

TABLE 4A.2 Repressive Control of the Enzymes
of the Isoleucine—Valine—Leucine Pathway
(See Fig. 4A.1)

| Enzymes | Corepressor |
|----------------|-------------------------------|
| d, e, f, g, h | Isoleucine + valine + leucine |
| i, j, jj, k, l | Leucine |

SUGGESTIONS FOR FURTHER READING

- ALBERTS, B., AND OTHERS, *Essential Cell Biology*, Garland Publishing, Inc., New York, 1998. (This book is an up-to-date, highly readable, relatively short text. The same authors have written a more detailed text, *Molecular Biology of the Cell*.)
- BLACK, J. G., *Microbiology: Principles and Applications* 3d ed., Prentice Hall, Upper Saddle River, NJ, 1996.
- COOK, P. R., The Organization of Replication and Transcription, *Science* 284:1790–1795, 1999.
- ELGARD, L., M. MOLINARI, AND A. HELENIUS, Setting the Standards: Quality Control in the Secretory Pathway, *Science* 286:1882–1888, 1999. (Also two related articles on quality control in post-translational processing and translation follow.)
- KELLY, M. T., AND T. R. HOOVER, Bacterial Enhancers Function at a Distance, *ASM News* 65:484–489, 1999.
- KOLTER, R., AND R. LOSICK, One for All and All for One, *Science* 280:226–227, 1998. (This article is an overview related to a more detailed article on quorum sensing in biofilms; see DAVIES, D. G., AND OTHERS, *Science* 280:295–298, 1998.)
- MADIGAN, M. T., J. M. MARTINKO, AND J. PARKER, *Brock Biology of Microorganisms*, 8th ed., Prentice Hall, Upper Saddle River, NJ, 1997.
- MORAN, L. A., K. G. SCRIMGEOUR, H. R. HORTON, R. S. OCHS, AND J. D. ROWN, *Biochemistry*, 2d ed., Prentice Hall, Upper Saddle River, NJ, 1994.
- STANIER, R. Y., AND OTHERS, *The Microbial World*, 5th ed., Prentice Hall, Englewood Cliffs, NJ, 1986.
- VON HIPPEL, P. H., An Integrated Model of the Transcription Complex in Elongation, Termination and Editing, *Science* 281:660–665, 1998.

PROBLEMS

- 4.1. Consider the aspartic acid pathway shown in Fig. 4A.1. Assume you have been asked to develop a high-lysine-producing mutant. What strategy would you pursue? (That is, which steps would you modify by removing feedback inhibition, and what changes in medium composition would you make over a simple mineral salts–glucose base medium?)
- 4.2. Why is *m*-RNA so unstable in most bacteria (half-life of about 1 min)? In many higher organisms, *m*-RNA half-lives are much longer (> 1 h). Why?
- 4.3. What would be the consequence of one base deletion at the beginning of the message for a protein?