

## Research Paper

# Glycyl-L-Glutamine Disposition in Rat Choroid Plexus Epithelial Cells in Primary Culture: Role of PEPT2

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**Purpose.** The purpose of this research was to determine the polarity and directionality of the PEPT2-mediated uptake and transepithelial transport of the neuropeptide glycyl-L-glutamine (GlyGln) in choroid plexus.

**Methods.** The transport kinetics of [<sup>3</sup>H]GlyGln was studied in neonatal rat choroid plexus epithelial cells in primary culture grown on laminin-coated Transwell filter inserts. Using a bicarbonate artificial cerebrospinal fluid (CSF) buffer (pH 7.4) at 37°C, GlyGln studies were performed as a function of time, substrate concentration, and the presence of potential inhibitors (at 1 mM).

**Results.** GlyGln (2 μM) accumulation was about three to four times greater when introduced from the apical (CSF-facing) as opposed to the basal (blood-facing) side of the cell monolayer, and transepithelial transport was about two times greater in the apical-to-basal direction. The apical uptake of radiolabeled GlyGln (2 μM) was inhibited significantly by dipeptides (i.e., unlabeled GlyGln and cysteinylglycine) and some neuropeptides (i.e., carnosine, *N*-acetylaspartylglutamate, kyotorphin), but was unaffected by amino acids (i.e., glycine, glutamine) as well as by [D-Arg<sup>2</sup>]-kyotorphin and glutathione. The concentration-dependent apical uptake of GlyGln (2–1000 μM) was characterized by a high-affinity process (i.e.,  $V_{max}$  of 72 pmol/mg/min;  $K_m$  of 136 μM), consistent with the properties of PEPT2. The intracellular hydrolysis of GlyGln was extensive, however, with only 40% of the dipeptide remaining intact after 1 h.

**Conclusions.** The results demonstrate that PEPT2 plays an important role in regulating the apical uptake of GlyGln at the blood–CSF interface. Once inside the cell, GlyGln is rapidly degraded to its constitutive amino acids for further processing.

**KEY WORDS:** choroid plexus; disposition; GlyGln; metabolism; PEPT2; transport; uptake.

## INTRODUCTION

Glycyl-L-glutamine (GlyGln) is a biologically active neuropeptide that inhibits the firing frequencies of neurons in the nucleus reticularis gigantocellularis of rat brain stem (1,2). It is also a potent inhibitor of the hypotension and respiratory depression produced by intracerebroventricular injection of morphine or β-endorphin in rats (1,2). GlyGln is incapable of crossing the blood–brain barrier (3) but is, instead, formed within the brain as a result of the proteolytic

cleavage of β-endorphin (4). Thus, GlyGln exerts its biological activity as a metabolic fragment of polypeptide biotransformation. The process by which the neuropeptide activity of GlyGln is terminated, however, is unclear since clearance mechanisms from the brain, including the cerebrospinal fluid (CSF), are poorly understood for naturally occurring di- and tripeptide substrates.

The close proximity of aminopeptidases and peptide transporters (PEPTs) in the brain makes the coupling of peptide metabolism and transport an attractive combination. In this regard, a number of peptidases have been identified in the choroid plexus (5). Likewise, a high-affinity proton-coupled oligopeptide transporter, PEPT2, has been identified as being expressed and functionally active at the apical membrane of choroid plexus whole tissue (6–8) epithelial cells in primary culture (9) and native tissue (10). As a result, it is possible that GlyGln, once formed, can be cleared by PEPT2 into the choroid plexus, where it can then be transported into the blood, sequestered in the tissue itself, or broken down into its constituent amino acids for further use.

Few studies have examined the uptake and transepithelial transport mechanisms of di- and tripeptides in the choroid plexus. This is particularly true for biologically active peptides. While the neuropeptides *N*-acetylaspartylglutamate

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**ABBREVIATIONS:** BCSFB, blood–CSF barrier; CSF, cerebrospinal fluid; CysGly, cysteinylglycine; DKTP, [D-Arg<sup>2</sup>]kyotorphin; GlyGln, glycyl-L-glutamine; GlySar, glycylsarcosine; GSH, glutathione; HPLC, high-performance liquid chromatography; KTP, kyotorphin; NAAG, *N*-acetylaspartylglutamate; PEPT, peptide transporter; PHT, peptide/histidine transporter.

(NAAG), cysteinylglycine (CysGly), GlyGln, kyotorphin (KTP), and carnosine inhibited the PEPT2-mediated uptake of radiolabeled glycylsarcosine (GlySar) in isolated rat choroid plexus (8), the direct study of dipeptide disposition in choroid plexus and the role of PEPT2 in this process yielded conflicting results. In particular, the accumulation and transcellular transport of GlySar (a synthetic model dipeptide) were three to four times higher when introduced from the apical as opposed to basal side of rat choroid plexus epithelial cells in primary culture (9). Likewise, carnosine (a naturally occurring neurodipeptide) was preferentially taken up from the apical as opposed to basolateral membrane of epithelial cell monolayers (11). In contrast to GlySar, however, basolateral efflux was limited and, as a result, the transepithelial flux of carnosine was not distinguishable from that of paracellular diffusion alone. Instead, carnosine accumulated substantially within choroid plexus cells, achieving concentrations 135 times higher than that of the extracellular medium.

Given the limited understanding of how neuropeptides are handled at the blood-CSF interface, including peptide disposition once in the cell, the polarity and directionality of GlyGln uptake and transport were examined in rat choroid plexus primary cell cultures. Our results indicate that although PEPT2 facilitates the apical uptake of GlyGln into choroidal epithelial cells, this neuropeptide (in contrast to GlySar and carnosine) is rapidly metabolized to its constituent amino acids for further processing.

## MATERIALS AND METHODS

All studies were carried out in accordance with the Guide for the Care and Use of Laboratory Animals as adopted and promulgated by the U.S. National Institutes of Health (NIH publication No. 85-23, revised in 1985).

### Materials

[<sup>3</sup>H]GlyGln (10 Ci/mmol; radiochemical purity 97.7%) was purchased from Moravak Biochemicals (Brea, CA) and [<sup>14</sup>C]mannitol (53 mCi/mmol) from American Radiolabeled Chemicals (St. Louis, MO). Unlabeled glycine and glutamine (i.e., amino acids), CysGly (i.e., dipeptide precursor of glutathione), and GlyGln, carnosine, NAAG, KTP, [D-Arg<sup>2</sup>]kyotorphin (DKTP) and glutathione (GSH) (i.e., neuropeptides) were purchased from Sigma (St. Louis, MO). Other chemicals were obtained from standard sources and were of the highest quality available.

### Intracellular Accumulation and Transepithelial Transport of GlyGln in Cell Cultures

Choroid plexus epithelial cells in primary culture were obtained from the lateral ventricles of 1- to 2-day-old Sprague-Dawley rats using the method of Strazielle and Gherzi-Egea (12). This method was validated and applied previously by our group in studying the uptake and transport kinetics of GlySar (9) and carnosine (11) at the blood-CSF barrier. In brief, choroid plexus cells were collected and seeded on laminin-coated Transwell-Clear filter inserts (12

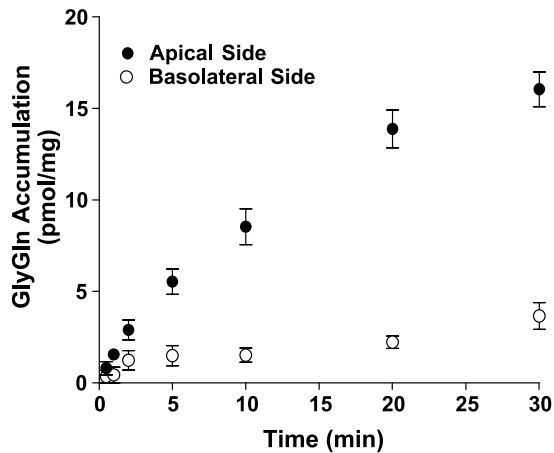
mm diameter, 0.4 μm pore size; Costar Plastics, Cambridge, MA). The culture medium was changed every 48 h after seeding and consisted of Dulbecco's minimum essential medium (DMEM)/F-12 (1:1) supplemented with 10% (v/v) fetal bovine serum, 2 mM glutamine, 25 μg/ml of gentamicin, 5 μg/ml of insulin, 5 μg/ml of transferrin, 5 ng/ml of sodium selenite, 10 ng/ml of epidermal growth factor, 2 μg/ml of hydrocortisone, and 5 ng/ml of basic fibroblast growth factor. Cells were grown in a sterile incubator at 37°C, 95% relative humidity, and 5% CO<sub>2</sub>. Cultures were inspected visually for epithelial growth (i.e., cobblestone appearance) on a regular basis, and confluence occurred 5–7 days post-seeding. Trans-epithelial electrical resistance measurements were performed prior to experimentation (using a Millicell-ERS; Millipore Corp., Bedford, MA) as a measure of monolayer integrity. Values of ≥150 ohm · cm<sup>2</sup> were reached at 10–14 days post-seeding, and were considered sufficient for experimentation.

Uptake and transport studies were performed in a bicarbonate artificial CSF buffer that consisted of (in mM): 127 NaCl, 20 NaHCO<sub>3</sub>, 2.4 KCl, 0.5 KH<sub>2</sub>PO<sub>4</sub>, 1.1 CaCl<sub>2</sub>, 0.85 MgCl<sub>2</sub>, 0.5 Na<sub>2</sub>SO<sub>4</sub>, and 5.0 glucose (pH 7.4), bubbled with 5% CO<sub>2</sub> and 95% O<sub>2</sub>. Cell monolayers were preincubated apically (0.4 ml) and basolaterally (1.2 ml) with this buffer at 37°C for 10 min. The buffer was removed and fresh buffer containing [<sup>3</sup>H]GlyGln and [<sup>14</sup>C]mannitol, with and without inhibitors, was added to the apical side (0.4 ml) or basolateral side (1.2 ml); control buffer (no isotope) was added to the opposite side. The cells were then incubated for the indicated period of time at 37°C. To measure transepithelial transport, an aliquot (100 μl) of the buffer was taken from the opposite side and the radioactivity was counted. To measure intracellular accumulation, the media were aspirated at the end of the incubation period and the monolayers rapidly washed four times on both sides with ice-cold buffer. The filters with monolayers were detached from the chambers and cells were solubilized in 0.5 ml of 0.2 M NaOH and 1% sodium dodecyl sulfate (SDS). The radioactivity of the collected buffer and of the solubilized cells was determined by liquid scintillation counting. The protein content of the solubilized cell monolayers was determined using a protein assay kit (Bio-Rad Laboratories, Hercules, CA) with bovine serum albumin as the standard.

GlyGln uptake data were corrected for filter binding and extracellular content, as described previously (7). The trans-epithelial transport of GlyGln was corrected for paracellular flux, as estimated by mannitol (9).

### Stability of GlyGln

The stability of GlyGln was evaluated in the apical, basolateral, and intracellular compartments of choroid plexus epithelial cells. Following the preparation of cell monolayers, as described previously, 2 μM of [<sup>3</sup>H]GlyGln was incubated in the apical chamber for 5, 15, 30, and 60 min at 37°C. At the end of incubation, media from the apical and basolateral compartments were aspirated and saved for analysis. Cell monolayers were washed four times with ice-cold buffer, the filters detached, and 0.5 ml of ice-cold Milli-Q water (Millipore Corp.) added prior to sonication for 10 min. Cell lysates were then treated with an equal volume of acetonitrile, vortex-mixed for 5 s, and centrifuged at 12,000 rpm



**Fig. 1.** Intracellular accumulation of GlyGln as a function of time in choroid plexus epithelial cells. Studies were performed with 2  $\mu$ M each of [ $^3$ H]GlyGln (0.04  $\mu$ Ci) and [ $^{14}$ C]mannitol (0.02  $\mu$ Ci; a marker for extracellular content) in bicarbonate artificial CSF buffer (pH 7.4) at 37°C. Data are expressed as means  $\pm$  SE ( $n = 8$ ). The apical uptake of GlyGln was significantly greater than basolateral uptake at all times, except at 0.5 and 1.0 min.

for 5 min at 4°C. The supernates were concentrated under cryo-vacuum (SpeedVac refrigerated concentrator; Savant Instruments Inc., Farmingdale, NY) and analyzed by high-performance liquid chromatography (HPLC). The stability of GlyGln was also evaluated in lysates prepared from choroid plexus whole tissue. For these preparations, 14–16 choroid plexuses were pooled from 7–8 neonatal rats and sonicated (1-s burst  $\times$  3) in ice-cold artificial CSF buffer. Tissue lysate (7.2 mg protein per 180  $\mu$ l of buffer) was preincubated for 5 min at 37°C and [ $^3$ H]GlyGln added to the mixture. GlyGln was then incubated continuously in tissue lysate (at pH 7.4 and pH 5.0) and samples (25  $\mu$ l) were obtained at 2, 5, 15, 30, and 60 min at 37°C. Samples were then centrifuged at 12,000 rpm for 5 min at 4°C and analyzed by HPLC. GlyGln stability was determined by its recovery and by the appearance of glycine and glutamine following incubation.

Glycine, glutamine, and GlyGln were detected using an HPLC system consisting of a pump (Waters, Model 510, Milford, MA), a reversed-phase column (Hypersil ODS, C-18, 5  $\mu$ m, 250 mm  $\times$  4.6 mm, Alltech, Deerfield, IL) and a radiochromatography detector (FLO-ONE 500TR, Packard Instrument Co., Meriden, CT). The mobile phase was comprised of 0.01 M phosphate buffer (pH 2.0) and 0.1% heptafluorobutyric acid, and pumped isocratically at 1 ml/min. Retention times for glycine, glutamine, and GlyGln were 4.07, 4.23, and 9.30 min, respectively, under ambient conditions. Peaks were identified by injecting known standards of radiolabeled glycine, glutamine, and GlyGln (Moravak Biochemicals).

#### Data Analysis

For kinetic studies, the concentration-dependent uptake of GlyGln was best fit to a Michaelis-Menten relationship such that:  $V = V_{\max} \cdot S / (K_m + S)$ , where  $V_{\max}$  is the maximal rate of saturable uptake,  $K_m$  is the Michaelis constant, and  $S$

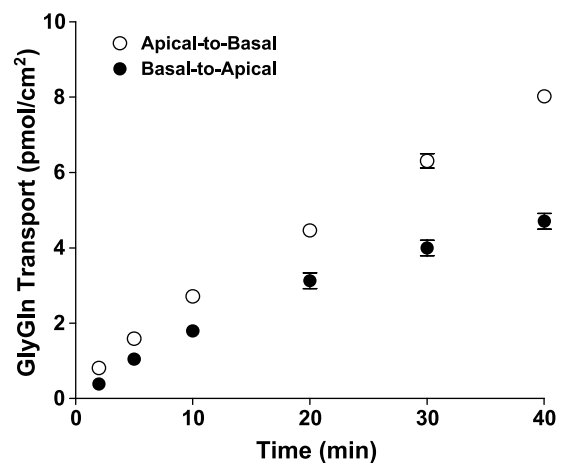
is the substrate concentration. To evaluate if more than one class of transporters was operational for GlyGln uptake, a Woolf-Augustinsson-Hofstee transformation was performed on the saturable component where:  $V = V_{\max} - K_m(V/S)$ .

Data are reported as means  $\pm$  SE (unless otherwise indicated), with data from each experiment determined in duplicate or triplicate. Statistical comparisons were performed with ANOVA (GraphPad Prism, v3.0; GraphPad Software, Inc., San Diego, CA) and *post hoc* pairwise comparisons made with Dunnett's test. A probability of  $p \leq 0.05$  was considered statistically significant. Linear and nonlinear regression analyses were performed with SigmaPlot 8.0 (SPSS Inc., Chicago, IL) and a weighting factor of unity. The quality of fit was determined by evaluating the coefficient of determination ( $r^2$ ), the coefficient of variation (CV%) of parameter estimates, and by visual inspection of the residuals.

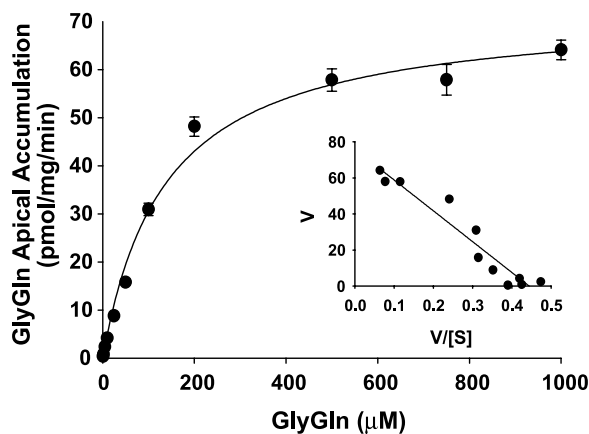
## RESULTS AND DISCUSSION

### Time Course of GlyGln Accumulation and Transepithelial Transport

As shown in Fig. 1, the uptake of GlyGln was substantially greater when introduced from the apical than from the basolateral surface of the cell monolayers (i.e., about three to four times). Similarly, the apical-to-basolateral flux of GlyGln was greater than that of GlyGln in the reverse direction (i.e., about two times) (Fig. 2). Given the strong preference of GlyGln for apical uptake, along with PEPT2 being localized to the apical surface of choroid plexus epithelial cells, the kinetic properties ( $V_{\max}$ ,  $K_m$ ) and specificity ( $\pm$  potential inhibitors) of GlyGln were further characterized at this membrane surface alone. For these analyses, initial rates were determined at 2 min, a time in which GlySar was still in the linear region of uptake.



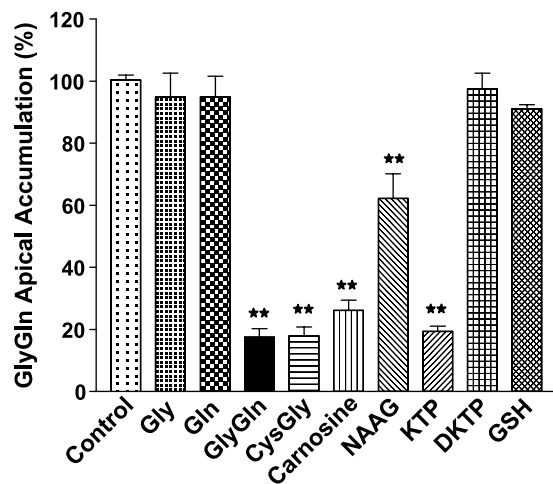
**Fig. 2.** Transepithelial transport of GlyGln as a function of time in choroid plexus epithelial cells. Studies were performed with 2  $\mu$ M each of [ $^3$ H]GlyGln (0.04  $\mu$ Ci) and [ $^{14}$ C]mannitol (0.02  $\mu$ Ci; a marker for paracellular transport) in bicarbonate artificial CSF buffer (pH 7.4) at 37°C. Data are expressed as means  $\pm$  SE ( $n = 16$ ). The apical-to-basolateral transport of GlyGln was significantly greater than basolateral-to-apical transport at all times, except at 2.0 min.



**Fig. 3.** Concentration-dependent uptake of GlyGln in choroid plexus epithelial cells. The 2-min apical uptake of [ $^3\text{H}$ ]GlyGln (0.04–1.2  $\mu\text{Ci}$ ; 2–1000  $\mu\text{M}$  total drug) was evaluated in bicarbonate artificial CSF buffer (pH 7.4) at 37°C; [ $^{14}\text{C}$ ]mannitol (0.02–0.4  $\mu\text{Ci}$ ; 2  $\mu\text{M}$  total compound) was also present as a marker for extracellular content. The experimental data are expressed as means  $\pm$  SE ( $n = 4$ –8). The predicted curve was generated using the mean parameters for  $V_{\text{max}}$  and  $K_m$ , as determined by nonlinear regression ( $r^2 = 0.991$ ) and reported in the text. The inset is a Woolf–Augustinsson–Hofstee plot of the transformed data [uptake ( $V$ ), pmol/mg/min vs. uptake per concentration ( $V/[S]$ ),  $\mu\text{l}/\text{mg}/\text{min}$ ], which is linear ( $r^2 = 0.926$ ). Error bars are omitted in the inset for clarity.

#### Concentration-Dependent Apical Uptake of GlyGln

The apical uptake of GlyGln was probed over a buffer concentration range of 2–1000  $\mu\text{M}$ . As shown in Fig. 3, GlyGln was taken up by choroid plexus epithelial cells in a concentration-dependent manner with a  $V_{\text{max}}$  of  $72.5 \pm 2.7$  pmol/mg/min and a  $K_m$  of  $137 \pm 18$   $\mu\text{M}$  ( $r^2 = 0.991$ ). Only one transport system was involved in the apical uptake of GlyGln, as suggested by the single linear slope during

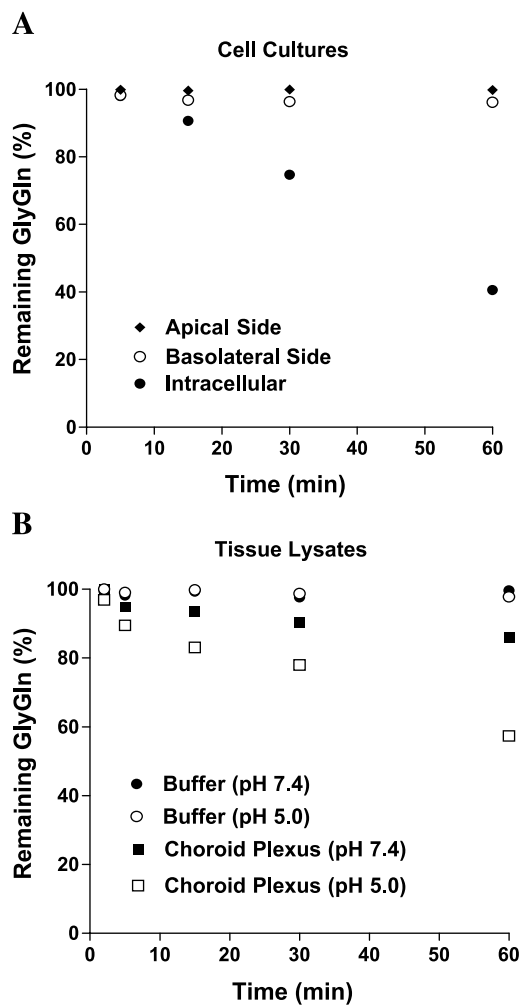


**Fig. 4.** Effect of potential inhibitors on the apical uptake of GlyGln in choroid plexus epithelial cells. The 2-min apical uptake of 2  $\mu\text{M}$  [ $^3\text{H}$ ]GlyGln (0.04  $\mu\text{Ci}$ ) was evaluated in bicarbonate artificial CSF buffer (pH 7.4) at 37°C; 2  $\mu\text{M}$  [ $^{14}\text{C}$ ]mannitol (0.02  $\mu\text{Ci}$ ) was also present as a marker for extracellular content. The experimental data are expressed as mean  $\pm$  SE ( $n = 4$ –9). \*\* $p < 0.01$  as compared to control values.

a Woolf–Augustinsson–Hofstee transformation of the data (see insert,  $r^2 = 0.926$ ). These values are consistent with that of PEPT2 in choroid plexus whole tissue (i.e., high-affinity interactions with  $K_m$  values of 39–260  $\mu\text{M}$  for peptides/mimetics) (5) and, in particular, with that of GlySar (i.e.,  $V_{\text{max}}$  of 31 pmol/mg/min,  $K_m$  of 60  $\mu\text{M}$ ) (9) and carnosine (i.e.,  $V_{\text{max}}$  of 73 pmol/mg/min,  $K_m$  of 34  $\mu\text{M}$ ) (11) in choroid plexus primary cell cultures.

#### Specificity of Apical Uptake for GlyGln

The specificity of GlyGln's apical uptake was probed using a wide range of potential amino acid, dipeptide, and neuropeptide inhibitors (Fig. 4). The constitutive amino acids of GlyGln were without effect, confirming the stability of GlyGln during the incubation process. As expected, GlyGln



**Fig. 5.** (A) GlyGln stability as a function of time in choroid plexus epithelial cells. Studies were performed in bicarbonate artificial CSF buffer (pH 7.4) at 37°C in which monolayers were incubated apically with 2  $\mu\text{M}$  of [ $^3\text{H}$ ]GlyGln. Stability was assessed in the apical and basolateral chambers, and in the intracellular compartment. Data are expressed as the mean of two experiments ( $\leq 10\%$  difference between studies). (B) GlyGln stability as a function of time in choroid plexus whole tissue lysates (pH 7.4 and pH 5.0). Data are expressed as the means of two preparations (with each preparation containing 14–16 choroid plexuses from 7–8 animals). Control studies were also performed in buffer alone (pH 7.4 and pH 5.0).



exhibited self-inhibition (i.e., reduced accumulation of radiolabel in the presence of unlabeled dipeptide) along with inhibition by the dipeptide CysGly (about 80–85%). With respect to neuropeptides, GlyGln was inhibited by carnosine (74%), NAAG (38%), and KTP (81%), but was not inhibited by DKTP or GSH. Thus, it appears that the apical uptake of GlyGln into choroid plexus cells may be influenced by dipeptide fragments and/or neuropeptides that share the same pharmacophore for PEPT2. For example, DKTP (L-Tyr-D-Arg) is a stereoisomer of KTP (L-Tyr-L-Arg) in which the L-configuration is preferred for transport. Likewise, free amino acids that are one (i.e.,  $\alpha$ -position) or at most two (i.e.,  $\beta$ -position) methylene groups from the backbone carbonyl group have been demonstrated to exhibit a high-affinity transport by PEPT2 (13). GSH, with a free amino acid in the  $\gamma$ -position would, therefore, have a much reduced or nonexistent uptake.

### Stability of GlyGln

For those primary cell culture studies in which 2-min uptakes were determined (i.e., kinetic and inhibition experiments), GlyGln was found to be stable during the incubation period. As shown in Fig. 5A, 99.8% and 96.2% of the label remained intact in the apical and basolateral compartments, respectively, at 60 min. In contrast, the intracellular compartment retained 97.9% of the intact label at 5 min, 90.6% at 15 min, 74.7% at 30 min, and only 40.6% after 60 min of incubation. Given the substantial metabolism of GlyGln at 60 min, yet the insignificant amount of degradation product found in apical or basolateral chambers, it appears that the constituent amino acids remain in the cell for further processing. To further understand the intracellular degradation of GlyGln, additional experiments were performed in lysates prepared from choroid plexus whole tissue. As shown in Fig. 5B, only about 15% of the drug was metabolized in pH 7.4 tissue lysate after 60 min of incubation, whereas about 43% of GlyGln was metabolized in pH 5.0 tissue lysate over the same time period. In contrast, control experiments (GlyGln added to buffer) showed no degradation of GlyGln as a function of pH alone. These results suggest that GlyGln may be preferentially metabolized in lysosomes, where a pH of about 5.0 in its interior is optimal for the activity of hydrolytic enzymes (14).

PEPT2 is a member of the proton-coupled oligopeptide transporter family along with PEPT1 and the peptide/histidine transporters (PHTs) PHT1 and PHT2 (15–18). Immunolocalization studies (9,10) have confirmed that whereas PEPT2 protein is expressed widely throughout the brain, including the apical membrane of choroid plexus, PEPT1 protein is absent in this tissue. Functional studies, using inhibition of radiolabeled dipeptide by excess L-histidine, have consistently failed to support a role for the peptide/histidine transporters, PHT1 and PHT2, at the plasma membrane of choroid plexus tissue (6–8) and primary cell cultures (9,11). Moreover, studies in wild type and PEPT2 null mice have clearly demonstrated that PEPT2 is responsible for about 80–95% of the uptake of GlySar (19,20), 5-aminolevulinic acid (20), carnosine (11), and cefadroxil (21) in choroid plexus whole tissue. Despite our current understanding that PEPT2 plays a predominant role in the choroid plexus uptake

of peptides/mimetics and peptide-like drugs, the disposition of specific neuropeptides, as opposed to model synthetic dipeptide probes, at the blood–CSF barrier (BCSFB) is still uncertain.

In the present study, we studied the uptake, trans-epithelial transport, and cellular metabolism of GlyGln, a dipeptide fragment of  $\beta$ -endorphin proteolysis, in choroid plexus epithelial cells in primary culture. We found that GlyGln is preferentially taken up and transported across choroid plexus epithelium by a PEPT2-mediated process at the apical surface of the cell membrane. We also found that GlyGln is rapidly degraded intracellularly, probably by lysosomes, into its constitutive amino acids. The ability to provide free glycine and glutamine for utilization by the cellular machinery might have significant value. In this context, glycine could combine with GluCys to form the antioxidant glutathione (22), and glutamine could be used in protein synthesis, as a nitrogen donor for the synthesis of purines, pyrimidines, nucleotides, and amino sugars, and in the regulation of acid–base balance (23). The importance of GlyGln as a source of free amino acids for these potential functions needs to be determined, however. Taken as a whole, PEPT2 serves as a conduit for a wide range of di- and tripeptides, neuropeptides, and peptide-like drugs to access the choroid plexus cell from the CSF. The disposition of these substrates, however, may vary once in the cell. While some peptides may be effluxed into the blood, as found for the hydrolysis- and peptidase-resistant dipeptide GlySar (9), others may accumulate in the choroid plexus, as observed for carnosine (11). In contrast, GlyGln is metabolized intracellularly by the choroid plexus, where it can then be recycled.

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