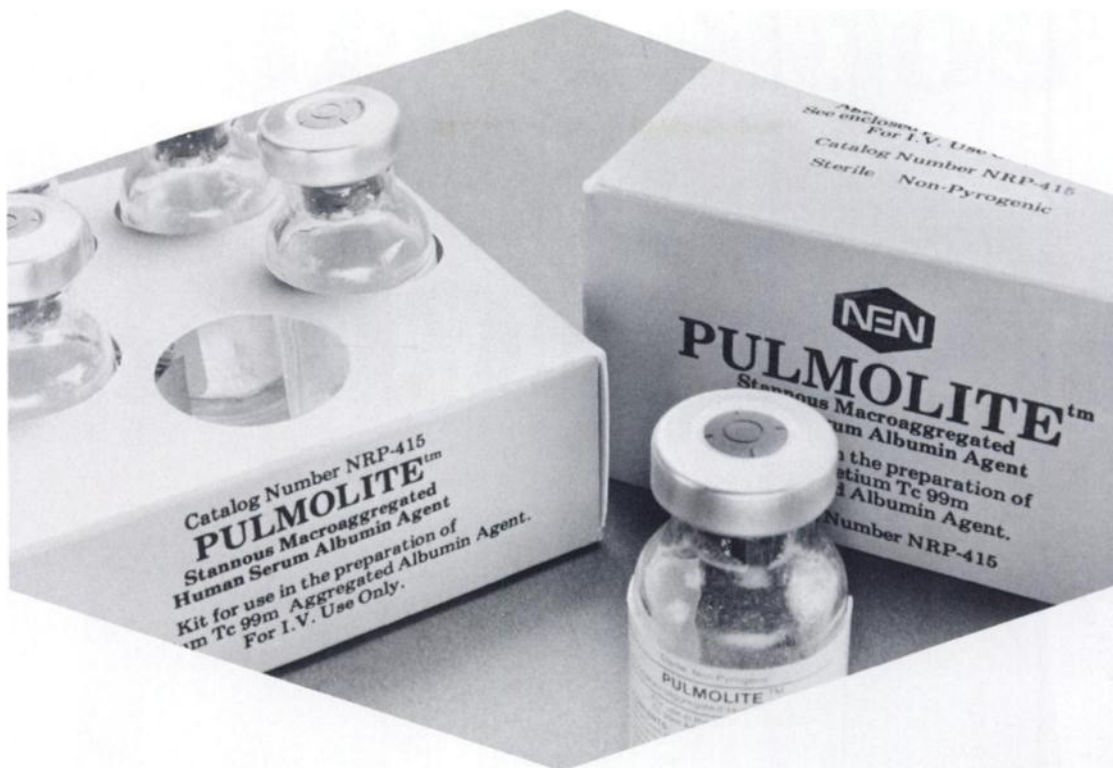
**ADVERSE REACTIONS.** Although no adverse reactions have been reported using NEN Technetium Tc 99m Aggregated Albumin, rare instances of hemodynamic or idiosyncratic reactions to other preparations of Tc 99m labeled macroaggregated albumin have been recorded.

**DOSAGE AND ADMINISTRATION.** The recommended intravenous dose range for the average patient (70 kg) is 2 to 4 millicuries.

For ease and accuracy in dispensing the prepared agent, it is recommended that prior to reconstitution, concentrated Pertechnetate Sodium Tc 99m be further diluted to a minimum volume of 5ml with fresh, preservative-free, sterile Sodium Chloride Injection (U.S.P.).

The patient dose should be measured by a suitable radioactivity calibration system immediately prior to patient administration. Re-suspend particles in syringe immediately prior to injection by repeated inversion of the syringe. (If blood is drawn into syringe, any unnecessary delay prior to injection may lead to clot formation in situ). Do not back-flush the syringe. Slow injection is recommended, and for optimum results, imaging should begin as soon as possible after injection.





# 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 TcO<sub>4</sub>

**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 5-75 $\mu$ m, and none larger than 150 $\mu$ m

Contact your NEN Representative or Customer Service for further details

## **NEN** New England Nuclear Radiopharmaceutical Division

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

# People Pictures for Clinical Clarity

```
L = LEAVE SYSTEM
H = HELP

ACTION? R;

NEXT PROTOCOL NAME? LIVER;

CURRENT PATIENT IS: ROBINSON-CRUSOE D SS56789
THIS PROTOCOL COLLECTS THREE STATIC VIEWS, THE FIRST
IS THE ANTERIOR, THE NEXT THE RIGHT LATERAL AND THE
LAST THE POSTERIOR.
NORMALISATION IS DONE WITH THE 'DIV' CORR. MTX.

STUDY NUMBER (12)? 127/741
NAME (30)? JONES D;
NUMBER (SS ETC) (14)? SS345;

CURRENT PATIENT IS: JONES D SS345

ADD, DELETE, LIST OR SELECT ? ;

NOW POSITION THE PATIENT SUPINE WITH THE LIVER AND
SPLEEN VISIBLE ON THE PERSISTENCE 'SCOPE.

TYPE CR TO GO ? ;
HIT SPACE BAR TO STOP EARLY
AGAIN, RESTART, KILL OR STOP? ;
VIEW NUMBER = 1; NUMBER OF MATRICES = 1

NOW POSITION THE PATIENT LYING ON THE LEFT SIDE.

TYPE CR TO GO ? ;
HIT SPACE BAR TO STOP EARLY
VIEW NUMBER = 2; NUMBER OF MATRICES = 1
IF YOU REQUIRE A POSTERIOR VIEW ANSWER YES
CONTINUE PROTOCOL ? Y;

NOW POSITION THE PATIENT PRONE WITH THE LIVER AND
SPLEEN VISIBLE ON THE PERSISTENCE 'SCOPE.

TYPE CR TO GO ? _
```

Typical protocol control dialogue (customer prepared) on monochrome display

At last! varicam,  
A sophisticated  
gamma camera  
computing system  
which not only  
provides a dynamic

capability but  
more significant  
static images  
without requiring  
computer  
expertise.



**varian associates**

611 Hansen Way, Palo Alto, California  
94303, USA. Telephone: (415) 493-4000

Molesey Rd, Walton-on-Thames, England.  
Telephone: (093 22) 28971 Telex: 261351



# From Abbott: a masterpiece of engineering... at a modest \$5795.

The Auto-LOGIC™ 50/121 Gamma Counting System represents an artful blend of advanced electronics and quality craftsmanship...at a price that's readily affordable.

Abbott designed the Auto-LOGIC System to get the job done—rapidly and efficiently—maximizing accuracy while minimizing tech time. The economical Auto-LOGIC 50/121 System is compact in size, big on performance and easy on your budget. Just compare:

50-sample capacity, 4.5 second sample cycle time, simplified pushbutton controls, automatic printout, automatic shut-off and more. So much more, in fact, that you'd have to look at systems costing twice as much to get comparable performance features.



Abbott Laboratories  
**Diagnostics Division**  
North Chicago, IL 60064  
800/323-9100



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Chicago, Illinois, 1975

**Auto-LOGIC 50/121:**  
state of the art.





# TI 201 opens up the heart

With TI 201 you enter a new era in the diagnostic  
evaluation of myocardial diseases.  
Myocard scintigraphy can now be carried out on a routine basis.  
Philips-Duphar supplies TI 201 wherever you are, whenever you want.

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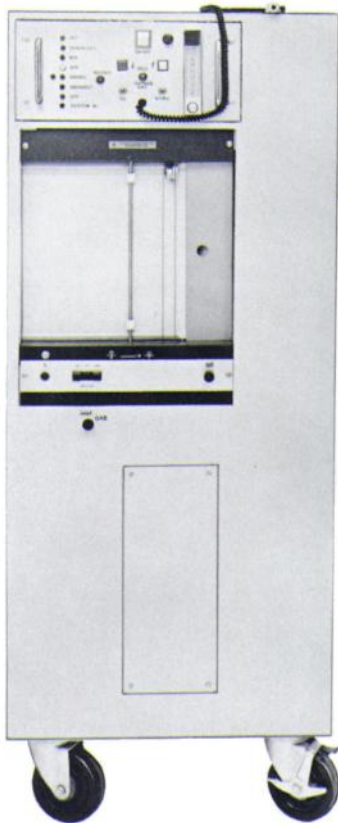


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PHILIPS-DUPHAR B.V. CYCLOTRON AND ISOTOPE LABORATORIES, PETTEN, HOLLAND.



*If you haven't found what you're looking for in a* **XENON-133**  
**LUNG FUNCTION UNIT**

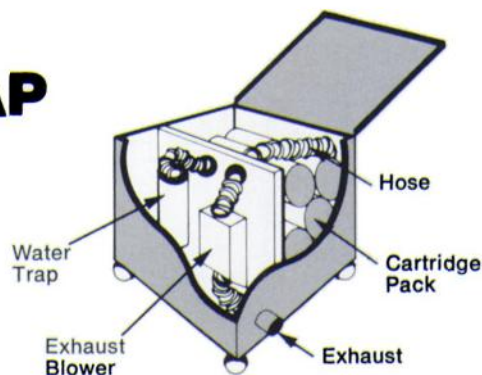


here are **3** good reasons to consider ours

- 1. SPIROMETER SYSTEM:** No bellows...just resistance-free breathing for your patients, regardless of the severity of their respiratory problem.
- 2. DIRECT BOLUS INJECTION:** No dead air space... your patient receives the direct, full bolus of xenon-133 exactly when desired.
- 3. RE-USE OF XENON GAS:** No costly waste, no matter what patient problem arises. You control the xenon flow throughout the system. It's always available for re-use during the same patient study.

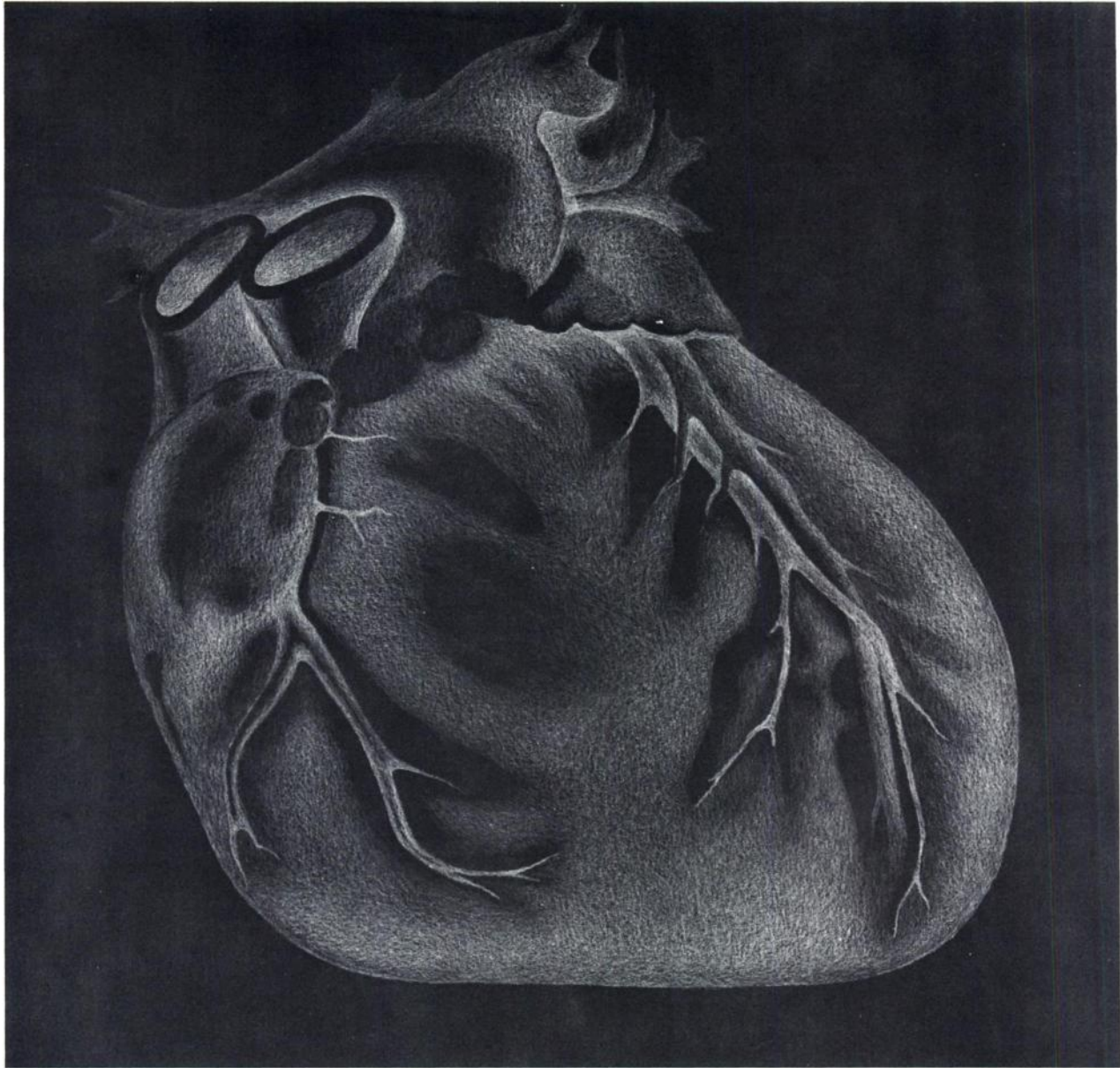
And **2** reasons to consider our  
**"NONEX" XENON GAS TRAP**

- 1.** Compatible with any xenon-133 gas handling system.
- 2.** Disposable 5-cartridge tandem filter removes all radioactive xenon from exhaled air. Outlasts single-cartridge units.



**NUCLEAR ASSOCIATES, INC.**  
 Subsidiary of  
**RADIATION-MEDICAL PRODUCTS CORP.**  
 100 VOICE ROAD, CARLE PLACE, N.Y. 11514

For full details,  
 ask for Bulletin  
 125-B

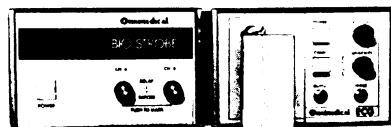


# Bio-Strobe: Cardiac Dual Gating

The immediate benefit to Cardiovascular Nuclear Medicine is simultaneous dual gating: ejection fraction calculations in one examination. Secondly, with no movement artifact, the Bio-Strobe achieves superior left ventricular function studies. And, in monitoring ventricular activities, it will isolate the T & P waves. More-

over, being a physiological gate, the Strobe has phonocardiogram capability, especially useful in cases like Bundle Branch Block. Modularity permits the use of the Strobe to gate any physiological parameter and the independent

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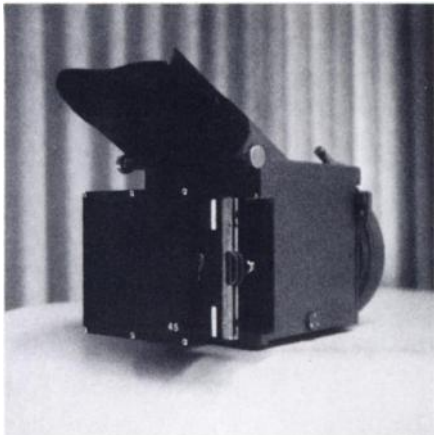
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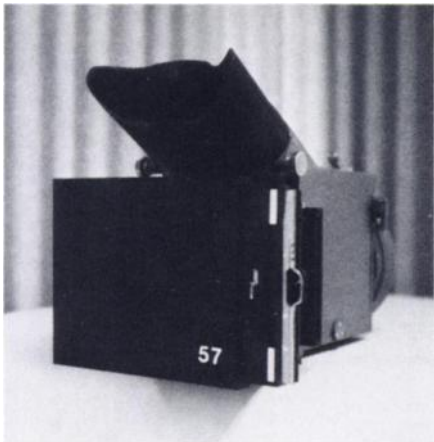
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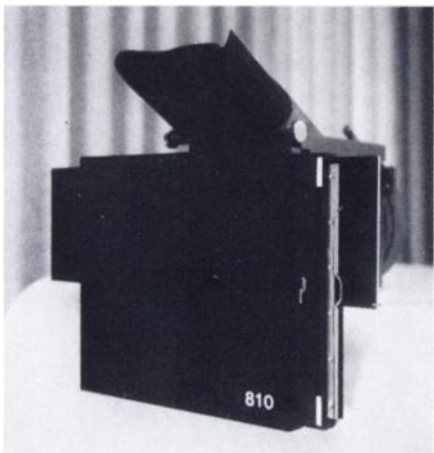
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**NUCLEAR MEDICINE TECHNOLOGIST:** Immediate opening, 1200-bed private hospital with medical school affiliation in Texas Medical Center. Well equipped expanding laboratory with 4 scintillation cameras and computer. Contact Staff Employment Manager, St. Luke's Episcopal & Texas Children's Hospital, 6621 Fannin, Suite 2R515, Houston, Tex. 77025. An Equal Opportunity Employer. M/F.

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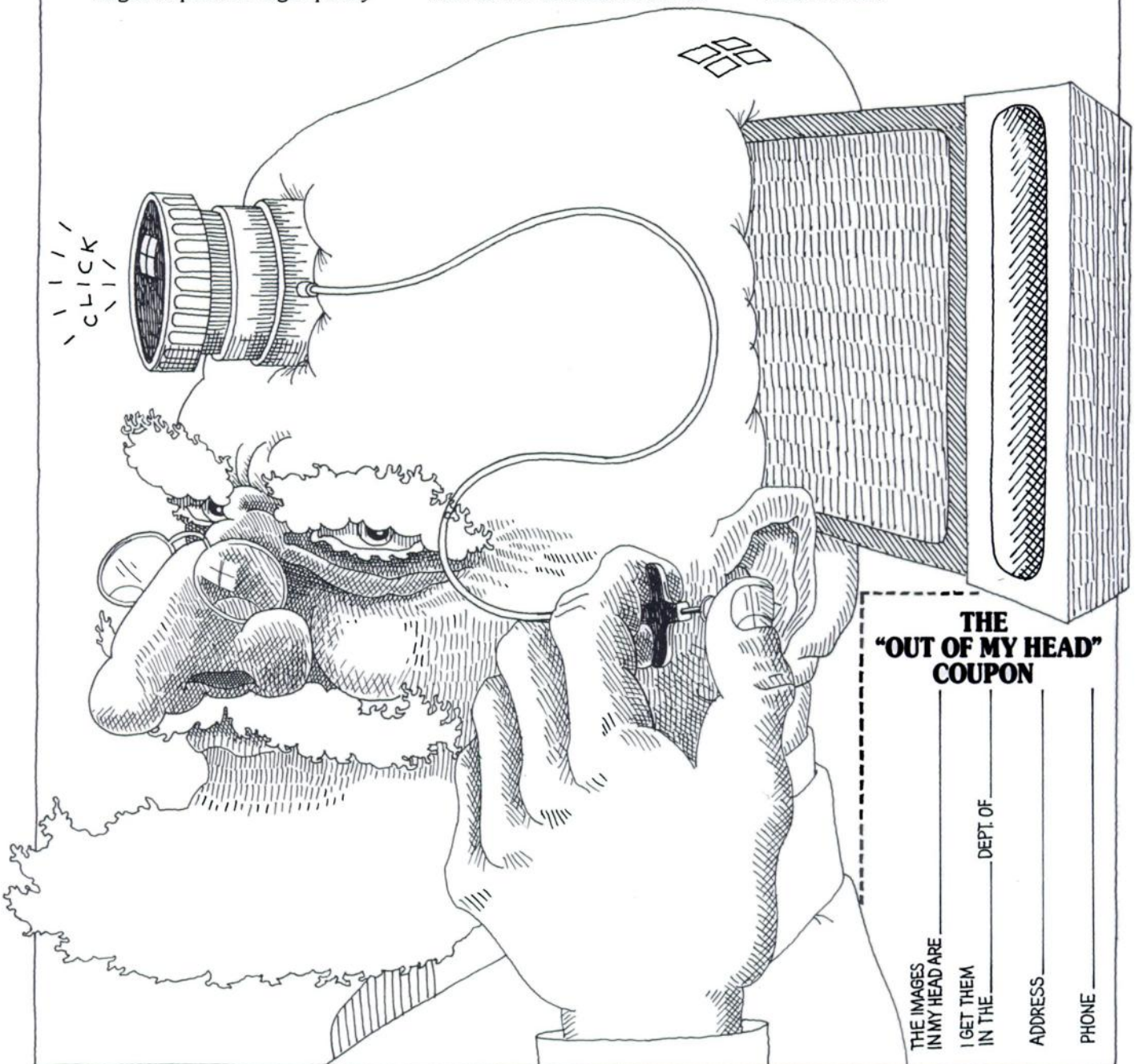
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For further information and applications for July, 1976, contact Howard Dworkin, M.D., Chief, Nuclear Medicine Department, William Beaumont Hospital, Royal Oak, Michigan 48072.

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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 1/8 page, \$90 for 1/4 page, \$165 for 1/2 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.

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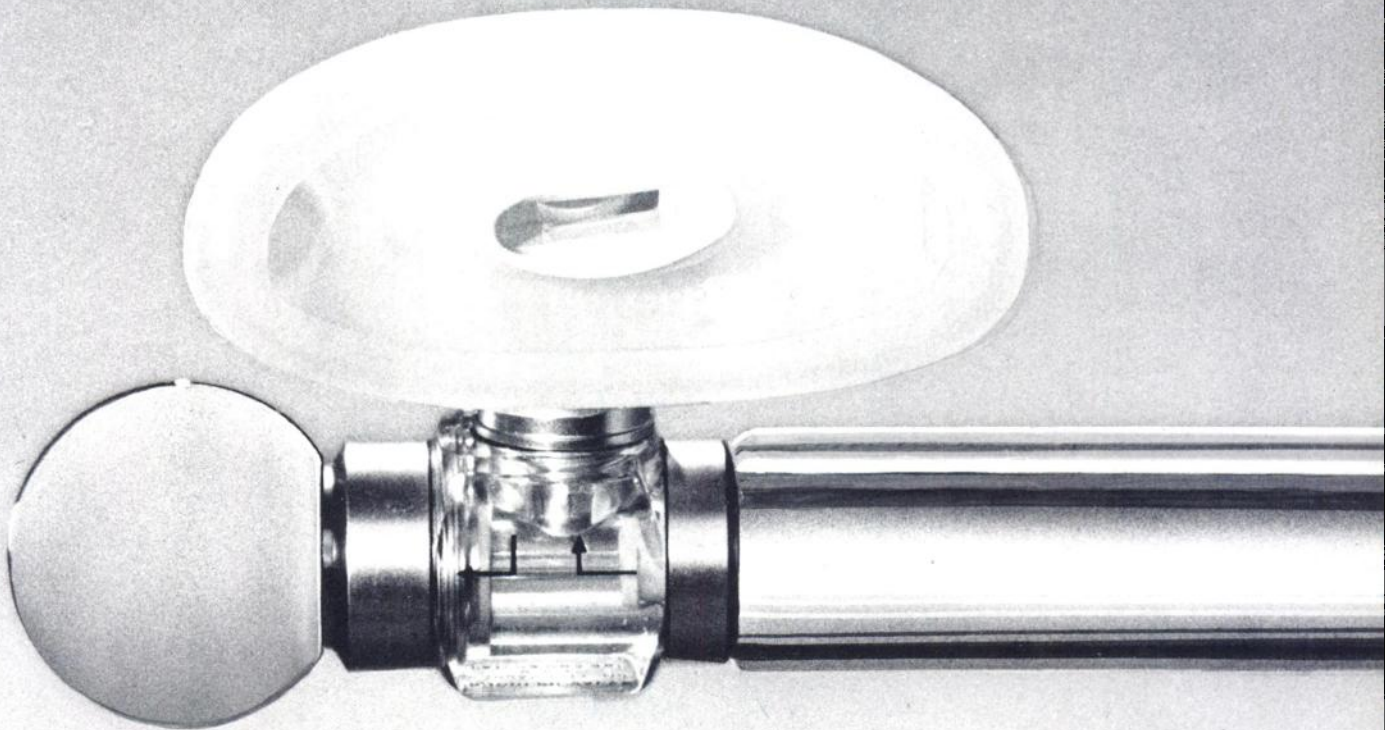
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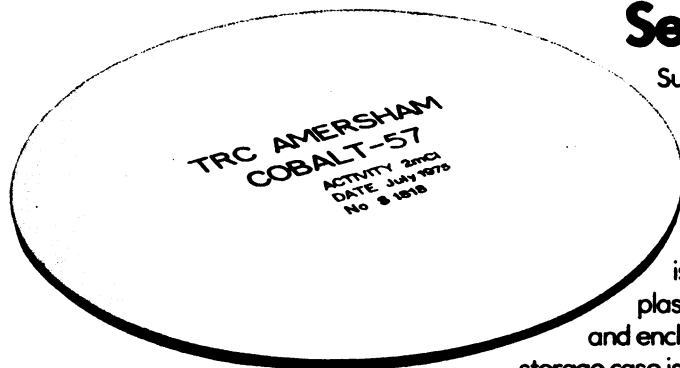
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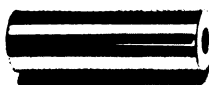
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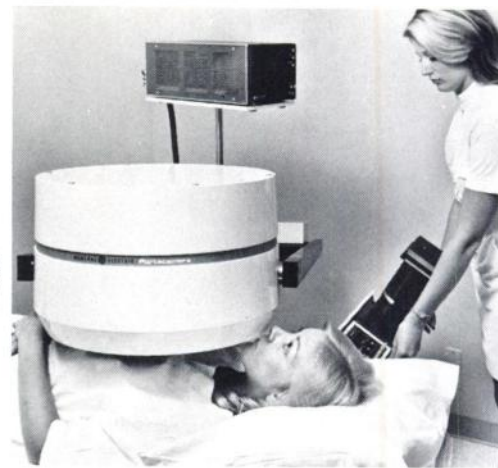


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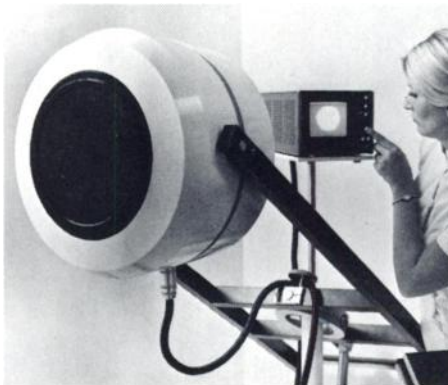


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- I am currently doing RIA work.
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## <sup>125</sup>I Folate

100 tube kit	\$ 70.00
200 tube kit	\$100.00

First to introduce <sup>125</sup>I Folate procedure.  
This procedure requires only 10 $\mu$ l of serum. Incubation Time: 45 minutes.

## Vitamin B<sub>12</sub> (<sup>57</sup>Co)

components for 200 tubes	\$100.00
--------------------------	----------

## <sup>125</sup>I Digoxin Kit

100 tube kit	\$ 55.00
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Sample Size: 50 $\mu$ l serum – Incubation Time: 30 minutes  
Separation: Charcoal – Sensitivity: 50 pg

## T<sub>4</sub>-RIA KIT

100 tube kit	\$ 70.00
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Specific T<sub>4</sub> Antiserum – No Extraction/No Evaporation  
Patient Sample: 5 $\mu$ l of serum – Incubation Time: 1 hour

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100 tube kit	\$ 21.00
200 tube kit	\$ 35.00

Patient Sample: 100 $\mu$ l – Incubation Time: 30 minutes

## T.S.H. KIT

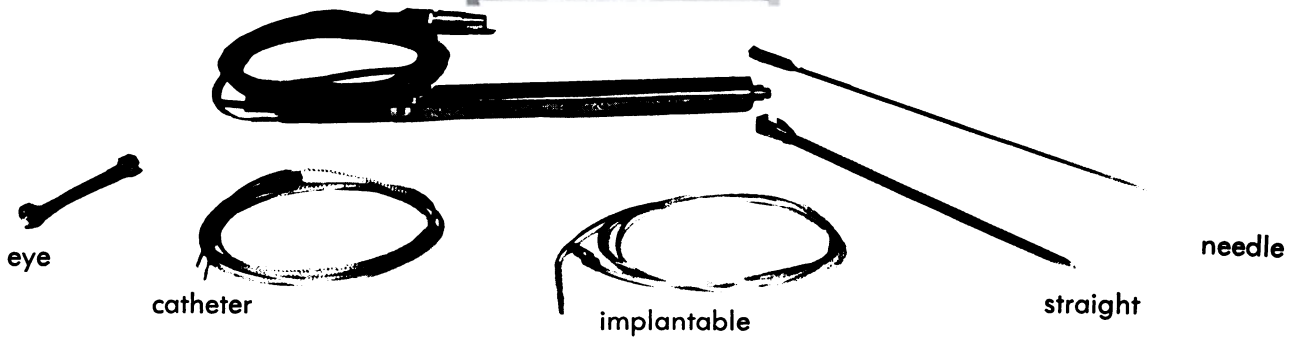
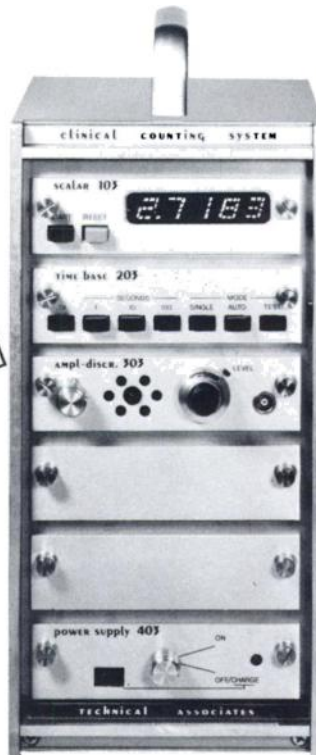
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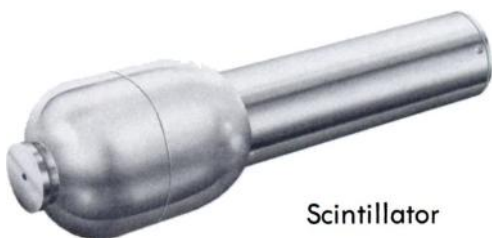


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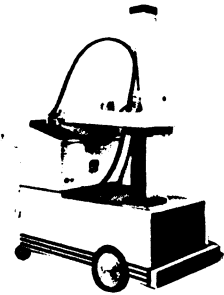


# mobility and dependability

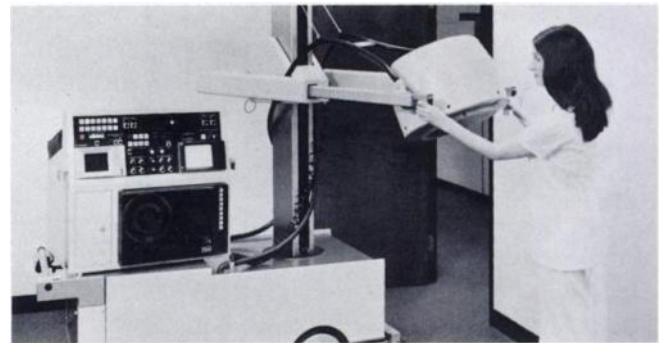


# with no loss in resolution

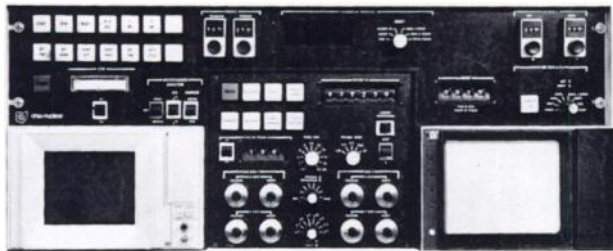
Wherever the need arises, in ICU, CCU, the Emergency Room, or within the NM Department, the Series 120 Mobile Camera is immediately available to generate high quality diagnostic information. And like all Ohio-Nuclear equipment, it is simple to operate.



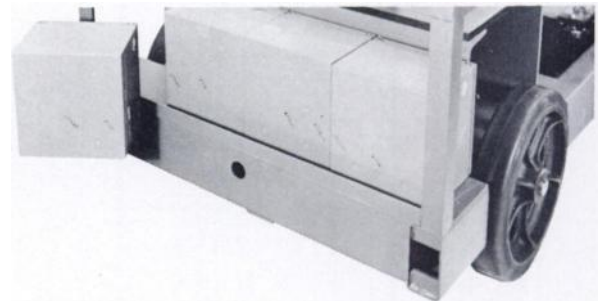
**Mobility.** The self-propelled Series 120 will travel at about 150' per minute, and negotiate a 10% incline under its own power, or it will creep for accurate patient positioning, all while maintaining full HV power to its photomultiplier tubes. This permits operation as soon as the unit is in place.



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PDR For Radiology and Nuclear Medicine is compiled with the same painstaking care and meticulous attention to detail that earns Physicians' Desk Reference its place as the traditional source of physician information. We wouldn't have it any other way.

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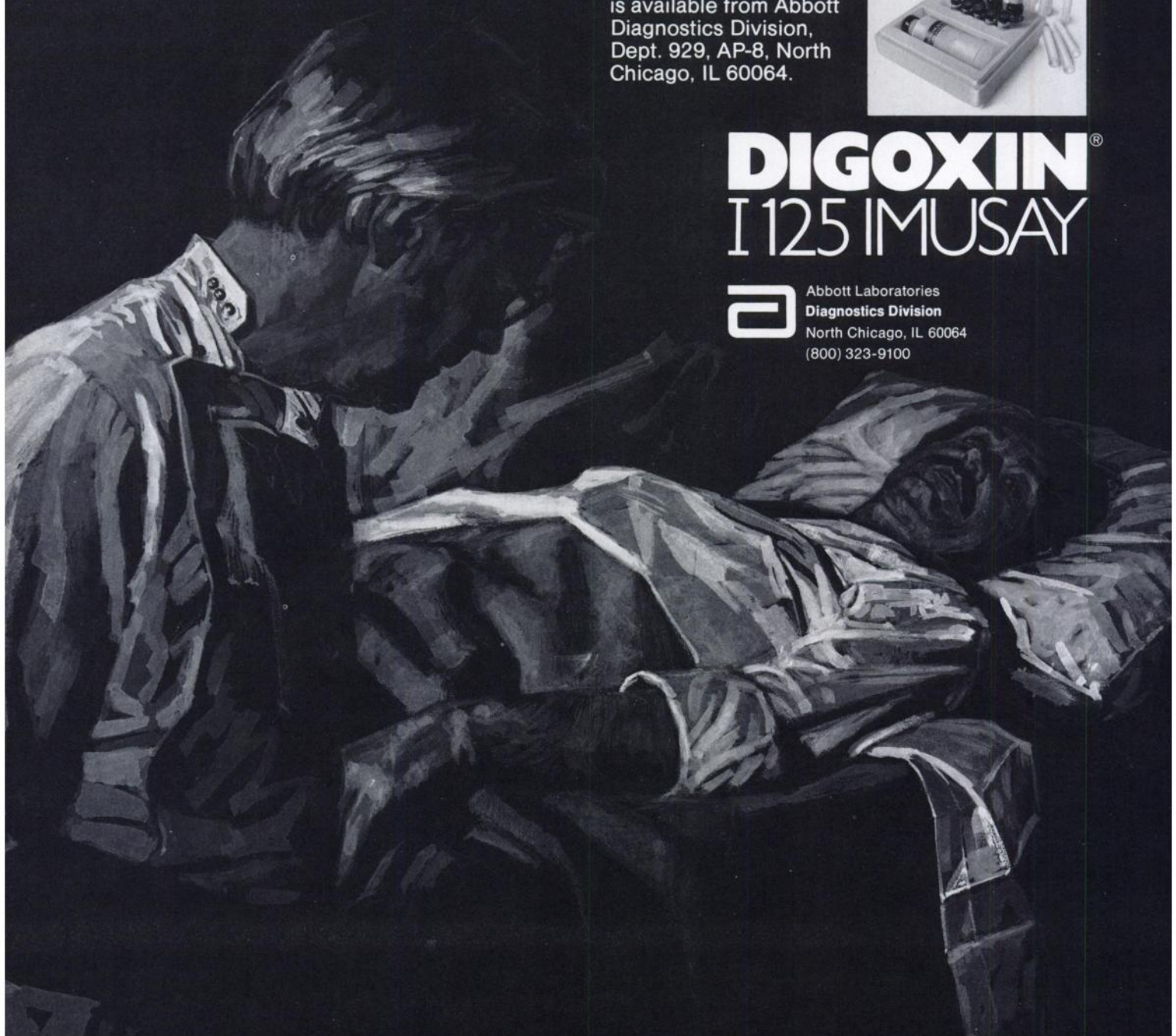
Important background information on cardiac glycosides prepared by a leading authority — Dr. Thomas W. Smith — is available from Abbott Diagnostics Division, Dept. 929, AP-8, North Chicago, IL 60064.



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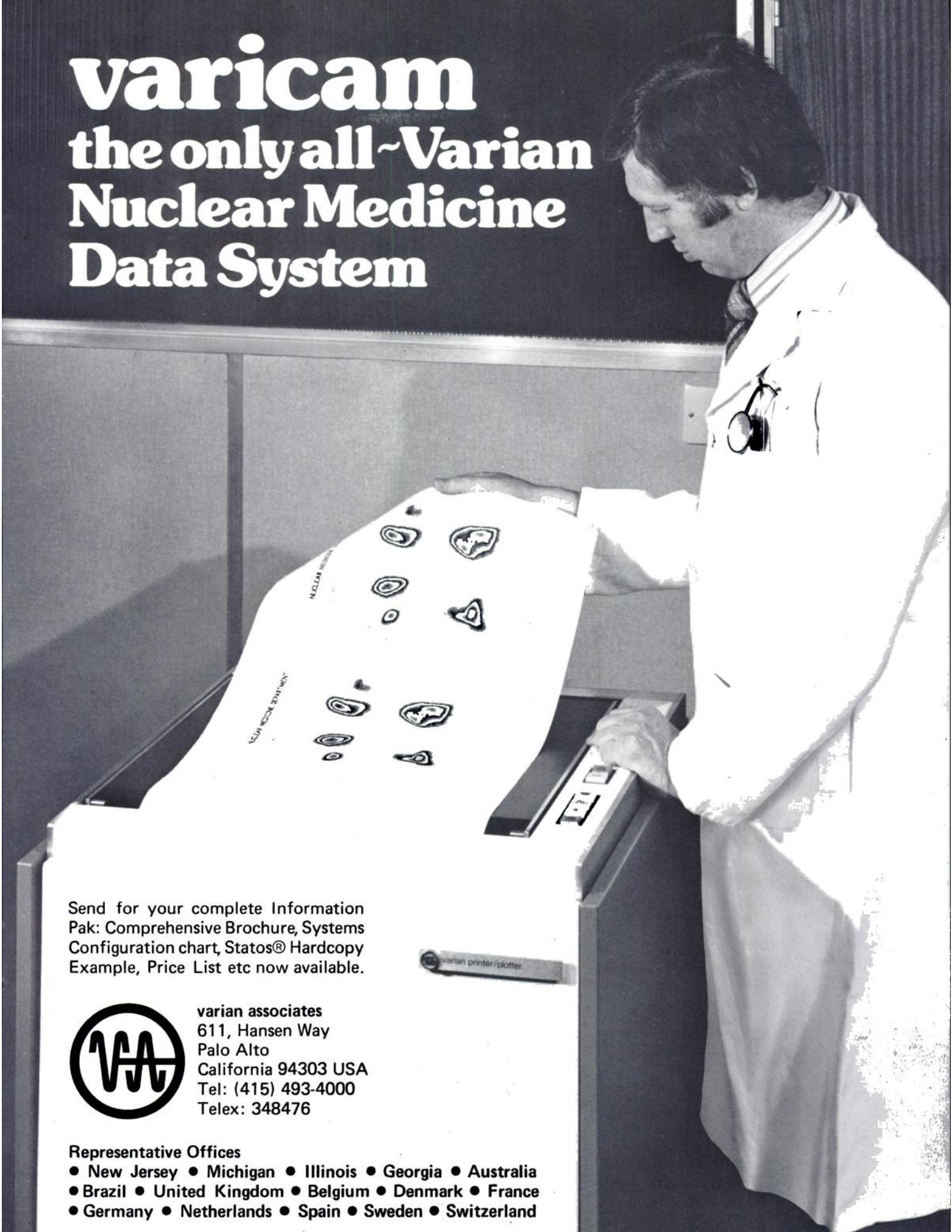
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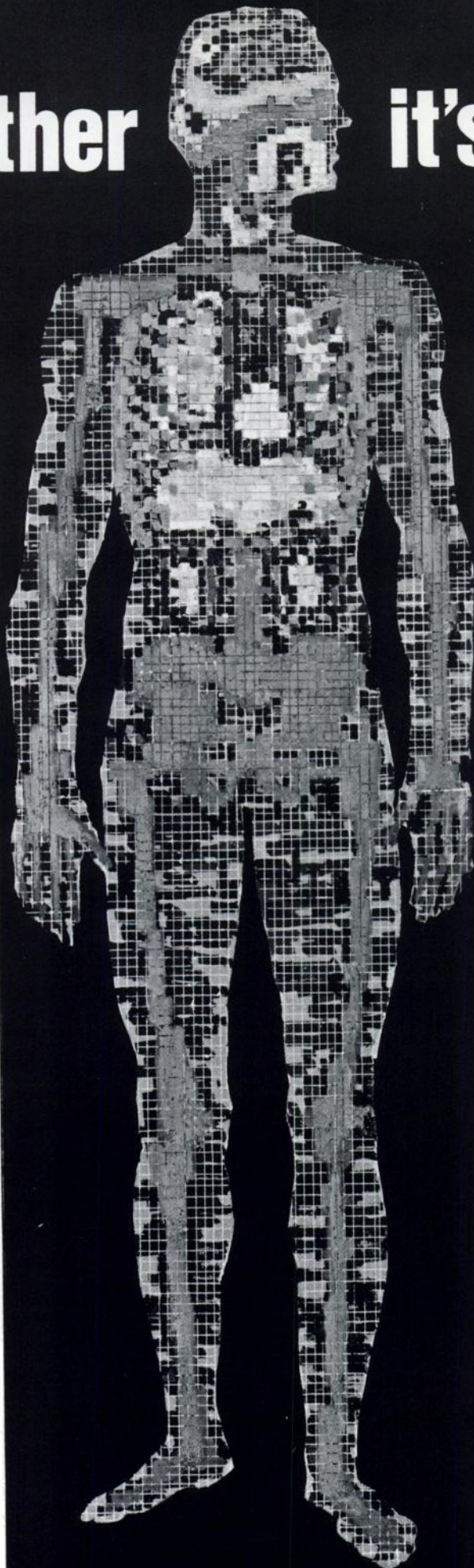
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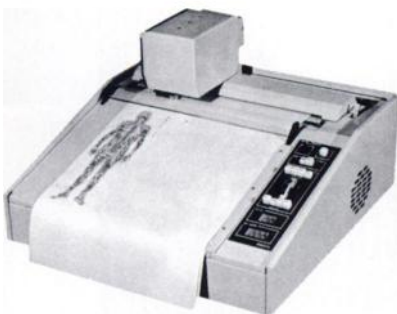
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Introducing the next generation of cortisol determinations — GammaCoat by Clinical Assays — the first solid phase Cortisol RIA. The greatly simplified extraction procedure, a test tube coated with cortisol — specific antibody and a <sup>125</sup>I cortisol derivative tracer brings accurate RIA cortisol determinations within reach of every clinical laboratory. A special additive is used to minimize the effects of variable serum proteins on the assay.

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4. Add tracer.  
Incubate 45 minutes.
5. Aspirate and wash.
6. Count the coated tubes.

The whole procedure takes less than two hours. Centrifugation and decanting are completely eliminated.

A <sup>3</sup>H Cortisol RIA with dextran coated charcoal separation is also available.

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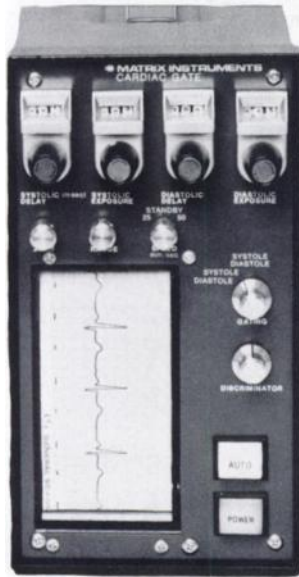
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# State of the art in cardiac and respiratory synchronization.

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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.

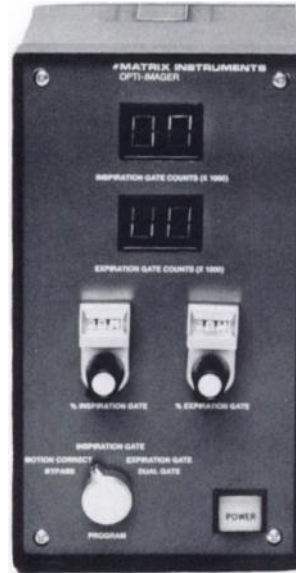
The dual gating operation mode allows recording of both end-systole and end-diastole simultaneously in a split screen two image format.

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**



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, inspiration plateau and expiration plateau images are recorded. The dual gating operation mode allows recording of both inspiration and expiration plateau images simultaneously in a split screen two frame format. Dual scalers record the number of counts in each image.

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**Endocrine sciences T3-38 and T4-15 thyroid hormone antisera offer:**

**Increased sensitivity and specificity**

**Assay times less than 5 hours\***

**Low sample volume requirements:**

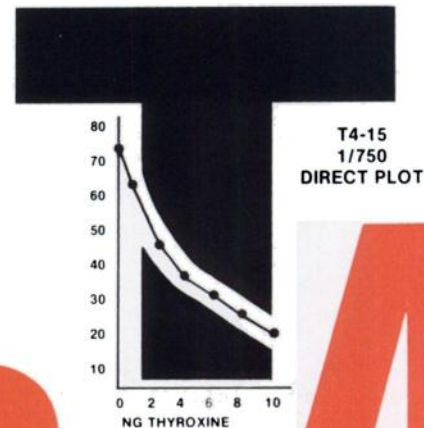
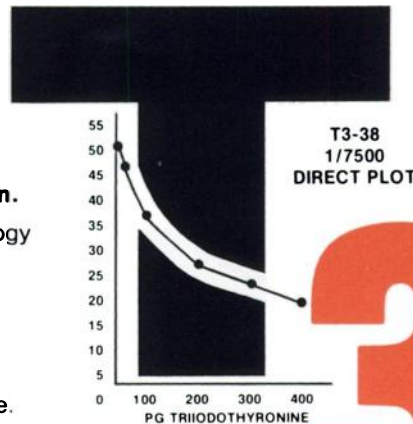
**Only 0.1 ml for T3-38\*  
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**Stability: Freeze-dried antisera are stable indefinitely if stored at -10°C, after reconstitution.**

Proven Endocrine Sciences methodology supplied with each antisera.

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# 3

# 4

**T3-38 and T4-15** are specific, high-affinity reagents developed for the radioimmunoassay of triiodothyronine (T3) and thyroxine (T4). Tested through routine use in our own clinical laboratories for over a year, T3-38 and T4-15 have been used in a simple RIA to determine T3 and T4 **directly** in plasma. The higher sensitivity and specificity of these antisera used in direct RIA offer distinct advantages over methods involving extraction and competitive protein binding. Increased sensitivity alone allows more precise measurement of T3 and T4 at critical lower physiological concentrations. Greater accuracy and precision are attained through elimination of errors associated with extraction and other sample processing.

**Sensitivity:** Standard curves normally obtained with T3-38 at a dilution of 1/7500 and T4-15 at a dilution of 1/750 are shown. Range and sensitivity of each curve were selected to measure generally encountered physiological concentrations of each hormone using sample volumes indicated above. The range of each can be adjusted to meet individual requirements by varying the dilution of the respective antiserum.

**Specificity:** T3-38 and T4-15 demonstrate very low cross-reactivity.

Multiple sample sizes with either T3-38 or T4-15 exhibit consistent linearity.

Hormone levels obtained in direct plasma RIA using T3-38 or T4-15 and those obtained after solvent extraction show no significant differences.

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Comparison of RIA using T4-15 with competitive protein binding:

Mean plasma T4 by RIA	9.5 ug%
Mean plasma T4 by CPB	9.0 ug%

## DIRECT PLASMA RIA

Today there is no better way to measure thyroid hormone levels in plasma than by radioimmunoassay, but RIA is only as reliable as the antiserum employed.

Clinical and research laboratories have been using Endocrine Sciences specific thyroid hormone antisera for more than a year now with complete confidence. Why? Because our T3 and T4 antisera were developed to meet exacting standards of specificity and sensitivity. Our customers know that each batch of T3 and T4 antiserum undergoes extensive quality control testing before its shipment. Users of our T3 and T4 antisera also know that our biggest customer is Endocrine Sciences Clinical Services Laboratory where these antisera must meet our own rigid standards daily.

Our antisera and reagents are offered as components rather than kits, because we believe in allowing more sophisticated users greater flexibility in methodology without incurring the additional expense of unnecessary reagents. Optimal sensitivity and reliability are easily attained using recommended procedures, thus eliminating the variability associated with most RIA kits. Check our specifications, then contact us for complete technical bulletins or to arrange for shipment.

Other Endocrine Sciences quality RIA reagents including T3 and T4 free plasma,  $^{125}$ I hormones, and purified bovine serum albumin are also available. Inquiries should be directed to our products division.

\* Based on use of RIA procedure similar to that provided by Endocrine Sciences.

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# New England Nuclear Radiopharmaceuticals

**INDICATIONS:** Inhalation of Xenon Xe 133 gas has proved valuable for the evaluation of pulmonary function and for imaging the lungs. It may also be applied to assessment of cerebral flow.

**CONTRAINDICATIONS:** To date, no known contraindications to the use of Xenon Xe 133 gas have been reported.

**WARNINGS:** This radiopharmaceutical should not be administered to pregnant or lactating women 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 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. Expired Xenon Xe 133 gas should be controlled in a manner that is in compliance with the appropriate governmental agency regulations.

Xenon Xe 133 adheres to some plastics and rubber and should not be allowed to stand in tubing or respirator containers. Such unrecognized loss of radioactivity from the dose for administration may render the study non-diagnostic. Xenon Xe 133 gas delivery systems, i.e., respirators or spirometers, and associated tubing assemblies must be leakproof to avoid loss of radioactivity into the laboratory environs not specifically protected by exhaust systems.

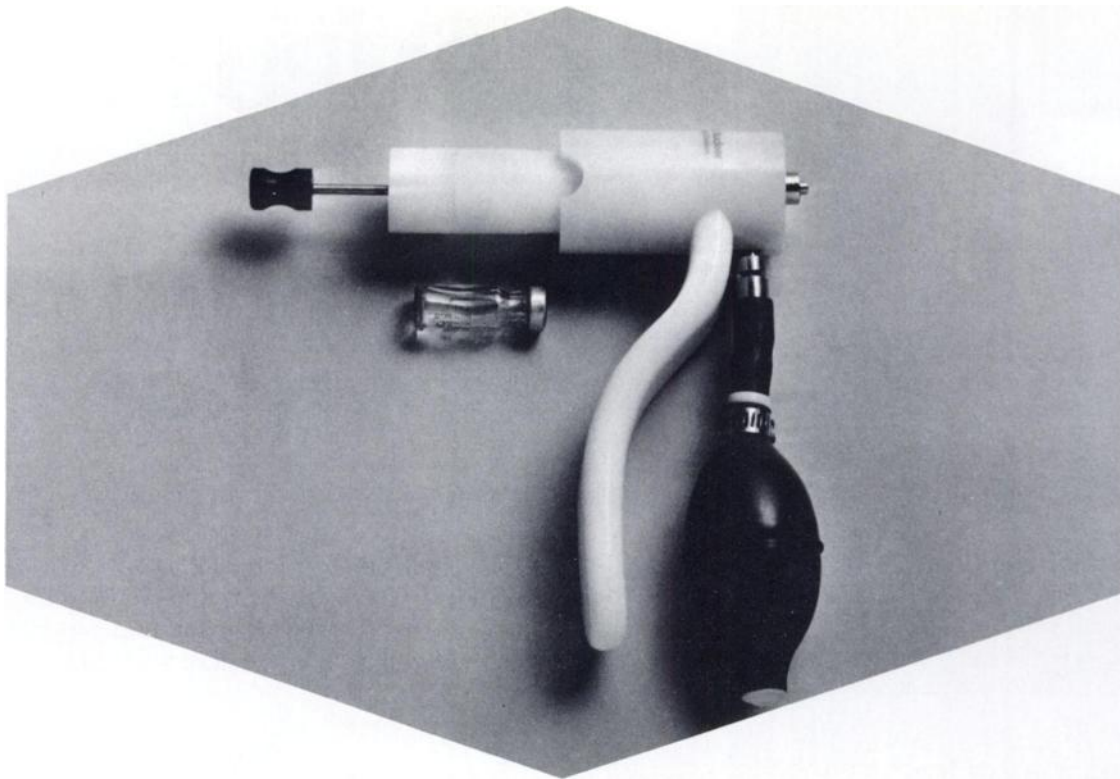
**ADVERSE REACTIONS:** To date, no adverse reactions based on the use of Xenon Xe 133 gas have been reported.

**DOSAGE AND ADMINISTRATION:** Xenon Xe 133 gas is administered by inhalation from closed respirator systems or spirometers.

The suggested activity range employed for inhalation by the average adult patient (70 kg) is:

Pulmonary function including imaging: 2-30mCi in 3 liters of air.  
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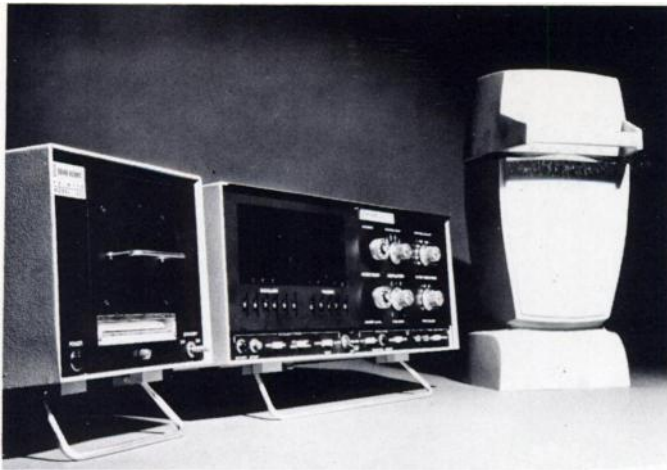
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Reconstitution of Phosphotec with sterile sodium pertechnetate-<sup>99m</sup>Tc results in an aqueous solution of Technetium 99m-Stannous Pyrophosphate Complex.

**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 <sup>99m</sup>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 <sup>99m</sup>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 <sup>99m</sup>Tc-sodium pertechnetate is added, adequate shielding of the resulting preparation must be maintained.

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 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.

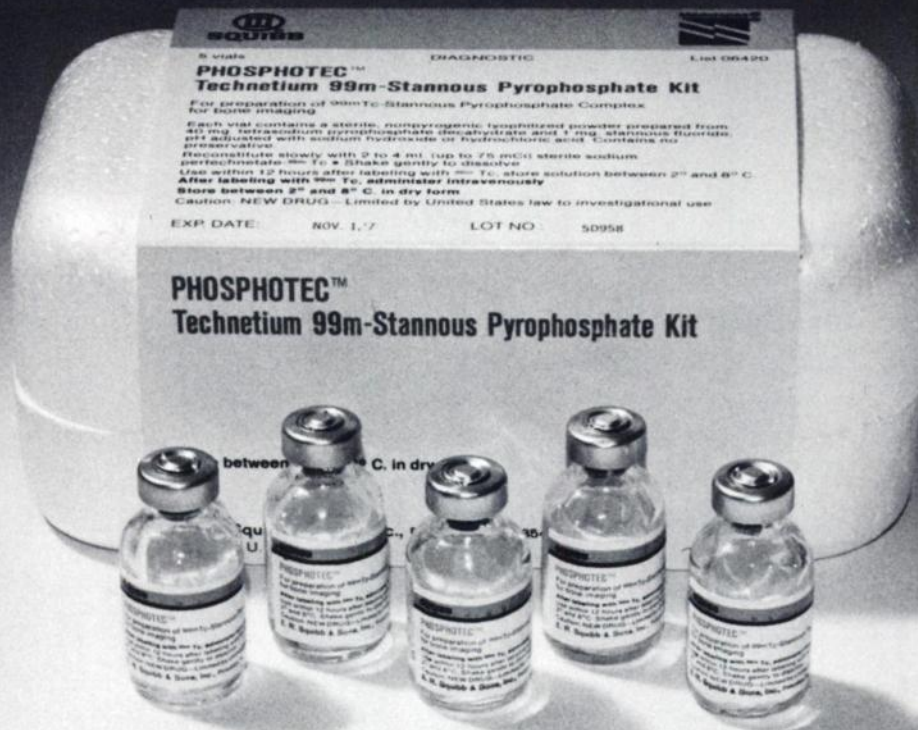
Use the preparation within 12 hours after labeling with <sup>99m</sup>Tc.

**ADVERSE REACTIONS:** At present, adverse reactions have not been reported following the administration of <sup>99m</sup>Tc-stannous pyrophosphate complex.

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

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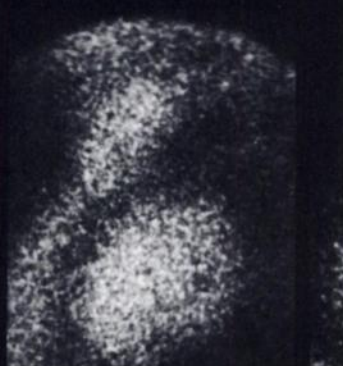
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RAO, DIASTOLE



RAO, SYSTOLE



LAO, DIASTOLE

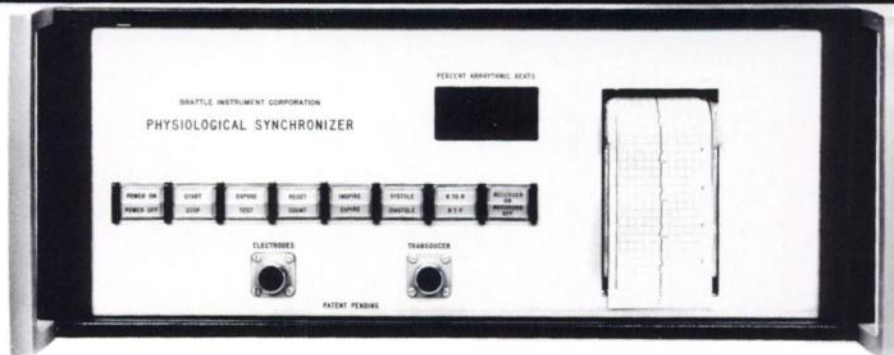


LAO, SYSTOLE

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  $^{99m}\text{Tc}$ -labelled Human Serum Albumin. The agent was prepared using the New

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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.

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