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Ogm Jaf Pkey Emulator V5.exe free iso.m oved.to.zenshare.org/w/j/f0/e75993649de 1168c543bbc14c2a9a6e8\$59.49.95.95Im proving the thermodynamic stability of an enzymatic catalyst by site-directed mutagenesis. A series of mutants of a thermostable triosephosphate isomerase (TIM) with improved stability were designed and characterized. Site-directed mutagenesis was employed to introduce charged residues in surface areas that may influence the stability of the protein. These mutants displayed significant increases in stability. The resistance to thermal denaturation was monitored by differential scanning calorimetry (DSC) and the unfolding equilibrium constant for folding was determined as a measure of the thermal stability of each mutant. The T61A, T63A, D64A, and E65A mutants displayed up to a 70-fold increase in thermal stability as compared to wildtype TIM. These results indicate that an

increase in the number of charged residues in the surface of TIM can improve the stability of the enzyme, presumably by decreasing the exposure of hydrophobic regions of the protein to water. However, increased thermal stability also increased the catalytic activity of TIM. Although these mutants were thermodynamically stable, they had an altered catalytic mechanism. I am a bit confused about the nomenclature of astrocytes and their roles in disease. "Classic" astrocyte marker(s) such as GFAP, S100β, vimentin have been established and have contributed to a better understanding of astrocyte phenotypes. In contrast, the field has recently turned to secreted proteins as useful markers of astrocyte populations. For example, S100β, a calcium-binding protein is expressed by both classic and non-classic astrocytes and is useful as an astrocyte marker. It is more specific than

GFAP for distinguishing astrocytes in high-density cultures, but is expressed in low numbers in adult brain. S100β is especially useful for identifying astrocytes in acute brain slices where it is readily detectable with simple immunostaining procedures. However, this marker does not always reveal differences in astrocyte populations, and it is not expressed by reactive astrocytes, and its expression persists in mature astrocytes (

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