EMFs are similar to the "stress response" used by all cells in reaction to harmful stimuli in the environment (1). Readers may wish to refer to volume 250 of the Advances in Chemistry Series (2) for a peerreviewed, balanced coverage of the issues.

Martin Blank

Department of Physiology and Cellular Biophysics, Columbia University College of Physicians and Surgeons, 630 West 168 Street, New York, NY 10032, USA

References

- M. Blank, O. Khorkova, R. Goodman, *Bioelectro-chem. Bioenerget.* **31**, 27 (1993); *ibid.* **33**, 109 (1994); R. Goodman *et al.*, *ibid.*, p. 115.
- M. Blank, Ed., *Electromagnetic Fields: Biological Interactions and Mechanisms* (American Chemical Society, Washington, DC, 1995).

Medical Imaging

The article by James Glanz "Computer processing gives imaging a sharper view" (News, 8 Sept., p. 1338) asserts that "the most important imaging medium of all is turning out to be the computer." The field of medical imaging has certainly made major advances since the discovery of x-rays by Roentgen some 100 years ago. The basis of medical imaging, however, is to use intrinsic differences in some physical property of the patient, such as the linear attenuation coefficient for x-rays or the acoustic impedance for ultrasonic waves, and generate an image that may distinguish normal from pathologic tissue. Simply put, it is the appropriate matching of the physics of the measurement process to the physical property of the tissue that determines the sensitive and overall quality of the final image. Although it is true that computers are being used more and more in medical imaging systems, in our opinion it is important to not lose sight of the fact that the underlying physics of the imaging modality is what dictates the diagnostic capability provided by the images, and ultimately the contribution to medical care.

> Stephen J. Riederer Richard L. Ehman Diagnostic Radiology, Mayo Clinic, Rochester, MN 55905, USA

Corrections and Clarifications

In the Research Article "Crystal structure of the MATa1/MAT $\alpha2$ homeodomain heterodimer



The world of SCIENCE On-line: Now you can access these exclusive features on the SCIENCE World Wide Web home page with just a click of your mouse:

• SCIENCE Electronic Marketplace: The latest scientific product information from top companies.

• SCIENCE GLOBAL CAREER NETWORK: On-line classified advertising.

• Beyond the Printed Page: Special interactive projects, important data and more.

• SCIENCE On-line: SCIENCE Table of Contents, the SCIENCE Editorial, This Week in SCIENCE available the same day that the printed version is published!

SCIENCE WWW Address: http://www.aaas.org



Sophisticated cyclic AMP and cyclic GMP Analogues Available!

bound to DNA" by T. Li et al. (13 Oct., p.

262), panels B and D in figure 6 (p. 267) were

In the correction on page 621 of the 4 August

The ScienceScope item "Peregrine falcon: Saved

or endangered?" (21 July, p. 291) should have

stated that the subspecies Falco peregrinus ana-

tum is being considered for reclassification or

removal from the endangered species list by

the U.S. Fish and Wildlife Service. Other

subspecies have been delisted or are not cur-

rently protected by the Endangered Species

Letters to the Editor

(at science_letters@aaas.org), fax (202-

289-7562), or regular mail (Science,

1333 H Street, NW, Washington, DC

20005). Letters will not be routinely ac-

knowledged. Full addresses, signatures,

and daytime phone numbers should be

included. Letters should be brief (300 words or less) and may be edited for

reasons of clarity or space. Beginning in

October 1995, our previous policy of

consulting with all letter authors before

publication will be discontinued.

Letters may be submitted by e-mail

issue, B. J. R. Philogene's name was mis-

inadvertently interchanged.

spelled.

Act.

- cyclic nucleotide based inhibitors of cAMP-/cGMPdependent protein kinases (cAK/cGK)
- hydrolysis-resistant activators of cAK/cGK for long-term activation experiments
- rapidly metabolizable structures for pulse type activation
- potent activators of cyclic nucleotide dependent ion channels
- analogues with different cGK isozyme selectivity ($I\alpha/\beta$)
- pairs of analogues with opposite site selectivity for preferential activation of either type I or type II of the cAK isozymes
- pairs of synergistic analogues with high activation potential
- analogues with extremely high membrane-permeability
- affinity gels with PDE-stable ligands for cAK, cGK & PDE
- main metabolites of PDE-sensitive structures
- common cAMP/cGMP analogues
- fluorescent structures
- reactive intermediates



- LIFE SCIENCE INSTITUTE -

Forschungslabor und Biochemica - Vertrieb GmbH

Head office: Flughafendamm 9a, P.O. Box 10 71 25 D-28071 Bremen, Germany Phone: 49 (0) 421 59 13 55 Fax: 49 (0) 421 59 47 71 US / Canada distributor: Ruth Langhorst Int'l. Marketing 7514 Girard Ave. Suite 1-411 La Jolla, CA 92037 Phone: (619) 457-1573 Fax: (619) 456-0810 In Japan contact: Wako Pure Chemical Ind., Ltd. Phone: Tokyo: 03-270-8571 Osaka: 06-203-3741

Circle No. 6 on Readers' Service Card