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SCIENCE ONLINE

- THIS WEEK IN SCIENCE **EDITORIAL Donald Kennedy**
 - **More Questions About** Research Misconduct
- **EDITORS' CHOICE**
- **NETWATCH**
- **CONTACT SCIENCE**
- 119 **NEW PRODUCTS**



34 Behind the **Bell Labs inquiry**

61 Two paths to thymidylate

NEWS

NEWS OF THE WEEK

- 26 PALEOANTHROPOLOGY: Were 'Little People' the First to Venture Out of Africa?
- 27 CANCER RISKS: Acrylamide in Food: **Uncharted Territory**
- 27 **HOMELAND SECURITY: Scientists Pan Plans** for New U.S. Agency
- 29 **GENOMICS CENTERS: Disease Gene Research** Heats Up in the Desert
- 29 SCIENCESCOPE
- 30 **CONDENSED-MATTER PHYSICS: Spintronics** Innovation Bids to Bolster Bits
- 30 MALARIA: Ecologists See Flaws in **Transgenic Mosquito**
- **▼**31 94 **FISHERIES RESEARCH: Mixed Schools a Must** for Fish?
- COMPUTER SCIENCE: Collective Effort Makes 33 the Good Times Roll

NEWS FOCUS

15

19

20

- 34 BELL LABS: Winning Streak Brought Awe, and Then Doubt
- MATHEMATICS: Graph Theory Uncovers the 38 Roots of Perfection
- 39 **IMAGING TECHNOLOGY: Beautiful Bioimages** for the Eyes of Many Beholders
- 41 SEISMOLOGY: Data Treasures of the Test **Ban Treaty**
- 44 **SOLAR SYSTEM: Comet Chasers Get Serious**
- 47 **RANDOM SAMPLES**



Comet close-ups

SCIENCE'S COMPASS

49 LETTERS

Kids, TV Viewing, and Aggressive Behavior P. H. Klopfer; S. N. Bakshi; R. Hockey. Response J. G. Johnson, P. Cohen, E. M. Smailes, S. Kasen, J. S. Brook. Smallpox Transmission Risks: How Bad? L. H. Kahn. Calling It Something Other Than Cloning D. A. Jones; E. Meyer. Response B. Vogelstein, B. Alberts, K. Shine.

HISTORICAL ESSAY

55 **PORTRAITS OF SCIENCE: From the Modern** Synthesis to Lysenkoism, and Back? U. Hossfeld and L. Olsson

BOOKS ET AL.

- 57 PALEONTOLOGY: Mammoths, Sabertooths, and Hominids 65 Million Years of Mammalian Evolution in Europe J. Augustí and M. Antón, reviewed by T. Flannery
- 58 ZOOLOGY: Encyclopedia of Marine Mammals W. F. Perrin, B. Würsig, J. G. M. Thewissen, Eds., reviewed by R. J. Schusterman
- 59 Browsings

PERSPECTIVES

▼60 75,78 81	PLANETARY SCIENCE: Tip of the Martian Iceberg? J. Bell
▼61	BIOCHEMISTRY: DNA Building Block
105	Reinvented A. G. Murzin
▼62	NANOTECHNOLOGY: Tools for the
72	Biomolecular Engineer C. M. Niemeyer
▼63	CANCER: Addiction to Oncogenes—the
102	Achilles Heal of Cancer I. B. Weinstein



RESEARCH

	BREVIA	▼ 85
65	Low-Temperature Thin-Film Deposition and Crystallization S. Park, B. L. Clark, D. A. Keszler, J. P. Bender, J. F. Wager, T. A. Reynolds, G. S. Herman	26
	REPORTS	
67	Pattern Formation in Homogeneous Polymer Solutions Induced by a Continuous-Wave Visible Laser R. Sigel, G. Fytas, N. Vainos, S. Pispas, N. Hadjichristidis	89 91
70	Probing and Controlling the Bonds of an Artificial Molecule A. W. Holleitner, R. H. Blick, A. K. Hüttel, K. Eberl, J. P. Kotthaus	√ 94
▼72 62	Sequence-Specific Molecular Lithography on Single DNA Molecules K. Keren, M. Krueger, R. Gilad, G. Ben-Yoseph, U. Sivan, E. Braun	31 96
▼75 60 78 81	Global Distribution of Neutrons from Mars: Results from Mars Odyssey W. C. Feldman, W. V. Boynton, R. L. Tokar, T. H. Prettyman, O. Gasnault, S. W. Squyres, R. C. Elphic, D. J. Lawrence, S. L. Lawson, S. Maurice, G. W. McKinney, K. R. Moore, R. C. Reedy	99
▼78 60 75 81	Maps of Subsurface Hydrogen from the High Energy Neutron Detector, Mars Odyssey I. Mitrofanov, D. Anfimov, A. Kozyrev, M. Litvak, A. Sanin, V. Tret'yakov, A. Krylov, V. Shvetsov, W. Boynton, C. Shinohara, D. Hamara, R. S. Saunders	▼ 102 63
▼81 60 75	Distribution of Hydrogen in the Near Surface of Mars: Evidence for Subsurface Ice Deposits W. V. Boynton, W. C. Feldman,	₩ 10: 61
78	S. W. Squyres, T. H. Prettyman, J. Brückner, L. G. Evans, R. C. Reedy, R. Starr, J. R. Arnold, D. M. Drake, P. A. J. Englert, A. E. Metzger,	108
	I. Mitrofanov, J. I. Trombka, C. d'Uston, H. Wänke, O. Gasnault, D. K. Hamara, D. M. Janes, R. L. Marcialis, S. Maurice, I. Mikheeva, G. J. Taylor, R. Tokar, C. Shinohara	110





- A New Skull of Early *Homo* from Dmanisi, Georgia A. Vekua, D. Lordkipanidze, G. P. Rightmire, J. Agusti, R. Ferring, G. Maisuradze, A. Mouskhelishvili, M. Nioradze, M. Ponce de Leon, M. Tappen, M. Tvalchrelidze, C. Zollikofer
- Rooting the Eukaryote Tree by Using a Derived Gene Fusion A. Stechmann and T. Cavalier-Smith
- Rapid Regulation of Light Harvesting and Plant Fitness in the Field C. Külheim, J. Ågren, S. Jansson
- Sustaining Fisheries Yields Over Evolutionary Time Scales D. O. Conover and S. B. Munch

An Essential Role of N-Terminal Arginylation in Cardiovascular Development Y. T. Kwon, A. S. Kashina, I. V. Davydov, R.-G. Hu, J. Y. An, J. W. Seo, F. Du, A. Varshavsky

- **Target-Selected Inactivation of the Zebrafish** *rag1* Gene E. Wienholds, S. Schulte-Merker, B. Walderich, R. H.A. Plasterk
- 02 Sustained Loss of a Neoplastic Phenotype by Brief Inactivation of MYC M. Jain, C. Arvanitis, K. Chu, W. Dewey, E. Leonhardt, M. Trinh, C. D. Sundberg, J. M. Bishop, D.W. Felsher
- 05 An Alternative Flavin-Dependent Mechanism for Thymidylate Synthesis H. Myllykallio, G. Lipowski, D. Leduc, J. Filee, P. Forterre, U. Liebl
- 08 Alternatively Spliced TCR mRNA Induced by Disruption of Reading Frame J. Wang, J. I. Hamilton, M. S. Carter, S. Li, M. F. Wilkinson
- 110 Reversion of B Cell Commitment upon Loss of Pax5 Expression I. Mikkola, B. Heavey, M. Horcher, M. Busslinger
- 114 Activation of *Drosophila* Toll During Fungal Infection by a Blood Serine Protease P. Ligoxygakis, N. Pelte, J.A. Hoffmann, J.-M. Reichhart

102

Switching oncogenes off and on



COVER 81

The south polar region of Mars is highly enriched in hydrogen (blue) in the form of water ice buried less than a meter beneath the surface, as revealed by the Mars Odyssey Gamma-Ray Spectrometer measurements of gamma-ray and epithermal neutron flux. The topographic relief (exaggerated) is from the Mars Orbiter Laser Altimeter. Three reports in this issue focus on martian hydrogen distribution. [Image: G. Shirah, Goddard Space Flight Center; W. Boynton and W. Feldman]



85 Diversity at Dmanisi

A transcription factor that

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CONTENT HIGHLIGHTS AS OF 5 JULY 2002

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White Collar-1, a Circadian Blue Light Photoreceptor, Binding to the frequency Promoter A. C. Froehlich, Y. Liu, J. J. Loros, J. C. Dunlap

White Collar-1, a DNA Binding Transcription Factor and a Light Sensor Q. He, P. Cheng, Y. Yang, L. Wang, K. H. Gardner, Y. Liu

In the fungus Neurospora crassa, a component of the circadian clock, White Collar-1, is revealed to be a FAD-utilizing photoreceptor as well as a transcription factor.



Use of Ionic Liquids for π -Conjugated **Polymer Electrochemical Devices** W. Lu *et al*.

Combining conducting polymers with ionic liquids is shown to lead to a lifetime extension of actuators and other electrochemical devices.

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GLOBAL: From Benchtop to Blackboard—Scientists in Schools Edited by J. Austin

Many nations have a shortage of science teachers and an abundance of young scientists who are eager to teach, but filling those jobs with those scientists isn't always easy.

- GERMANY: The Never-Ending Debate S. Steghaus-Kovac A continuing discussion on the effects that university reform will have on the careers of young scientists.
- UK: Biotechnology in Britain—Growth and Opportunities E. Pain At the BiologyInBusiness network, Chris Lowe, director of the Institute of Biotechnology, paints an optimistic picture of the UK biotech industry.
- EUROPE: Who Cares About Slovak Ph.D. Students? G. Gregušová Despite poor stipends and the prospect of low wages upon qualification, Ph.D.'s are increasingly popular in Slovakia.
- US: Postdoc Parents See Room for Improvement J. C. Reddy A physician-scientist describes how parenthood turned her into an agent for change.
- AUSTRALIA: Contract Versus Tenure-Which Way to Go? T. J. Piva The pros and cons of contract and tenure positions in Australian academia.

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- Marked for Death? R. J. Davenport Scientists expose proteins damaged in Alzheimer's disease.

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signal transduction knowledge environment

Review: "Crosstalk" in Abscisic Acid Signaling N.V. Fedoroff How integrated signaling from multiple pathways orchestrates plant responses to stress.

Protocol: Screening for Gene Regulation Mutants by Bioluminescence Imaging V. Chinnusamy, B. Stevenson, B. Lee, J.-K. Zhu A method for identifying plants with defective stress responses.

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THIS WEEK IN Science

RecA Resists for Nanowiring

One approach to wiring together nanometer-scale electronics relies on using DNA as a template-the ends of a long DNA molecule are designed to hybridize to components bearing complementary strands, and the DNA is then metallized. Keren et al. (p. 72; see the Perspective by Niemeyer) show that the major protein responsible for homologous recombination in Escherichia coli, RecA, can be used to pattern metallized DNA. RecA was polymerized on single-stranded DNA molecules, which were then hydridized to double-stranded DNA substrate molecules. RecA could be used either lithographically to create nonmetallized gaps in the DNA, or to perform recombination reactions that created branched networks.

More from Dmanisi

Studies at Dmanisi, Georgia, have shown that early *Homo* migrated out of Africa by 1.75 million years ago. Vekua *et al.* (p. 85; see edited by Phil Szuromi

Subsurface Martian Water Since February 2002, the Gamma-Ray Spectrometer, Neutron Spectrometer, and High Energy Neutron Detector on board the 2001 Mars Odyssey mission have been used to map the flux

of gamma rays and secondary neutrons, which are attributed to hydrogen concentrations in the top meter of the martian surface. Results from just 30 to 60 days of data presented by Feldman et al. (p. 75; see the cover), Mitrofanov et al. (p. 78), and Boynton et al. (p. 81) indicate that, at high latitudes (>60°) near both poles, a putative water ice-rich layer exists about 30 to 60 centimeters below the surface (see the Perspective by Bell). This layer, which is tens of centimeters in thickness, may contain 20 to 35% by weight of water ice, and this ice would fill in most of the estimated porous regolith at this depth. Nearer the equator, the depth of this layer increases to about 100 centimeters, and the amount of possible water ice in the layer decreases. Thus, Mars may have collected and concentrated a significant amount of water just below the surface at both poles. The presence of water is consistent with a wetter past, seasonally modulated water transport, current gully formation by subsurface water-ice erosion, and the potential for a higher concentration of water than expected beneath the martian surface.

And in Brevia ...

A technique developed by Park *et al.* (p. 65) allows refractory oxides, such as zirconium oxide, to be grown as crystalline thin films at low temperatures (~130°C) and on plastic substrates.

news story by Balter and Gibbons) now report a new hominid skull and mandible from this site. Together, these fossils represent the largest collection of individuals from any one site older than about 800,000 years ago. The specimens may even have been from a common group. Regardless, their diversity may provide a view of the variation in morphology of early hominid species. This newly discovered hominid has a relatively small brain, implying that evolution in brain size may not have determined when migration out of Africa occurred.

Getting to the Root of It All

Despite decades of phylogenetic research, the position of the root of the eukaryotic tree has remained controversial. Stechmann and Cavalier-Smith (p. 89) have used a gene fusion that is almost certainly derived and compared it to the ancestral bacterial condition to show that the root must lie below the divergence point between opisthokonts (animals, fungi, and Choanozoa) and biciliated eukaryotes (which include plants, chromists, and almost all protozoa). This new evidence is congruent with a simple interpretation of the evolution of the eukaryotic cytoskeleton and a basic divergence in mitochondrial structure, and is the most substantial advance to date in clarifying the position of the root and the nature of the first eukaryote. An On/Off Switch for Oncogenes

Among the many new strategies being developed for cancer therapy are drugs that inhibit the function of oncogenes that are involved in tumorigenesis. Such drugs might be toxic if used chronically, yet they might be ineffective in the short term because of possible reactivation of the oncogene once treatment is stopped. To investigate the latter possibility, Jain et al. (p. 102; see the Perspective by Weinstein) used a sophisticated mouse genetic model to examine what happens to MYC-induced tumors when the MYC oncogene is briefly inactivated and subsequently reactivated. Surprisingly, they found that transient MYC inactivation leads to permanent loss of the neoplastic phenotype. When MYC was removed, osteogenic sarcoma cells differentiated into bone cells; re-expression of MYC did not restore the cells' tumorigenic potential but, rather, caused the cells to undergo apoptosis. These results suggest that brief inactivation of an oncogene may permanently

change the epigenetic context of a tumor cell so that it cannot revert to its original malignant behavior.

Quantum Dot Molecules

Quantum dots, like molecules, have discrete energy levels that can be tuned by varying their size and composition. Holleitner *et al.* (p. 70) have controlled the coupling between two quantum dots with a technique based on cotunneling of electrons through each of the dots. They then probe the development of energy spectra analogous to the formation of molecular levels.



Fisheries management is based mainly on population dynamics, but in response to predation, it is conceivable that populations might evolve rapidly, which could lead to diminished yields. Conover and Munch (p. 94) harvested large, small, or randomly sized members of an exploited fish, the Atlantic silverside (*Menidia menidia*), from captive populations over four generations. Harvesting large individu-



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CONTINUED FROM 9 THIS WEEK IN SCIENCE

als initially increased overall catch but quickly led to lower yields, as slow-growing phenotypes began to dominate. The authors suggest that creating no-take reserves or enforcing a maximum size limit may preserve the genetic variability in wild populations.

Second-Sourcing Thymidylate

It might be assumed that mechanisms for the biosynthesis of essential DNA precursors would be conserved in all organisms. However, Myllykallio *et al.* (p. 105) show that a large class of proteins that do not have sequence similarity with the classic thymidylate synthase (ThyA) synthesizes thymidylate by a different mechanism. These enzymes (ThyX) are mainly limited to microbial genomes that lack ThyA and are an important drug target as they occur in several human pathogens. In a Perspective, Murzin comments on a rare confluence of events in which the functional characterization is complemented by the recent structural characterization of ThyX by the Joint Center for Structural Genomics.

Compensating for Nonsense

Mutations that introduce premature termination codons (nonsense) can affect alternative splicing of transcripts by disrupting exonic splicing enhancers. Wang *et al.* (p. 108) have examined a T cell receptor (TCR) gene that acquires nonsense codons at high frequency during normal T cell development. In contrast to previous explanations, they show that nonsense-associated altered splicing (NAS) is enhancer independent. Instead, it appears that nonsense codons up-regulate alternatively spliced messenger RNA by a translation-like mechanism.

Sticking with It

Specification of cell lineage requires early decisions that, in some cases, may be reversible. The transcription factor Pax5, is required for entry into the B cell lineage via the repression of non–B cell genes and establishment of the B cell–specific gene profile. After conditionally inactivating the *Pax5* gene, Mikkola *et al.* (p. 110) observed that the developmental program of committed early B cells could be reset. As a result, other myeloid and lymphoid lineages could be derived from these cells in vivo and in vitro. This type of persistent control over early lineage preservation may prove an important general principal in upholding cell fate decisions.



Taking Degradation to Heart

The ubiquitin pathway of protein degradation has gained prominence as a major regulatory network in mammalian cells. Arginine is commonly conjugated to the amino-terminus of proteins during ubiquitin-dependent degradation, but the physiological functions of arginylation are unknown. Kwon *et al.* (p. 96) generated mice genetically deficient in one of the Arg-tRNA-protein transferases catalyzing this modification (ATE-1) and found that the embryos die from defects in heart development and angiogenesis. The authors also discover a possible mechanism for the early cardiac defects: Amino-terminal cysteine is oxidized prior to its arginylation by ATE-1, suggesting that the amino-terminal arginylation may function as an oxygen sensor.

Breaking the Mold

Drosophila protect themselves against bacterial and fungal infection by secreting antimicrobial peptides from the fat body, the insect equivalent of the liver. One mechanism required for initiating this in response to Gram-positive bacteria and fungi is provided by the Toll pathway, which is activated via binding of a proteolytically cleaved form of its ligand Spaetzle. Using mutagenesis screening, Ligoxygakis *et al.* (p. 114) identified mutants for a serine protease gene named *persephone*, which suppressed the Tolldependent antifungal response. Unlike other activators of the Toll pathway, *persephone* has no inherent microbial recognition domain, which suggests that it depends on an upstream fungal receptor to initiate its cleavage of Spaetzle.



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Simplify your protein expression work with the **RTS ProteoMaster Instrument**. This compact instrument supports all three RTS *E. coli* HY Kits – **RTS 100**, **RTS 500**, and **RTS 9000**.

The Rosette Nebula C Robert Gendler



RTS ProteoMaster Instrument with crystals and X-ray diffraction pattern of phosphoserine phosphatase, produced using the RTS 500 *E. coli* HY Kit.

Product	Description	Cat. No.	Pack Size
RTS Linear Template Generation Set	generate PCR templates for expression in RTS 100	3 186 237	1 set (96 reactions)
RTS 100 <i>E. coli</i> HY Kit	express up to 20 µg protein/reaction in just four hours	3 186 148 3 186 156	1 kit (24 x 50 µl reactions) 1 kit (96 x 50 µl reactions)
RTS 500 <i>E. coli</i> HY Kit	generate up to 5 mg protein/reaction in 24 hours	3 246 817 3 246 949	1 kit (2 x 1 ml reactions) 1 kit (5 x 1 ml reactions)
RTS 9000 <i>E. coli</i> HY Kit	scale-up to 50 mg protein/reaction in 24-36 hours	3 290 395	1 kit (1 x 10 ml reaction)
RTS ProteoMaster Instrument	supports all RTS kits	3 064 859	1 instrument plus inserts

For detailed information, visit **www.proteinexpression.com** or contact your Roche sales representative.

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The PTC-100[™] Thermal Cycler

The first MU Research Peltier-effect thermal cycler delivers precision and ease of use in a compact and reliable instrument. The novel Peltier heating and cooling technology establishes a widespread reputation for innovation and versatility. Today, the updated PTC-100 continues to deliver reliability and economy.

1988



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2000



The DNA Engine Opticon[™] Real-Time System

The development of fluorescence-based assays has radically changed DNA analysis and the novel Opticon" detector accessory enables quantitative and melting-curve analysis as well as other precision temperature-controlled measurements on the flagship DNA Engine thermal cycler. Its unique rugged design uses LED excitation, and PMT detection to deliver high sensitivity with no moving parts.

^{arts.} 2001



The MiniCycler[™] Thermal Cycler

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The DNA Engine Tetrad[™] Thermal Cycler

The DNA Engine family grows, adding an instrument which incorporates four DNA Engine cyclers in a single minimal footprint. MJ also designs and introduces today's standard configuration for 384-well blocks and vessels. Together, these innovations create a single "Genome Engine", with a capacity sixteen times that of a 96-well system, which becomes the standard in high-throughput DNA sequencing, and the thermal-cycling backbone of the Human Genome Project.



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FAMRI ANNOUNCES THE SECOND YEAR CLINICAL RESEARCH AWARDS

In October 1991, Miami attorneys, Stanley and Susan Rosenblatt, brought a class action suit against the tobacco industry seeking damages on behalf of flight attendants and their survivors, for the diseases and death that have been caused by their exposure to secondhand tobacco smoke in airline cabins. The October 1997 settlement, after four months of trial, among other substantial benefits to class members, established an endowment fund of \$300 million that has supported a not-for-profit research foundation, the Flight Attendant Medical Research Institute (FAMRI). The Mission of FAMRI is to sponsor scientific and medical research for the early detection, treatment, and cure of diseases and medical conditions associated with exposure to secondhand tobacco smoke. FAMRI is governed by a Board of Trustees with the majority of flight attendants. A Medical Advisory Board of highly qualified, internationally recognized clinical scientists, chaired by former United States Surgeon General Julius Richmond, M.D. and a Lay Advisory Board of dedicated concerned citizens assist the Governing Board in decision-making. FAMRI has contracted the American Institute of Biological Sciences to conduct the peer review of proposals for the three clinical research awards detailed below. More information about FAMRI and the awards, including the Requests for Applications will be available on the web at: http://www.famri.org after May 1, 2002. Other communications and queries should be directed to: Beth Kress, FAMRI Executive Director, 201 S. Biscayne Blvd., Suite 1310, Miami, FL 33131. E-mail famri@bellsouth.net; phone, 305-379-7007; fax, 305-577-0005.

YOUNG CLINICAL SCIENTIST AWARD (YCSA)

The purpose of the FAMRI YCSA is to help prepare and support new clinical investigators with a M.D. or Ph.D. as they begin their careers as independent researchers. The program is limited to the development of young researchers in smoking-related disorders. FAMRI is particularly interested in helping to provide the bridge between the clinic and the laboratory for the critical translation of basic research findings into diagnostic and therapeutic approaches. The YCSAs are being offered to two groups of scientists: research fellows and junior faculty members.

CLINICAL INNOVATOR AWARD (CIA)

FAMRI established the CIA to stimulate novel medical and clinical scientific research studies on the effects of exposure to secondhand tobacco smoke. While considerable government and non-government funding is available to support established mainstream biomedical research projects, funds for high-risk projects are generally quite limited. With the CIA, FAMRI hopes to foster innovative breakthroughs and creative collaborations. The CIA is available to clinical investigators with a M.D. or Ph.D.

CENTER OF EXCELLENCE (CoE)

FAMRI's CoE will be the centerpiece to linking physicians and scientists from various disciplines into multidisciplinary programs in patient care and research. The aims of FAMRI's Centers of Excellence are to enhance the knowledge base relating to exposure to second hand tobacco smoke, to serve as a new source for more effective approaches to detection, diagnosis, and therapy for diseases associated with such exposure, and to serve as principal deliverers of medical advances to those suffering from such exposure, especially flight attendants. FAMRI has developed this award program on an institutional basis, striving for comprehensive research plans, including the entire range of research endeavors from basic to clinical research, as well as community outreach.

FAMRI does not support individuals who

or institutions that are currently receiving funds from the tobacco industry or its affiliates. Full disclosure of any prior tobacco industry funding is required of all applicants.



The McNeil Research Grant Awards are designed to sponsor original research on: The mechanism of action of acetaminophen or New indications and uses for acetaminophen.

A maximum of \$60,000 per grant per year will be made available. Extensions of one additional year (maximum grant \$120,000) will be considered. Selected applicants must be willing to publish results in a peer-reviewed journal or present findings at a national scientific meeting. Investigators will be required to submit interim reports on the project's progress. Proposals must be received post-marked, or submitted online, by September 30, 2002. All applicants will be notified of their status by November 30, 2002.

For details of the procedures and conditions of the competition visit www.tylenolgrants.com

Or contact:

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Go to www.amershamscienceprize.org to obtain the mandatory entry form.

And take your step. Avraham Yaron did and won our regional prize for all other countries in 2000. Then in 2001, he was published for the first time in *Science*—"Sorting of Striatal and Cortical Interneurons Regulated by Semaphorin-Neuropilin Interactions,"*Science* **293**:872-875 (2001).

* For the purpose of this prize, molecular biology is defined as "that part of biology which attempts to interpret biological events in terms of the physico-chemical properties of molecules in a cell" (McGraw-Hill Dictionary of Scientific and Technical Terms, 4th Edition).

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