American Association for the Advancement of Science



3 JANUARY 1992 Vol. 255 **B** Pages 1–132 \$6.00

Introducina

Stratagene's Eagle Eye" Still Video System

IMAGE INTEGRATION DEVICE for exposure time control

IMAGE STORAGE DEVICE for high resolution data storage

THERMAL PRINTER for publication quality documentation

CUSTOMIZED VIDEO CAMERA for image capture

ZOOM LENS for ease of use and maximum resolution

INTERFERENCE FILTER for high signal to noise ratio



At a fraction of the cost of instant photography



RECORDED STILL VIDEO MAGING Vs. CONVENTIONAL PHOTOGRAPHY

A: Still video image captured at F/stop 1.2 and integration of 10 frames (0.33 second). B: Photo using Polaroid Type 667 film (ISO 3000) exposed for 1 second at F/stop 8

50 ng of lambda/Hind III size markers along with various dilutions of pUC 18 supercoiled DNA were loaded on a 5 mm thick 0.8% agarose, 1X TAE slab gel and electrophoresed. The gel was stained in 0.5 mg/ml ethidium bromide for 30 minutes, then destained for 45 minutes in deionized water. All images were generated using the same transilluminator (302 nm). Lane 1: lambda/Hind III markers; Lane 2: 4 ng pUC 18 DNA; Lane 3: 2 ng pUC 18 DNA; Lane 4: 1 ng pUC 18 DNA; Lane 5: 0.5 ng pUC 18 DNA; Lane 6: 0.25 ng pUC 18 DNA.

S tratagene's advanced Eagle Eye[™] still video system uses state-of-the-art still video imaging technology to enhance the sensitivity, increase the speed, and reduce the cost associated with gel documentation. The Eagle Eye still video system increases the limit of detection of DNA to 0.25 ng, a four-fold improvement over conventional instant photography. The system's high resolution quick focus



zoom lens eliminates the tedious chore of adjusting the camera focal length associated with all other systems. Publication quality thermal prints are produced in seconds at a fraction of the cost of instant photography, and additional prints are obtained by the push of a button. This system is also ideal for the documentation of protein gels, TLC plates and more. Using the optional still video storage device, images can be stored on still video floppy disks for archiving and processing. Dimensions: $41"(H) \times 16"(D) \times 26"(W)$.

Please Contact Stratagene for a Distributor Near You.

Corporate Headquarters: 800-424-5444 Fax: 619-535-5430 Telex: 9103809841 *Germany:* Stratagene GmbH Telephone: (06221) 40 06 34 Telefax: (06221) 40 06 39 United Kingdom: Stratagene Ltd. Telephone: (0223) 42 09 55 Telefax: (0223) 42 02 34 Telex: 81417 INNCEN G

Circle No. 40 on Readers' Service Card

Quantikine

For the precise Quantitation of Cytokines in fluids...

CYTOKINE

The Tool to Accurately Measure Cytokines

Immunoassay vs. Bioassay

The usual assay for a specific cytokine is based on its ability to affect the growth or differentiation of an actively growing, indicator cell line.

However, since most of these indicator cell lines can respond to many growth or inhibitory stimuli (including other cytokines), precise detection and quantitation of a specific cytokine in such complex fluids as: cell culture medium, serum, plasma, urine and synovial fluid (see *e.g.* J. of IMMUNO. **145**:8, 1990 pp 2514-2519) proves difficult and often imprecise.

Now with the Quantikine[™] series of "sandwich" immunoassay, a precise, accurate quantitation of a given human cytokine can easily be made in a variety

of complex biological fluids. Each Quantikine kit has been

formulated under exacting standards to ensure that it is:

Sensitive -detects less than 10 pg/ml.

Rapid -results in less than 4.5 hours.

Specific -unaffected by the presence of other cytokines.

Calibrated -to WHO (National Institute for Biological Standards and Control) interim reference standards.

FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES.



IRPC

To place an order or request product information, call us at 1-800-328-2400.

In Europe contact:

British Bio-technology, Ltd. 4-10 The Quadrant, Barton Lane Abingdon, Oxon OX14 3YS Telephone: 0865 781045 3792956 Fax: 0235 53342 Telex: 838063 BIOTEC G In Japan contact:

Funakoshi Co., Ltd. 9-7, Hongo 2-Chome Bunkyo-ku, Tokyo 113 Telephone: 81 3 5684 1616 Fax: 81 3 5684 1633 Telex: J28489 FUNA

Your Source for Cytokine Reagents



GM-CSF

IL-7

nal Detectable Dose

Minimal Detectable Dose.

R&D Systems 614 McKinley Place N.E. Minneapolis, MN 55413 In Minnesota: (612) Fax:(612)379-6580 Telex: 750627

4.5 pg/ml

Circle No. 82 on Readers' Service Card

American Association for the Advancement of Science

4



ISSN 0036-8075 3 JANUARY 1992 VOLUME 255 NUMBER 5040

	7	This Week in Science
Editorial	9	Achievable New Year's Resolutions
Letters	10	Science and the Press: J. E. BISHOP; D. E. KOSHLAND, JR. ■ Coverage of the "Gallo Case": J. CREWDSON; J. COHEN
ScienceScope	19	1992 preview: The year in funding, big projects, and science policy
News & Comment	20	Conservation Biology in the Fast Lane
	22	Soviet Environment Slips Down the Agenda
	24	Can There Be a Better Grade of "Pork?"
	25	Plant Biotechnology Explored in Indianapolis
	26	NRC Faults Science Behind Ozone Regs
	27	Briefings: Twinkle Twinkle Little LED ■ Where HUGOing? ■ Invertebrates Need Love Too ■ Open Freezer at NCI
Research News	28	Microbes From 20,000 Feet Under the Sea Superbugs in Waiting: Some Cautionary Tales
	30	A Fall Harvest of Earth Science in San Francisco: Loma Prieta's Long Reach Was a Matter of Mirrors Are Earthquakes a Ticking Clock for Los Angeles? A Conundrum at Steens Mountain
:	32	Catching the Rhythm of The Bacterial Twist
:	33	Twin Study Links Genes to Homosexuality
	34	Anything Goes at the Cell Biology Meeting: New Evidence Found for a Nuclear Matrix ■ Fruit Fly Learning Research Mushrooms ■ New Clues to How Bacteria Get Into Cells
Perspective	39	Mechanisms of Transcriptional Timing in Drosophila: C. S. THUMMEL
Articles	41	America's Children: Economic Perspectives and Policy Options: V. R. FUCHS AND D. M. REKLIS
	46	Fermi Surfaces, Fermi Liquids, and High-Temperature Superconductors: W. E. PICKETT, H. KRAKAUER, R. E. COHEN, D. J. SINGH
	54	Molecular Code for Cooperativity in Hemoglobin: G. K. ACKERS, M. L. DOYLE,

54 Molecular Code for Cooperativity in Hemoglobin: G. K. Ackers, M. L. Doyle, D. Myers, M. A. DAUGHERTY

SCIENCE (ISSN 0036-8075) is published weekly on Friday, except the last week in December, by the American Association for the Advancement of Science, 1333 H Street, NW, Washington, DC 20005. Second-class postage (publication No. 484460) paid at Washington, DC, and additional mailing offices. Copyright © 1991 by the American Association for the Advancement of Science. The title SCIENCE is a registered trademark of the AAAS. Domestic individual membership and subscription (51 issues): \$87 (\$47 allocated to subscription.) Domestic institutional subscription (51 issues): \$195. Foreign postage extra: Mexico, Caribbean (surface mail) \$50; Other countries (air assist delivery) \$95. First class, airmail, student and emeritus rates on request. Canadian rates with GST available upon request, GST #1254 88122. Change of address: allow 6 weeks, giving old and new addresses and 11-digit account number. Postmaster: Send change of address to Science, P.O. Box 2033, Marion, OH 43305–2033. Single copy sales: \$6.00 per issue prepaid includes surface postage; Guide to Biotechnology Products and Instruments, \$20. Bulk rates on request. Authorization to photocopy material for internal or personal use under circumstances not falling within the fair use provisions of the Copyright Act is granted by AAAS to libraries and other users registered with the Copyright Clearance Center (CCC) Transactional Reporting Service, provided that the base fee of \$1 per copy plus \$0.10 per page is paid directly to CCC, 27 Congress Street, Salem, Massachusetts 01970. The identification code for Science is 0036-8075/83 \$1 + .10. Science is indexed in the Reader's Guide to Periodical Literature and in several specialized indexes.

The American Association for the Advancement of Science was founded in 1848 and incorporated in 1874. Its objectives are to further the work of scientists, to facilitate cooperation among them, to foster scientific freedom and responsibility, to improve the effectiveness of science in the promotion of human welfare, to advance education in science, and to increase public understanding and appreciation of the importance and promise of the methods of science in human progress.



COVER The calculated surfaces in momentum space ("Fermi surfaces") for the charge carriers in the high-temperature superconductor YBa₂Cu₃O₇. Charge carriers can be electron-like or hole-like; blue indicates low-velocity, high-mass carriers, and red indicates high-velocity, low-mass carriers. This Fermi surface, first calculated theoretically, has recently been confirmed by several experimental spectroscopies. See page 46. [Image by R. E. Cohen with AVS 3.0 software; image printed on a Kodak XL7700 at the Naval Research Laboratory Connection Machine Facility]

RA	m	nrte	
		orts	

64	Orientational Ordering of Polymers by Atomic Force Microscope Tip-Surface Interaction: O. M. LEUNG AND M. C. GOH
66	Luminescent Colloidal Silicon Suspensions from Porous Silicon: J. L. HEINRICH, C. L. CURTIS, G. M. CREDO, K. L. KAVANAGH, M. J. SAILOR
68	Strontium Isotopic Composition of Estuarine Sediments as Paleosalinity- Paleoclimate Indicator: B. L. INGRAM AND D. SLOAN

- 72 Production of Isotopic Variability in Continental Basalts by Cryptic Crustal Contamination: A. F. GLAZNER AND G. L. FARMER
- 74 The Antiquity of Oxygenic Photosynthesis: Evidence from Stromatolites in Sulphate-Deficient Archaean Lakes: R. BUICK
- 77 Interspecific Brood Parasitism in Blackbirds (Icterinae): A Phylogenetic Perspective: S. M. LANYON
- 79 Activation of T Cells by a Tyrosine Kinase Activation Domain in the Cytoplasmic Tail of CD3 ε: F. LETOURNEUR AND R. D. KLAUSNER
- 82 Torsional Rigidity of Positively and Negatively Supercoiled DNA: P. R. SELVIN, D. N. COOK, N. G. PON, W. R. BAUER, M. P. KLEIN, J. E. HEARST
- 85 Interaction of p107 with Cyclin A Independent of Complex Formation with Viral Oncoproteins: M. E. EWEN, B. FAHA, E. HARLOW, D. M. LIVINGSTON

Technical Comment

Book Reviews

Products & Materials

Oncoproteins: M. E. EWEN, B. FAHA, E. HARLOW, D. M. LIVINGSTON
87 Interaction Between Human Cyclin A and Adenovirus E1A-Associated p107 Protein: B. FAHA, M. E. EWEN, L.-H. TSAI, D. M. LIVINGSTON, E. HARLOW
90 The Updating of the Representation of Visual Space in Parietal Cortex by Intended Eye Movements: J.-R. DUHAMEL, C. L. COLBY, M. E. GOLDBERG
92 Enumerating Buckminsterfullerane Isomers: G. THIMM AND W. E. KLEE; M. SAUNDERS
93 Taking Society's Measure, *reviewed by* D. L. SILLS A Physicist on Madison Avenue; Science à la Mode, J. SILK The Molecular Biology of Plastids, C. W. BIRKY, JR. Prices of Books Books Received

 97 Clampless Filtration Manifold System ■ Automated Capillary Zone Electrophoresis System ■ DNA Sequencing Gel Accessory ■ Three-Dimensional Rendering Software ■ Grant Accounting Software ■ Fast Blot-Developer System ■ Literature

Information to Contributors is found on pages 36 to 38.

Board of Directors Donald N. Langenberg

Retiring President, Chairman

Leon M. Lederman President

F. Sherwood Rowland President-elect Mary Ellen Avery Francisco J. Ayala Eugene H. Cota-Robles Robert A. Frosch Joseph G. Gavin, Jr. Florence P. Haseltine Jean'ne M. Shreeve Warren M. Washington William T. Golden

Treasurer Richard S. Nicholson Executive Officer Editorial Board Charles J. Arntzen Elizabeth E. Bailey David Baltimore William F. Brinkman E. Margaret Burbidge Pierre-Gilles de Gennes Joseph L. Goldstein Mary L. Good Harry B. Gray John J. Hopfield F. Clark Howell Paul A. Marks Yasutomi Nishizuka Helen M. Ranney Robert M. Solow Edward C. Stone James D. Watson

Board of Reviewing Editors

John Abelson Frederick W. Alt Don L. Anderson Stephen J. Benkovic David E. Bloom Henry R. Bourne James J. Bull Kathryn Calame Charles R. Cantor C. Thomas Caskey Dennis W. Choi Ralph J. Cicerone John M. Coffin Bruce F. Eldridge Paul T. Englund Fredric S. Fay

Douglas T. Fearon Harry A. Fozzard Victor R. Fuchs Theodore H. Geballe Margaret J. Geller Roger I. M. Glass Stephen P. Goff Corey S. Goodman Stephen J. Gould Ira Herskowitz Eric F. Johnson Stephen M. Kosslyn Konrad B. Krauskopf Charles S. Levings III Harvey F. Lodish Richard Losick Anthony R. Means Mortimer Mishkin Roger A. Nicoll William H. Orme-Johnson III Stuart L. Pimm Yeshayau Pocker Dennis A. Powers Ralph S. Quatrano Erkki Ruoslahti Thomas W. Schoener Ronald H. Schwartz Terrence J. Sejnowski Thomas A. Steitz Richard F. Thompson Robert T. N. Tiian Emil R. Unanue Geerat J. Vermeij Bert Vogelstein Harold Weintraub Zena Werb George M. Whitesides Owen N. Witte William B. Wood Keith Yamamoto

3 JANUARY 1992

TABLE OF CONTENTS 5

EUKARYOTIC TRANSCRIPTION

We make the grade like no one else.

HeLaScribe" Nuclear Extract in vitro Transcription Grade

- First commercially available mammalian nuclear extract qualified for transcription
- Highly active for specific in vitro transcription
- Accurate initiation for promoter analysis
- Successfully tested on CMV, Ad2MLP, HTFR, and DHFR promoters*

CMV = Cytomegalovirus Immediate Early gene; Ad2MLP = Adenovirus 2 Major Late Promoter; HTFR = Human Transferrin Receptor gene; DHFR = Dihydrofolate Reductase gene.

Available as a complete transcription system or as a stand-alone extract

HeLaScribe[™] Nuclear Extract is another exciting addition to Promega's expanding line of products for Eukaryotic Transcription Regulation which already includes:

- Transcription Extracts
- Transcription Factors
- Gel Shift Assay Systems
- Primer Extension System
- Footprinting Systems

... by Promega.



800-356-9526

To find out more about HeLaScribe Nuclear Extract – *in vitro* Transcription Grade, please call us toll free to request Technical Bulletin 123. For information on our line of Eukaryotic Transcription products, please request the Eukaryotic Transcription folder.

O Technical Builettic

Circle No. 83 on Readers' Service Card



America's children

merican children appear to be worse off than their parents were as children, both materially and culturally. During the 1980s, material conditions of children of lower income parents deteriorated the most, according to census data. Policy discussions have focused either on government mandates on employers to provide benefits to parents or on direct government transfers. Fuchs and Reklis (p. 41) point out that employer-mandated programs would not benefit most poor children, who live in households that would be unaffected by the programs. Paid parental leave, for example, would provide the greatest dollar benefits to children with the highest paid parents. The authors argue that transfer of benefits from households without children to those with them, such as through tax credits, will be required.

Fermi liquid

The high-temperature oxide superconductors are metallic phases at room temperature; Pickett *et al.* (p. 46; cover) argue that recent experimental results show that the normal state is a Fermi liquid, or a band metal, rather than a doped insulator. They discuss some features of the normal state that still need to be resolved, such as low-energy excitations, as well as how these results constrain theoretical explanations of the superconducting state.

Hemoglobin code

etrameric hemoglobin is the classical model for understanding allosteric regulation. Ackers *et al.* (p. 54) review recent experimental data that reveal a previously unrecognized symmetry feature in the switching between the tense and the relaxed states. Switching occurs whenever binding of a molecule at the heme site forms a tetramer with at least one ligand-binding subunit on each half of the molecule.

Orienting polymers

Periodic nanometer-scale patterns can be induced in polystyrene films with the atomic force microscope (AFM). The AFM is normally used to image surfaces by recording vertical displacements of a softly sprung tip as it is dragged over a surface. Leung and Goh (p. 64) used relatively large tip forces (10^{-7} newtons) to increase the interaction between the tip and the surface. Repeated scans produced periodic furrowed patterns in the direction perpendicular to the scanning.

Glowing silicon

uminescent silicon, made by etching silicon wafers to a high porosity, can be dispersed by ultrasound to produce a colloid, which in turn can be put into a polymer matrix to create luminescent films. Silicon does not normally luminesce, but etching silicon in hydrofluoric acid can produce a porous network of small structures that emits red-orange to yellow light after ultraviolet irradiation. The emission presumably arises from quantumconfinement effects. Because porous silicon is itself rather fragile, the preparation of colloids and films by Heinrich et al. (p. 66) may provide more convenient options for exploiting its properties.

Brood parasitism

Theories about the coevolution of brood parasite species that use another species, or host species, to raise its young, suggest that a generalist strategy, in which many species are hosts, would be used at first, and that specifically targeting one host species would evolve later. Not so for cowbirds, a brood parasite that uses other blackbirds to raise its young. Lanyon (p. 77) studied an 852-base pair region of the mitochondrial cytochrome b gene of six cowbird species and 20 other blackbirds. The North American cowbird, *Molothrus alter*, which para-

Edited by Phillip D. Szuromi

sitizes more than 200 host species, appears to be the youngest taxon of the group studied. The use of many hosts may reduce the probability of their developing effective defenses against parasitism.

Cell cycle complexes

omplexes containing several regulatory proteins are implicated in the control of the cell cycle. The oncoprotein encoded by the adenovirus early region 1A (E1A) interacts with the retinoblastoma protein (pRB) (a tumor suppressor) and p107, a cellular protein that is similar to pRB. Also associated with E1A are p33^{cdk2} and cyclin A, two proteins that interact to form a protein kinase that is likely to function in regulating progression through the cell cycle. Faha et al. (p. 87) show that p107 associates with cvclin A in the absence of E1A or pRB. Ewen et al. (p. 85) report that although p107 and pRB bind to similar sets of cellular proteins, cyclin A interacts only with p107. Cyclin A binding requires a discrete sequence of p107 distinct from the domains shared with pRB at which E1A and other proteins appear to bind.

Intended movements

ovement of the eye shifts the position of images of station-Lary objects on the retina; Duhamel *et al.* (p. 90) show that single neurons in the parietal cortex of the monkey can use information about intended movements of the eye to maintain the accuracy and continuity of visual representations. The receptive field of some parietal neurons shifted before an eye movement occurred, and almost all of the parietal neurons responded when eye movement brought the site of a previously used stimulus into the receptive field, even though the stimulus itself was absent. The shift in the receptive field updates the coordinates of remembered stimuli to maintain a dynamic link between successive retinal images.

When we introduced a complete line of factors and cytokines, the response was overwhelming.

Improved growth. Improved function. Cells respond to Boehringer Mannheim.

The ultimate goal: reproduce *in vivo* cell growth and function, *in vitro*. Today, anything that helps attain

that goal is in demand.

In response, we've introduced a complete line of natural and recombinant factors and cytokines. A line your cells respond to with improved growth and function.

Circle No. 51 on Readers' Service Card

BOEHRINGER

CORPORATION

And Boehringer Mannheim factors and cytokines respond to *your* needs, too.

They're function-tested to assure performance.

They're easy to use with convenient pack sizes and complete instructions. And they're control-dated to

guarantee activity.

You get Boehringer Mannheim quality. Backed by technical support and service. All available in a rapidlygrowing line of factors and cytokines.

Boehringer Mannheim Corporation Biochemical Products 9115 Hague Road P.O. Box 50414 Indianapolis, IN 46250-0414 Orders: 800 262 1640 Technical Service: 800 428 5433

Respond now for your free catalog.

Take the first step today. Get your free Cell Culture catalog. Call our Technical Services Department at **800-428-5433.**

For research use on

"...simplify, simplify."

– Henry David Thoreau

Let BIOSIS' Life Science Network simplify your search for information.

Searching for life science information used to be anything but easy. Then came the *Life Science Network*. Offering you access to nearly 80 recognized life science databases, the *Life Science Network* also provides something equally valuable:

Simplicity.

How? An easy-to-use, menu-driven system does away with the need to learn separate, complicated search languages or database names. Just enter your search terms, and the *Life Science Network* does the rest – it even scans several relevant databases at once!

To top it all off, the *Life Science Network* offers a simple, uniform price schedule to help you track expenses, plus extensive "help" services and a customized software package – LSN Lifeline[™] – for easy dial up, log on and downloading!

Free Video

Find out about the *Life Science Network*! Order your free *Life Science Network* VHS video tape* from BIOSIS today! Contact BIOSIS, Marketing Department S192SS, 2100 Arch Street, Philadelphia, PA 19103-1399 or call toll free 1-800-523-4806 (USA except PA) or 215-587-4800 (worldwide); Telex 831739;

Fax (215) 587-2016.

*for distribution in the U.S. only

The Life Science Network is sponsored by BIOSIS, serving the life science community worldwide. BIOSIS is a registered trademark of Biological Abstracts, Inc.



GENOME MAPS 1991 Send in your order for a reprint of the Genome Maps 1991, featured in the 11 October issue of Science Magazine. This colorful 21" x 32" foldout wall chart has two key features. In one section it highlights progress in the Human Genome Project—localization of genes

Circle No. 4 on Readers' Service Card

32" foldout wall chart has two key features. In one section it highlights progress in the Human Genome Project—localization of genes and markers on the chromosomes as well as sequencing effects. In addition, because of the importance of model systems in biology and medicine, the chart summarizes mapping and sequencing achievements in one of the classic model systems, *Drosophila melanogaster*.

Order a copy of the Map for your friends, and family by completing the coupon. Please make checks payable to Science (US funds only).



	'etal number andara	1@\$2.00					
	Total number ordered @ \$8.00 Subtotal						
Subtotal For shipment to California, add applicable sales tax.							
							Postage & Hand
In the US	\$1.50			E .			
International Air	r \$5.00						
International Sur	rface \$2.00						
Method OF PAY	MENT						
Visa	MasterCard	Check en	closed				
Card #:		Ex	<pre>cp:</pre>				
NAME:							
ADDRESS:							
	S		ZIP:				
Send Orders to:	Corrine Harris 1333 H St., N.W., Washington, D.C. 202 326-6527 (ph	. 20005	2-0816 (fax)				



HOW TO ACHIEVE ENHANCED CHARACTERIZATION OF BIOMOLECULES.

The Electrospray System from Finnigan MAT simplifies tedious sequencing processes, and lets you produce accurate and intelligent data in a fraction of the time.

Picomole and femtomole sensitivity in molecular weight determination, coupled with structural elucidation achieved in hours—not days or weeks—makes the Electrospray System a powerful tool.

The Electrospray System combines electrospray ionization (ESI) with our high-performance TSQ^{IM} 700 mass

spectrometer to provide molecular weight determination of biomolecules, such as peptides and proteins with mass accuracy of 0.01%.

And the innovative Finnigan MAT data processing software extracts meaningful information and presents it in a format tailored for the biochemist, letting you spend more time on science and less time crunching numbers.

To seek higher intelligence in high mass analysis, call a Finnigan MAT office listed below or FAX (408) 433-4823.



A subsidiary of Thermo Instrument Systems, Inc.

California (408) 433-4800 • Georgia (404) 424-7880 • Ohio (513) 891-1255 • Illinois (708) 310-0140 • New Jersey (201) 740-9177 • Maryland (301) 698-9760 Germany 421-54931 • UK 442-233555 • France 1-6941-9800 • Italy 6-601-1742 • Netherlands 838-527266 • Sweden 08-680-0101 • Japan (03) 3372-3001

Pure mRNA in Minutes...

...Directly from Small or Large Samples of Cells or Tissue.

FastTrack™ and MicroFastTrack™ set the industry standard in high quality mRNA isolation.

MicroFastTrack™*: 20 Reactions

- Ideal for PCR, Northerns and cDNA synthesis
- Isolation from samples ranging in size from 10-3×106
- cells or 10-250mg of tissue. – Reproducible yields of high quality mRNA.

FastTrack™*: 6 Reactions

- mRNA isolation for Northerns, cDNA, library construction, PCR, microinjection, RNA protection studies and *in vitro* translation.
- Isolation from samples ranging in size from 10⁷-10⁸ cells or 0.4-1.0 gram of tissue.
- Fast, efficient recovery of large amounts of polyA + RNA from a variety of sources.

Both systems offer:

- High yields of intact mRNA with low ribosomal contamination.
 Eliminate the need for total RNA isolation or the use of toxic
- The most cost effective means of generating high quality
- mRNA.
- Consistency, convenience and the fastest isolation time.

For the very best in direct mRNA isolation FastTrack™ and MicroFastTrack™ are the choice of thousands of research labs worldwide. When the quality of your mRNA is important, turn to the original source for purity, reliability and convenience; turn to Invitrogen.





3985 • B Sorrento Valley Blvd. • San Diego, CA 92121 (619) 597-6200 Phone • (619) 597-6201 Fax

RITISH BIOTECHNOLOGY LTD, UK - TEL: 44-235529449 • AMS BIOTECHNOLOGY UK LTD, UK - TEL: 44-993822786 • BDH INC., CANADA - TEL: 800-268-0310 • BIO-TRADE, AUSTRIA - TEL: 43-2228284694 • CELBIO, ITALY - TEL: 39-24048646 • FUNAKOSHI PHARMACEUTICALS, JAPAN - TEL: 81-356841622 • ITC BIOTECHNOLOGY GMBH, GERMANY - TEL: 06221-303907 • KEBO LABS AB, SWEDEN - TEL: 46-86213400 • MEDOS COMPANY PTY LTD, AUSTRALIA - TEL: 61-38089077

*patent pending. mRNA model courtesy of BIOSYM

Circle No. 72 on Readers' Service Card

AAAS☆92

The AAAS Annual Meeting

6–11 February 1992 Hyatt Regency Chicago

Don't miss this 158th national gathering of the scientific community. At AAAS☆92, you'll take part in a cross-disciplinary exchange of unmatched caliber. This is the consummate synergy of the sciences, where you can explore the ways in which the sciences interconnect and learn how advances in other fields impact upon your own.

Here's what you'll find:

- More than 200 scientific sessions and 40 topical lectures organized into 22 concurrent "tracks" (see box at right). Each track represents a virtual "meeting within a meeting." You can follow a single track throughout the meeting or pick and choose sessions from various tracks.
- In-depth, three-day seminars on cognitive neuroscience and on molecular modeling and computational chemistry.
- + Plenary lectures by distinguished scientists.
- ✦ An employment exchange in which employers and job candidates can meet for face-to-face interviews.
- ✦ A full series of poster sessions where researchers can discuss their research one-on-one with colleagues.
- Exhibits by science publishers, computer hardware and software companies, on-line information services, and other sciencerelated organizations.
- A unique opportunity to network with your colleagues from across disciplines and from every corner of the scientific community.

Register on site at the Hyatt Regency Chicago!

For a complete program, including prices and registration information, see the 15 November 1991 issue of *Science* or call the AAAS Meetings Office at 202-326-6450.

Meeting Tracks

Molecular Genetics and Evolution

Native American Origins

Climate and Global Change

Medicines and Technologies of the Future

Ethics and Research Policies

Crisis in Health Care

Fantastic Voyages: From Columbus to the Cosmos

Energy for the 21st Century

Feeding the World

Waging War Against Pollution

Mathematics, Communication, and Information Processing

Psychology and Child Behavior

Patterns of Life in Urban and Rural America

Industry and the Changing Workforce

Environmental Modeling and Policy

Science and Math Education: Striving for Excellence

Preserving World Peace

Physics: From Fermi to the Future

Long-Term Research in U.S. National Parks

International Issues

Scientists and Journalists

Science for Everyone

Seminars

Cognitive Neuroscience

Molecular Modeling and Computational Chemistry

American Association for the Advancement of Science

The Best System For Maximum Protein Expression...



Is a Full Magnitude Better!

MaxBac, the only complete baculovirus expression kit, allows high level eukaryotic expression of functionally active protein. Invitrogen, the leader in baculovirus technology, has made recent improvements that make baculovirus expression a full magnitude better, including:

High Expressing Insect Cells – Invitrogen offers a number of insect cell lines, including High 5 (BTI-TN-5B1-4) that has been shown to give up to one log higher levels of protein expression compared to Sf9 cells.

levels of protein expression compared to Sig cells.

■ Transfection Quality Linear Viral DNA – Allows simplified plaque screening, decreases the number of wild type plaques and results in recombinant plaque frequencies of up to 50%.

■ Vectors for Screening and Purification – Impart blue color to recombinant plaques, making plaque identification easier than ever before. Vectors are available for one step purification of recombinant proteins without knowledge of the proteins' biochemical properties.

PCR* Primers for Recombinant Verification – Allows identification of recombinants from plaques or viral preparations and determination of wild type virus contamination in a recombinant plaque.

Put Invitrogen's MaxBac Kit, the most powerful baculovirus expression system, or our **Custom Baculovirus Expression Services** to work in your lab by calling toll free.





3985•B Sorrento Valley Blvd. • San Diego, CA 92121 Phone 619-597-6200 Fax 619-597-6201

TISH BIOTECHNOLOGY LTD, UK - TEL: 44-235529449 • BDH INC., CANADA - TEL: 800-268-0310 • BIO-TRADE, AUSTRIA -TEL: 43-2228284694 • CELBIO, ITALY - TEL: 39-24048646 • FUNAKOSHI CO., LTD., JAPAN - TEL: 81-356841622 • ITC BIOTECHNOLOGY GMBH, GERMANY - TEL: 06221-303907 • SWITZERLAND- TEL: 155-5044 • KEBO LABS AB, SWEDEN - TEL: 46-86213400 • MEDOS COMPANY PTY LTD, AUSTRALIA - TEL: 61-38069077

*PCR is covered by patent issued to Cetus Corporation

Circle No. 73 on Readers' Service Card



Thermostability, Fidelity & Versatility From The Ocean Depths



Heat stability of various thermal stable DNA polymerases. All were incubated at 95°C under standard assay conditions.

For high fidelity, thermostability up to 100°C, primer extensions consistently in the 5-10 Kb range, thermal cycle DNA sequencing and high temperature dideoxy DNA sequencing, discover New England Biolabs' complete range of cloned thermostable Vent[™] DNA polymerases:

- Vent[™]_R DNA Polymerase
- Deep Vent_R[™] DNA Polymerase
 Vent_R[™] (exo⁻) DNA Polymerase

For thermal cycle DNA sequencing, choose our new easy-to-use system:

■ CircumVent[™] DNA Sequencing Kit



Tube worms at 2010 meters around a hydrothermal vent where the native organism containing Deep Vent[™] DNA Polymerase was isolated.



Vent[™] DNA Polymerase

High fidelity thermostable DNA polymerase with fidelity levels up to 20 fold higher than Taq DNA polymerase due to a $3' \rightarrow 5'$ proofreading exonuclease activity.

- Discovered and cloned at NEB
- High fidelity
- Primer extensions up to 13 Kb
- Stable up to100°C
- Half life of 6.7 hours at 95°C

Deep Vent[™] DNA Polymerase NEB's second thermostable DNA polymerase with a 3' \rightarrow 5' proofreading exonuclease activity. Originally isolated from an ocean submarine thermal vent at 2010 meters, Deep Vent_B[™] DNA Polymerase is even more thermostable than Vent_R[™] DNA polymerase at temperatures of 95°C to 100°C.

- Discovered and cloned at NEB
- Superior stability up to 100°C Half life of 23 hours at 95°C



Reversion frequency reflects error rate in DNA synthesis and was measured by the opal codon reversion assay. Kunkel et al. (1987) Proc. Natl. Acad. Sci. USA 84, 4865-4869. Mattila P. et al. (1991) Nucleic Acids Res. 19, 4967-4973.

Vent_R[™] (exo⁻) DNA Polymerase Genetically modified form (exo^-) of the cloned Vent_R^w DNA polymerase with fidelity levels up to 2-fold higher than Taq DNA polymerase. Preferred form for hightemperature dideoxy DNA sequencing and thermal cycle DNA sequencing.

- Discovered and cloned at NEB
- High temperature dideoxy sequencing
- Primer extensions up to 13 Kb
- Stable up to 100°C

CircumVent[™] DNA Sequencing Kit The only thermal cycle sequencing kit that uses the Vent_R[™] (exo⁻) DNA polymerase.

- Easier than conventional methods Fast - no need to collapse double stranded plasmids, eliminates centrifugation steps and independent priming steps. You save valuable research time.
- Requires only nanograms of template (only femtomoles when using ³²P end labelled primers)
- Allows direct sequencing from colonies, plaques, cosmids, or DNA fragments isolated from agarose gels
- Diminished secondary structure effects due to high temperature reaction
- Compatible with radiolabel or chemiluminescent detection

New England Biolabs Inc. 32 Tozer Road, Beverly, MA 01915 USA 800-NEB LABS (US and MA) Tel. (508) 927-5054 Fax (508) 921-1350 New England Biolabs Ltd., Canada Tel. (800) 387-1095 (416) 672-3370 Fax (416) 672-3414 New England Biolabs GmbH, Federal Republic of Germany Tel. (06196) 3031 Fax (06196) 83639

DISTRIBUTORS: AUSTRALIA GENESEARCH Tel. (075) 94 0299 / FINLAND, SWEDEN, DENMARK, USSR FINNZYMES (Finland) Tel. (0) 420-8077 / FRANCE OZYME Tel. (1) 30 57 0025 / INDIA BIOTECH INDIA Tel. (542) 311473 / ISRAEL GAMIDOR Tel. (03) 535-1205 / ITALY C.A.M.Bio Tel. (02) 38103171 / JAPAN DAIICHI PURE CHEMICALS CO. LTD. Tel. (03) 3272-0671 / KOREA KORAM BIOTECH Tel. (02) 556-0311 / THE NETHERLANDS WESTBURG Tel. (033) 95 00 94 / NEW ZEALAND BIOLAB SCIENTIFIC Tel. (09) 418-3039 / NORWAY ING. F. HEIDENREICH Tel. (02) 22 04 11 / PEOPLE'S REPUBLIC OF CHINA CUBC Tel. (1) 256 -1627 / PORTUGAL ISODER TEI. (01) 647208 / SPAIN LANDERDIAGNOSTICO TEL. (01) 594 08 06 / SWITZERLAND FLOW LABORATORIES TEI. (061) 4814713 / TAIWAN LONG CHAIN INTERNATIONAL TEI. (02) 552-2605 / UK CP LABORATORIES TEI. (0279) 758200



Circle No. 20 on Readers' Service Card

Information For Contributors

Science is a weekly, peer-reviewed journal with offices in Washington, DC, and London that publishes research in every field of scientific endeavor. Submitted manuscripts should be intelligible to readers in a variety of disciplines and should be brief and clearly written.

The guidelines below describe our manuscript selection, review, and publication process. Please follow these guidelines in preparing your manuscript to ensure speedy handling by our editorial offices.

Manuscript Preparation

Use double-spacing throughout the text, tables, figure legends, and references and notes, and leave margins of at least 2.5 centimeters. Put your name on each page and number the pages starting with the title page.

Titles and subheadings should be descriptive clauses, not complete sentences or questions. The maximum length is 76 characters and spaces for general articles, and 76 to 114 characters and spaces for research articles and reports.

Abstracts should explain to the general reader why the research was undertaken and why the results should be viewed as important. The abstract should convey the paper's main point and outline the results or conclusions.

Text. A brief introduction describing the paper's significance should be intelligible to readers in different disciplines. Technical terms should be defined. All tables and figures should be cited in numerical order.

Figures and tables should be submitted on separate pages from the text. For each figure submit three high-quality prints, laser prints, or original drawings no larger than 22 by 28 centimeters (8 $\frac{1}{2}$ by 11 inches). On the back of every figure write the first author's name and the figure number and indicate the correct orientation. Photocopies of figures are not acceptable; transparencies, slides, or negatives cannot be used because they cannot be sent to reviewers. Papers that include a large number of figures or tables and a small amount of text may present layout problems. In preparing the manuscript, try to maintain a balance between text length and illustrations.

On acceptance of a paper, authors requesting the use of color will be asked to pay \$600 for the first color figure or figure part and \$300 for each additional figure or figure part to help defray the cost of obtaining color separations. There will be an additional charge for color figures in the reprints.

Cover illustration suggestions may be included with the manuscript. Submit prints, not slides, negatives, or transparencies. After an image is chosen for use on the cover, a positive transparency will be required.

Informed consent. Investigations on humans must include a statement indicating that informed consent was obtained after the nature and possible consequences of the studies were explained.

Animal welfare. Authors using experimental animals must include a statement that their care was in accordance with institutional guidelines. For animals subjected to invasive procedures, include the anesthetic, analgesic, and tranquilizing agents used, as well as the amounts and frequency of administration.

Uncertainties and reproducibility. Evidence that the results are reproducible and the conditions under which this reproducibility (replication) was obtained should be explicitly stated. The effect of limitations in experimental conditions on generalizability of results should be discussed. Uncertainties should be stated in terms of variation expected in independent repetitions of the experiments; they should include an allowance for possible systematic error arising from inadequacies in the assumed model and other known sources of possible bias. Probabilities from statistical tests of significance should not replace the reporting of results and associated uncertainties.

Permissions to reprint illustrations or tables from other publications must be obtained in writing by the author. The written permission must include complete citation from the copyright owner (usually the publisher) to reprint such illustrations in *Science*. Papers are not sent to the printer until copies of all permission letters have been received.

Copyright law requires that we obtain copyright transfer from authors of each paper published in *Science*. Copyright forms are sent to all authors prior to accep-

Inquiries to our London office should be directed to Alun Anderson at 071-494-0062; fax, 071-494-0063. tance and must be signed and returned to the editorial office immediately. U.S. government employees sign the section of the form stating exemption from copyright laws. Alterations to or substitutions for our form are not acceptable.

Categories of Signed Papers

General Articles (3000 to 5000 words or three to five printed pages) are expected to review new developments in one field that will be of interest to readers in other fields; describe a current research problem or a technique of interdisciplinary significance; or discuss some aspect of the history, logic, policy, or administration of science. Readers should be able to learn from a general article what has been firmly established and what are unresolved questions or future directions. Many general articles are solicited by the editor, but unsolicited articles are welcome. Both solicited and unsolicited articles undergo review.

General articles should include a note giving the authors' names, titles, and addresses; an abstract (50 to 100 words); an introduction that outlines for the general reader the main point of the article; and brief subheadings to indicate the main ideas. The reference list should not be exhaustive; a maximum of 50 references is suggested.

Research Articles (up to 4000 words or four printed pages) are expected to contain new data representing a major breakthrough in a field. The article should include an author note, abstract, introduction, and sections with brief subheadings. A maximum of 40 references is suggested.

Figures and tables together with their legends should occupy about one printed page for General Articles and Research Articles.

Reports (up to 2500 words or three printed pages) are expected to contain important research results. Addresses for all authors should be listed on the title page and the corresponding author should be indicated by an asterisk. Reports should include an abstract (no more than 100 words) and an introductory paragraph. A maximum of 30 references is suggested. *Figures and tables together with their legends should occupy no more than one of the pages.*

Policy Forum provides a platform for scientists to present in-depth discussions of policy issues relevant to science. Whenever possible, Policy Forums representing opposing sides are presented in the same issue.

Perspectives analyze recent advances in fast-breaking fields and express opinions as to the impact the developments will have on future research. Perspectives should be ei-

Checklist for Submission

Manuscripts should be addressed to the Editor, Science, 1333 H Street, NW, Washington, DC 20005. Submit three copies together with a letter giving

- the names and telephone numbers of the authors.
- the title of the paper and a statement of its main point.
- the names, addresses, telephone numbers, and fields of interest of four to six persons outside your institution who are qualified to referee the paper.
- the names of colleagues who have reviewed the paper.
 the total number of words (including text, references, and figure and table
- legends) in the manuscript.
- a statement that the material has not been published and is not under consideration for publication elsewhere.

Also include with your manuscript:

- any paper of yours that is in press or under consideration elsewhere and includes information that would be helpful in evaluating the work submitted to Science.
- written permission from any author whose work is cited as a personal communication, unpublished work, or work in press but is not an author of your manuscript.
- for manuscripts based on crystallographic data, two copies of the coordinates.

By submitting a manuscript, an author accepts the responsibility that all those listed as authors of a work have agreed to be so listed, have seen and approved the manuscript, and are responsible for its content.

ther one or two published pages.

Letters are selected for their pertinence to material published in *Science* or because they discuss problems of general interest to scientists. Letters about material published in *Science* may correct errors, provide support or agreement, or offer different points of view, clarifications, or additional information. Personal remarks about an author are inappropriate. Letters may be reviewed by outside consultants. Letters selected for publication are intended to reflect the range of opinions received. The author of the paper in question is usually given an opportunity to reply.

All letters are acknowledged by postcard; authors are notified if their letters are to be published. Preference is given to short letters (250–500 words). Letters accepted for publication are frequently edited and shortened in consultation with the author.

Technical Comments (up to 500 words) may criticize articles or reports published in *Science* within the previous 6 months or may offer useful additional information. Minor issues should be resolved by private correspondence. The authors of the original paper are asked for an opinion of the comment and are given an opportunity to reply in the same issue if the comment is published. Comments and replies are subject to the usual reviewing and editing procedures. Priority disputes may undergo extensive review and are published only when action is recommended.

Book and Software Review selections are made by the editors. Instructions and length specifications accompany items to be reviewed when they are sent to the reviewers, who are chosen by the editors.

Manuscript Review and Selection

Before being reviewed in depth, most papers are rated for their interest and overall suitability by a member of the Board of Reviewing Editors. Papers submitted in disciplines for which there is no appropriate member of the Board of Reviewing Editors may be screened by editorial staff in consultation with outside experts. Papers that are not highly rated are mailed back to the authors within about 2 weeks; the title page and abstract from one copy are retained for our files.

Approximately 35% of submitted papers are reviewed in depth by two or more outside referees. Reviewers are telephoned prior to being sent a paper and are expected to decline to review if they are not qualified or there is a possible conflict of interest. Reviewers are expected to return their comments within 2 weeks and are instructed that the manuscript is a privileged document that is not to be disseminated or exploited. It is the policy of *Science* that reviewers are kept anonymous.

When the review process is complete, the manuscript and reviewers' comments are discussed by the editors at a weekly meeting. Manuscripts are evaluated in terms of their technical merit as well as their merit in relation to other papers under consideration.

In selecting papers for publication, the editors give preference to those of novelty and general significance that are well written, well organized, and intelligible to scientists in different disciplines. An attempt is made to balance the subject matter in all sections of *Science*. Membership in the AAAS is not a factor in selection.

Authors are notified of acceptance, rejection, or need for revision, usually within 8 to 10 weeks. Accepted papers are edited to improve accuracy and clarity and to bring them within the specified length limits.

Papers cannot be resubmitted over a disagreement on interest level or relative merit. If the author can demonstrate that a paper was rejected on the basis of serious reviewer error, resubmission will be considered.

Conditions of Acceptance

When a paper is accepted for publication in *Science*, it is understood that

• any materials and methods necessary to verify the conclusions of the experiments reported will be made available to other investigators under appropriate conditions.

• sequence and crystallographic data will be offered for deposit to the appropriate data bank and the identifier code will be sent to *Science* for inclusion in the published manuscript (coordinates should be released no later than 1 year after publication).

• the author or authors agree to transfer copyright of the paper to *Science*; and the paper will remain a privileged document and will not be released to the press or the public before publication.

• if there is a need in exceptional cases to publicize data in advance of publication, the AAAS Office of Communications (202-326-6440) must be consulted.

Authors may provide a copy of their manuscript on disc upon acceptance. Specific instructions will be provided when the manuscipt is returned for revision.

Printing and Publication

Proofs and reprints. One set of proofs and an order blank for reprints are sent to the authors.

Scheduling. Papers are scheduled for publication after *Science* has received corrected proofs. Papers with tables or figures that present problems in layout, or with cover pictures, or that exceed the length limits may be subject to delay.

INFORMATION FOR CONTRIBUTORS 37

Science Style Sheet

Acknowledgments, including funding information, should be gathered into a brief statement at the end of the references and notes and will be edited to conform to Science style.

Equations and formulas should be typed with quadruple-spacing if they are to be set off from the text. Define all symbols and number all equations.

Figures. Most figures will be printed at a width of 5.9 cm (2.3 inches or 1 column) or 12.2 cm (4.8 inches or 2 columns). Some illustrations (for example, bar graphs, simple line graphs, and gels) may be reduced to a smaller width. Symbols and lettering should be large enough to be legible after reduction. Composite figures should be labeled A, B,C,.... If mounting is necessary, use cardboard.

Legends should be typed double-spaced in numerical order on a separate page. No single legend should be longer than one page. Nomenclature, abbreviations, symbols, and units used in a figure should match those used in the text. The figure title should be given as the first line of the legend.

Line drawings should be labeled on the ordinate and abscissa with the parameter or variable being measured, the units of measure, and the scale. Scales with large or small numbers should be presented as powers of 10. Definitions of symbols should usually appear in the figure legend and not in the figure. Simple symbols (circles, squares, triangles, and diamonds, solid or open) will best survive reduction.

Recommended symbols at the size they should appear after reduction:

Ο 🗆 🔺 Δ

Avoid the use of light lines, shading, and stippling. Use heavy lines or boxes for emphasizing or marking off areas of the figure, and use black, white, hatched, and cross-hatched designs in place of stippling in bar graphs and ball-and-stick molecular models. Authors using computer graphics should choose screens between 20 and 60%.

Halftones, such as electron micrographs, should be submitted as high-quality prints or orginals (do not send irreplaceable artwork). If possible, use scale bars in place of, or in addition to, magnifications. In gels, the lanes should be numbered and identified by number in the figure legend.

For color art please provide a positive slide, if possible, and a print or laser proof. Indicate positioning, lettering, and cropping limits on the print. For composite figures, send the original composite board rather than a print if the quality of the original is much better than that of the print. Do not send irreplaceable artwork.

Lettering in Helvetica font is preferable. Use boldface type for axis labels and for the labels A, B, C,... in composite figures; use italic type only as it would be used in the text (for example, for variables and genes). The first letter of each entry should be uppercase; otherwise, use uppercase letters as they would be used in the text (for example, for acronyms). Avoid wide variation in type size within a single figure. In the printed version of the figure, letters should be about 7 point (2 mm high).

Sequences may be reduced considerably so make sure the typeface in the original is clear. There should be about 130 characters (including spaces) per line for a sequence occupying the full width of the printed page and about 84 characters per line for a sequence occupying two columns.

References and notes are numbered in the order in which they are cited, first through the text and then through the table and figure legends. List a reference only one time. References that are always cited together may be grouped under a single number. Reference to unpublished data should be given a number in the text and placed, in correct sequence, in the references and notes. Use conventional abbreviations for well-known journals; provide complete titles for other journals. Do not use op. cit. See "Science Reference Style" (at right) for examples.

Symbols, abbreviations, and acronyms should be defined the first time they are used.

Tables should supplement, not duplicate, the text. They should be numbered in the order of their citation in the text. Each table should be generated on a separate page with its legend double-spaced above the table. The first sentence of the legend should be a brief descriptive title. Three horizontal lines are used in tables: at the top and bottom of the table and between the column headings and the table body. Vertical lines are not used between the columns.

Every vertical column should have a heading consisting of a title with the unit of measure in parentheses. Units should not change within a column. Centered headings of the body of the table can be used to break the entries into groups. (See the section on lettering for use of italic type and uppercase letters.)

Footnotes should contain information relevant to specific entries or parts of the

Science **Reference Style**

Journals

- 1. I. N. Tang, Atmos. Environ. 14, 819 (1980). [one author]
- J. C. Smith and M. Field, Proc. Natl. Acad. Sci. 2. U.S.A. 51, 930 (1964).
- J. C. Cheeseborough III, S. Trajmar, J.-T. Yang, З. EMBO J., in press. [three to five authors]
- G. Sunshine et al., Lancet i, 711 (1975). [more than five authors]
- M. Schmidt, Sci. Am. 251, 58 (November 1984). 5. [journal paginated by issue] 6.
 - J. Brown, ibid., p. 67.

Technical reports

- 1. D. E. Shaw, Technical Report No. CUCS-29-82 (Columbia University, New York, 1982).
- 2. F. Press, "A report on the computational needs for physics" (National Science Foundation, Washington, DC, 1981). [unpublished or access by title]
- 3 "Assessment of the carcinogenicity and mutagenicity of chemicals," WHO Tech. Rep. Ser. No. 546 (1974).

Proceedings

- Proceedings of the Fifth IEEE Pulsed Power 1. Conference, Arlington, VA, inclusive dates of meeting (publisher, publisher's location, year).
- Proc. IEEE 88, 452 (1968).
- Title of symposium published as a book, sponsoring organization, location of meeting, dates (publisher, location, year)

Paper presented at a meeting (not published)

1. M. Konishi, paper presented at the 14th Annual Meeting of the Society for Neuroscience, Anaheim, CA. 10 October 1984, [Sponsoring organization should be mentioned if it is not part of the meeting name.1

Theses and unpublished material

- 1. B. Smith, thesis, Georgetown University (1973).
- 2. J.A. Norton, unpublished material.

Books

- 1. A. M. Lister, Fundamentals of Operating Systems (Springer-Verlag, New York, ed. 3, 1984), pp. 7–11. [third edition]
- J. B. Carroll, Ed., Language, Thought and Reality: Selected Writings of Benjamin Lee Whorf (MIT Press, Cambridge, MA, 1956).
- R. Davis and J. King, in *Machine Intelligence*, E. Acock and D. Michie, Eds. (Wiley, New 3. York, 1976), vol. 8, chap. 3.
- D. Curtis et al., in Clinical Neurology of Development, B. Walters, Ed. (Oxford Univ. Press, New York, 1983), pp. 60-73. [et al. = more than five authors]
- 5. F. R. Sabier, Contributions to Embryology (Publ. 18, Carnegie Institution of Washington, Washington, DC, 1917), p. 61.
- Principles and Procedures for Evaluating the Toxicity of Household Substances (National Academy of Sciences, Washington, DC, 1977). [organization as author and publisher]

table. The sequence of symbols for footnotes is

*, †, ‡, §, II, ¶, #, **, ††, ‡‡,

Units of measure are given in metric. If measurements were made in English units, give metric equivalents.