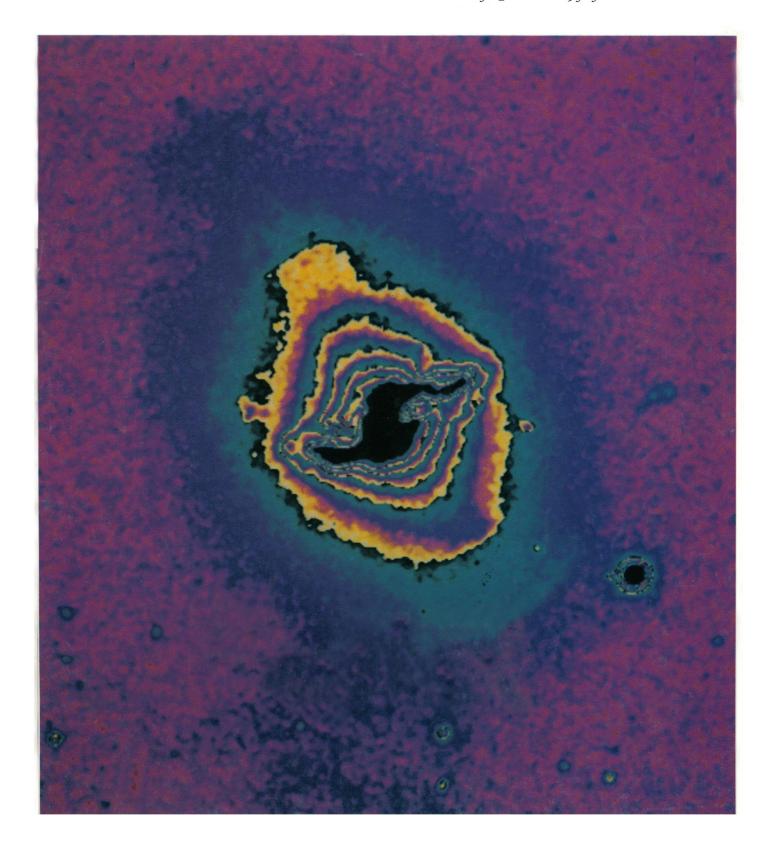
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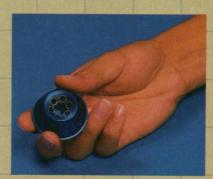
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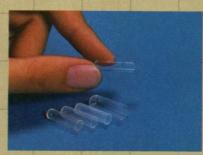
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Peculiar galaxy Arp 230, believed to be the remnant of a recent merger of two galaxies. The striking, sharp-edged ripples on either side of the center consist mostly of old stars that may have belonged to an intruding disk galaxy. Model calculations suggest that this intruder plunged into the main galaxy on a near-radial orbit. Gaseous debris, dust, and young stars form a disk seen here as black oval near the center. See page 227. [Photograph taken by F. Schweizer with a 4-meter telescope at Cerro Tololo, Chile, and computer-enhanced with Astronomical Image Processing System, Department of Terrestrial Magnetism, Carnegie Institution of Washington, Washington, DC 20015]

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This Week in SCIENCE

Galaxy dynamics

ALAXY formation is a continuing process (cover); its dynamics are becoming clearer through a combination of new space observations and computer-generated models (page 227). The complex orbits of galaxies apparently provide them with opportunities for close encounters with each other; they also interact with clouds of hot gases and satellites. Data gathered in 1983 by the Infrared Astronomy Satellite support the concept that collisions and mergers of galaxies, accompanied by star and quasar formation and the emission of great amounts of infrared energy, are regular and frequent events. Schweizer describes forces that may act during collisions and mergers, those which culminate in the "violent relaxation" that precedes the establishment of each new equilibrium configuration, and features of galaxies—such as tidal tails and ripples—that provide evidence for past merger events. These details of the origin, evolution, structure, and fate of galaxies suggest that not all galaxies were produced after a single Big Bang.

Industries, universities, and biotechnology

IOTECHNOLOGY companies invested about \$120 million in university biotechnology research laboratories in 1984, about 20 percent of the total dollar investment that year; this contrasts with the contribution (about 3 to 4 percent annually) of industries to other types of university research (page 242). Blumenthal et al. analyzed the relations of 106 biotechnology companies with university laboratories. Nearly half the companies supported university research in 1984. Benefits to the companies were seen in the abundance of patent applications (two to four times as many applications per dollar invested in university as in inhouse research) and trade secrets generated, which may translate into financial gain. Most arrangements were for 2 years or less and could have the longrange effect of shifting some university research from basic toward applied studies. Relations of university researchers with firms and with government sponsors differ. The generation of trade secrets may jeopardize a fundamental scientific and academic value, the free exchange of information. It is considered desirable that the government, which is the major sponsor of biotechnology research in universities, remain so.

Acid fog

YDROXYM ET HANESULFON ATE ion (HMSA) is a strong acid that can form in fog and cloud water in regions with sulfur dioxide air pollution (page 247). The fog and cloud droplets may then transport the pollutant to distant sites. HMSA forms when the pH of water droplets is at intermediate values and when sulfur dioxide is abundant; it is then best preserved in droplets by acidification and serves as an important reservoir of reduced sulfur. Munger et al. used ionpairing chromatography to measure HMSA in fog droplets collected at two sites in the southern San Joaquin Valley of California. There the oil industry generates the precursors for HMSA formation, and the humidity is high enough for droplets to form. Concurrently, the agriculture industry generates ammonia, which neutralizes the droplets and promotes decay of HMSA. Droplets from the site closer to the source of sulfur dioxide pollution had more HMSA, formaldehyde, and reduced sulfur than droplets at the more distant site, but at both sites there was evidence for the pollution-generated compound.

Ice and the martian terrain

¬Low of surface materials on Mars ◀ aided by ground ice may have led over time to the softening of the planet's terrain (page 249). Squyres and Carr used images collected by the Viking Orbiter to map landforms on Mars. Ground ice and water, generally thought to have once been widely distributed on Mars, appear to have migrated to higher, colder latitudes. Characteristic features called lobate debris aprons, concentric crater fill, and terrain softening formed as a result of ice. Their distribution supports the idea that ice creep varied with climate and that ice was unstable in the warmer equatorial regions but stable at higher latitudes where temperatures are below the frost point. Thus, with a cooling of the martian climate, desiccation has taken place at the equator while a stable ice layer has formed at the higher latitudes. Another explanation for the terrain softening—deposition of sediment by wind is considered a less likely cause of the distinctive patterns observed in the martian terrain.

Rearranged genes in leukemia

 \blacksquare ENES for β_1 interferon and for a Jamong those repositioned on chromosomes of patients with acute monocytic leukemia (page 265). Hybridization techniques showed that the interferon genes, normally clustered on human chromosome 9, were split: the α gene cluster remained on chromosome 9, while the β_1 gene was translocated to chromosome 11. The c-ets-1 proto-oncogene, normally found on chromosome 11, was found on chromosome 9. Exactly how these genes in their new locations participate in the malignant transformation of acute monocytic leukemia is not known. Diaz et al. speculate that when the oncogene is translocated near the interferon cluster, regulatory sequences that activate the interferon genes may also activate the oncogene. Although proliferation of monocytic cells is not likely to represent overactive genes of the interferon system (since this system typically produces antiproliferative effects), proliferation could occur if the activated oncogene stimulates an alternative pathway that generates tumor cells.

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The Social Process of Science

eing well-informed about science is not the same thing as understanding science." Therewith, James Bryant Conant discounted the popular interest in science roused by the thunderclap that ended World War II. Now as then, in a world transformed by the application of scientific knowledge, people put that knowledge in the same category with what they know by revelation or other received authority. What needs to be understood is how, scientifically, we come to know what we know.

Scientists know nothing for certain. The advancement of science is a social process, a public process, and yet an intensely private one. Societies that would enjoy its material benefits must understand science in both its aspects.

"The truth of an idea," William James perceived at the turn of this century, "is not a stagnant property inherent in it. Truth happens to an idea . . .[by] the process of . . . its verification." In the private process, the scientist must face the singular loneliness of the sovereign. He can accept no authority but his own conscience and judgment.

The work proceeds in ways very different from that suggested by its impersonality in formal publication. "The process I want to call scientific," the physicist Percy Bridgman wrote, "is a process that involves the continual apprehension of meaning . . . accompanied by the running act of checking to be sure that I am doing what I want to do, and of judging correctness or incorrectness. This checking and judging and accepting . . . are done by me and can be done for me by no one else. They are as private as my toothache, and without them science is dead."

The intensity of this private process—its toothache—is raised by the fact that it is integrally public. It is intended for publication. Without publication, science is dead.

Upon publication, verification of the work proceeds. As the sociologist Robert Merton has observed, "Only after the originality and consequence of the work have been attested by significant others can the scientist feel reasonably comfortable about it." Merton's term for this public process is "'Communism'.... The substantive findings of science are the product of social collaboration and are assigned to the community The scientist's claim to 'his' intellectual 'property' is limited to that recognition and esteem which, if the institution functions with a modicum of efficiency, are roughly commensurate with the significance of the increment brought to the common fund of knowledge."

The remarkable fact, established by the open public record of science, is that this social process functions with high efficiency. It is not that scientists are more dedicated, honest, and selfless than other citizens; they are disciplined to behave that way by their collaboration. Error and fraud are exposed sooner rather than later by the communal process of verification. With equal reliability the consensus of the community distinguishes the significant from the trivial. This is the more remarkable considering what Merton calls "the basic uncertainty of genuinely independent originality in science.'

Thus Niels Bohr was once prompted to observe of a radical and baffling proposal by the aging Werner Heisenberg: "Yes, it is crazy, but it is not crazy enough!"

It has been said that science will flourish only in a society that cherishes its norms. The reason, openness, tolerance, and respect for the autonomy of the individual that distinguish the social process of science, however, are norms desirable in every human community. They describe a world in which, we can agree, all of us want to live.

Happily, the social process of science brings along the means to realize its values. For it finds convincing verification in the technologies it begets. During the past four centuries science has been liberating increasing numbers, now nearly one-third, of mankind from toil and want and even from submission to received authority. No national constitution written in this century has failed to hold out the promise, at least, of political and economic democracy. The people of the world—if nations can keep the peace—may see this revolution in the condition of man fulfill its promise in the next century.—GERARD PIEL,* Chairman, Scientific American, 415 Madison Avenue, New York 10017, and President, AAAS.

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^{*}This editorial is excerpted from a public lecture given at Moscow State (Lomonsov) University, Moscow, U.S.S.R., on 25 November 1985, on the occasion of the conferral by the university of the degree Doctor Honoris Causa.

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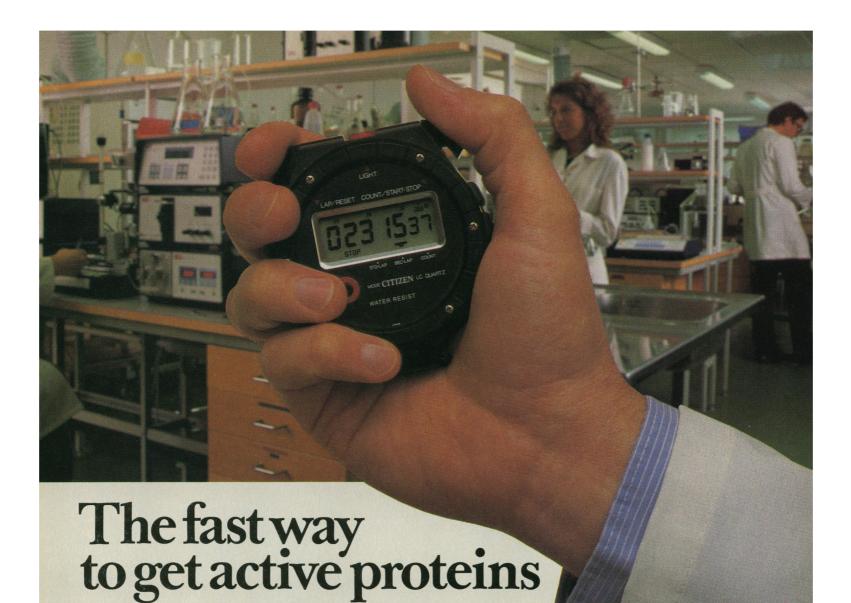
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Zeiss Instrumentation for the New Biology

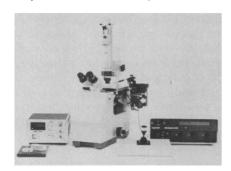
Observe, analyze, quantify, heat, manipulate and inject with advanced systems from Zeiss. Here are some of them:

Faster, More Accurate Micromanipulation/ Microinjection

For faster, more accurate results in bioresearch involving micromanipulation and microinjection, Zeiss has combined a Zeiss inverted microscope with an electrically-operated micromanipulator, a high speed microinjector with three adjustable injection pressures, and a controlled temperature stage, all in a single compact workstation. The systems are available with a choice of the Zeiss IM, IM 35 or ICM 405 Invertoscopes.

Equipped with DC motor controls, the **Micromanipulator** offers exceptionally smooth, responsive and vibration-free X, Y, and Z movement at speeds ranging from 0 to 0.1 mm/sec. For positioning of X-Y movements parallel to the stage plate a special holder permits precise mounting of the micromanipulator at any elevation to the right or left of the stage plate.

With three adjustable injection pressures for holding, injecting and cleansing, the **Microinjector 5242** increases injection rate from the 100 cells a day typical of manual systems to 500 cells per hour. Other features: dosages in the picoliter range; an LED display of number of injections, pressure, mode and automatic or manual time; an acoustic signal when the injection sequence has been completed.



The **Heating Stage M** has built-in transistors (no electrical interference) and a temperature range of 3°C above ambient to approximately 50°C, with a stability of better than 0.2°C. The control unit monitors both programmed temperature and actual temperature for the heating stage and preheating plate.

The new Zeiss Micromanipulation/ Microinjection Systems provide the researcher in physiology, cell biology, neurobiology with a complete workstation capable of transfering materials into the living cell with new levels of speed and accuracy.

EM 902: New Tool for Biomedical Research

The Zeiss EM 902 is an advanced analytical TEM that enables biomedical researchers to localize specific elements in the ultrastructure of tissues and cells. In employing a new technique, Electron Spectroscopic Imaging (ESI), the EM 902 operates on the principle that electrons in various elements nitrogen, oxygen, phosphorus, calcium and sodium – exhibit different levels of energy loss. A fully integrated electron energy-loss spectrometer registers the electron energy-loss spectra (EELS) of a specimen, producing a high-resolution distribution map of the elemental components. The new TEM achieves an unprecedented point resolution as low



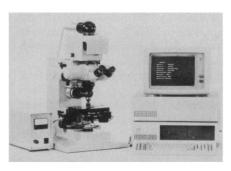
Extremely simple to operate, the EM 902 is suitable for research and routine applications. Besides the ESI technique, the EM 902 can be used for contrast enhancement on thin unstained sections, on thick sections up to 1 µm (the chromatic aberration equals that of a 1 million volt TEM), and for unlimited conventional electron microscopy.

New UMSP Microscope Photometer System

Flexible in design and easy to use, the new Zeiss UMSP System for highresolution quantitative microscopy offers the biological researcher maximum problem solving capabilities.

The UMSP System teams the Zeiss UEM microscope with a modified IBM PC/XT, and user-friendly, state-of-the-art software developed for spatial and spectral analysis. Data acquisition is accomplished through either solid state camera or photomultiplier tube (PMT) depending on the specific application. The system can be configured with other Zeiss microscopes such as the Universal and Standard 16.

Powerful applications software such as the biologically-oriented Image Scan enable a researcher to measure the



grey level distribution and thereby generate a density map of a single cell. Simplicity in design of all hardware and software gives the user clear, logical and functional sequences for easy use and adaptable control of the system.

Video-Enhanced Microscopy for Submicron Observation

New techniques of video-enhanced microscopy from Zeiss allow the researcher to detect and record intra- and intercellular details at the submicron level. The recommended systems combine the famous Zeiss Axiomat™ microscope with a choice of compatible video units.

The unparalleled mechanical stability of the Axiomat microscope and its internally corrected planapochromatic lens produce the excellent image quality at high magnification required by techniques such as the patented Allen Video-Enhanced Contrast (AVEC) method.

Features such as zoom capability can vary the image transfer as desired, optimizing the size of the specimen on the TV monitor with no loss of resolution. The image is further enhanced by electronic manipulation of the video signal. The instrument's single set of objectives covers the range of most commonly utilized optical techniques: brightfield, darkfield, phase contrast, POL and DIC.

In the inverted configuration for biological applications, the Axiomat microscope is connected to a video system by an optical interface developed by

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Tales of Chemists:

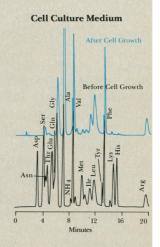
The Amino Acid Story



"Ion exchange is really specific for analyzing amino acids in complex samples like cell culture media. The problem is that it's just too slow. So I investigated prederivatization HPLC. It was faster, but now I had interferences to contend with. Not to mention all the time I ended up spending on sample preparation.

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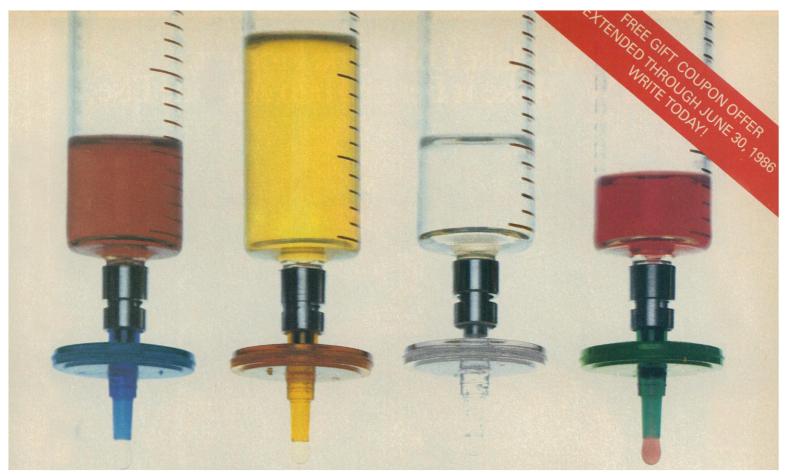
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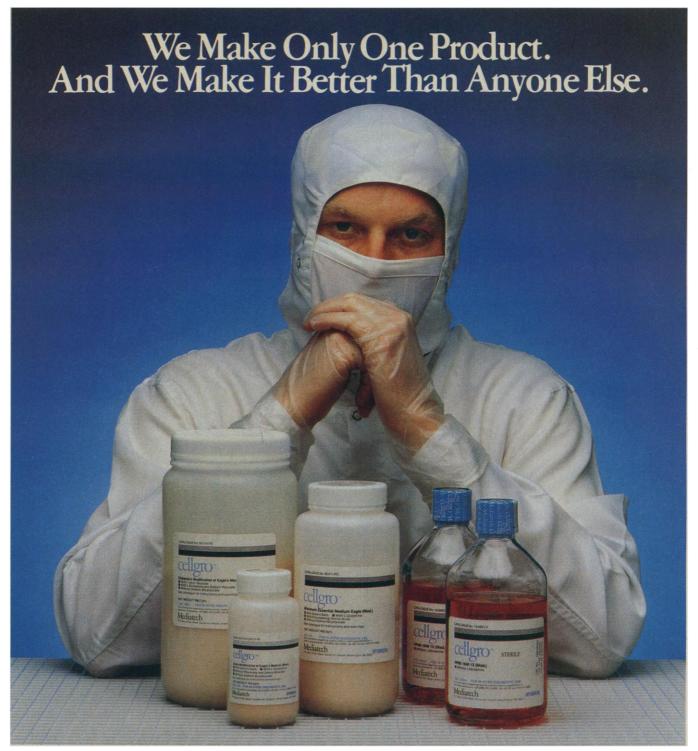
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Paris, April 7-9

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Dexamethasone-Suppressible Hyperaldosteronism

Rome, June 5-6

Scientific Organization: M.I. New (USA)

Corticosteroids and Peptide Hormones in Hypertension

Mannheim, Sept. 6-7

Scientific Organization: E.G. Biglieri (USA), F. Mantero (I) and P. Vecsei (D)

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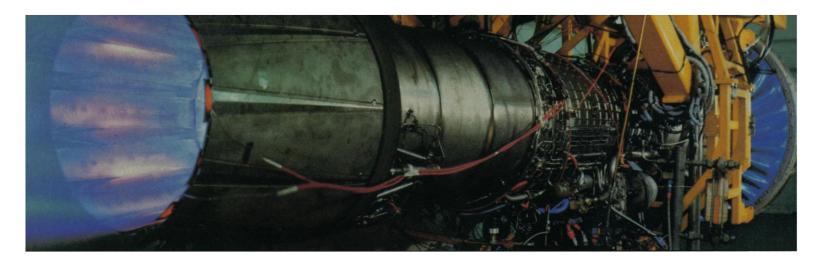
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Scientific Organization: E. Diczfalusy (S) and M. Bygdeman (S)



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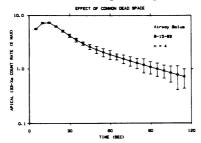


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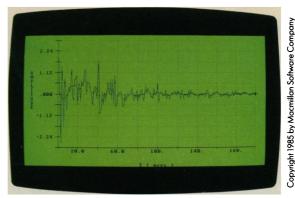
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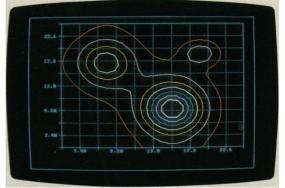
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