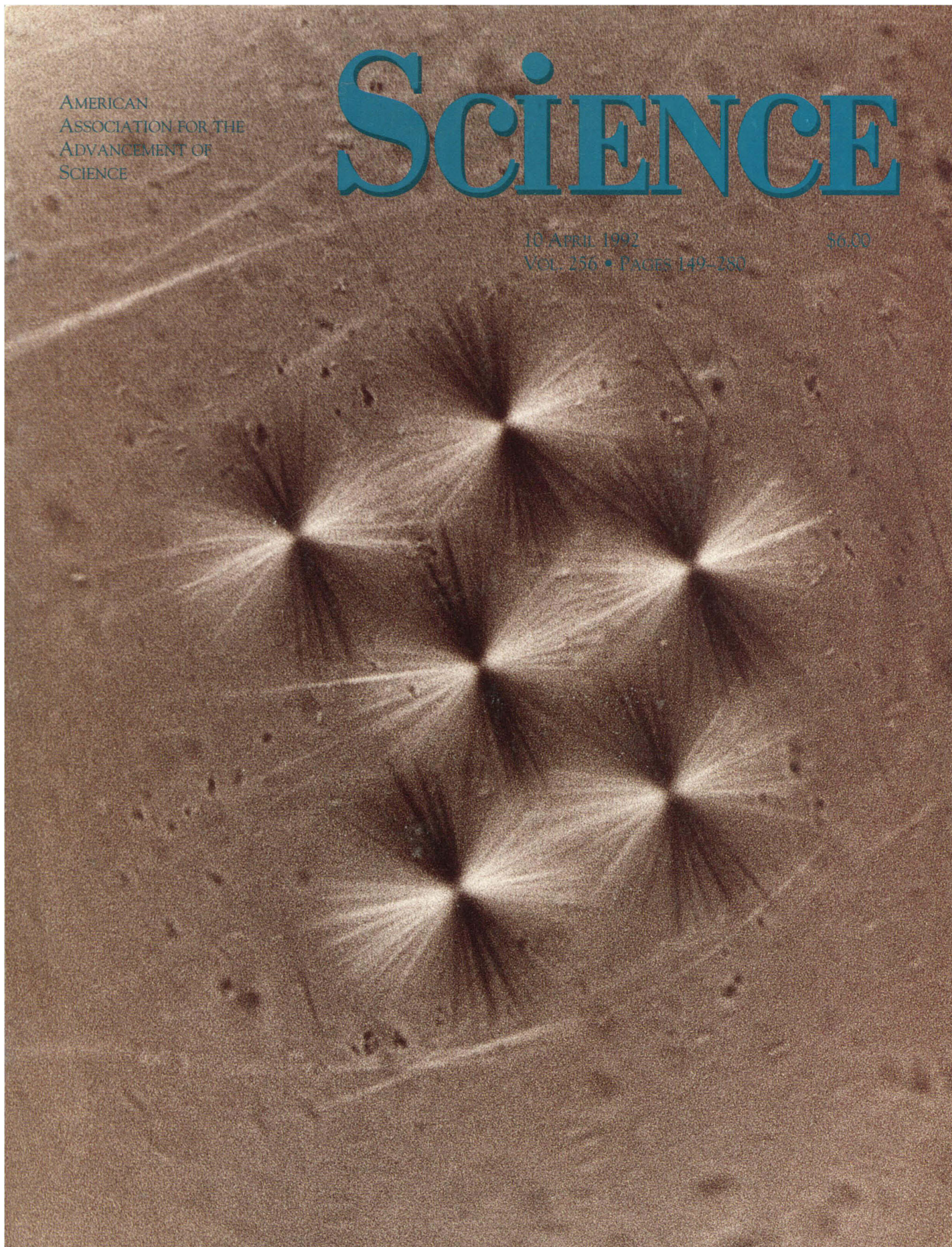


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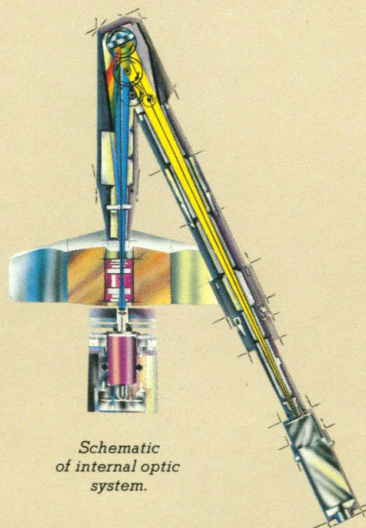




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THE ORIGINAL DNA THERMAL CYCLER

88

DATE 12-12-91

SUBJECT λ DNA AMPLIFICATION

PERIOD ENDING Q2

PAGE 47

		Operating budget:	April	May	June	Total
PCR MIX	VOLUME (μ L)					
H ₂ O	61.5	costs:				
10X BUFFER	10	facturing labor	\$57,600	\$60,500	\$63,400	\$181,500
dATP	2	materials	53,800	56,400	59,200	169,400
dCTP	2	ting supplies	6,500	6,900	7,300	20,700
dGTP	2	r labor and parts	7,300	12,400	6,500	26,200
dTTP	2	r heat, light	4,200	4,500	4,800	13,500
AMPLITAQ	0.5	l direct costs	129,400	140,700	141,200	411,300
PRIMER #1	5					
PRIMER #2	5					
BACTERIOPHAGE λ DNA	10 \leftarrow (λ DNA DILUTED)	vision	5,500	5,500	5,500	16,500
	100 μ L	ort labor	28,500	28,500	28,500	85,500
PIPETTE MASTER		axes	8,700	8,700	8,700	26,100
MIX INTO REACTION			20,500	20,500	20,500	61,500
TUBE. ADD 50 μ L		ood costs	63,200	63,200	63,200	189,600
MINERAL OIL.		ontrollable costs	192,600	203,900	204,400	600,900
AMPLIFY.		verhead	72,000	72,000	72,000	216,000
PCR PROTOCOL		total cost	\$264,600	\$275,900	\$276,400	\$816,900
DENATURE:						
94°C - 1 MINUTE						
ANNEAL:						
37°C - 1 MINUTE		er shifts	3	3	3	3
EXTEND:			20	21	22	63
72°C - 2 MINUTES		asses per shift	33	33	33	33
- 25 CYCLES		er of equipment	35	35	34	-
		duction:				

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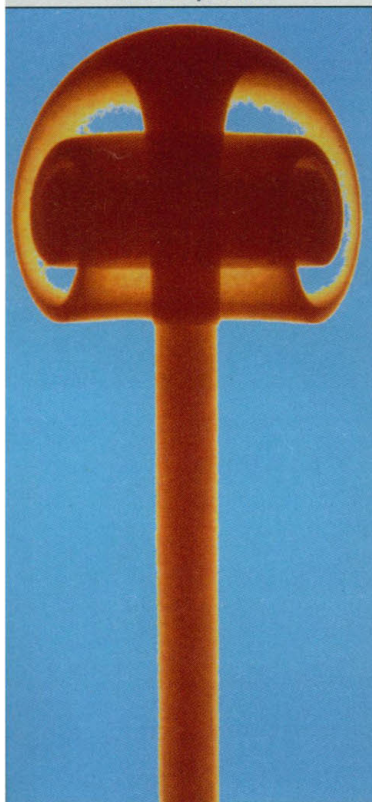
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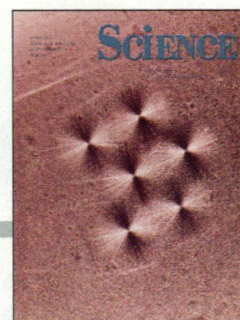
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COVER

Polarized light micrograph of asters (systems of microtubules emanating from centrosomes; these microtubules contribute to the mitotic spindle during cell division) in cytoplasmic lysates prepared from oocytes of

the Atlantic surf clam, *Spisula solidissima*. A method of duplicating centrioles outside of the confines of the cell has now been developed. See page 219. [Photomicrograph by Robert E. Palazzo]



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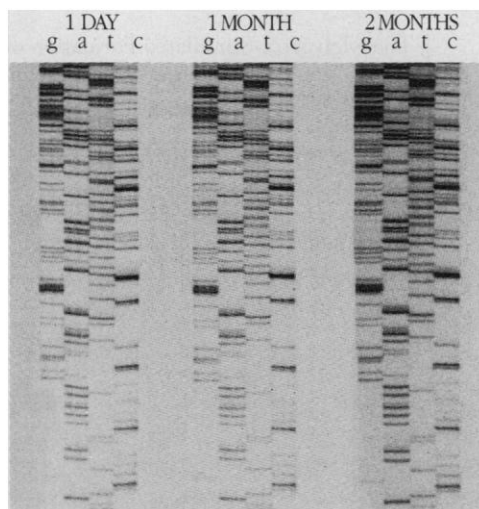
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Darwin's finches

Interbreeding or hybridization is relatively common among many species of birds, but the fate of hybrid offspring is not well known. Grant and Grant (p. 193) have documented the frequency of hybridization between three species of Darwin's finches on a small island in the Galápagos chain through a number of generations and assessed the fitness consequences. Surprisingly, the hybrids in this natural environment in the years studied showed higher overall fitness than conspecifics by combined measures of survival and breeding success.

Come together

Site-specific recombination in bacteriophage λ is mediated by higher order protein-DNA complexes that bring together distant DNA sites on the chromosome of *Escherichia coli*. Kim and Landy (p. 198) studied the excision of bacteriophage λ DNA from its *E. coli* host by comparing synergistic effects between pairs of mutants with marginally impaired function. A map is proposed for the association of the two prophage DNA sites, *attL* and *attR*, with copies of the *Int* protein, which can bind two DNA strands, and with the DNA bending proteins IHF, Xis, and Fis, which facilitate long-range tethering.

Triton's ionosphere

Initial analysis of Voyager observations suggested that Triton's extensive ionosphere was produced by precipitation of electrons when Triton passed through Neptune's magnetic equatorial plane. The principal ion present was thought to be N^+ . A photochemical model of

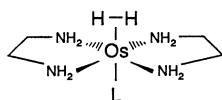
Triton's atmosphere by Lyons *et al.* (p. 204) that accounts for reactions among H, C, and N species shows that the extensive ionosphere could be produced simply by solar radiation if the principal ion in the ionosphere was C^+ rather than N^+ .

Solar diamonds

Many primitive meteorites contain trace amounts of small (about 2 nanometers across) diamonds whose chemical and isotopic nature indicate that they were produced at low pressures as circumstellar grains and predate formation of the solar system. Russell *et al.* (p. 206) report the presence of similar appearing diamonds from the enstatite chondrite Abee, except that these diamonds lack the chemical and isotopic characteristics of the presolar diamonds and thus likely formed within the solar nebula.

Versatile NMR probe

An osmium complex, $[\text{en}_2\text{Os}(\eta^2\text{-H}_2)]^{2+}$ (where en is ethylenediamine), binds readily



to a wide variety of biological molecules (ligand L) in aqueous solution. Li and Taube (p. 210) show that the nuclear magnetic resonance (NMR) peak associated with the dihydrogen ligand, which is bound in a side-on fashion to the metal, appears in the clear part of the NMR spectral window between 0 and -20 parts per million. This peak can serve as a probe for the binding of nucleotides, RNA, amino acids, peptides, and phospholipids.

Backbone engineering

Synthesis of large proteins by chemical means is attractive because unusual modifications can be made to proteins, such as in the backbone chain. Schnölzer and Kent (p. 221) synthesized a fully active, backbone-engineered form of HIV-1 protease by ligating two large peptide segments. The segments were modified so that they formed a thioester bond between Gly⁵¹ and Gly⁵² in each half of the dimer. The solubility of these unprotected peptide segments allowed the synthesis to proceed in high yield.

HMG1 and cisplatin

Cisplatin [$\text{cis-Pt}(\text{NH}_3)_2\text{Cl}_2$] is an effective anticancer drug, whereas its trans isomer is not, even though both bind to DNA and block replication. Pil and Lippard (p. 234) show that DNA damaged by cisplatin specifically binds the high-mobility group protein HMG1. Such binding is not seen with therapeutically inactive platinum compounds. The cytotoxicity of cisplatin may result because binding of HMG1 might prevent repair, or perhaps HMG1 and related proteins are made unavailable for transcription.

Muscles minus myoD?

Members of the myoD family of DNA binding proteins appear to exert control over the formation of muscle cells in vertebrates, but experiments in the nematode *Caenorhabditis elegans* by Chen *et al.* (p. 240) suggest that involvement of these proteins may not be necessary in invertebrates. Only one gene, *hlh-1*, was found in a search for myoD homologs in *C. elegans*. This gene is expressed in body-

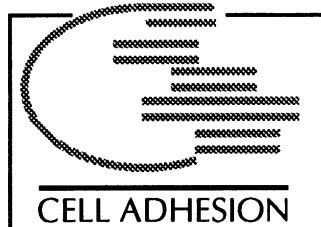
wall muscle cells and their precursors. Several gamma-ray-induced mutants that were homozygous for deficiencies in the *hlh-1* gene still differentiated to form muscle cells and myofibrils. Myogenic activity may be initiated through a myoD-independent pathway.

Lithium in development

The teratogenic effects of lithium on the development of early embryos are well known, but poorly understood. Maslanski *et al.* (p. 243) injected lithium into a cell in the vegetal region of early frog (*Xenopus laevis*) embryos, which disrupts the normal development of the mesoderm. Co-injection of specific isomers of inositol counteracted the effects of lithium. Lithium may inhibit the polyphosphoinositide cycle, which may normally contribute to the signal transduction pathways that result in induction of mesoderm.

Activating actin

Polymerization of actin is increased in cells stimulated by growth factors or chemoattractants. Shariff and Luna (p. 245) report that in the slime mold *Dictyostelium discoideum*, diacylglycerol (DG), which is produced from phosphatidylinositol biphosphate during mitogenesis or chemotaxis, stimulates nucleation of actin filaments at the plasma membrane. The effect of DG on actin polymerization required one or more proteins tightly associated with the plasma membrane. Protein kinase C is activated by DG, but it did not appear to mediate the effect of DG on actin polymerization (see news story by Travis, p. 177).



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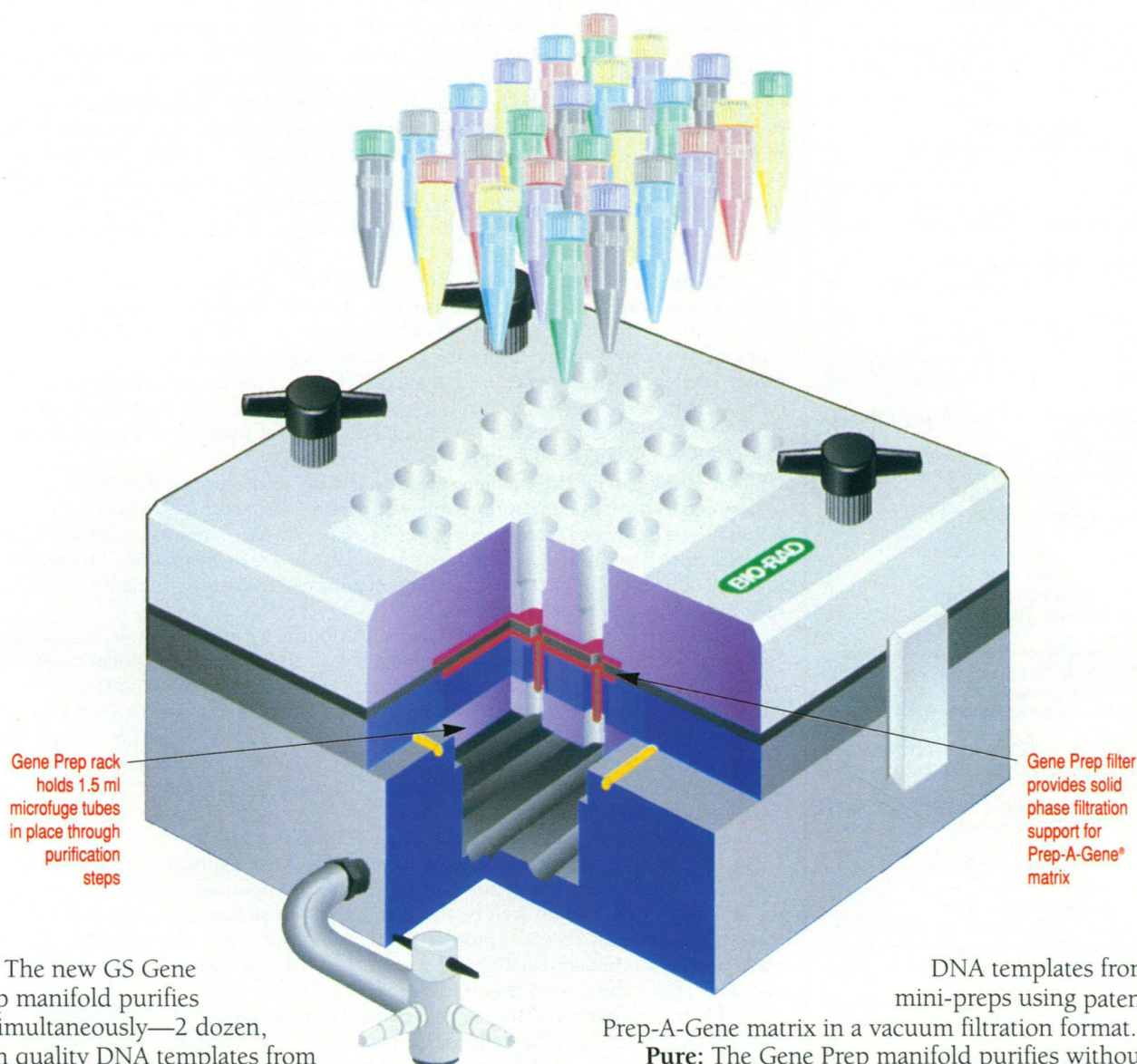
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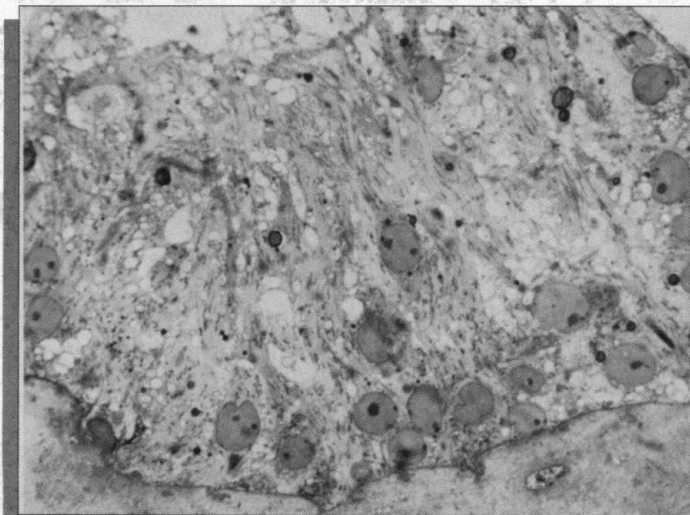
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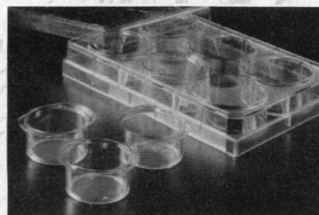


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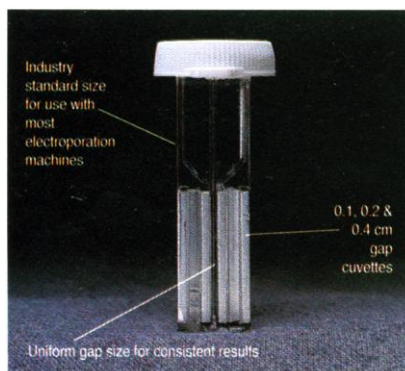
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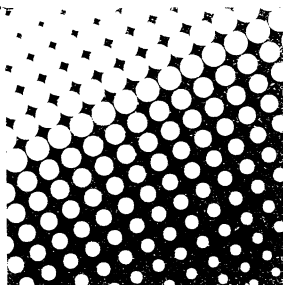
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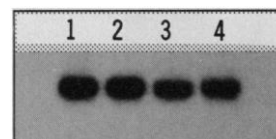
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TWINNING PROGRAM WITH ROMANIA

The National Academy of Sciences is accepting proposals for research programs which link individual U.S. scientists with Romanian counterparts in fields which are normally supported by the National Science Foundation. These twinning programs should have the strong support of the scientists' home institutions and should lead to long-term sustained linkages.

Twinning programs require a 2 to 4-year commitment beginning in 1993. One month of travel support per year in each direction will be available for the participating scientists.

Applications will be accepted from individuals who are a) United States citizens; b) native residents of a possession of the United States; or c) green card holders or permanent residents of the United States. They must:

- be engaged in research careers (or research and teaching careers) and be in possession of a PhD degree;
- be affiliated with an educational or research institution in the United States;
- have existing contacts with Romanian researchers and/or institutions.

U.S. twinning program participants will receive grants to support their travel between their residences and Romania, their living and travel expenses within Romania, and the living and travel expenses in the United States for their Romanian twinning partners.

Applicants should submit 5 copies of each of the following:

- curriculum vitae
- list of publications
- 1 page single-spaced description & justification of proposed research
- letter of support from the Dean or other senior official of the applicant's institution
- correspondence with Romanian twinning partner

Each applicant must arrange for two letters of recommendation to be sent to the National Academy of Sciences.

Application materials should be sent to the address listed below postmarked no later than May 30, 1992.

Selection criteria will emphasize scientific achievements and ability and benefits from research carried out in Romania.

All candidates will be notified of final decisions by July 30, 1992.

Inquiries and application materials should be addressed to:

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National Academy of Sciences FO 2014
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fax: (202) 334-2614
Attn: Elisa Chait

Announcement for the AAAS Black Church Health Connection Project

To boost the number of blacks who are scientifically literate and who understand how the body functions and how it can be abused, AAAS is conducting a three-year project to develop a set of participatory biology activities and an accompanying training program.

Scientists and health professionals are invited to develop a set of participatory activities that will emphasize basic concepts of biology, as well as how the body functions and can be abused by addictive drugs, tobacco, and alcohol. Activities, which will be used in a church setting by school-aged children as well as adults, should also emphasize problem-solving skills and creative thinking. Scientists and health professionals who submit activities selected and used in this project will receive \$100. The deadline for submitting activities is May 22, 1992. For detailed information on guidelines for submitting materials, contact **Audrey B. Daniel at (202) 326-6670 or write to: Audrey B. Daniel, AAAS Directorate for Education and Human Resources Programs, 1333 H Street, NW, Washington, DC 20005.**

Funded by the Alcohol, Drug Abuse, and Mental Health Administration (ADAMHA), National Institutes of Health, U.S. Department of Health and Human Services, Washington, DC

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*PCR is covered by U.S. Pat. #'s 4,683,202 and 4,683,195 issued to Cetus Corporation.

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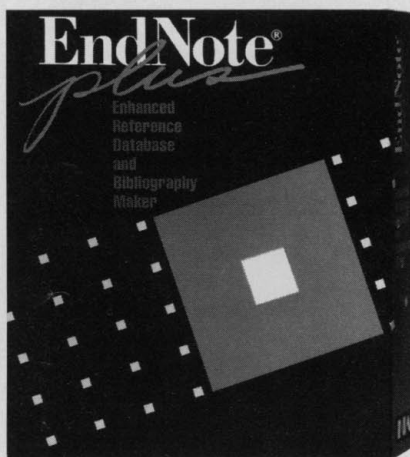
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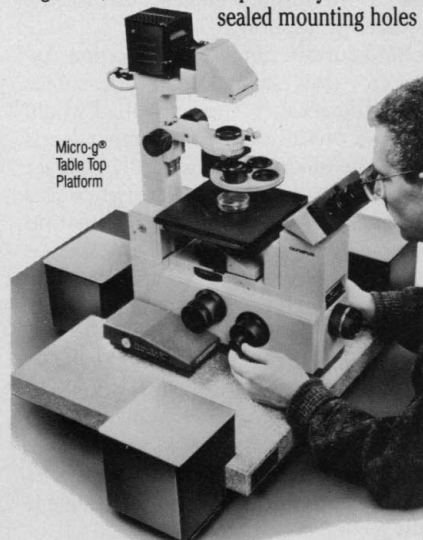
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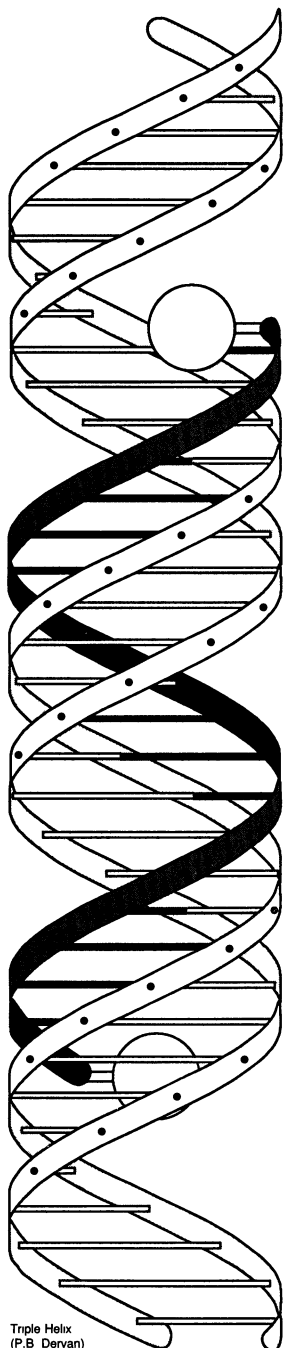
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