Table II: Kinetic Parameters for Amino Acid Transport under Various Conditions^a

| | | pH 5.5 | | | | | |
|---------------|------|---------------------------|---|-------------------|---------------|--|-----------------|
| amino acid | pI | K _m (app) (μM) | V_{max} [nmol/(min mg of protein)] driving force | | | pH 7.5° | |
| | | | | | | $K_{\mathbf{m}}(\text{app}) V_{\mathbf{max}} [\text{nmol/(min}]$ | |
| | | | $\Delta \mu_{H^+}^b$ | $\Delta \Psi^{c}$ | ΔpH^d | (μM) | mg of protein)] |
| proline | 6.10 | 0.7 | 1.6 | 0.7 | 1.2 | 0.7 | 1.9 |
| glycine | 5.97 | 2.0 | 0.9 | 0.6 | 0.6 | 2.5 | 0.6 |
| alanine | 6.02 | 4.4 | 1.7 | 1.4 | 0.9 | 4.4 | 1.1 |
| serine | 5.68 | 9.9 | 9.3 | 4.9 | 6.0 | 9.9 | 14.4 |
| threonine | 6.53 | 3.9 | 3.1 | 1.6 | 1.3 | 3.7 | 1.6 |
| tyrosine | 5.65 | 1.5 | 2.7 | 1.5 | 0.9 | 1.2 | 0.7 |
| tryptophan | 5.88 | 0.3 | 0.9 | 0.3 | 0.7 | 0.3 | 0.7 |
| phenylalanine | 5.98 | 1.0 | 3.6 | 1.6 | 2.3 | 1.0 | 3.0 |
| glutamate | 3.22 | 9.2 | 1.0 | 0.8 | 0.5 | 9.2 | 0.5 |
| aspartate | 2.97 | 6.8 | 1.8 | 0.9 | 1.4 | 6.8 | 1.8 |
| lysine | 9.74 | 0.5 | 0.6 | 0.5 | 0.3 | 1.0 | 0.7 |

^a Experiments were performed as outlined in Methods and in Figures 1 and 6-9 by using the amino acids listed at the following specific activities and ranges of concentration: [U-¹⁴C]proline (240.6 mCi/mmol) at 0.16-8.1 μM; [U-¹⁴C]glycine (123 mCi/mmol) at 0.4-10.7 μM; [U-¹⁴C]alanine (171.2 mCi/mmol) at 1.2-26.5 μM; [U-¹⁴C]serine (147 mCi/mmol) at 2.7-26.4 μM; [U-¹⁴C]threonine (185.2 mCi/mmol) at 0.5-20.8 μM; tyrosine (452 mCi/mmol) at 0.22-2.2 μM; [U-¹⁴C]tryptophan (17.1 mCi/mmol) at 0.1-1.95 μM; [U-¹⁴C]phenylalanine (464 mCi/mmol) at 0.15-8.9 μM; [U-¹⁶C]glutamate (263 mCi/mmol) at 1.5-21.4 μM; [U-¹⁶C]aspartate (208 mCi/mmol) at 0.18-18.5 μM; and [U-¹⁶C]ysine (280.9 mCi/mmol) at 0.14-21.4 μM. Kinetic parameters were determined by plotting initial velocity, V, as a function of V/S, where S represents solute concentration. Values for V_{max} were estimated from the y intercept of the plots, and the apparent K_m was estimated from the slope of the functions. Assays performed in the absence of ionophores. Assays performed in the presence of nigericin at a final concentration of 0.2 μM.