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MBP and Innate Immunity

I enjoyed the timely and informative Research News article by Clare Thompson about mannose-binding protein (MBP) (21 July, p. 301). MBP is a lectin found in serum, and quite a number of observations point to its importance for the innate immune defense against a variety of microorganisms. This defense does not depend on priming of the immune system, as is characteristic for acquired immunity, for example, after vaccinations.

Many important investigators and their contributions to this field are mentioned by Thompson. However, credit is not given to some Japanese groups who made decisive contributions. Kawasaki's group discovered and characterized the lectin in liver and serum (1). Later they and another Japanese group demonstrated its capacity to activate the complement system (2), crucial for the antimicrobial activity of the lectin. It was also Japanese investigators who characterized a new enzyme necessary for this activation (3). This information showed that MBP is the initiator of a novel complement activation pathway, referred to as the lectin pathway.

The Japanese named the protein "mannan-binding protein," as its discovery and purification were dependent on its calcium-dependent binding to the polysaccharide mannan, which is extracted from baker's yeast. The binding to carbohydrates classified the protein as a lectin, and its biological functions require its binding to carbohydrates on microorganisms. Mannan is composed mainly of polymers of the monosaccharide mannose, and American and British researchers, who entered the field later, renamed the protein "mannose-binding protein." They used the monosaccharide mannose instead of mannan for the purification of the lectin, but there seems to be no scientific reason for maintaining this new name, as it clearly is the same lectin in Japan and elsewhere. The biological function of the protein is mediated through binding to oligo- and polysaccharides rather than monosaccharides. Examination of binding to a variety of monosaccharides even shows that mannose is not the monosaccharide bound with highest affinity by MBP.

We all know the importance of names; I suggest that one should acknowledge the crucial contributions of the Japanese researchers, as well as the right of the discoverer to name new entities, and return to using the original name.

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References

1. N. Kawasaki, R. Etoh, I. Yamashina, *Biochem. Biophys. Res. Commun.* **81**, 1018 (1978); Y. Kozutsumi, T. Kawasaki, I. Yamashina, *ibid.* **95**, 658 (1980).
2. K. Ikeda, T. Sannoh, N. Kawasaki, T. Kawasaki, I. Yamashina, *J. Biol. Chem.* **262**, 7451 (1987); Y.-H. Ji, M. Matsushita, H. Okada, T. Fujita, M. Kawakami, *J. Immunol.* **141**, 4271 (1988).
3. M. Matsushita and T. Fujita, *J. Exp. Med.* **176**, 1497 (1992); A. Takahashi, Y. Takayama, H. Hattuse, M. Kawakami, *Biochem. Biophys. Res. Commun.* **190**, 681 (1993).

EMF Studies

Gary Taubes' News article "Another blow weakens EMF-cancer link" (29 Sept., p. 1816) deals with a controversy regarding EMF (electromagnetic field)-stimulated transcription in HL60 cells that is still being debated at scientific meetings and in scientific journals. There appear to be differences in the cells used by scientists on both sides of the issue, as well as differences in extraction techniques, that could account for the different results.

Many experimental studies showing EMF-stimulated changes in biosynthesis suggest that some concern is warranted. Of particular significance is the observation that changes in protein synthesis caused by

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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 peer-reviewed, balanced coverage of the issues.

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References

1. M. Blank, O. Khorkova, R. Goodman, *Bioelectrochem. Bioenerget.* **31**, 27 (1993); *ibid.* **33**, 109 (1994); R. Goodman *et al.*, *ibid.*, p. 115.
2. 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 ad-

vances 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.

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Corrections and Clarifications

In the Research Article "Crystal structure of the MATa1/MATa2 homeodomain heterodimer

bound to DNA" by T. Li *et al.* (13 Oct., p. 262), panels B and D in figure 6 (p. 267) were inadvertently interchanged.

In the correction on page 621 of the 4 August issue, B. J. R. Philogene's name was misspelled.

The ScienceScope item "Peregrine falcon: Saved or endangered?" (21 July, p. 291) should have stated that the subspecies *Falco peregrinus anatum* 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 currently protected by the Endangered Species Act.

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