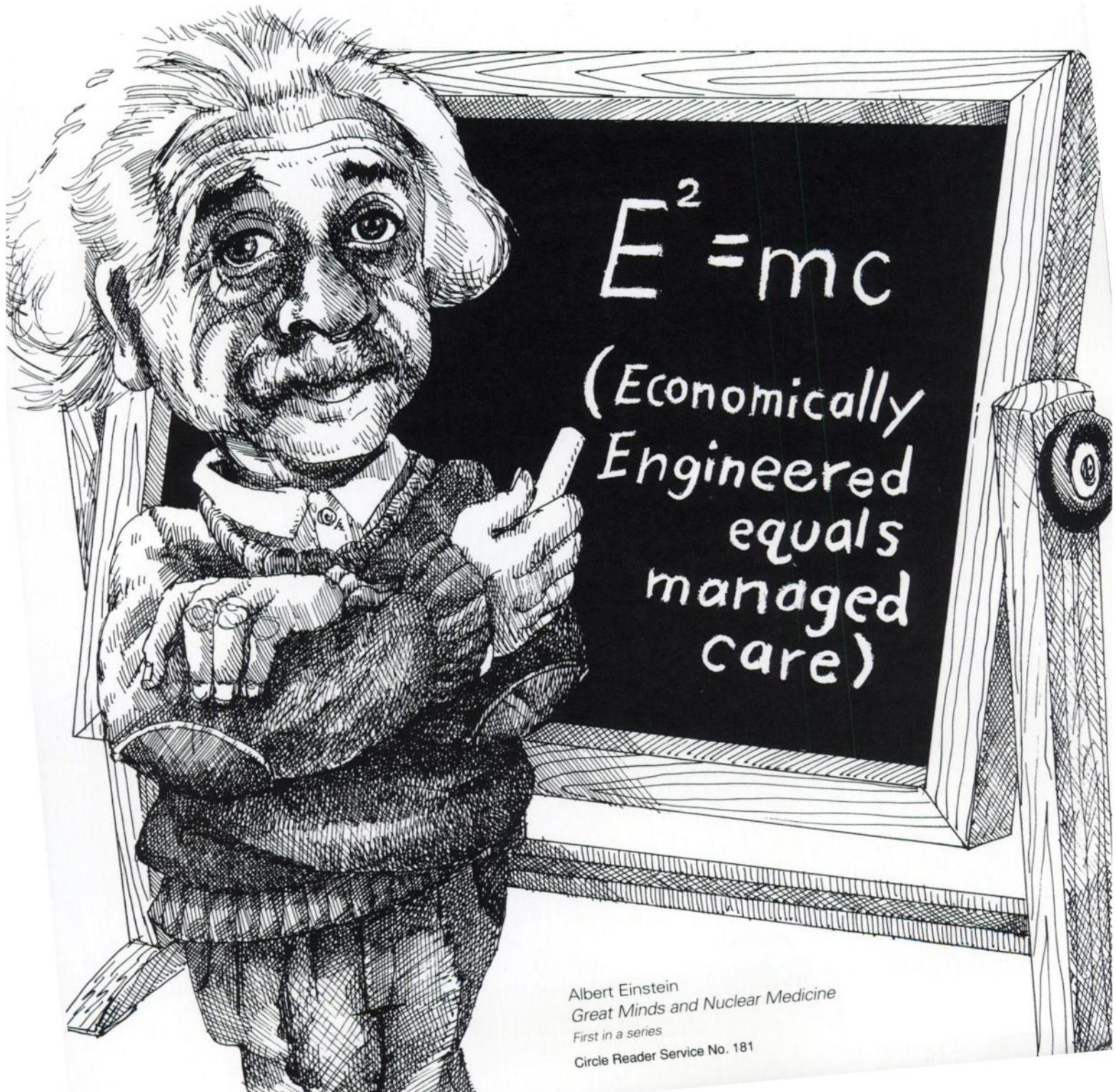


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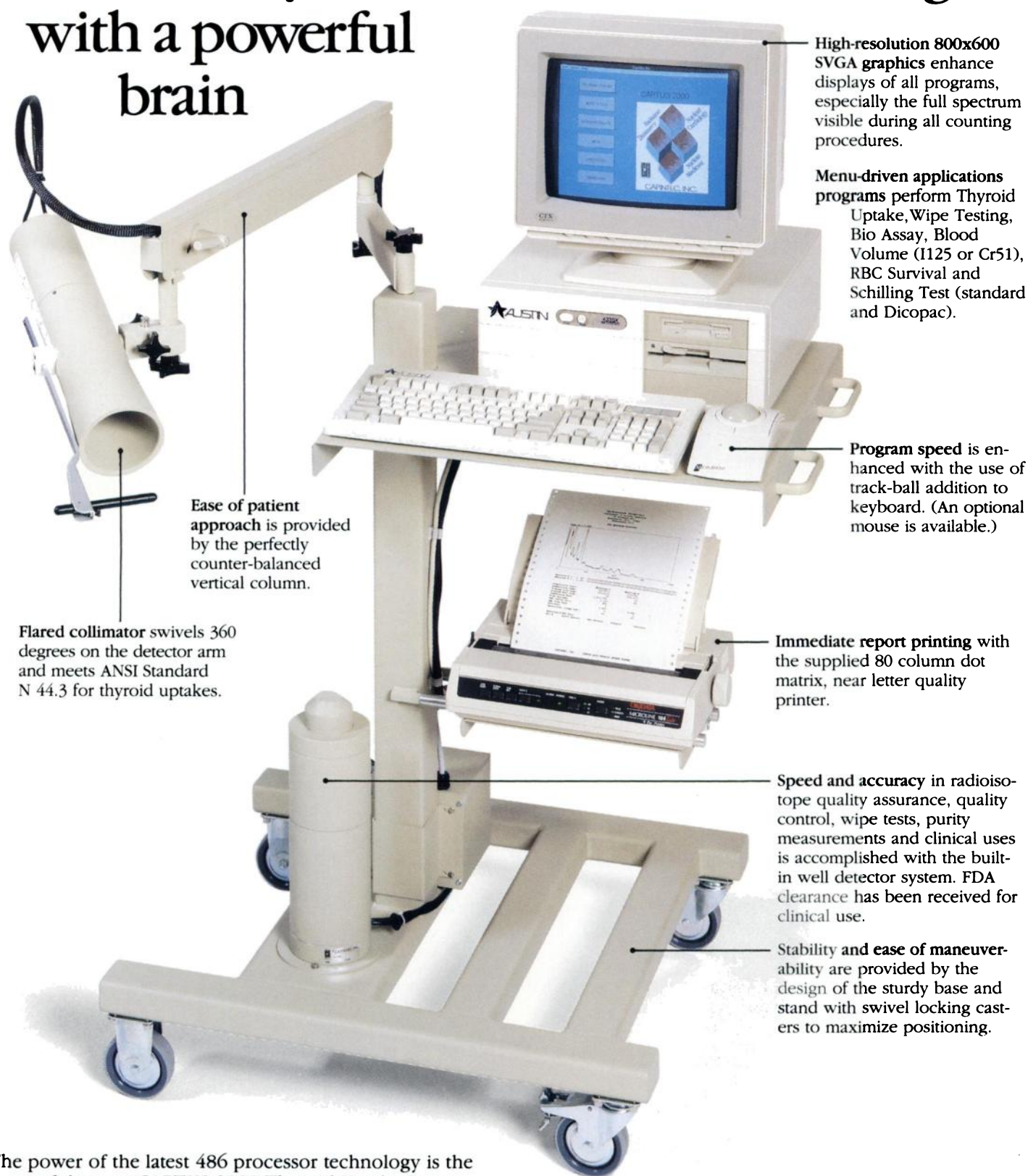
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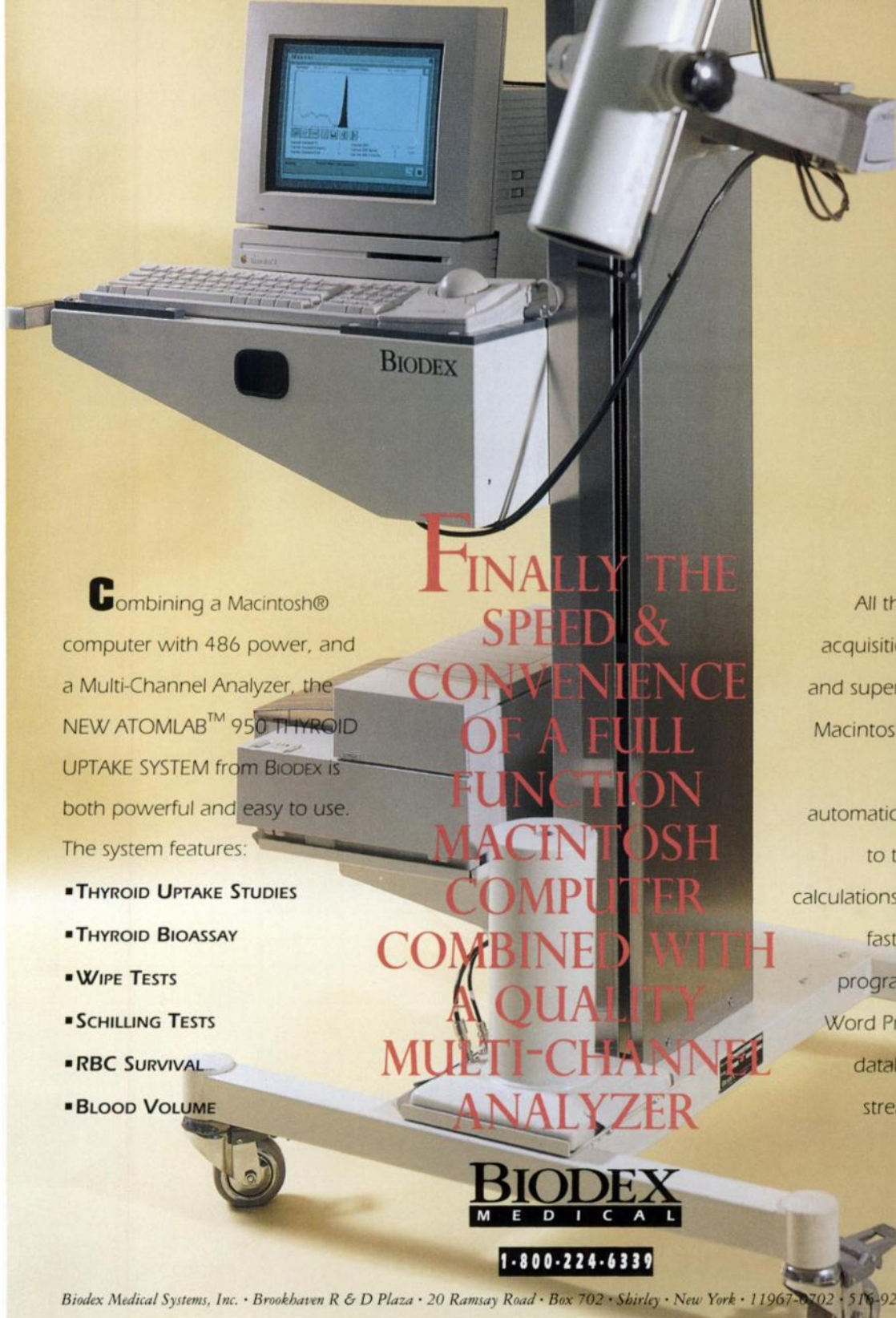
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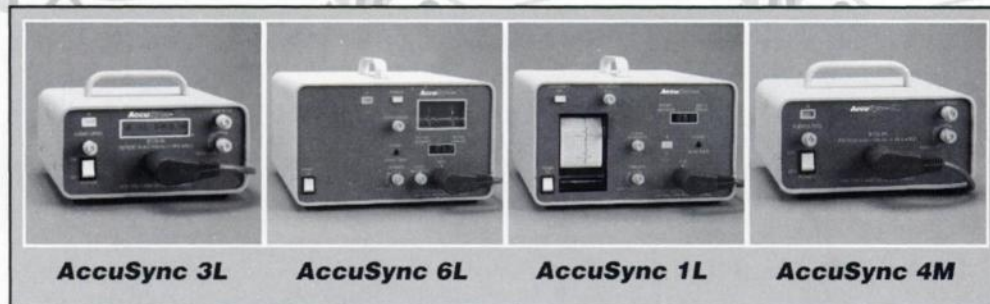
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The AccuSync 5L, our top model (featured at left) includes CRT monitor (visual) and Strip Chart Recorder (hard copy).

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Model	Strip Chart	CRT Monitor	HR/R-R Int	Trigger
5L	•	•	•	•
6L		•	•	•
1L	•		•	•
3L			•	•
4M				•

Accessory and optional products available:

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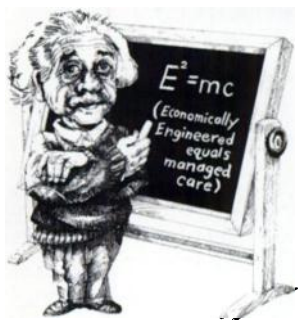
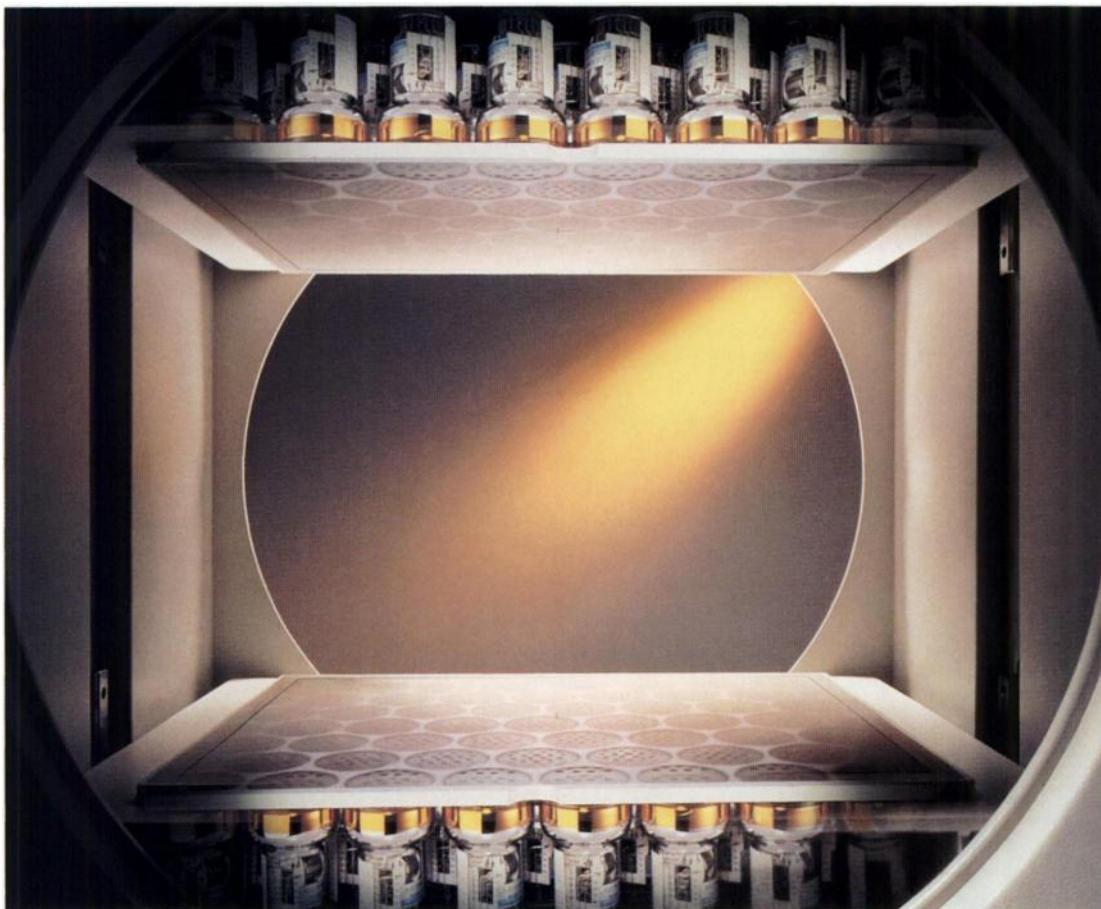
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Breakthrough in cancer detection...

ONCO SCINT[®] CR/OV

Satumomab Pendetide (1mg/2mL)

A diagnostic tool
that can assist
decision making
in patients with
colorectal or recurrent
ovarian adenocarcinoma

Please see last page for brief summary of prescribing information.

ONCOscINT[®] CR/OV

Satumomab Pendetide (1mg/2mL)

To enhance decision making in the management of patients with

The first monoclonal antibody-based
in determining both the location and

Reveals malignancy with tumor-targeted accuracy—

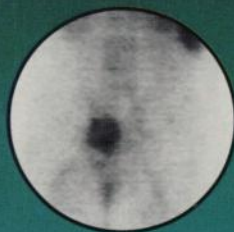
specifically targets the cell-surface antigen, TAG-72, common to colorectal and ovarian adenocarcinomas,^{1,2} making it possible—with the use of existing nuclear medicine equipment—to detect extrahepatic abdominal and pelvic lesions.^{3,4}

Provides information beyond the scope of standard diagnostic modalities—when interpreted in conjunction with a review of information obtained from other appropriate tests

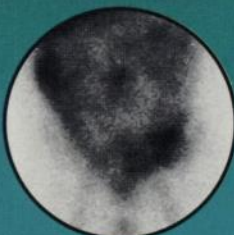
Found to be beneficial in these difficult situations:



- determining the source of a rising serum tumor marker in patients with an otherwise-negative workup^{2,4,5}




- determining extrahepatic abdominal and pelvic involvement in patients thought to have isolated and resectable recurrence^{2,4}



- differentiating disease from postsurgical or postradiation anatomic changes⁴

OncoScint is a registered trademark of CYTOGEN Corporation.

Please see last page for brief summary of prescribing information.



colorectal or recurrent ovarian cancer...

Imaging agent effective extent of disease

Assists decision making in patient management²⁻⁵—

enhanced medical/surgical management in
difficult colorectal^{3,4} and recurrent ovarian
cases.^{2,5}

Excellent safety profile*—

with generally minor and transient side
effects occurring in less than 4% of patients
studied (most frequently reported: fever,
chills and clinically insignificant changes in
blood pressure).²

*See Adverse Reactions section of brief summary on
following page.

For further information, please call
1-800-833-3533.

ONCO SCINT[®] CR/OV

Satumomab Pendetide (1mg/2mL)

Tumor-targeted cancer detection

OncoScint® CR/OV Kit (satumomab pendetide)

Kit for the Preparation of indium In 111 satumomab pendetide
For Intravenous Use Only
Brief summary of prescribing information

INDICATIONS AND USAGE

OncoScint® CR/OV-In (indium In 111 satumomab pendetide) is a diagnostic imaging agent that is indicated for determining the extent and location of extrahepatic malignant disease in patients with known colorectal or ovarian cancer. Clinical studies suggest that this imaging agent should be used after completion of standard diagnostic tests when additional information regarding disease extent could aid in patient management. The diagnostic images acquired with OncoScint® CR/OV-In should be interpreted in conjunction with a review of information obtained from other appropriate tests.

OncoScint® CR/OV-In is not indicated as a screening test for ovarian or colorectal cancer.

Administration of OncoScint® CR/OV-In may result in falsely elevated values from *in vitro* immunoassays, including tests for carcinoembryonic antigen (CEA) and CA 125. Because this interference may persist for months, the clinical laboratory should investigate for assay interference in patients who develop elevated CEA or CA 125 subsequent to imaging with OncoScint® CR/OV-In (see *Drug/Laboratory Test Interactions*).

Due to insufficient safety and efficacy data regarding repeat administration of this product, this imaging agent is limited to single use only.

CONTRAINDICATIONS

OncoScint® CR/OV-In (indium In 111 satumomab pendetide) should not be used in patients who are hypersensitive to this or any other product of murine origin or to indium In 111 chloride.

WARNINGS

Allergic reactions, including anaphylaxis, can occur in patients who receive murine antibodies. Although serious reactions of this type have not been observed in clinical trials after OncoScint® CR/OV-In (indium In 111 satumomab pendetide) administration, medications for the treatment of hypersensitivity reactions should be available during administration of this agent.

PRECAUTIONS

General The components of the kit are sterile and pyrogen free and contain no preservative. OncoScint® CR/OV-In (indium In 111 satumomab pendetide) should be used within 8 hours after radiolabeling. It is essential to follow the directions for preparation carefully and to adhere to strict aseptic procedures during preparation of the radiolabeled product.

Each OncoScint® CR/OV kit is a unit of use package. The contents of the kit are to be used only to prepare OncoScint® CR/OV-In; unlabeled OncoScint® CR/OV should NOT be administered directly to the patient. After radiolabeling with indium-111, the entire OncoScint® CR/OV-In dose must be administered to the patient for whom it was prescribed. **Reducing the dose of either component may adversely impact imaging results, and, therefore, is not recommended.**

The contents of the kit are not radioactive. However, after the indium In 111 chloride is added, appropriate shielding of OncoScint® CR/OV-In must be maintained. Care should be taken to minimize radiation exposure to patients and medical personnel, consistent with proper hospital and patient management procedures.

In addition, radiopharmaceuticals should be used only by physicians and other professionals who are qualified by training and experience in the safe use and handling of radionuclides.

Information for Patients Murine monoclonal antibodies are foreign proteins, and their administration can induce human anti-murine antibodies (HAMA). While limited data exist concerning the clinical significance of HAMA, the presence of HAMA may interfere with murine-antibody based immunoassays, could compromise the efficacy of diagnostic or therapeutic murine antibody-based agents, and may increase the risk of adverse reactions. For these reasons, patients should be informed that the use of this product could affect the future use of other murine-based products, including OncoScint® CR/OV-In, and should be advised to discuss prior use of murine-antibody based products with their physicians (see *Heterologous Protein Administration*).

Heterologous Protein Administration Murine monoclonal antibodies (MAbs) are heterologous proteins, and their administration can induce human anti-murine antibodies (HAMA).

OncoScint® CR/OV-In has been shown to induce HAMA to murine IgG after single administration in about 40% of patients in tumor imaging trials. HAMA levels became negative (undetectable or < 400 ng/mL) in approximately half of such patients at 4 to 12 months after infusion.

While limited data exist concerning the clinical significance of HAMA, detectable HAMA levels can alter the clearance and tissue biodistribution of MAbs. In patients who develop persistently elevated serum HAMA levels, the efficacy of diagnostic or therapeutic murine antibody-based agents may be compromised.

When considering the administration of OncoScint® CR/OV-In to patients who have previously received murine antibody-based products, physicians should be aware of the potential for HAMA to alter clearance and biodistribution. The quality or sensitivity of the imaging study may be compromised. Therefore, prior to administration of murine antibodies, including OncoScint® CR/OV-In, the physician should review the patient history to determine whether the patient has previously received such products.

Preliminary data are available from repeat-administration studies of OncoScint® CR/OV-In in 69 patients who have received 105 repeat doses. However, there are insuffi-

cient data to determine the safety and efficacy of this product after repeat administration (see ADVERSE REACTIONS).

Drug/Laboratory Test Interactions: The presence of HAMA in serum may interfere with two-site murine antibody-based immunoassays, including assays for carcinoembryonic antigen (CEA) and CA 125. When present, this interference generally results in falsely high values. If HAMA is known or suspected to be present, the clinical laboratory should be notified and appropriate measures taken to avoid this interference. These methods include the use of non-murine immunoassays, or HAMA removal by adsorption, blocking, or heat inactivation.

Carcinogenesis, Mutagenesis, Impairment of Fertility Long-term animal studies have not been performed to evaluate the carcinogenic or mutagenic potential of OncoScint® CR/OV-In or to evaluate its effect on fertility in males or females.

Pregnancy Category C Animal reproduction studies have not been conducted with OncoScint® CR/OV-In. It is also not known whether OncoScint® CR/OV-In can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. OncoScint® CR/OV-In should not be administered to a pregnant woman unless, in the opinion of the physician, the information to be gained outweighs the potential risks. MAB B72.3 has been shown to react with fetal gastrointestinal tissues.

In general, examinations using radiopharmaceuticals in women of childbearing potential should be performed during the first few days (approximately 10) following the onset of menses.

Nursing Mothers and/or Lactating Women It is not known whether OncoScint® CR/OV-In is excreted in human milk and, if so, for how long. Because many drugs are excreted in human milk, caution should be exercised when OncoScint® CR/OV-In is administered to a nursing woman. OncoScint® CR/OV-In has not been administered to lactating females and therefore should not be administered to nursing mothers unless, in the opinion of the physician, the information to be gained outweighs the potential risk. In such cases, formula feedings should be substituted for breast feedings.

Pediatric Use The safety and effectiveness of OncoScint® CR/OV-In in children have not been established.

ADVERSE REACTIONS

After administration of over 500 single i.v. doses of OncoScint® CR/OV-In (indium In 111 satumomab pendetide) in clinical trials, adverse reactions were observed in less than 4% of patients. No deaths attributable to OncoScint® CR/OV-In administration were reported. The most common adverse reaction was fever, which occurred in less than 2% of patients. Other adverse reactions, each of which occurred in less than 1% of patients, are listed in order of decreasing frequency: chills, hypotension, hypertension, rash, pruritus, sweating, nausea, arthralgia, asthenia, chest pain, headache, hypothermia, pain, flushing, diarrhea, confusion, dizziness, nervousness, crying, and angioedema. Although causality was not determined, an isolated occurrence of reversible thrombocytopenia was observed in a patient who received OncoScint® CR/OV-In in clinical trials.

Additional adverse reactions after 105 repeat administrations of OncoScint® CR/OV-In in 69 patients included two reports of fever, one complaint of abdominal pain, and two cases of non-serious, readily reversible reactions characterized primarily by flank pain.

OVERDOSAGE

The maximum amount of OncoScint® CR/OV-In (indium In 111 satumomab pendetide) that can be safely administered has not been determined. In clinical trials, single doses of 20 mg of OncoScint® CR/OV-In were administered to 64 patients with various types of epithelial carcinomas; the type and frequency of adverse reactions at this dose were similar to those observed with lower doses.

DOSE AND ADMINISTRATION

The dose of OncoScint® CR/OV (satumomab pendetide) is 1 mg radiolabeled with 5 mCi of indium In 111 chloride. Each dose is administered intravenously over 5 minutes and should not be mixed with any other medication during its administration. The patient dose of the radiolabel should be measured in a dose calibrator prior to administration.

Each OncoScint® CR/OV kit is a unit dose package. After radiolabeling with indium-111, the entire OncoScint® CR/OV-In dose should be administered to the patients. **Reducing the dose of either component may adversely impact imaging results, and is, therefore, not recommended.**

HOW SUPPLIED

The OncoScint® CR/OV kit (NDC No. 0044-0579-01) for the preparation of indium-111 labeled OncoScint® CR/OV includes one vial containing 1 mg of satumomab pendetide per 2 mL of sodium phosphate buffered saline and one 2 mL vial of sodium acetate buffer solution, 0.5 M. These solutions are sterile and pyrogen free and contain no preservative. Each kit also includes one sterile 0.22 µm Millex® GV filter, prescribing information, and two identification labels.

U.S. Patent Nos. 4,671,958 and 4,741,900

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Revised 12/30/92

References 1. Nabi HA, Doerr RJ. Radiolabeled monoclonal antibody imaging (immunoscintigraphy) of colorectal cancers: current status and future perspectives. *Am J Surg*. 1992;163:448-456. 2. Data on file. CytoGen Corporation, Princeton, NJ. 3. Doerr RJ, Abdel-Nabi H, Krag D, et al. Radiolabeled antibody imaging in the management of colorectal cancer: results of a multicenter clinical study. *Ann Surg*. 1991;118-124. 4. Collier RJ, Abdel-Nabi H, Doerr RJ, et al. Immunoscintigraphy with ¹¹¹In-CYT-103 in the management of colorectal carcinoma: a comparison with computed tomography. *Radiology*. 1992;185:179-186. 5. Surwit EA, Childers JM, Krag DN, et al. Clinical assessment of ¹¹¹In-CYT-103 immunoscintigraphy in ovarian cancer. *Gynecol Oncol*. 1993;48:285-292.

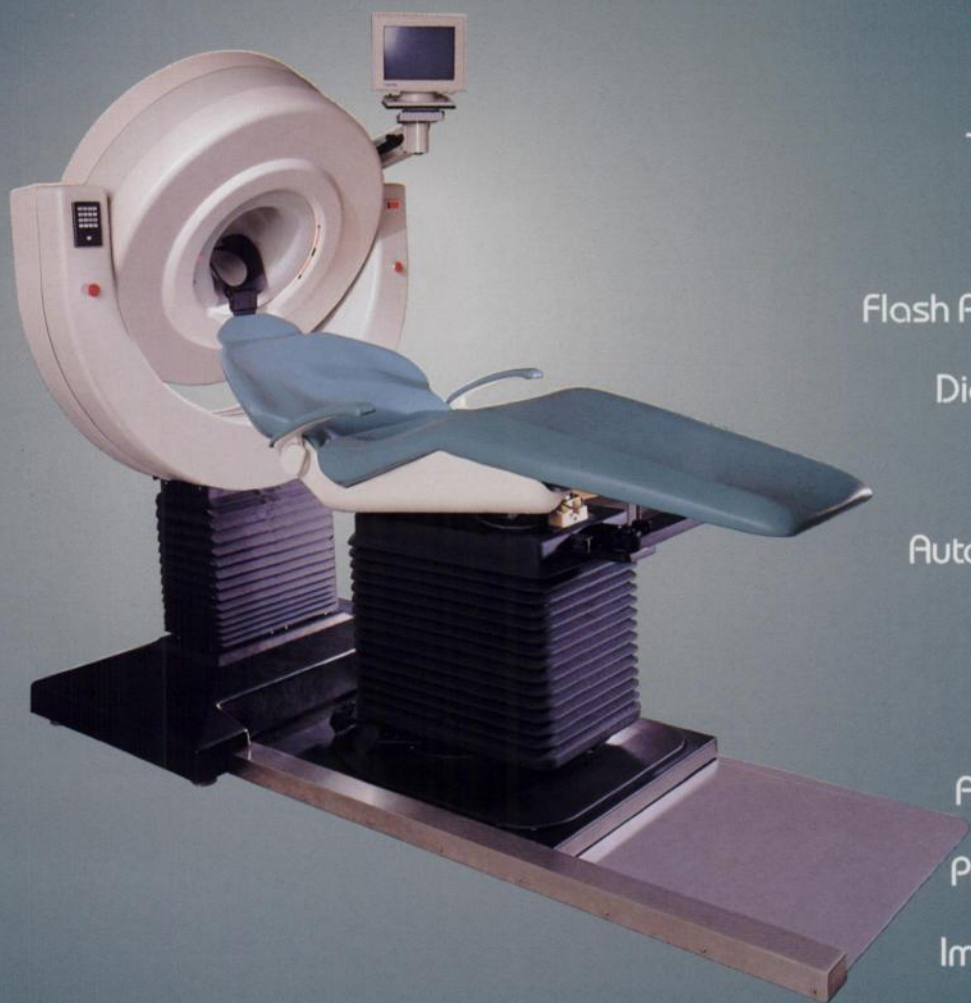
To assist decision making
in the management of patients with colorectal
or recurrent ovarian cancer...

ONCOSCINT® CR/OV
Satumomab Pendetide (1mg/2mL)

Effective in determining
both the location and extent
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Please refer to complete prescribing information before using OncoScint CR/OV.

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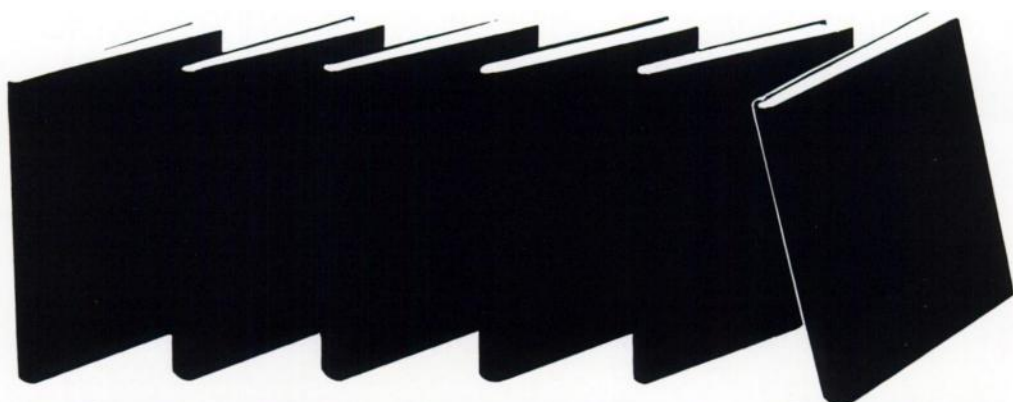
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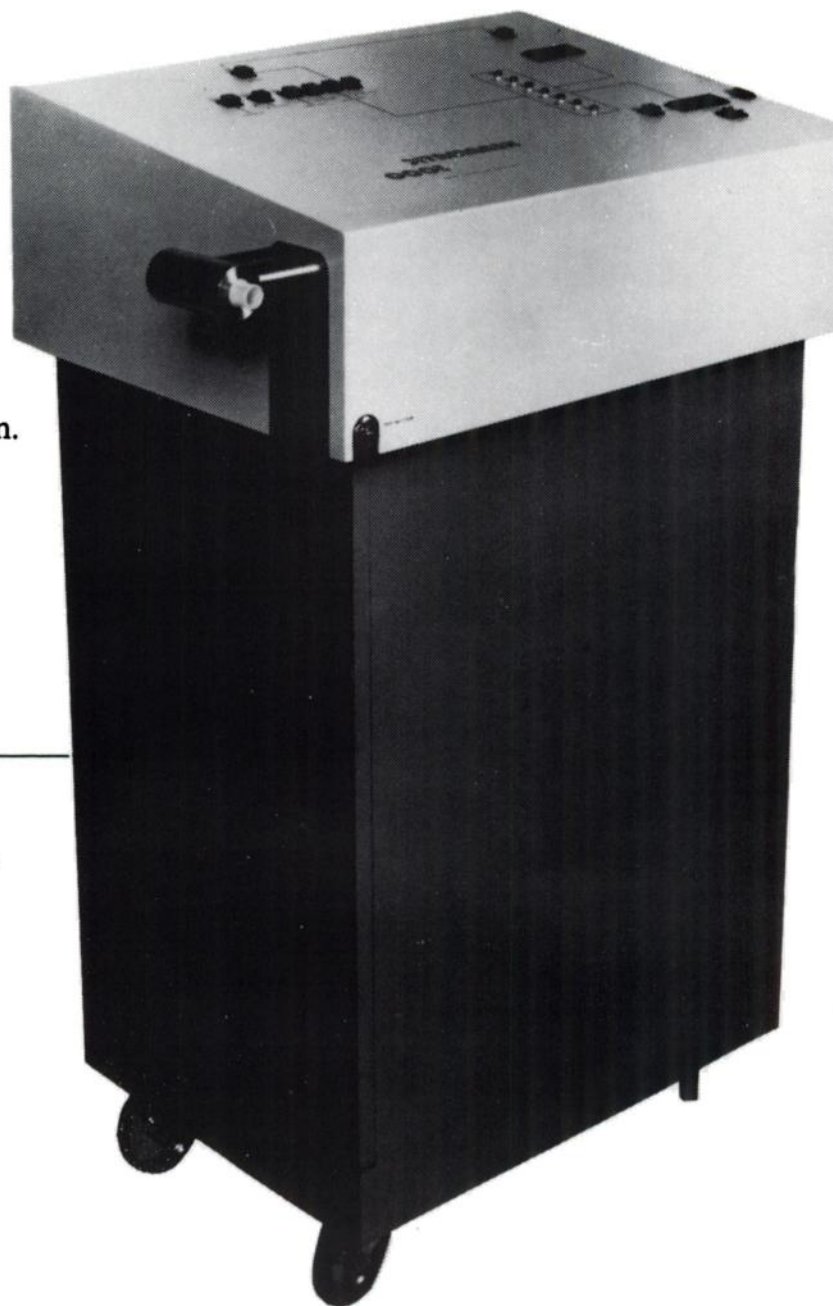
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Department of Radiology
Section of Nuclear Medicine



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This program is sponsored by the Medical College of Wisconsin.

TUITION:

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CREDIT:

The Medical College of Wisconsin is accredited by the Accreditation Council for Continuing Medical Education to sponsor continuing medical education for physicians.

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Funding Announcements: June 6, 1994
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Funding Availability: July 1, 1994

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Refresher and state-of-the art continuing education courses in chemistry, physics, quality assurance, cardiovascular nuclear medicine, PET, SPECT and NMR will supply up-to-the-minute approaches and procedures for all clinical settings.

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An extensive display of scientific posters and exhibits will augment the presentation.

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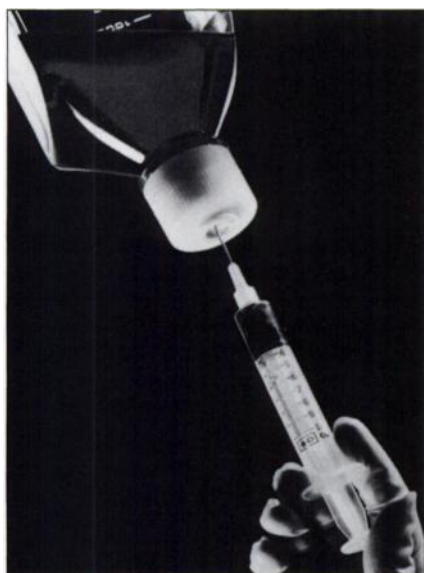


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Each description of the products below was condensed from information supplied by the manufacturer. The reviews are published as a service to the professionals working in the field of nuclear medicine and their inclusion herein does not in any way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

Septum Closure



A unique closure system suitable for use with any bottle or container with a 38-430 neck has been introduced by Nalge Company. The Nalgene autoclavable septum closure features a thermoplastic elastomer septum for aseptic injection of reagents or sample withdrawal without compromising the sterility or integrity of the container's contents. Nalgene also manufactures suitable containers for use with the new septum closure such as the Nalgene culture vessel and culture vessel mixing system, media bottles and other square bottles. The closure is designed for use with an 18-gauge or smaller needle and is for laboratory use only (not for in vitro diagnosis or parenterals). **Nalge Company, P.O. Box 20365, Rochester, NY 14602. (716) 264-3985.**

Gridded Ionization Chamber

Ordela, Inc. has introduced the model 8210A gridded ionization chamber for easier screening of soil, water, air filter and smear samples by rapid, direct alpha spectrometry. Sample preparation is easy, quick and does not

require toxic, labor-intensive radiochemistry or special laboratory equipment. Soil samples are finely ground, mixed with solution, sprayed onto 10" diameter stainless steel planchets and dried under heat lamps. The sample is then placed in the sliding tray assembly and inserted into the chamber via a sealed, horizontal access port. Alpha resolutions of 50 keV FWHM are possible for ^{230}Th in thin samples and 50-65-keV resolutions FWHM are possible for alpha energies from ^{230}Th to ^{24}Cm . Water samples of 5-15 ml can be directly sprayed onto the planchets and counted. Air filter samples and smears up to 10" in diameter can be directly counted with no sample preparation and without introducing inconsistencies due to sample trimming. The gridded ionization chamber is ruggedly constructed of mild steel and is completely nickel-plated for low-background environmental counting and ease of decontamination. The grid is specially fabricated for ease of replacement or decontamination. **Ordela, Inc., 1009 Alvin Weinberg Dr., Oak Ridge, TN 37830. (615) 483-8675.**

Nonlatex Gloves

A new nonlatex, nonvinyl examination and surgical glove has been introduced by SmartPractice. The new gloves are called Tactyl 1 and are made from Tactylon™, a new synthetic material that provides protection against latex allergy reactions. Tactyl 1 gloves offer all the benefits of natural latex elasticity, tactility and barrier protection along with the

added value of problem-free wear for latex-sensitive individuals and contain none of the chemical irritants and latex proteins that are the source of most adverse skin reactions associated with latex. These latex-free gloves resist deterioration caused by oxidation, light and ozone exposure and provide enhanced reliability against tearing, cracking and pinpoint holes. Tactyl 1 gloves are available in sizes ranging from 5 1/2 to 9. **SmartPractice, 3400 East McDowell, Phoenix, AZ 85008-7899. (602) 225-9090.**

Infrared Laser Imaging Film

Agfa has introduced a new high-speed laser printer film called Scopix LT-IR. The new film is designed for optimum performance in infrared laser imagers and provides excellent hard copy results for CT scans, MR images, ultrasound and other nuclear medicine modalities. The film is available in two versions; LT-IRB, coated on a blue-tinted base, and LT-IRC, coated on a clear base. These films feature Agfa's SEL (split emulsion layer) technology which incorporates two separate emulsion layers. The first layer provides low contrast in the toe of the H & D curve while minimizing cross-scan line visualization caused by laser writing mechanics. The second layer ensures high shoulder contrast and high D max for crisp images and ultra-sharp alphanumerics. **AGFA, Miles, Inc., 100 Challenger Rd., Ridgefield Park, NJ 07660-2199. (201) 440-2500.**

Macintosh-based Thyroid Uptake System



The first Mac-based thyroid uptake system has been introduced by Biodex Medical Systems, Inc. The Atomlab 950, which was officially unveiled at the Society of Nuclear Medicine's 40th Annual Meeting in Toronto in June, provides a full display screen, point and click pull-down menus and comprehensive patient management capabilities. The Atomlab 950 allows complete, unlimited patient and test data stor-

age and retrieval. Reports are easily generated to include all data, thus eliminating the need for a manual report to supplement the standard test result printout. With 4 MB of RAM and an 80-MB hard drive, the Macintosh LC III can enhance the overall capabilities of the nuclear medicine department by adding word processing, spreadsheet and database programs. Compliance is simple with the improved Wipe Test mode. Various wipe sites can be individually identified, tested and stored to create a single compliance report for multiple wipes. In addition to thyroid uptake, thyroid bioassay, wipe tests, Schilling tests and administration/QA, the new system offers in vitro tests including blood volume and red cell survival. Spectrum analysis is displayed during acquisition with detailed spectrum analysis tools to manipulate channels and values. **Biodex Medical Systems, Inc., Brookhaven R&D Plaza, P.O. Box 702, Shirley, NY 11967-0702. 1-800-224-6339.**

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The 1994 examination will be given Saturday, June 4, 1994, in Orlando, Florida, in conjunction with the 41st Annual Meeting of The Society of Nuclear Medicine.

The examination is written and consists of two parts. Part 1 (approximately 3.5 hr) assesses knowledge of basic aspects of Nuclear Medicine Science. Part 2 (approximately 2.5 hr) examines in depth the knowledge of a predetermined subspecialty area of the candidate's choice including:

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The examination fee is \$450 (\$400 refundable if you do not qualify).

For applications and more information please contact:

**Christine Santos, Associate Coordinator
American Board of Science in Nuclear Medicine
The Society of Nuclear Medicine
Department of Meetings Services
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(212) 889-0717 Fax: (212) 545-0221.**

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MEDICAL DIRECTOR, NUCLEAR MEDICINE. Prince George's Hospital Center, a 450-bed hospital located in Cheverly, Maryland, a suburb of Washington, DC, is seeking a Medical Director for its Nuclear Medicine Department. Candidates for the position will have completed a two-year fellowship in Nuclear Medicine and be board certified by the American College of Nuclear Medicine, Radiology, or Pathology. A minimum of two

years clinical experience post-residency is required and administrative/managerial experience is preferred. Candidates should submit their resume and letter of interest by January 31, 1994 to: Faranak Sotoudeh, MD, Chairperson, Nuclear Medicine Search Committee, Prince George's Hospital Center, 3001 Hospital Dr., Cheverly, Maryland 20785. Candidates should direct any questions to: Louis Morgan, PhD, CNMT, Administrative/Technical Director, Nuclear Medicine Department, (301) 618-2283.

NUCLEAR MEDICINE PHYSICIAN. Northern California—The Kaiser Permanente Medical Center in Santa Clara, CA is seeking a BC/BE Nuclear Medicine physician for a career opportunity with the nation's leading HMO. All aspects of nuclear medicine services are provided to a 250,000 member prepaid patient population. The ideal candidate should have experience in clinical management of thyroid disease and the performance of treadmill tests. Board certification/eligibility in internal medicine is preferred. Academic opportunities are available. Competitive salary, generous benefits and a comprehensive retirement program. For more information, send CV with cover letter to: Diane Butler, The Permanente Medical Group, Inc., Physician Recruitment, 1814 Franklin, 4th Floor, Oakland, CA 94612. EOE.

Radiation Safety

VA Medical Center, San Francisco, CA. The Medical Center holds a NRC License of broad scope, performs both diagnostic and therapeutic nuclear medicine, and has one of the largest research programs in the VA system with approximately 45 biomedical research laboratories and 400 users of radioactivity. Candidates must have at least a year's full-time experience in a medical radiation safety program, meet the RSO qualifications in 10 CFR 35, and be familiar with U.S. NRC and DOT regulations. An advanced degree, especially a doctorate, in a relevant field and certification by the ABR, ABMP or ABSNM are desirable. Salary is dependent upon qualifications. The VA is an equal opportunity employer. Selectee may be subject to drug testing. Please send your CV and references to: Ralph R. Cavalieri, M.D., Chief, Nuclear Medicine Service (115), VA Medical Center, 4150 Clement Street, San Francisco, CA 94121.

Senior Radiochemist

The Division of Nuclear Medicine at the Hospital of the University of Pennsylvania is seeking a senior radiochemist for its radiochemistry laboratory. The Division of Nuclear Medicine has strong research and clinical programs in both

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Technical Coordinator

The Department of Radiology at the Columbia-Presbyterian Medical Center in New York is searching for a full-time Technical Coordinator for our new PET facility. The successful candidate will function as a key member of our PET team and will have various responsibilities. Requirements include a Bachelor's degree, Master's preferred; AART (N) and/or CNMT registration or comparable background required. Strong background in computer science (SUN/UNIX) preferred. Familiarity with NYC BRC regulations, medical health physics, and medical use of radiopharmaceuticals preferred. Previous patient imaging and PET imaging experience desired. Excellent communication skills required. Salary commensurate with experience. Please send resume to: Ms. Joan Massler, Columbia-Presbyterian Medical Center, Department of Nuclear Medicine, Milstein Building 3-224, 177 Fort Washington Avenue, NY, NY 10032. Columbia-Presbyterian Medical Center is an affirmative action/equal opportunity employer.

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NUCLEAR MEDICINE PHYSICIAN, ABNM-certified. Ample experience in basic research, academics, government, and private practice. Solid background in all areas of nuclear medicine. Seeking position, preferably private practice, but will consider offers in other settings, including corporate appointments. Call 301-390-0984 or write: P.O. Box 101, The Society of Nuclear Medicine, 136 Madison Ave., New York, NY 10016.

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West Virginia University is located in Morgantown, West Virginia, a pleasant university community near many major urban centers. The environment is ideal for raising a family. West Virginia University is an affirmative action/equal opportunity employer. Women and minority candidates are strongly encouraged to apply. Send letters of interest, CV and three references to: Howard H. Kaufman, MD, Professor and Chairman, Department of Neurosurgery, PO Box 9183, Health Sciences Center, West Virginia University, Morgantown, WV 26506.

Review of applicants will begin February 1, 1994, and will continue until the position is filled.

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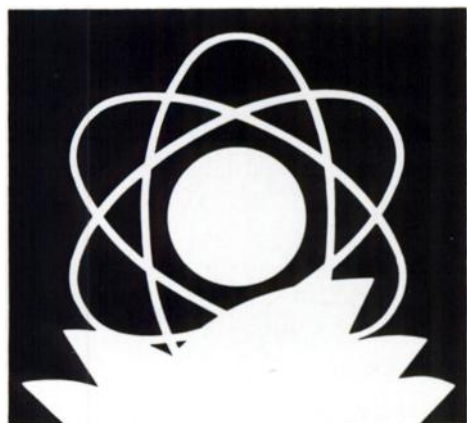
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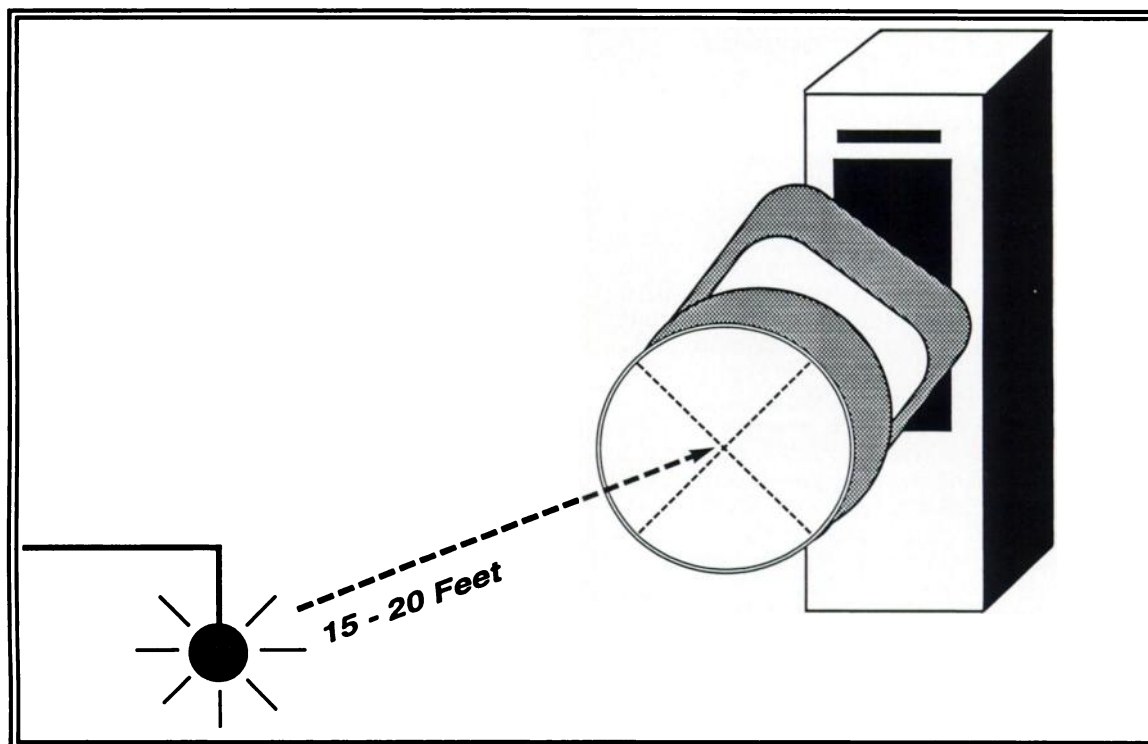
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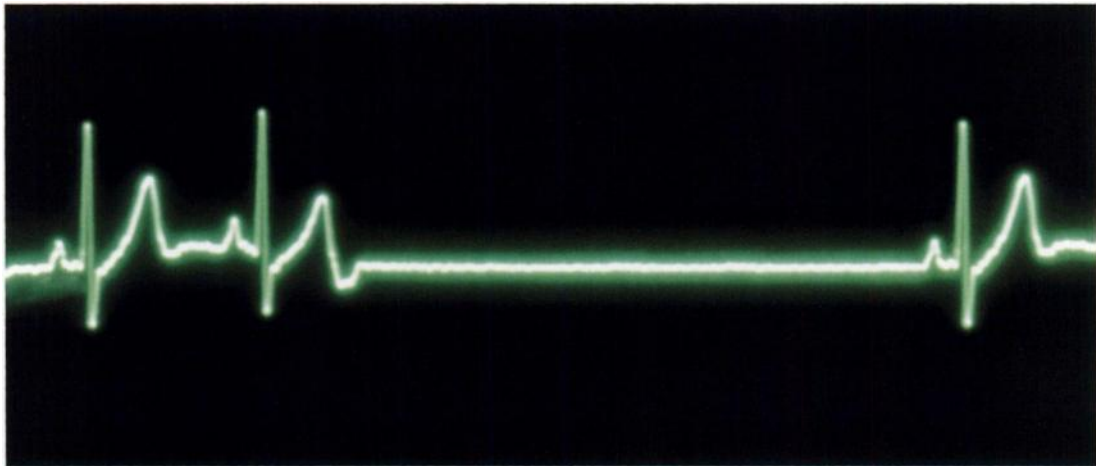
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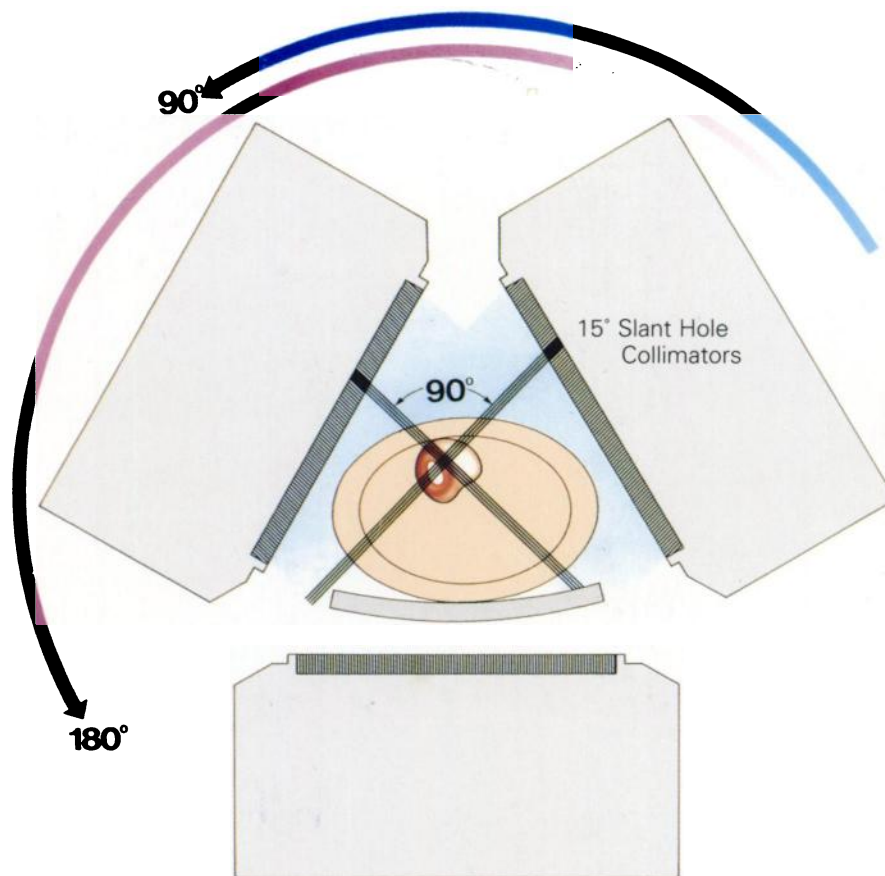
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