

# Southerns/Northerns: Electrophoresis, Blotting, and Crosslinking in 2.5 Hours Instead of 30.

Stratagene has streamlined agarose gel electrophoresis and blotting. The system decreases the time required, from sample loading to prehybridization ten fold.

## STRATAGENE METHOD—TIME 2.5 HOURS

2 HOURS 15 MIN 15 MIN 30 SECONDS

Size Fractionate DNA Stain Gel Transfer To On Agarose Gel And Pretreat A Solid Support

12 HOURS 4 HOURS 12 HOURS CONVENTIONAL METHOD—TOTAL TIME 30 HOURS

#### VAGE™ System

The VAGE<sup>™</sup> vertical agarose/acrylamide gel electrophoresis system allows the casting of agarose or acrylamide gels in the unit. Nucleic acids can be electrophoresed through a 3 mm, 0.8 % vertical agarose gel in less than two hours with excellent resolution (Figure 1).

Because the gels are thin, staining, depurination, and denaturation can be accomplished in 15 minutes.

Crosslink DNA To

2 HOURS

A Solid Support

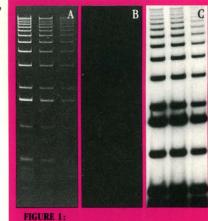
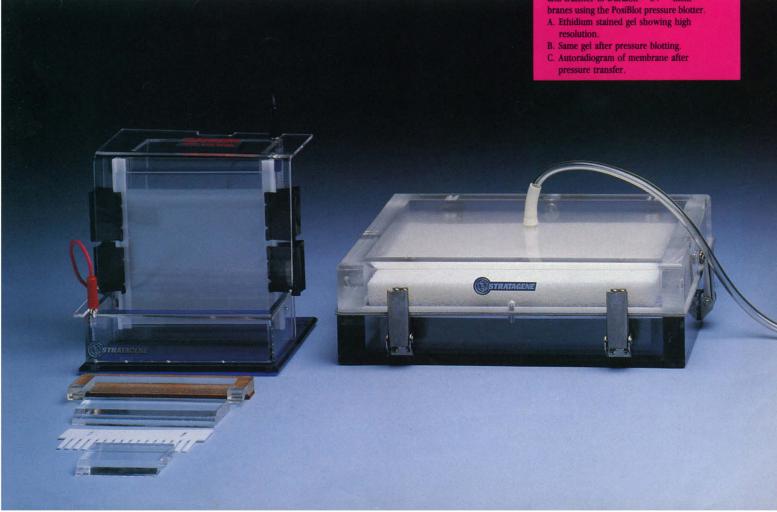


Figure Legend: Fractionation of end labeled DNA markers on 3mm thick 0.8% agarose by the VAGE apparatus and transfer to Duralon—UV<sup>TM</sup> membranes using the PosiBlot pressure blotter



#### PosiBlot<sup>™</sup> Pressure Blotter

# Buo01 Buo01 Buo01 Buo01 Buo01 Buo01 Buo01 Buo08 Buo08 Buo08 Buo08 Buo08 Buo08 Buo08 Buo08 Buo09 Buo08 Buo09 Bu

#### FIGURE 2:

Figure Lengend: <sup>32</sup>P end-labeled lambda Hind III markers were electrophoresed in 0.8 % agarose. The DNA was then transferred to a nylon membrane with a vacuum blotter at 30mm Hg below atmospheric or with the PosiBlot pressure blotter at 100mm Hg above atmospheric. Both transfers were carried out for 15 minutes. As can be seen, pressure blotting transferred significantly more DNA in the same period of time, especially in the higher molecular weight range (largest band is 23 kilobases).

The PosiBlot<sup>™</sup> positive pressure blotter permits the transfer of nucleic acids in 1/3 the time of vacuum blotters and 1/50 the time of capillary blotting (Figure 2). Pressure blotting does not dehydrate gels as do other methods. This allows the use of substantially higher

pressure differentials, compared with vacuum blotting, without gel collapse. The PosiBlot apparatus reduces blotting time to 15 minutes.

#### Stratalinker™ UV Crosslinker

The Stratalinker™ UV Crosslinker fixes nucleic acids to solid supports such as nitrocellulose or nylon membranes, in less than one minute. This compares favorably to vacuum baking, which requires 2 hours. The Stratalinker actually monitors the ultra violet energy flux and deactivates the light source upon reaching the user-programmed energy level (Figure 4). Figure 3 shows an autoradiogram of a human genomic Southern blot performed using the VAGE, PosiBlot and Stratalinker all in 2.5 hours.

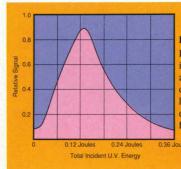


FIGURE 4:

Figure Legend: The effects of altering the incident energy for crosslinking nucleic acids to nylon membranes. The significant drop in signal intensity at energy levels below and above 0.12 Joules demonstrates the limited optimal range for UV treatment.

Stratagene offers a full selection of nitrocellulose, reinforced nitrocellulose and nylon membranes. Each membrane is stringently lot tested to ensure consistency when performing Northern and Southern blotting. Please call Technical Services for detailed information on Stratagene's time saving blotting systems and membranes.



#### FIGURE 3:

Figure Legend: Autoradiogram showing the resolution of 2.8 and 1.3 Kb Msp I RFLP alleles revealed by a cystic fibrosis human DNA probe using the VAGE, PosiBlot and Stratalinker all in 2.5 hours



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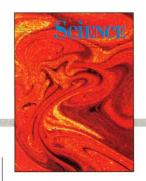
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Fractal pattern formed by small particles floating on a fluid undergoing complicated motion. High particle density (yellow-blue) indicates regions of past compressive surface flow or downwelling. Low particle density (black-red) indicates past upwelling. The pat-

tern geometrically summarizes the long-term, chaotic behavior of typical elements of the fluid surface; it is the strange attractor of a random dynamical system. See page 335. [Image: John C. Sommerer]



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#### THIS WEEK IN SCIENCE

edited by PHIL SZUROMI

#### Space shuttle astronomy

Because Earth's atmosphere strongly absorbs ultraviolet light, astronomical objects that emit in this wavelength range must be observed with telescopes in space. The space shuttle Columbia carried the Astro-1 ultraviolet observatory into space in December 1990 for a 9-day mission. Davidsen (p. 327) reviews results from one of the instruments aboard the shuttle, the Hopkins Ultraviolet Telescope, which obtained spectrophotometric data on 77 objects at wavelengths from 912 to 1850 angstroms. Among the findings were evidence for a hot corona around the Milky Way, new information about galactic nuclei, and a measurement of the properties of the ionized interstellar medium.

#### Metal nanocrystals in carbon particles

During the synthesis of carbon nanotubes, polyhedral carbon nanoparticles (about 20 to 40 nanometers in diameter) also form. These particles can be thought of as nested giant fullerenes that surround an internal cavity. Ruoff et al. (p. 346) describe the synthesis of carbon nanoparticles in which the internal cavity is filled with single crystals of a metal, the  $\alpha$ phase of LaC2. This carbide is metallic but is hygroscopic at room temperature and readily converts to La<sub>2</sub>O<sub>3</sub>. High-resolution transmission electron microscopy revealed that this carbon-encapsulated form of LaC<sub>2</sub> is stable for many days under ambient conditions.

1\_

#### After the flood

Evidence for catastrophic flooding is observed on Earth, where

enormous floods have resulted from rupture of the ice dams of large glacial lakes, as well as on other planets, notably Mars. Baker et al. (p. 348) briefly review known events and describe a cataclysmic flood in the Altay Mountains in Siberia. The flows inferred from the morphology of landforms produced by the flood rival those associated with the Channeled Scabland region of the northwestern United States. Peak flows are inferred to have exceeded 18 million cubic meters per second, flow depths were 400 to 500 meters, and flow velocities may have ranged up to 45 meters per second.

#### **Gecko rivalry**

Invading species frequently out compete native species, but the mechanisms are often unknown. Petren et al. (p. 354) investigate the mechanisms of the displacement of an asexual native gecko lizard species by an invading sexual species in the Pacific Islands. They find that light on buildings is a major factor in this competition because it attracts insects, the lizards' food source. The smaller asexual native geckos avoid the larger sexual geckos at these concentrated food sources, which leads to a decline in the condition of the asexual native geckos.

#### **Open for business**

The formation of an RNA polymerase-promoter complex suitable for transcription initiation requires conformational changes in both RNA polymerase and the promoter DNA. Suh *et al.* (p. 358) find that two different open complexes exist when RNA polymerase is complexed with the  $\lambda$  phage promoter Pr. Only the open complex formed in the presence of magnesium, however, has single-stranded DNA at the transcription start site.

#### Transporting tRNA

Export of transfer RNA (tRNA) molecules from the nucleus can be slowed by specific base mutations. Singh and Green (p. 365) isolated a 37-kilodalton protein that binds to wild-type tRNA<sup>Met</sup> but not to mutants that have reduced export activity. This protein is the glycolytic enzyme glyceraldehyde-3-phosphatedehydrogenase (GAPDH). In addition to being a cytoplasmic enzyme that catalyzes the reaction between its cofactor, nicotinamide adenine dinucleotide (NAD+) and carbohydrates, GAPDH is also present in the nucleus, where it selectively binds tRNAs. Binding of tRNA and NAD+ is competitive, and these molecules may share a binding site.

#### Physical basis of a fractal

Fractals have been discerned in everything from broccoli to the coastline of England, but few studies have rigorously connected the fractal dimensions calculated to the underlying physical principles of plant growth or shore erosion. Moreover, the mathematical fractals (known as strange attractors) that characterize many dynamic systems are abstract geometric entities. Sommerer and Ott (p. 355; cover) describe a strange attractor in physical space—tracer particles floating on a fluid surface. The particles produce a physical fractal pattern when the fluid flow is pulsed; the authors can then establish a quantitative connection between the fractal dimension of the pattern and the process that produced it.

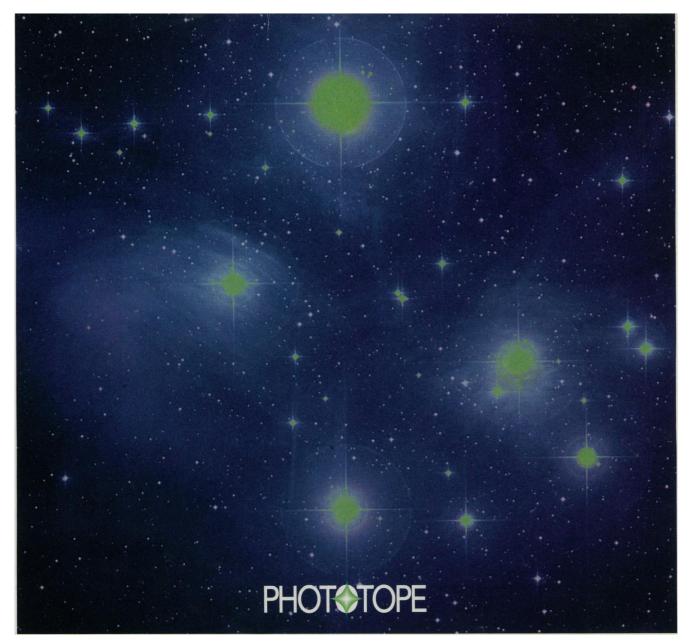
#### Detecting tumors

Some tumors can escape attack by the immune system even though they express potentially antigenic peptides; for example, if the costimulatory molecule B7 is not also expressed, then an effective T cell response against the tumor may not be elicited. Townsend and Allison (p. 368) transfected B7 into a murine melanoma cell line that expresses major histocompatibility molecules but is only weakly immunogenic. Mice implanted with melanoma cells that expressed B7 either did not develop tumors or developed tumors that regressed over time. Immunization with the B7 tumors also protected mice against the growth of implanted tumors that did not contain B7. This response was mediated by CD8+ cells. Appropriate costimulation can apparently "jump start" the cytotoxic T lymphocyte response without exogenous help from CD4+ cells (see news story by Travis, p. 310).

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#### Delivering nerve growth factor

Nerve growth factor (NGF) is a potential therapeutic agent for diseases in which cholinergic neurons degenerate, such as Alzheimer's disease, but the blood-brain barrier (BBB) prevents the delivery of this protein from the blood stream. Friden et al. (p. 373) conjugated NGF to an antibody to the rat transferrin receptor, which is responsible for delivering iron across the BBB. Injection of this NGF-antibody conjugate into rats increased the survival of both cholinergic and noncholinergic neurons that had been transplanted from the medial forebrain into the anterior chamber of the eye.



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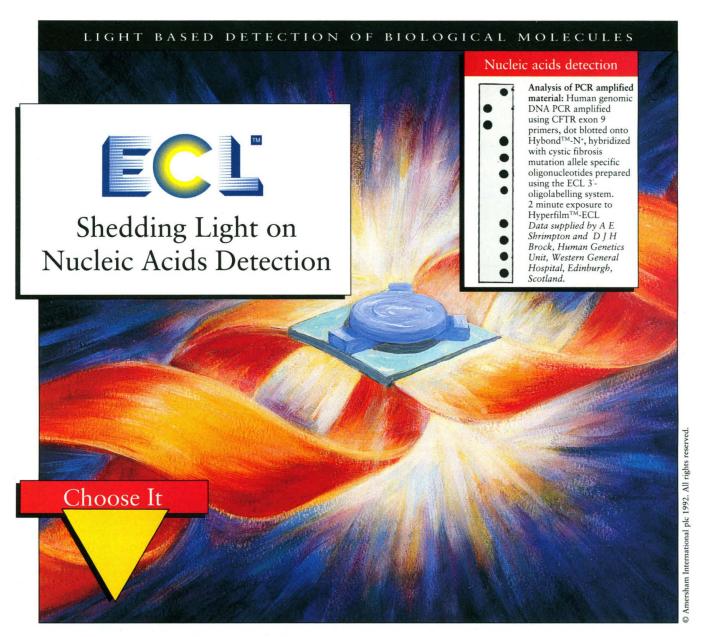
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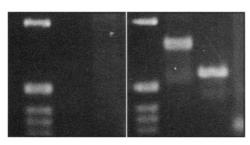
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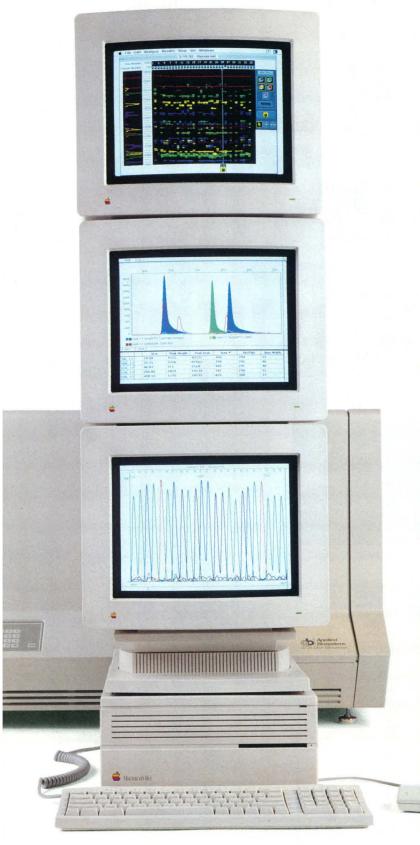
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<sup>\*</sup> PCR is covered by U.S. Patent No. 4,683,202 issued to Cetus Corporation.

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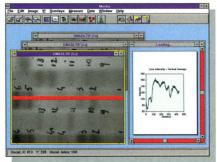
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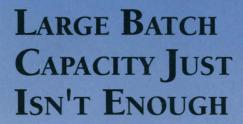
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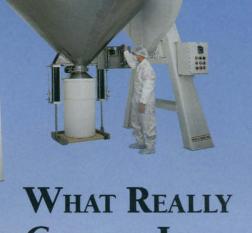
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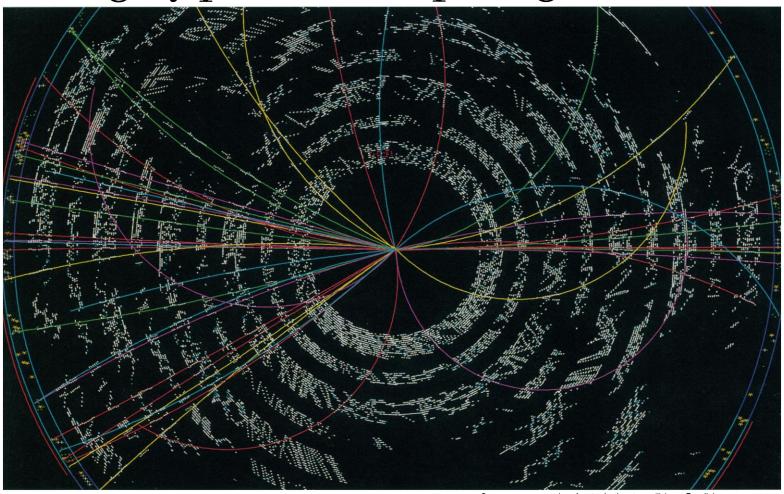
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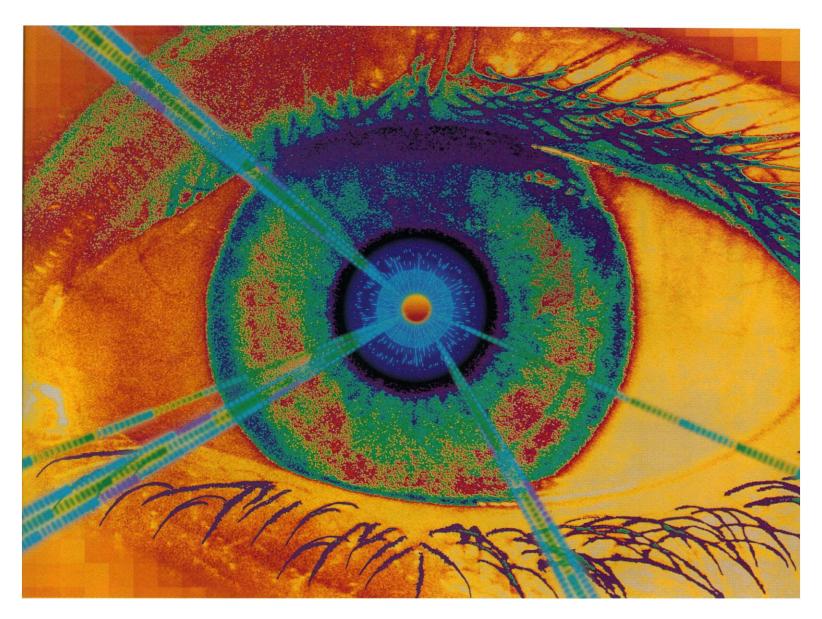
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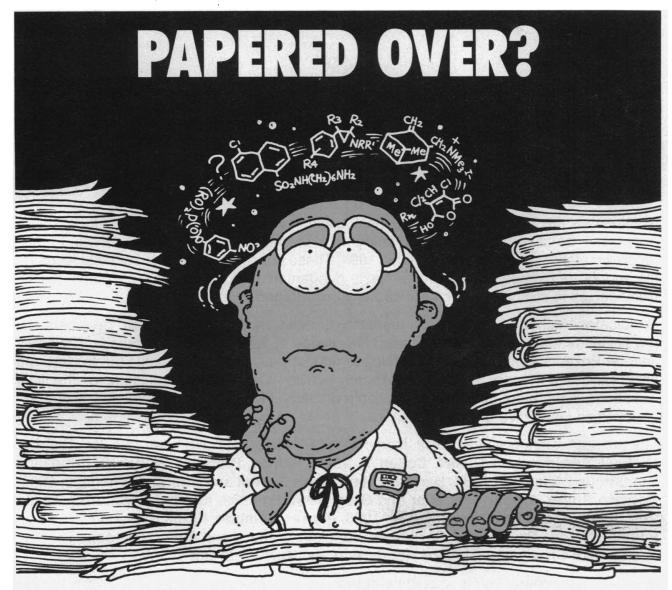
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Candidates (either individuals or research groups) must be nominated by scientists, physicians or institutions having detailed knowledge of the candidates' research. The Louis Jeantet Prize for Medicine is not intended to honour past accomplishments but to help and encourage the winners' continued research activity. Candidates shortlisted for the final selection will therefore be asked to provide a research project to which the financial support of the Prize could give decisive impetus.

The previous winners of the Louis Jeantet Prize for Medicine have been Luc Montagnier, Michael Berridge and Désiré Collen in 1986, Sidney Brenner, Walter Gehring and Dominique Stehelin in 1987, Bert Sakmann, John Skehel and Rolf Zinkernagel in 1988, Roberto Poljak, Walter Schaffner and Greg Winter in 1989, Nicole Le Douarin, Harald Von Boehmer and Gottfried Schatz in 1990, Pierre Chambon, Frank Grosveld and Hugh Pelham in 1991, Paul Nurse, Christiane Nüsslein-Volhard and Alain Townsend in 1992, Jean-Pierre Changeux, Richard Henderson and Kurt Wüthrich in 1993.

The following general points should be noted:

- 1. The Prize is limited to researchers working in European countries, members of the Council of Europe. The candidates need not, however, be themselves nationals of any of these countries.
- 2. Applications must be submitted, confidentially, on the official forms only. These are obtainable from:

The Secretary of the Science Committee
The Louis Jeantet Foundation for Medicine
P.O. Box 277
CH—1211 GENEVA 17
Switzerland

Further information will be sent with the nomination form.

3. The deadline for applications is February 15, 1993.

The name(s) of the winner(s) of the 1994 Louis Jeantet Prize for Medicine will be announced in January 1994. The Prize Ceremony will take place in Geneva (Switzerland) in April 1994.



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