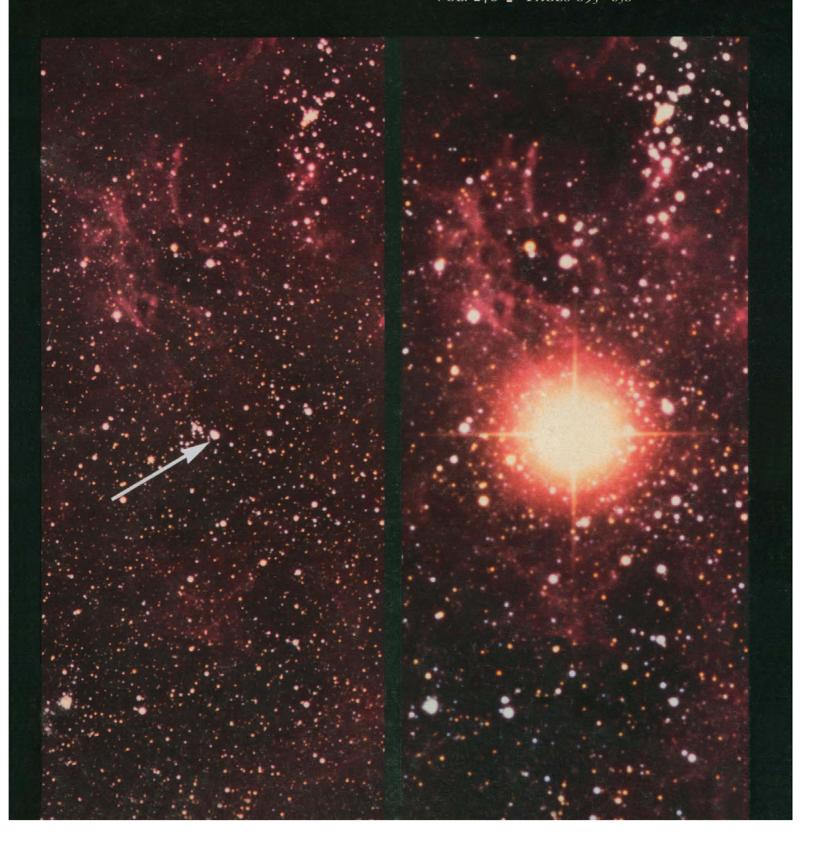
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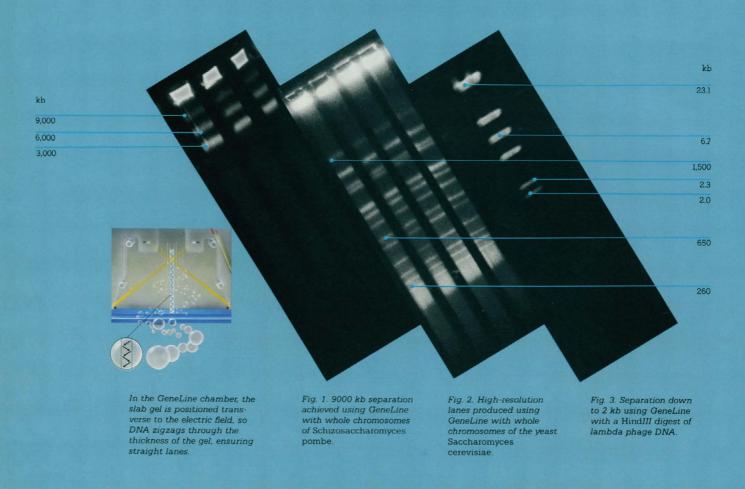
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Grasp 3.0	Blue Start     33       Clickart Publications     37       Colormate     56       Decision Map     108       Dollars & Sense     112       Double Helix     273       Easy 3D     112       In Talk 3.0     146       Laser Author     44       Lazerstart 2.5     71
Grasp 3.0	Blue Start     33       Clickart Publications     37       Colormate     56       Decision Map     108       Dollars & Sense     112       Double Helix     273       Easy 3D     112       In Talk 3.0     146       Laser Author     44       Lazerstart 2.5     71       MAC Vision     224       Mainstreet Filer     75
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Grasp 3.0	Blue Start         33           Clickart Publications         37           Colormate         56           Decision Map         108           Dollars & Sense         112           Double Helix         273           Easy 3D         112           In Talk 3.0         146           Lazer Author         44           Lazerstart 2.5         71           MAC Vision         224           Mainstreet Filer         75           Microsoft Chart         78           Microsoft Excel         296
Grasp 3.0	Blue Start         33           Clickart Publications         37           Colormate         56           Decision Map         108           Dollars & Sense         112           Double Helix         273           Easy 3D         112           In Talk 3.0         146           Laser Author         44           Lazerstart 2.5         71           MAC Vision         224           Mainstreet Filer         75           Microsoft Chart         78           Microsoft Excel         296           Microsoft Works         198           Microsoft Multiplan         118
Grasp 3.0	Blue Start         33           Clickart Publications         37           Colormate         56           Decision Map         108           Dollars & Sense         112           Double Helix         273           Easy 3D         112           In Talk 3.0         146           Laser Author         44           Lazerstart 2.5         71           MAC Vision         224           Mainstreet Filer         75           Microsoft Chart         78           Microsoft Excel         296           Microsoft Works         198           Microsoft Multiplan         118
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Macroworks												٠,	26
Pin Point													. 66
Power Print													30
Newsroom										1	v.		. 45
Print Magic								4			ě.		. 45
Print Shop Compar	io	n.						14					. 30
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# SCIENCE

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699 This Week in Science

### Editorial 701 World Competition in Biotechnology etters College Calculus: How Should It Be Taught?: M. M. GRELLER; A. RALSTON ■ The 3 K Microwave Background and Olbers' Paradox: P. MARMET ■ Biology Teaching: S. L. Weinberg News & Comment 710 Europe Seeks Strategy for Biology ■ Focus on the Genome Field Test Data Inadequate, OTA Says 713 Science Budget Squeeze and the Zero Sum Game 714 Anti-Acne Drug Poses Dilemma for FDA 715 Fresh Look at Acid Rain Toxic Waste R&D Effort Stalled OECD Sets Guidelines for Cooperation 717 NAS Elects New Members 718 Animals of Invention Research News Obstacles to an AIDS Vaccine 721 Mathematics at 100 Exploiting the Insights from Protein Structure A New Tool Maker in the Hominid Record? ■ One Species . . . One Tool Articles Segregation of Form, Color, Movement, and Depth: Anatomy, Physiology, and 740 Perception: M. Livingstone and D. Hubel 750 Supernova 1987A!: S. E. Woosley and M. M. Phillips The Design of Molecular Hosts, Guests, and Their Complexes: D. J. CRAM Research Articles Visualizing Gene Expression in Time and Space in the Filamentous Bacterium Streptomyces coelicolor: A. Schauer, M. Ranes, R. Santamaria J. Guijarro, E. LAWLOR, C. MENDEZ, K. CHATER, R. LOSICK Reports Direct Observation of the Femtosecond Excited-State cis-trans Isomerization in Bacteriorhodopsin: R. A. MATHIES, C. H. B. CRUZ, W. T. POLLARD, C. V. SHANK The Nature of the Interior of Uranus Based on Studies of Planetary Ices at High Dynamic Pressure: W. J. Nellis, D. C. Hamilton, N. C. Holmes, H. B. RADOUSKY, F. H. REE, A. C. MITCHELL, M. NICOL

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Supernova 1987A—an exploding star. On 23 February 1987, light and neutrinos reached Earth from the brightest supernova in almost 400 years. "Before (left) and after (right)" photos show the star that exploded and the supernova shortly after outburst. Photographs were taken using the 3.9-meter Anglo-Australian Telescope in New South Wales, Australia. See page 750. [Courtesy of David Malin, copyright 1987, the Anglo-Australian Telescope Board]

		781		canthropus robustus from or: R. L. Susman	Member 1, Swartkrans	: Fossil Evidence for			
		784	Proreceptor	stant Diabetes Due to a Processing: Y. Yosнім. H. Kuzuya, H. Імига	asa, S. Seino, J. Whit	taker, T. Kakehi,			
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		<i>7</i> 96	Middle Miss J. A. WATER	issippian Blastoid Extino s	ction Event: W. I. Aus	ICH, D. L. MEYER,			
		798 Directional Selection and the Evolution of Breeding Date in Birds: T. PRICE, M. KIRKPATRICK, S. J. ARNOLD							
		800	CS Peptides	Lipopeptide Vaccines: I G. H. Lowell, W. R. INGER, W. T. HOCKME	BALLOU, L. F. SMITH,				
Technica	al Comments	803 New Zealand Marine Terraces: Uplift Rates: C. M. WARD; W. B. BULL AND A. F. COOPER Guanylate Cyclase and the Adrenal Natriuretic Factor Receptor S. A. WALDMAN, D. C. LEITMAN, J. ANDRESEN, F. MURAD; R. K. SHARMA							
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Book Re	views	817	Analysis, R.	ucture and Performance L. RANDS Sedimental Geochemistry of Abyss	tion in the African Rifts	s, R. G. Coleman			
Software	Software Reviews  824 PC Software for Artificial Intelligence Applications: H. Epp, M. Kalin, D. Miller								
Products	s & Materials	832		Sample Handling ■ Microercomputer ■ Fast 3-D citerature					
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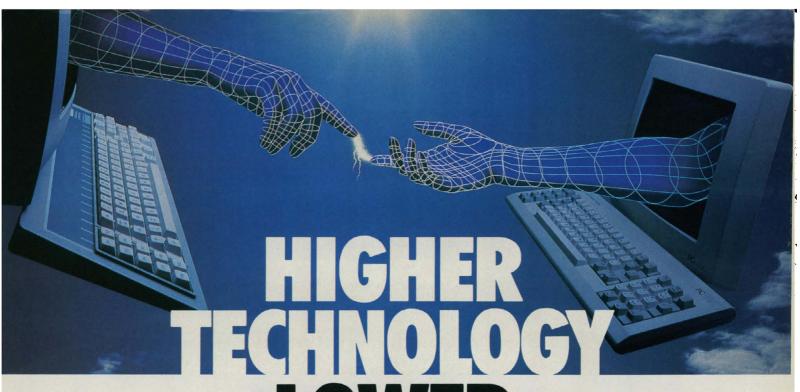
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# This Week in

# Science

# **Insight into vision**

visual image has color, depth, movement, form, and texture (page 740). Anatomic and physiologic studies in monkeys and perception studies in humans indicate that these components are processed by separate channels and that the segregated, parallel processing begins in the eye and continues as far into the brain as has been probed. There is also some supportive clinical evidence that distinct circuits process individual components of an image: stroke victims sometimes have selectively short-circuited vision, experiencing, for example, loss of only one segment of vision such as color vision. The experimental studies and the insights they provide into the structure, function, and evolution of the visual system are described by Livingstone and Hubel. It is enigmatic that visual images seem to be unified, when, in fact, processing keeps the parts of the whole separate from one another.

# Highlighting differentiation

TREPTOMYCETES are soil bacteria that form colonies in which physically distinct cells perform specialized functions (page 768). It takes several days for a colony to mature. Substrate mycelium cells initially burrow down into the medium on which the colony is growing; later aerial mycelium cells grow upward out of the substrate cells and eventually produce spores. When and where various genes are turned on and off within the colony during this complex differentiation process has now been studied by Schauer et al. through the use of a "reporter" gene cluster, luxAB. This cluster was transferred into streptomycetes cells next to different genes; as indigenous genes were expressed during differentiation, their promoters concurrently turned on nearby luxAB genes. The location of active genes was devilishly simple to identify because, when the luxAB gene cluster is turned on, the enzyme luciferase is produced and light is emitted: steps in morphologic differentiation are in this way spatially and temporally illuminated.

# **Fast-forward chemistry**

state-of-the-art laser system has made it possible to directly observe steps in a chemical reaction that goes from reactant to product within 500 femtoseconds (10<sup>-15</sup> seconds) (page 777). In the prototype experiment, the retinal prosthetic group of bacteriorhodopsin, a light-sensitive bacterial membrane protein that pumps protons through the membranes of Halobacterium halobium, was observed to isomerize from a trans to a cis configuration. Mathies et al. experimentally induced the chemical transformation with 60-femtosecond optical pulses; dynamic structural changes were followed spectrophotometrically with 6-femtosecond resolution. The spectroscopic signature (a direct monitor of a structural change) at 150 femtoseconds showed the prosthetic group twisting; the torsional displacement of the prosthetic group resulted in a permanent change in the molecule's geometry.

# **Inside Uranus**

TEW models of the interior of Uranus and new ways of testing these models have been developed since Voyager II took a closeup look at the distant planet in 1986 (page 779). Uranus consists of ice, gas, and rock and has a powerful but uneven magnetic field. Nellis et al. have investigated in the laboratory how methane, ammonia, and a mixture of water, ammonia, and isopropanol called synthetic Uranus—all thought to be prevalent on the planet—behave under simulated planetary conditions. During passage of a strong shock wave, the materials were molecularly ionized and dissociated. Conductivity was strong within certain high temperature and pressure ranges, accounting for (and constraining calculations of) the internal dynamo that produces the planet's magnetic field. All materials deep within the planet are likely to be dense and stiff; conditions in

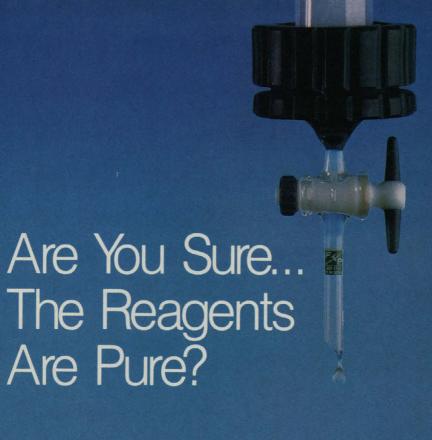
Uranus might thus be right for production of diamond-like carbon, nitrogen, and oxygen.

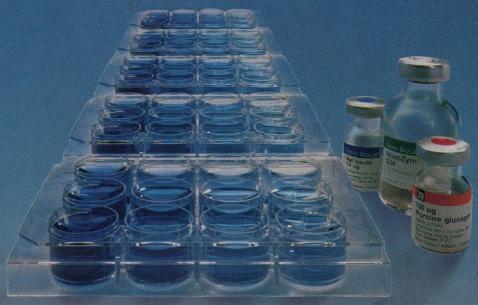
### Hand bones and tools

AND bones discovered at the 1.8-million-year-old Swartkrans site in South Africa indicate that the extinct hominid Paranthropus robustus could have made and used tools (page 781). Bone and stone artifacts suitable for digging or preparing foods were also found. It had not before been appreciated that P. robustus, like contemporaneous species Homo habilis and Homo erectus, had grasping fingers and a human-like thumb. However, the shapes, sizes, and muscle insertion sites of the 22 hand bones found indicate similarities to human hands and major differences from monkey and ape hands, which are adapted for power grasping during climbing. Paranthropus robustus had a small brain and large teeth, and the species' extinction has been attributed to a presumed combination of low intelligence and an inability to use tools and engage in other sophisticated behaviors; Susman suggests that, since it is now clear that Homo did not have a monopoly on the capacity to use tools, some other explanation must be found for the survival of Homo and not Paranthropus species (see also Lewin, page

### The bottom lines

**▼**HE explosion of the 10-millionyear-old blue star Sanduleak -69°202 into Supernova 1987A was astronomical in every way: the energy output in the first second exceeded by two orders of magnitude the energy output of the sun through its entire lifetime, supernova neutrinos from the star's collapse were detected deep within the earth, temperatures may have reached 10 billion kelvin in the exploding star's core, and astronomers and astrophysicists were afforded the opportunity of a lifetime—to test old theories and formulate new ones about how stars live and die (page 750).





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# World Competition in Biotechnology

irtually every developed country and many developing countries have targeted leadership in biotechnology as a national goal. In efforts to compete with the United States in applications of molecular biology to produce pharmaceuticals, would-be competitors are latecomers, with limited prospects. But in agriculture, the picture is quite different, and the United States could become second-rate. In the United States, generous support of biomedical research for decades led to great advances in molecular biology and to the training of a large number of talented people capable of advancing the biomedical sciences. Would-be competitors have limited reservoirs of expert personnel. Another favorable factor for the United States has been an abundance of venture capital, which was in short supply elsewhere. For some possible competitors the cost of development and clinical trials of pharmaceuticals (\$75 to \$100 million) needed to gain approval from the U.S. Food and Drug Administration is an inhibiting factor.

In contrast to federal support of biomedical research, funds for basic research in plant biology have been meager and the sums provided individual investigators have been tiny. Consequently, the knowledge base of the molecular biology of plants is limited. The level of conventional agricultural science in other countries is comparable to that here, and in some instances possibly superior. Expertise in the new biotechnology is widespread. Average yields of wheat per hectare in the Netherlands are more than twice those in the United States as are yields in the United Kingdom. There are mitigating factors, but these do not cancel the contrasts in yields. Our balance of agricultural trade has dwindled.

The crop surpluses in the United States have been used as an argument for curtailing research. However, if we move slowly in exploiting new biotechnology, we will lose competitiveness fairly rapidly. The time span and the investment required to introduce modified plants or symbiotic bioengineered microorganisms is small compared with that required for pharmaceuticals. Economics is one reason for pursuing vigorous development of plant biotechnology. Other benefits include diminished need for fertilizers and pesticides. Were productivity to be increased, less land would be required for crops, with less related soil erosion.

Major companies are devoting substantial funds to agricultural biotechnology. Their efforts are complemented by many new, small outfits. But progress has been greatly impeded by regulatory processes and legal actions. Although some caution in introducing new technologies is warranted, caution has been overdone. For example, there has been concern about introducing into the field a corn plant with a single altered gene. But Howard A. Schneiderman has pointed out that to convert a corn plant into a weed would require hundreds of genetic changes, because corn does not have a "weedy personality."\* Regulations should take into consideration the basic characteristics of plants into which a gene or genes are to be incorporated. Some of our crop varieties require human assistance for survival.

Another area in which progress is being impeded is in the introduction of beneficial soil microorganisms. A prejudice exists against organisms whose DNA has been modified by recombinant technology. But during most of this century, rhizobia (designed to enhance nitrogen fixation) have been added to millions of acres of agricultural soil. Roughly 10<sup>18</sup> rhizobia improved through mutation by chemicals or radiation are added each year. This release has not produced a negative environmental impact. Inoculants of selected mycorrhizae have greatly helped in restoring wastelands. In estimating potential hazards of introducing modified organisms, the gene's location in the genome is important. A gene located on a plasmid is much more likely to be transferred to another organism than is a gene that is part of a chromosome.

Advances in molecular biology have created great opportunities for advances in agriculture. The United States can persist in a policy of starving agricultural basic research and of overregulating biotechnology. Others may not follow such a path.

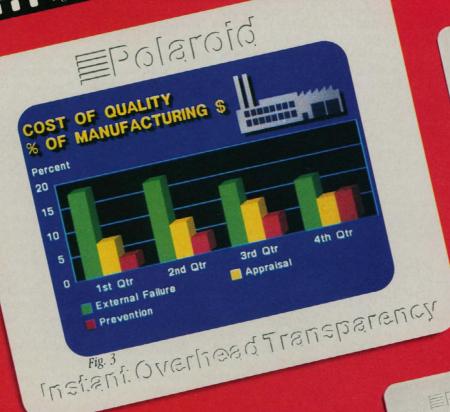
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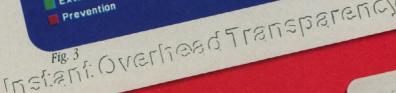
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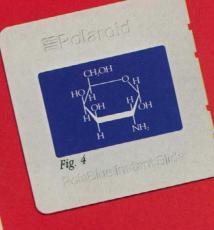
<sup>\*</sup>H. A. Schneiderman, "Biotechnology: A key to America's economic competitiveness in health care and agriculture," speech at the Second Annual American Society for Microbiology Conference on Biotechnology, San Diego, CA, June 1987.













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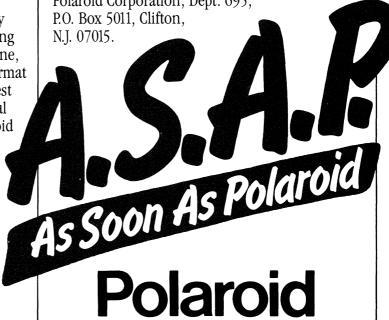
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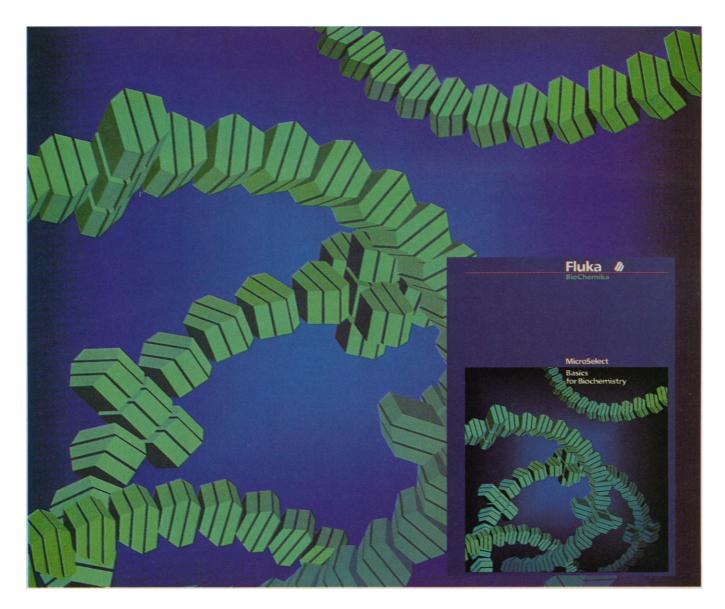
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### **Biology Teaching**

As a biology teacher with 28 years of experience, mostly in high school, and as a biology textbook author, I wish to join in the great frog debate: dissection versus frog integrity. My students did some dissection, yet I feel that requiring a student to sue in order to avoid participation is silly. If a student of mine occasionally objected to doing dissection, with or without specifying a reason, I retained my equanimity. The student simply retreated to the back of the room and took up another assignment. More often than not, with the rest of the class happily (and noisily) engaged, the dissenting student would edge forward to see what all the excitement was about. Sometimes the boy or girl would have second thoughts and ask to join in the dissection. When such an activity has to depend on legal sanctions, its educational value is likely

Why excitement at frog dissection? The reason is that I used live frogs, never pickled specimens of any kind. Just before class I pithed them—in a preparation room out of the students' sight. Thus the tissues and organs remained fresh and functioning. Excitement was generated by seeing the beating heart alternately flushing and paling. Often an enthralled boy or girl called me over and said in wonder: "Look, it's still beating!" Students marveled at an isolated heart continuing to pulsate in saline, at the intestine slowly writhing in peristalsis, at the gastrocnemius responding to nerve stimulation. These close-ups of glistening, functioning, living tissues generate a perception of life processes not matched by reading a textbook, by computer simulation, by audiovisual media, or by classroom lectures or discussions, however lively.

Yet dissections were not numerous in my course. I depended much more on behavioral studies of living animals, and to a lesser extent on studies of plants and microbes. Before the frog dissection lesson the students were given live frogs to observe in a variety of situations. An even more intriguing animal was the edible Burgundy snail, Helix aspersa. As science chairman in a high school on New York City's Lower East Side, with funds for supplies very limited, I had the task of providing 1500 biology students with living materials. But the Italian fish stores on Second Avenue provided snails at \$1 a pound (100 snails). With 2 pounds of snails and a few heads of lettuce, the student body had a fascinating lesson.

Frogs and snails, alas, are no longer available. Frogs have become scarce and expensive. *Salmonella* infections have restricted the importation of Burgundy snails, and when

they are available they cost far more than a penny apiece. But we should not despair. Resources of living materials are almost unlimited; the limiting factor is teacher ingenuity. If three or four beans are put in a jar of water and incubated overnight, on the next day the culture will be swarming with bacteria. Students seemed never to tire of watching the swarming culture, with spirilla and bacilli vigorously swimming about. Nor did I ever tire of watching the bacteria through the microscope.

These activities were not "experiments," and I did not call them that. We referred to them as "lab exercises." In a large high school class, with limited resources, a rigid time limitation, and a limited student background, it is difficult to do genuine, honest, scientific experiments. Many so-called student experiments are faked or fudged; or the results are determined a priori. The common pressure on students to get the "right answers" also destroys the scientific validity of the "experiments." In the kinds of activities described above there are no "right answers." Any function of an organism or of its organs that a student observed and accurately reported was the right answer.

Are these activities "science"? I think not. They are more in the nature of natural history or "pre-science." For this I offer no apology. James Watson began as a bird watcher. As a boy Charles Darwin collected beetles and birds' eggs. As the Chinese proverb puts it: "Tell me and I hear. Show me and I observe. Involve me and I understand." It is a standard axiom in education, honored more in the quotation than in the observance, that students learn by doing; learning is an active process. I tried to have my students "do," and many of them have gone into science as a profession or vocation.

Robert B. Eckhardt reports that he found his biology course dull (Letters, 18 Mar., p. 1361). Horrors! Living creatures are endlessly fascinating to people and especially to young people. Give students live animals or living tissues to work with and they cannot help but find biology interesting. More than a few of my former students have told me that they were first attracted to biology when they watched and tested the beating heart of a freshly pithed frog.

STANLEY L. WEINBERG Iowa Committee of Correspondence, Ottumwa, IA 52501

Erratum: In Mark Crawford's story "Budget crunch stalls Super Collider" (News & Comment, 1 Apr., p. 17) the caption accompanying the photograph was incorrect. The superconducting magnet pictured was 4.5 meters long, not 17 meters, as stated.

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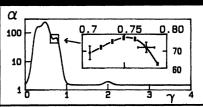
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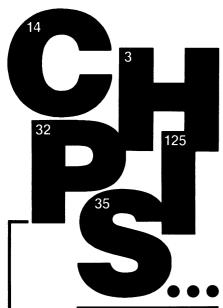


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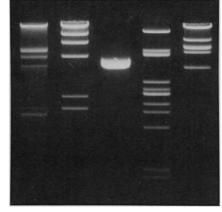
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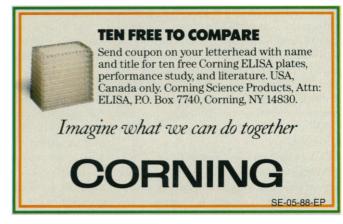
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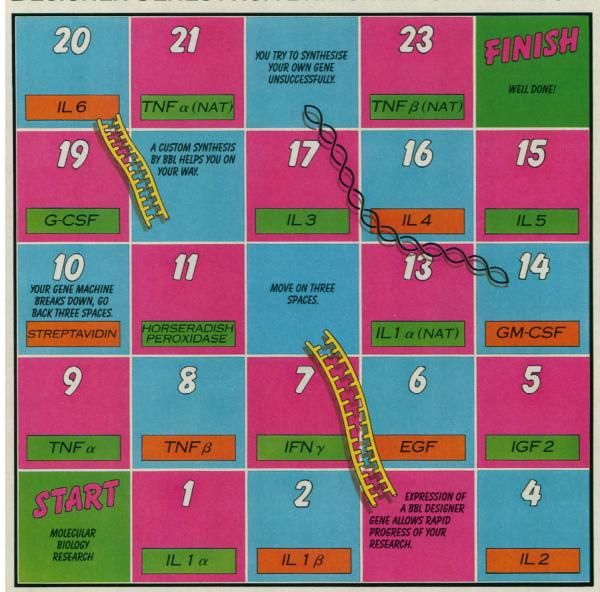
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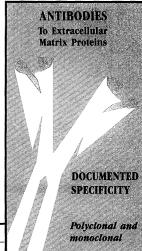
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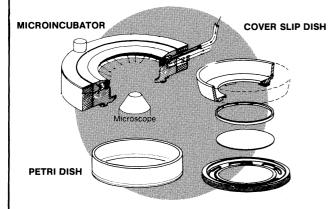
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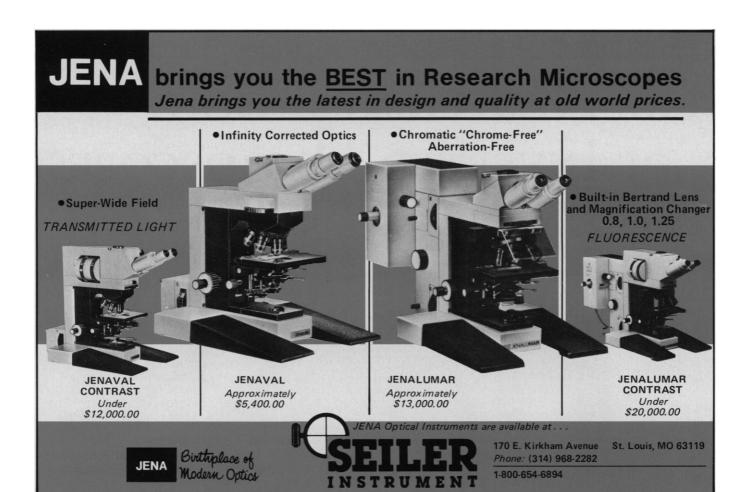
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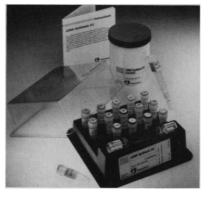


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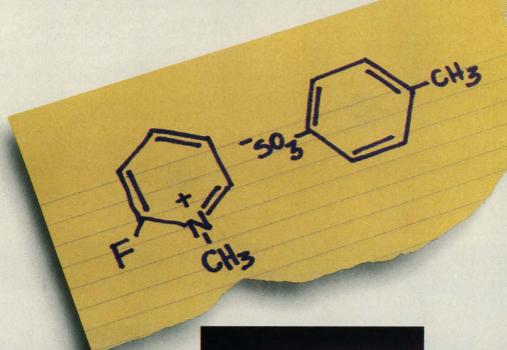
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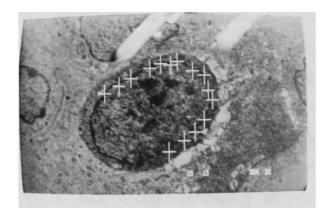
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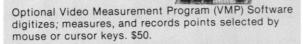


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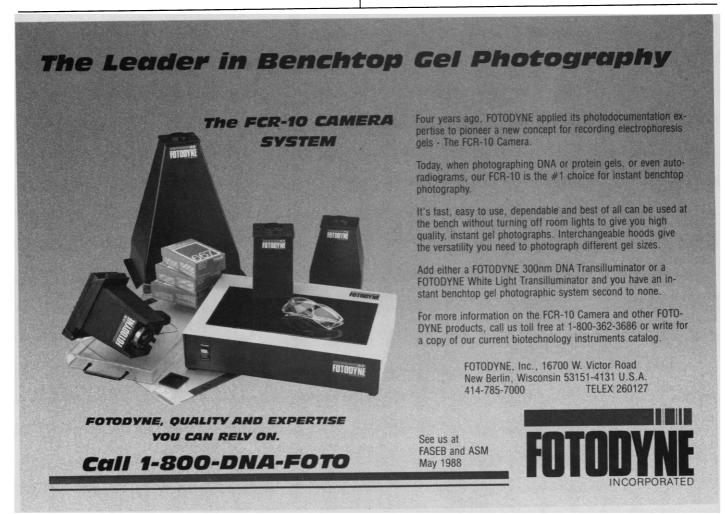
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### **CELL CULTURE**

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centrifuge tubes

centrifuges, cell-washing

centrifuges, continuous-flow

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centrifuges, hemocrit

centrifuges, micro

centrifuges, refrigerated: below 25,000 rpm

centrifuges, tabletop centrifuges, ultra

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chromatography, column: columns

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chromatography, HPLC supplies

chromatography, ion-exchange: HPLC specialty resins chromatography, ion-exchange: prepacked columns

chromatography, liquid: products

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electrophoresis equipment, gel

immunoelectrophoresis products

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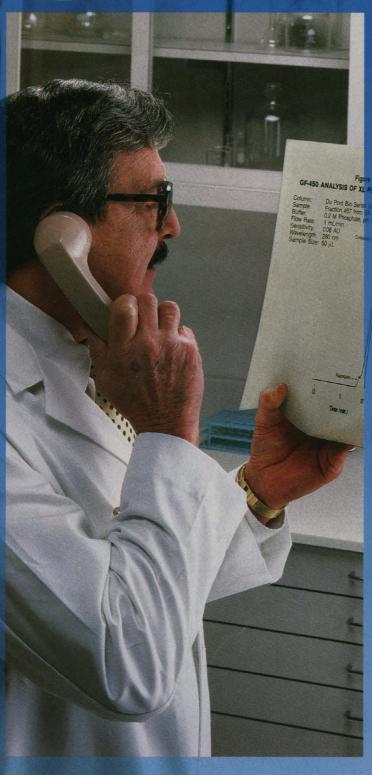
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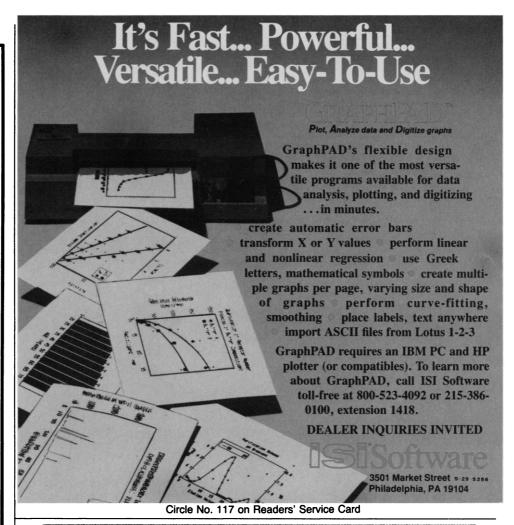
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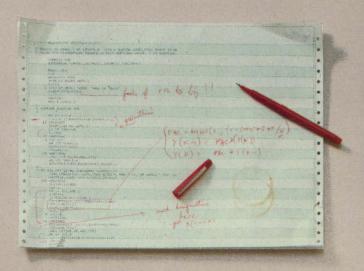
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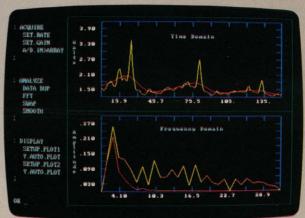
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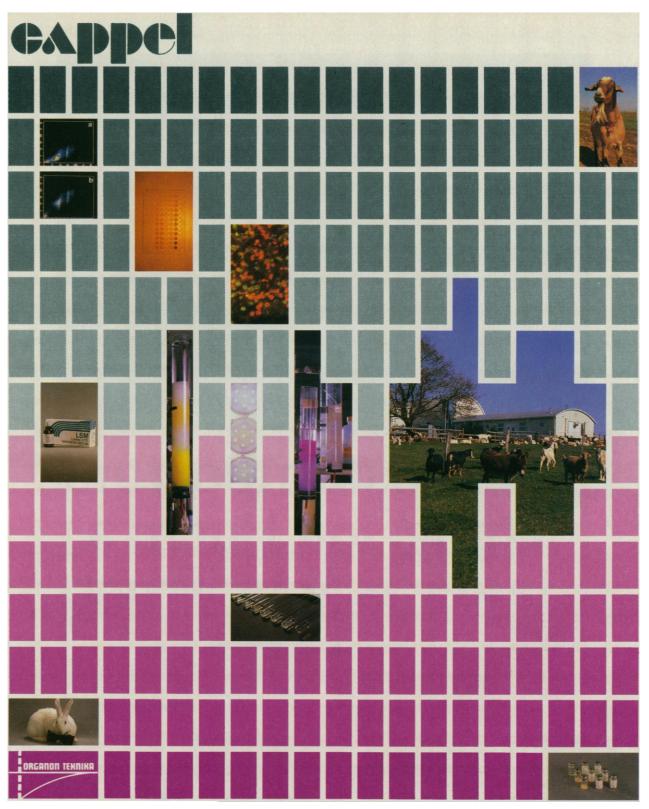
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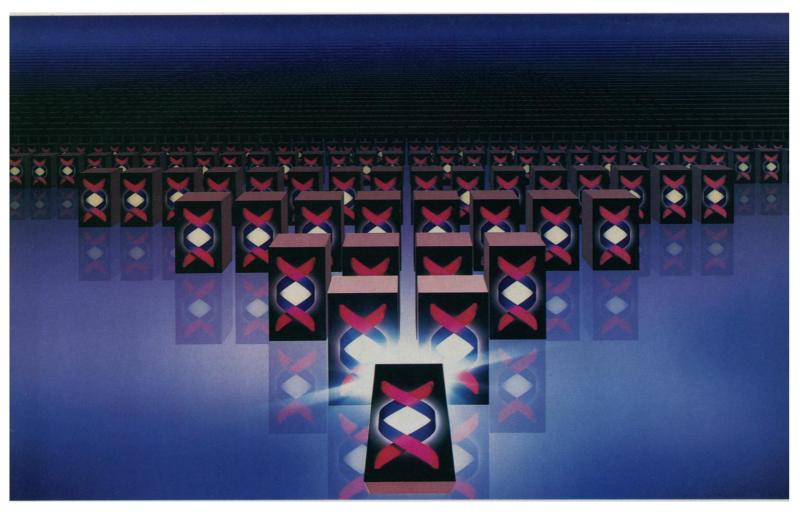
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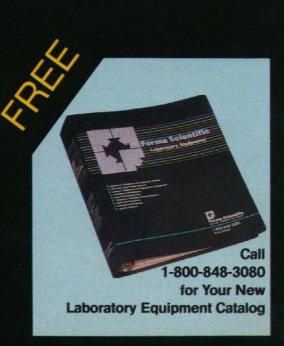
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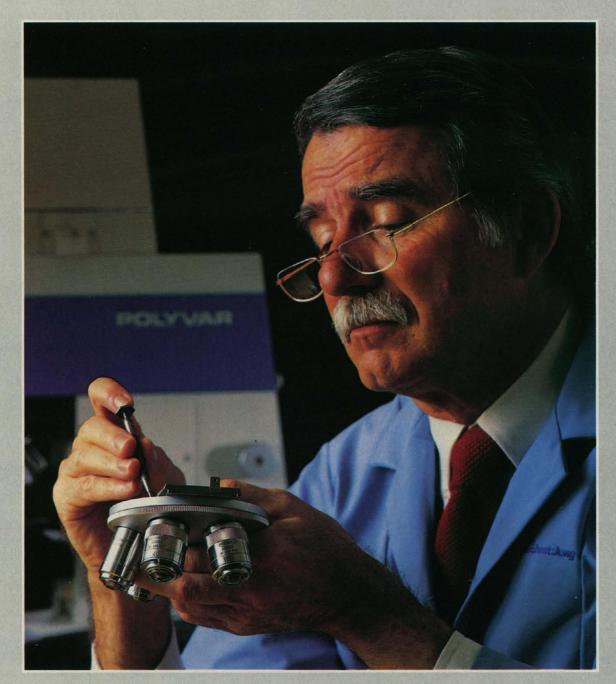
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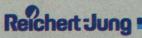


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