If you know get to know



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The in vitro test unmatched for reproducibility, convenience and accuracy.

Reproducible. Over 15 million tests conducted over the past eight years have made Triosorb® the standard of T-3 tests.

Convenient. The disposable Triosorb® Kit is ready for immediate use at room temperature making it one of the simplest, most convenient thyroid function tests available.

Accurate. Approximately 15 drugs and conditions produce misleading Triosorb®-T-3 test results, compared with over 200 factors which affect PBI.

* Also available as Triosorb®-131.



Tetrasorb-125 T-4 Diagnostic Kit

An improved, simplified method for measuring total serum thyroxine with diagnostic accuracy equal to or better than any currently used measures of thyroid function. Unlike other tests, exogenous iodines don't affect Tetrasorb® results.

one of these, them all.

The T-7 value completes the thyroid profile.

It's the Abbott method for determining the in vitro free thyroxine index.

T-7 is not a test but a numerical value derived from the multiplication of T-3 and T-4 test values. Because it is a product of two other numbers, the *T-7 value* will *move* only when both the T-3 and T-4 values move in the *same direction*. There are *only* two physiological conditions which cause this to occur, *hypothyroidism* and *hyperthyroidism*. With the exception of those patients receiving liothyronine or d-thyroxine therapy, all other factors which affect thyroid function tests will cause the T-3 and T-4 values to move in opposite directions, and the T-7 value to remain in the normal range.

When you provide the Abbott T-3, T-4 and T-7 values you furnish a complete thyroid profile with unparalleled clinical accuracy.

With LOGIC your final step is as easy as 1,2,3.

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- 2. Take a post-wash reading.

 Pre-set *timer* for the baseline established in step 1.
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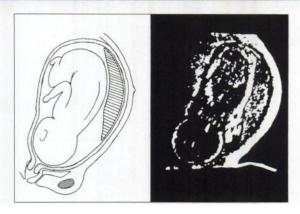
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PRECAUTIONS, ADVERSE REACTIONS: Care should be taken to administer the minimum dose consistent with patient safety and validity of data. The thyroid gland should be protected by prophylactic administration of concentrated iodide solution. Urticaria and acute cor pulmonate, possibly related to the drug, have occurred.

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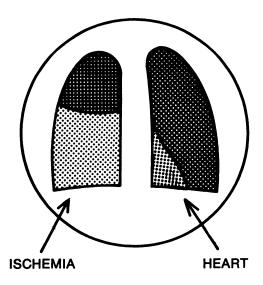
Volume 12, Number 2 ix



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Pulmonary Embolism?

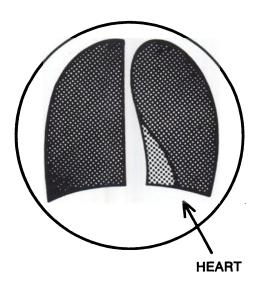




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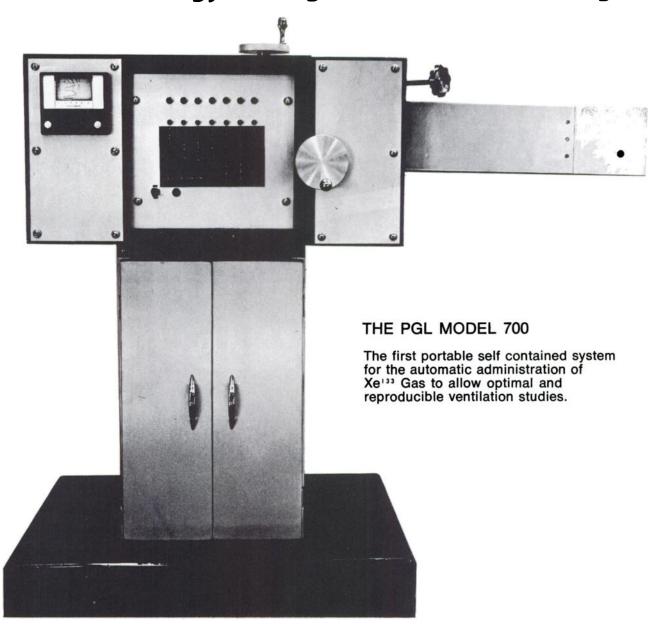




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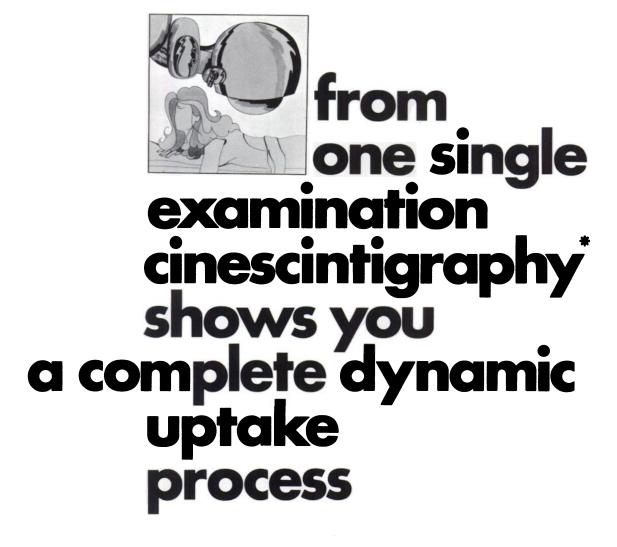


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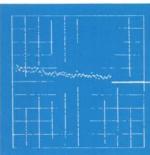


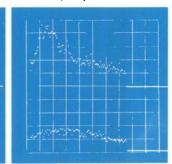


Below is a renogram

picture on which Replay of the digital magnetic tape gives, on the oscilloscope screen, 4 regions of interest the dynamic uptake curves for each region: activity versus time. have been selected Successive elementary images, corresponding to each point by light pen. of the curves, could also be displayed.









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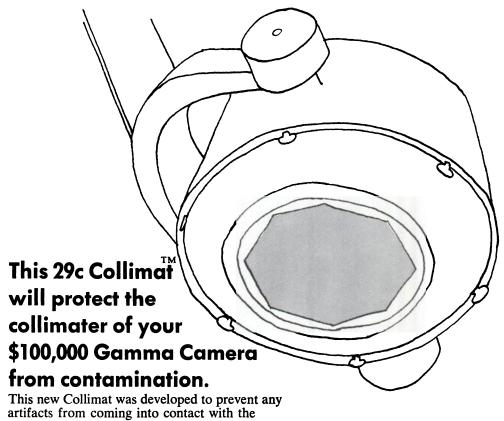
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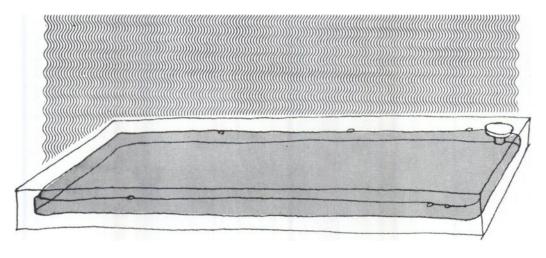
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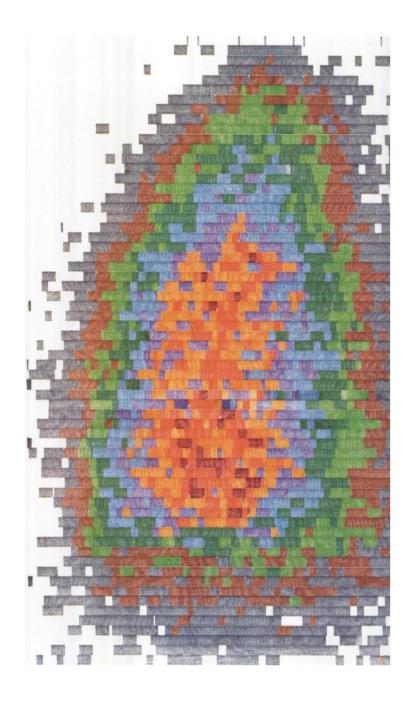
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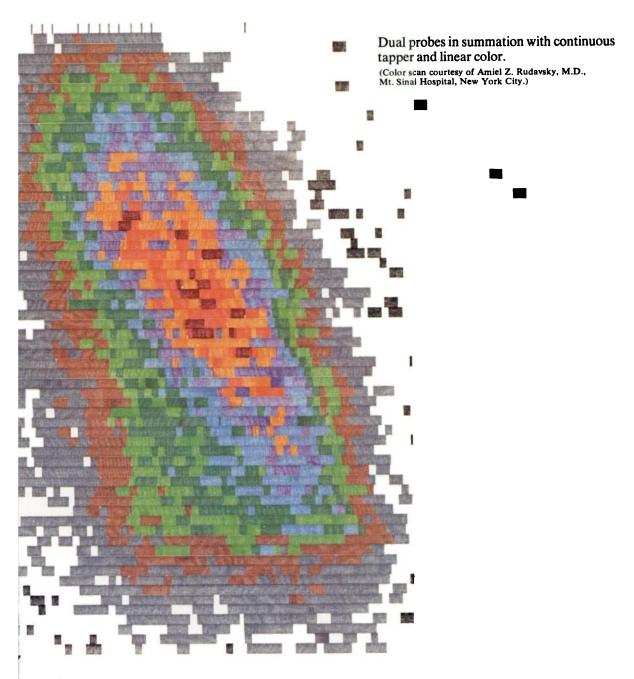
By simply inserting a plug, you can change a Raytheon imaging device from conventional linear color operation to the color dot contrast enhancement mode. Raytheon offers a wide variety of plugs to meet your clinical requirements for color contrast enhancement. The accompanying graph illustrates the results you can expect at various count rate activity levels.

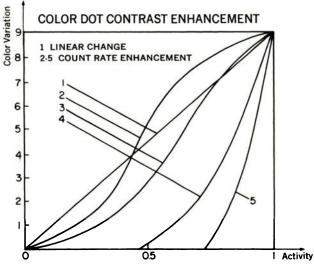
In addition, you have a choice of continuous color, another Raytheon exclusive, or conventional color recording with variable tapper frequency.

But there are a number of features of Raytheon

nuclear imaging devices that make them the most advanced units available today. For example: The scanning heads are completely flexible. Tomograms, oblique scans of normally masked areas, parallel-headed scanning for whole body applications, and conventional opposed-head scanning are some of the ways the heads can be manipulated.

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scanner will give you various levels of enhancement simultaneously.

With a Raytheon nuclear imaging device, you can also have a unit that can be updated to meet your future needs. You can convert a single 3" scanner to a single 5, dual 3, or dual 5 right in the hospital.

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We would like to provide you with additional details on Raytheon's family of nuclear imaging devices. Write or call Raytheon Company, Medical Electronics, 190 Willow St., Waltham, Mass. 02154.
Tel. 617-899-5949.

Raytheon's 12-minute, color film on nuclear imaging devices is available for your viewing. To arrange a convenient time to see this informative film, contact your nearest Raytheon sales representative. Or, get in touch with Raytheon Company, Medical Electronics, 190 Willow Street, Waltham, Mass. 02154. Telephone 617-899-5949.



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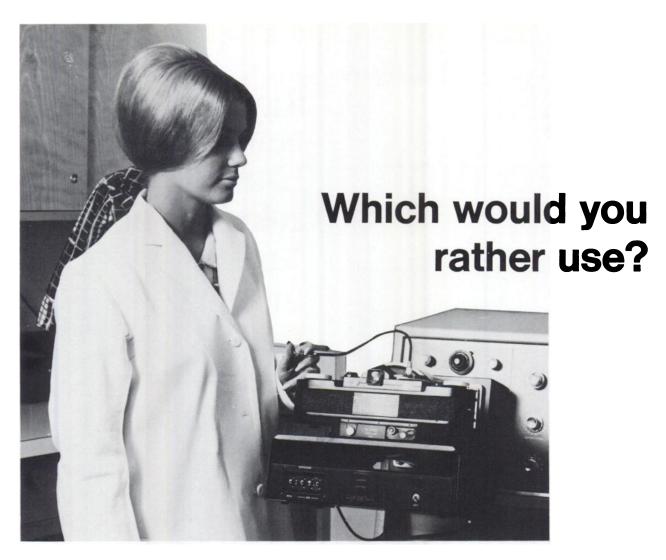
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Volume 12, Number 2 xxi



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FIGURE 1. SERIAL SCINTIPHOTOS. ANTERIOR VIEW.

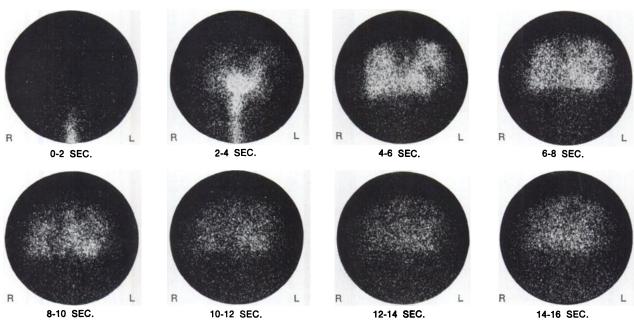


FIGURE 2. AREAS-OF-INTEREST. ANTERIOR VIEW.

R

FIGURE 3. PULMONARY DILUTION CURVES, ABNORMAL. Traced from original chart recordings for clarity of reproduction.

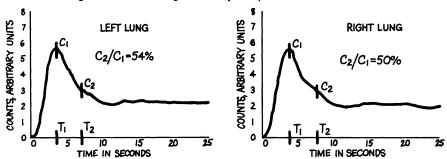
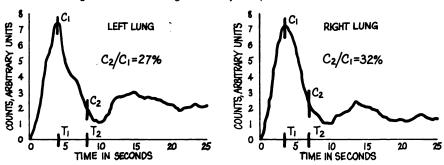


FIGURE 4. PULMONARY DILUTION CURVES, NORMAL. Traced from original chart recordings for clarity of reproduction.



The Cardiac Dynamic Study

A Dynamic Technique Using the Nuclear-Chicago Pho/Gamma® Scintillation Camera and Data-Store/Playback System

This study combines serial scintiphotos of the circulation of 99mtechnetium pertechnetate through the heart and lungs, photographed from the Pho/Gamma Scintillation Camera, with a time-concentration curve of the pulmonary circulatory dynamics using the Data-Store/Playback Accessory and a dual-channel ratemeter/dual-pen chart recorder.

SETTING UP. The patient is positioned beneath the Pho/Gamma detector so that the heart and lungs are included within the field of view. For adults, a central venous catheter is inserted and the tip is advanced to the superior vena cava. For children, a percutaneous femoral venous puncture is performed.

ISOTOPE AND DOSE. 50 microcuries/lb. of 99mTc pertechnetate are injected as a bolus. This is followed by a sterile saline "flush." It is imperative that the tracer be administered as a bolus for proper interpretation of the pulmonary dilution curve.

DATA ACCUMULATION. Since the 99mTc pertechnetate is injected so close to the heart, serial handpulled scintiphotos are started immediately. Each exposure is for 1-2 seconds and no more than eight films are necessary. Alternatively, the automaticsequencing 35mm camera may be used to obtain precisely timed sequential images.

The Data-Store/Playback Accessory plays an important role in the examination. The entire sequence is recorded in a high-resolution digital format (256 x 256 matrix) on the magnetic tape recording system. Subsequent replay of the tape allows reconstitution of the serial images at any desired frame rate and permits correction of film exposure factors to provide excellent scintiphotos. The study may be viewed on the system's variable-persistence oscilloscope during both original recording and upon tape replay.

The pulmonary dilution curves are obtained by choosing two separate areas-of-interest, one corresponding to the right lung field, the other to the left lung field. With this system's variable controls, these areas-of-interest may be rectangular or oval in shape. It is important, however, that these areas-of-interest correspond only to the lung fields, and no portion of the heart or great vessels should be included. Timeactivity curves are generated with the dual ratemeter/ recorder with a time constant of 0.5 seconds and a chart speed of 12 inches/minute.

CASE HISTORY. The clinical study on the opposite page is that of a seven-year-old child suspected of having a small left-to-right intercardiac shunt based on the characteristics of a systolic murmur. The child was not cyanotic. Following the diagnostic nuclearmedicine procedure, the patient was catheterized. A ventricular septal defect with a 1.2-to-1 left-to-right shunt was revealed as determined by standard dye dilution curves. In addition, there was a supervalvular obstruction of the pulmonary artery. Systemic pressures were observed in the right ventricle suggesting the diagnosis of an "Acyanotic Tetralogy of Fallot."

EVALUATION. The serial two-second images (Fig. 1) were produced upon replay of the Data-Store/Playback Accessory. The bolus of 99mTc pertechnetate is clearly seen in the inferior vena cava (0-2 sec.), having been injected into the right femoral vein. The tracer, thereafter, flows into the right atrium (2-4 sec.), then into the right ventricle and out through the pulmonary artery into both lung fields (4-6 sec.). Later frames show the return of the tracer to the left atrium, the left ventricle, and then out the aorta.

The pulmonary dilution curves were produced by adjusting the area-of-interest controls of the Data-Store/Playback Accessory, causing the areas-ofinterest to correspond to the right and left lungs as indicated by the intensified areas seen on the representative scintiphoto (Fig. 2). The resulting pulmonary dilution curves (Fig. 3) show a rapid rise in count rate to a peak count rate C_1 at time T_1 . T_1 - T_0 is the interval from time of rise onset to time of peak activity. At time T_2 ($T_2 - T_1 = T_1 - T_0$), count rate C_2 is determined from the curve. As shown, C_2 is 50 - 54% (C_2/C_1) of count rate C_1 . These curves are abnormal and suggest the possibility of a left-to-right shunt. Normally, C_2/C_1 is less than 40% as shown by normal curves (Fig. 4).

CONCLUSIONS. The diagnosis of a left-to-right shunt was confirmed in this case, both at cardiac catheterization and at surgery.

An abnormal pulmonary dilution curve, it should be noted, does not indicate the anatomical location of the defect, nor does it indicate the severity of the left-to-right shunt. This cardiac dynamic study should be considered only as a screening procedure. In the event of an abnormal radionuclide pulmonary dilution curve, further diagnostic procedures are indicated.

An exchange of information on topics



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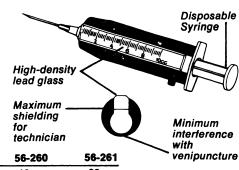
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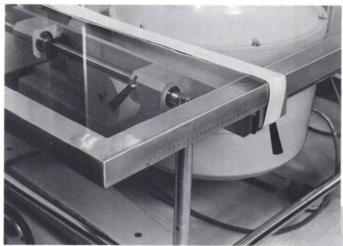
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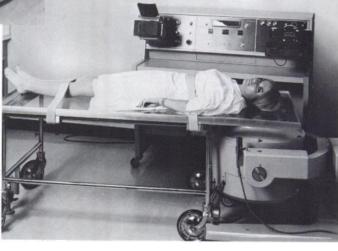
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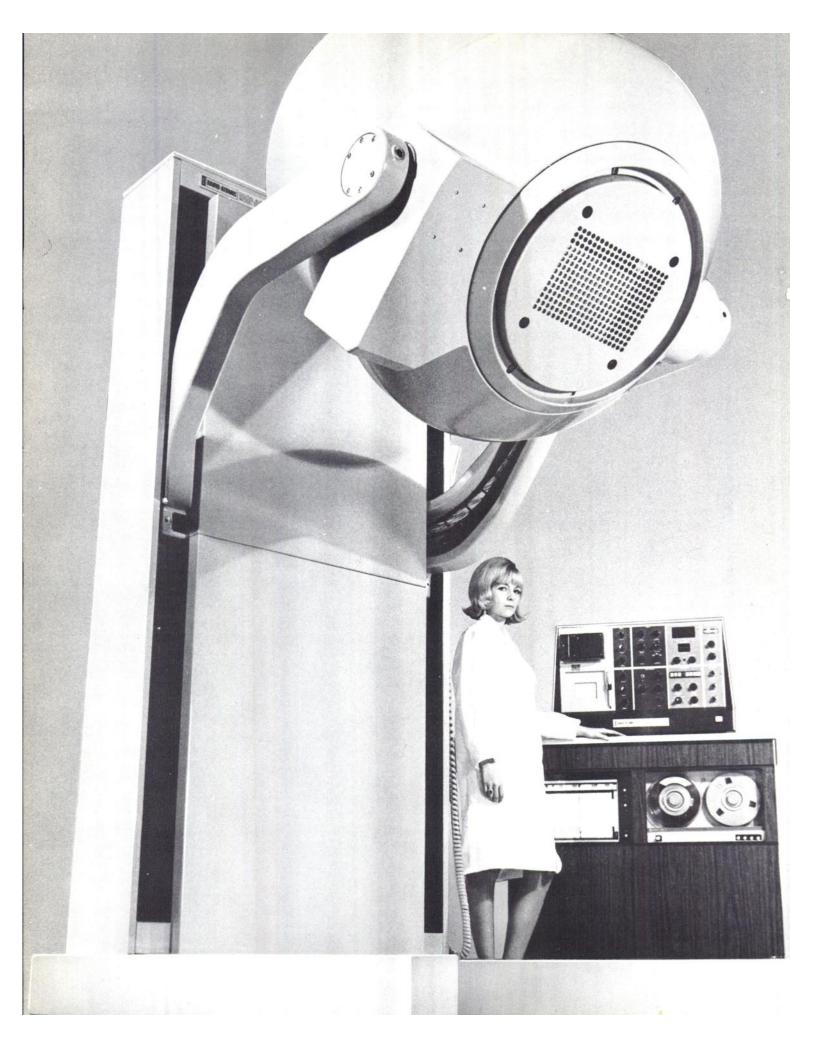
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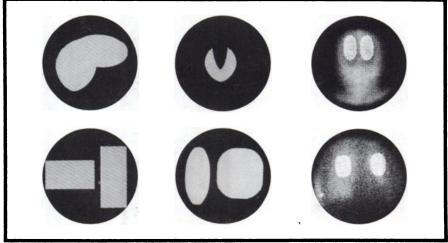
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