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Presumptive brain influx of large neutral amino acids and the effect of phenylalanine supplementation in patients with Tyrosinemia type 1

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Abstract

Introduction

Hereditary Tyrosinemia type 1 (HT1) is a rare metabolic disease caused by a defect in the tyrosine degradation pathway. Current treatment consists of 2-(2-nitro-4-trifluoromethylbenzoyl)-1,3-cyclohexanedione (NTBC) and a tyrosine and phenylalanine restricted diet. Recently, neuropsychological deficits have been seen in HT1 patients. These deficits are possibly associated with low blood phenylalanine concentrations and/or high blood tyrosine concentrations. Therefore, the aim of the present study was threefold. Firstly, we aimed to calculate how the plasma amino acid profile in HT1 patients may influence the presumptive brain influx of all large neutral amino acids (LNAA). Secondly, we aimed to investigate the effect of phenylalanine supplementation on presumptive brain phenylalanine and tyrosine influx. Thirdly, we aimed to theoretically determine minimal target plasma phenylalanine concentrations in HT1 patient to ensure adequate presumptive brain phenylalanine influx.

Methods

Data of plasma LNAA concentrations were obtained. In total, 239 samples of 9 HT1 children, treated with NTBC, diet, and partly with phenylalanine supplementation were collected together with 596 samples of independent control children. Presumptive brain influx of all LNAA was calculated, using Michaelis-Menten parameters ($K_m$) and $V_{max}$-values obtained from earlier articles.

Results

In HT1 patients, plasma concentrations and presumptive brain influx of tyrosine were higher. However, plasma and especially brain influx of phenylalanine were lower in HT1 patients.
Phenylalanine supplementation did not only tend to increase plasma phenylalanine concentrations, but also presumptive brain phenylalanine influx, despite increased plasma tyrosine concentrations. However, to ensure sufficient brain phenylalanine influx in HT1 patients, minimal plasma phenylalanine concentrations may need to be higher than considered thus far.

**Conclusion**

This study clearly suggests a role for disturbed brain LNAA biochemistry, which is not well reflected by plasma LNAA concentrations. This could play a role in the pathophysiology of the neuropsychological impairments in HT1 patients and may have therapeutic implications.

**Introduction**

Hereditary Tyrosinemia type I (HT1; McKusick 276700) is an inborn error of metabolism caused by a deficiency in the catabolic pathway of tyrosine. Due to a genetic defect in the enzyme fumarylacetoacetate hydrolase (FAH), several toxic products accumulate, causing liver failure, renal tubulopathy, rickets, cardiomyopathy, porphyria like syndrome, and hepatocellular carcinoma [1]. Since 1992, HT1 patients are treated with 2-(2-nitro-4-trifluoromethylbenoyl)-1,3-cyclohexanedione (NTBC) [2]. NTBC inhibits tyrosine catabolism upstream from the primary enzymatic defect (at the level of 4-OH-phenylpyruvate dioxygenase), preventing the formation of the toxic products, and thereby substantially improving clinical outcome [3]. NTBC treatment strongly increases plasma tyrosine concentrations, necessitating dietary restriction of tyrosine and its precursor phenylalanine [1,4]. Such combined treatment of NTBC and diet may still result in high plasma tyrosine concentrations, while phenylalanine concentrations are often low [5–8]. While previous studies have shown the possible importance of phenylalanine supplementation in HT1 patients, the minimum plasma phenylalanine target level remains to be established [5,8].

In the past few years, neurocognitive impairments have been observed in HT1 patients [6,9–13]. The pathophysiological mechanisms underlying these neurocognitive impairments are not fully understood. The metabolic derangement in HT1 with high plasma tyrosine and low phenylalanine concentrations is supposed to play a central role. However, the resulting transport of these amino acids across the blood-brain barrier (BBB) could be even more important, as this could directly influence brain amino acid and neurotransmitter concentrations [8,14].

At the BBB, all large neutral amino acids (LNAA) are primarily transported by the so-called L-type amino acid transporter 1 (LAT1) in a competitive manner [15–17]. The transport rate of each individual LNAA is determined by its plasma concentration, and the $K_m$-value (the plasma concentration at which the transport rate is 50% of its maximum), which differs for each LNAA [18,19]. Under physiological conditions, LAT1 is saturated for $>95\%$ by these LNAA [19]. Therefore, high plasma concentrations of a particular LNAA would not only increase brain influx of this LNAA but also outcompete brain uptake of the other LNAA [18–20].

The competitive transport mechanism has for example been shown to exist in phenylketonuria (PKU) mice [21] and maple syrup urine disease (MSUD) mice [22]. In these mice, Vogel et al. (2014) clearly showed that plasma amino acid concentrations in PKU and MSUD mice represent only a partial picture of brain amino acid homeostasis. Next to this, therapeutically,
the addition of a specific amino acid that interferes with the transport of the other amino acids at the BBB has shown to improve the outcome in some defects of amino acid metabolism such as guanidinoacetate methyltransferase deficiency, pyridoxine dependent epilepsy and MSUD [23–25]. Therefore, taking together these theoretical considerations, experimental, and clinical data, the competitive transport of amino acids across the BBB shows to be important in various inborn errors of amino acid metabolism [26].

Therefore, the aim of the present study was threefold. Firstly, we aimed to calculate how the plasma amino acid profile as observed in HT1 patients on NTBC and dietary treatment may influence the presumptive brain influx of all LNAA compared to healthy controls. Secondly, we aimed to investigate the effect of phenylalanine supplementation on presumptive brain phenylalanine and tyrosine influx in four of these HT1 patients. Thirdly, we aimed to theoretically determine minimal plasma phenylalanine concentrations in HT1 patients to ensure adequate presumptive brain phenylalanine influx.

Methods
Patients

In total, 239 plasma samples of nine HT1 patients (six males, three females) were obtained between 2002 and 2015, from diagnosis or first measurement after introduction of NTBC and diet (median: 0.53 years; range: 0.04–12.2 years) till the current age with a maximum of 18 years (median: 11.5 years; range: 2.8–18.0 years). Six patients were diagnosed primarily suffering from liver failure, among others characterized by severe coagulopathy due to decreased synthetic liver function. For these patients, data from the first period after diagnosis were excluded until coagulopathy had been restored (defined by prothrombin time <16 sec and activated partial prothrombin time <35 sec) to avoid samples with high methionine concentrations associated with liver failure. Three other patients were diagnosed due to neonatal screening or because of an affected sibling. For these patients, only the first measurement after diagnosis was excluded for adjustment to the NTBC and dietary treatment, if liver synthesis function was still normal. All patients were treated with NTBC (1–2 mg/kg/day) and a tyrosine and phenylalanine restricted diet in the University Medical Center Groningen or Maastricht University Medical Center. Four of these patients were treated with phenylalanine supplementation ranging from 10 to 32 mg/kg/day next to their regular NTBC and dietary treatment. Of all 239 plasma samples, 73 were obtained during phenylalanine supplementation.

In total, 596 anonymized amino acid profiles from independent control participants (mean: 4.8 years, all younger than 18 years old) were obtained. Control participants were patients who underwent analysis of plasma amino acid concentrations as part of the diagnostic process in the hospital, but who eventually did not have an inborn error of amino acid metabolism.

All HT1 patients and/or their parents or guardians gave written informed consent for retrospective analysis of their data, acknowledging that one patient had died and no informed consent was obtained. This study was approved by the Medical Ethical Committee of the University Medical Center Groningen.

Biochemical analyses

Plasma samples of both HT1 patients and control participants had been taken in the clinic, not taking into account the timing of plasma sampling and its relation to the last meal of the patient or control individuals. In these samples, concentrations of LNAA (except for tryptophan) and glutamine had been quantified in deproteinized plasma by cation-exchange high-performance liquid chromatography followed by post-column ninhydrin derivatization, using...
norleucine as an internal standard, on a Biochrom 20 or 30 analyser (Pharmacia Biotech, Cambridge, UK).

**Calculations of presumptive brain amino acid influx**

Plasma LNAA concentrations were used to calculate presumptive brain influx of individual LNAA. Brain influxes were calculated using classical Michaelis-Menten parameters ($K_m$) and $V_{\text{max}}$-values for each LNAA as reported by Smith et al. 2000, measured with an in situ rat brain perfusion technique [19]. For each LNAA, substrate inhibition ($K_{\text{app}}$) was calculated in the presence of competing amino acids using previously published equations [24].

$$K_{\text{app}} = K_m \left[1 + \sum_{i=1}^{n} \left(\frac{C_i}{K_i}\right)\right]$$

In this equation, $K_m$ ($\mu$M) is the substrate concentration at which the reaction rate is half of the $V_{\text{max}}$. $C_i$ is the current plasma concentration of each competitive amino acid ($\mu$M), and $K_i$ is the $K_m$ of these respective inhibiting amino acids ($\mu$M). The resulting $K_{\text{app}}$ shows the current inhibitory effect of all LNAA together. This value is used to determine the presumptive brain influx using the following equation: brain influx = $(V_{\text{max}})^{'(C)/(K_{\text{app}}+C)}$. In this equation, brain influx is the presumptive brain uptake of the LNAA (nanomoles per minute per gram of brain tissue (nmol/(min’g))), $V_{\text{max}}$ is the maximum transport velocity (nmol/(min’g)), and $C$ is the plasma concentration of the amino acid ($\mu$M).

In all NTBC and dietary treated HT1 patients, Z-scores for both plasma concentrations and presumptive brain influxes of individual LNAA were determined. Z-scores were calculated by subtracting the mean value of the control participants from the values of the patients, and dividing this difference by the standard deviation of the control values. Afterwards mean Z-scores for individual patients were calculated.

To assess the effect of phenylalanine supplementation on presumptive brain phenylalanine and tyrosine influx, we investigated the four HT1 patients with the lowest plasma phenylalanine concentrations who were subsequently treated without and with phenylalanine supplementation in addition to the regular NTBC and dietary treatment. Plasma phenylalanine and tyrosine concentrations, plasma phenylalanine/tyrosine (phe/tyr) ratios, and presumptive brain phenylalanine and tyrosine influx in four HT1 patients before and during phenylalanine supplementation were calculated.

Thereafter, for all HT1 patients, plasma phenylalanine concentrations that would probably result in sufficient brain phenylalanine influx (determined as mean, -1, and -2 standard deviation (SD) below the mean values of presumptive brain influx in control participants) were calculated, if all LNAA concentrations in the HT1 patients would remain the same. To do so, the previously used equation for brain influx was rewritten into: $C = (\text{brain influx} \times K_{\text{app}})/(V_{\text{max}} \times \text{brain influx})$. To estimate these minimum target plasma phenylalanine concentrations for HT1 patients, the mean brain influx as well as the -1 SD and -2 SD of the mean brain influx of control participants were used in the equation, together with the previously calculated individual $K_{\text{app}}$-values.

**Statistics**

The distribution of brain influx of LNAA was compared between HT1 patients and controls using multiple Mann Whitney U tests. Considering the fact that we have repeated observations within each patient and independent data of 596 controls, we used the following testing procedure. To test the null hypothesis that the observations of patients and controls originated from equal distributions, we repeatedly took one random observation from each patient and all observations from the controls. We calculated for each (sub)dataset the p-value, using the
Mann-Whitney U test. After 1000 tests for each LNAA, we evaluated the distributions of p-values. To evaluate the effect of phenylalanine supplementation in HT1 patients, means of plasma values and presumptive brain influx of phenylalanine and tyrosine before and after supplementation were calculated for each patient. Correlational analyses were performed in each individual patient to assess whether age was correlated with presumptive brain phenylalanine influx in HT1 patients. In control participants, linear regression analysis was done after logarithmic transformation to assess a possible correlation between age and presumptive brain phenylalanine influx. Linear mixed effect models with variance components for random effects were used to investigate a possible correlation between presumptive brain phenylalanine influx and plasma tyrosine concentrations, plasma phenylalanine concentrations, and phe/tyr ratios in HT1 patients without receiving phenylalanine supplementation. In all statistical tests, a (mean) p-value of \(<0.05\) was considered statistically significant. Analyses were conducted with the statistical program SPSS 22 (IBM, Chicago, Illinois).

### Results

#### Plasma concentrations and presumptive brain influx of individual LNAA in HT1 patients and controls

Plasma LNAA concentrations without phenylalanine supplementation were repetitively measured in all patients, ranging from 1 to 40 times. Table 1 shows mean plasma concentrations of all LNAA (except for tryptophan) and glutamine for HT1 patients and control participants calculated from the mean concentrations of each HT1 individual. Plasma phenylalanine concentrations were significantly lower (in 99% out of 1000 tests \(p<0.05\); mean \(p = 0.00462\)), while plasma tyrosine concentrations were much higher (in 100% out of 1000 tests \(p<0.05\); mean \(p = 2.67 \times 10^{-7}\)).

<table>
<thead>
<tr>
<th>LNAA</th>
<th>HT1</th>
<th>Controls</th>
<th>Mean P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phenylalanine</td>
<td>27 ±14</td>
<td>54 ± 16</td>
<td>0.00462</td>
</tr>
<tr>
<td></td>
<td>33 [16–38]</td>
<td>50 [43–61]</td>
<td></td>
</tr>
<tr>
<td>Tyrosine</td>
<td>384 ± 53</td>
<td>64 ± 23</td>
<td>2.67 \times 10^{-7}</td>
</tr>
<tr>
<td></td>
<td>382 [337–432]</td>
<td>60 [49–74]</td>
<td></td>
</tr>
<tr>
<td>Valine</td>
<td>226 ± 26</td>
<td>188 ± 54</td>
<td>0.199</td>
</tr>
<tr>
<td>Isoleucine</td>
<td>59 ± 8</td>
<td>55 ± 22</td>
<td>0.494</td>
</tr>
<tr>
<td></td>
<td>59 [52–64]</td>
<td>52 [41–64]</td>
<td></td>
</tr>
<tr>
<td>Leucine</td>
<td>115 ± 18</td>
<td>105 ± 39</td>
<td>0.454</td>
</tr>
<tr>
<td>Methionine</td>
<td>23 ± 6</td>
<td>21 ± 9</td>
<td>0.401</td>
</tr>
<tr>
<td>Histidine</td>
<td>81 ± 11</td>
<td>83 ± 22</td>
<td>0.572</td>
</tr>
<tr>
<td></td>
<td>79 [73–89]</td>
<td>81 [70–94]</td>
<td></td>
</tr>
<tr>
<td>Threonine</td>
<td>129 ± 37</td>
<td>118 ± 62</td>
<td>0.427</td>
</tr>
<tr>
<td>Glutamine</td>
<td>480 ± 46</td>
<td>517 ± 121</td>
<td>0.382</td>
</tr>
<tr>
<td></td>
<td>488 [446–507]</td>
<td>513 [451–578]</td>
<td></td>
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</table>

Plasma amino acid concentrations (μmol/l) in HT1 patients treated with NTBC and diet and controls. Data are presented as Mean ± SD and Median with IQR.

[https://doi.org/10.1371/journal.pone.0185342.t001](https://doi.org/10.1371/journal.pone.0185342.t001)
p<0.001) in HT1 patients compared to control participants. Plasma concentrations of all other LNAA were not significantly different between HT1 patients and controls.

Fig 1 shows the mean plasma concentrations and presumptive brain influx of the measured LNAA and glutamine in HT1 patients not receiving phenylalanine supplementation, expressed as Z-scores of values in control participants. Mean plasma tyrosine concentrations and presumptive brain tyrosine influx in HT1 patients were higher than values in control participants with Z-scores of 13.8 and 15.2, respectively. In contrast, mean plasma phenylalanine concentrations in HT1 patients tended to be slightly lower than in controls with a Z-score of -1.6, while mean presumptive brain phenylalanine influx was much lower with a Z-score of -4.2. Mean plasma concentrations and presumptive brain influxes of other LNAA were all between Z-scores of -0.9 and 0.7.

Effect of phenylalanine supplementation on presumptive brain phenylalanine and tyrosine influx in HT1 patients

Table 2 shows the biochemical effects of phenylalanine supplementation in four HT1 patients. Descriptive analyses show that, on phenylalanine supplementation, mean plasma phenylalanine and tyrosine concentrations increased in these four patients. The proportional increase of plasma phenylalanine concentrations was higher than the increase of plasma tyrosine concentrations in 2 patients, resulting in higher phe/tyr ratios in these patients. On phenylalanine supplementation, the increase in plasma phenylalanine concentrations tended to be accompanied by an increase of the mean presumptive brain influx of phenylalanine in 3 patients. Although plasma tyrosine concentrations increased on phenylalanine supplementation in all patients, presumptive brain influx of tyrosine only increased in two of these patients.

![Graph showing plasma concentrations and presumptive brain influx of individual LNAA in HT1 patients.](https://doi.org/10.1371/journal.pone.0185342.g001)

**Fig 1.** Plasma concentrations and presumptive brain influx of individual LNAA in HT1 patients. Mean plasma concentrations and presumptive brain influx with SD of individual LNAA in HT1 patients expressed as Z-scores (individual patient means) of values in control participants. Dashed lines represent Z-scores of -1 and +1.
Presumptive minimal target plasma phenylalanine concentrations in HT1 patients

In Fig 2, presumptive brain phenylalanine influx is plotted against age for both HT1 patients and controls. Presumptive brain phenylalanine influx in HT1 patients was significantly lower than in controls (in 100% out of 1000 tests \( p < 0.05 \); mean \( p < 0.001 \)). In individual HT1 patients, no clear effect of age on the presumptive brain phenylalanine influx could be seen, although age was significantly correlated with presumptive brain phenylalanine influx in two (out of nine) patients \((r = -0.448; p = 0.006 \text{ and } r = 0.562; p = 0.001)\). However, the correlation coefficient was opposite in both patients. In control participants, age was not significantly correlated with presumptive brain phenylalanine influx \((p = 0.100)\).

Linear mixed effect model analyses on presumptive brain phenylalanine influx and parameters of plasma amino acid (tyrosine and phenylalanine concentrations and phe/tyr ratios) biochemistry revealed that presumptive brain phenylalanine influx was most strongly correlated with plasma phenylalanine concentrations \((p < 0.001, \beta = 0.167, t = 23.57)\). Correlations between presumptive brain phenylalanine influx and plasma tyrosine concentrations and plasma phe/tyr ratios were less strong based on Akaike’s Information Criteria.

Given the strong correlations between presumptive brain phenylalanine influx and plasma phenylalanine concentrations, minimal plasma phenylalanine concentrations were calculated to ensure sufficient presumptive brain phenylalanine influx in HT1 patients. Mean presumptive brain phenylalanine influx in HT1 patients not receiving phenylalanine supplementation was calculated to be \(5.5 \pm 2.5 \text{ nmol/}(\text{min} \times \text{g})\) and \(13.5 \pm 1.9 \text{ nmol/}(\text{min} \times \text{g})\) (-1SD: 11.6; -2SD: 9.7 nmol/\(\text{min} \times \text{g}\)) in control participants. To reach mean, -1SD, or -2SD values of the presumptive brain phenylalanine influx of control participants, plasma phenylalanine concentrations in HT1 patients needed to be increased from 27 to 84 \(\pm 5\), 68 \(\pm 4\), and 53 \(\pm 3\) \(\mu\text{mol/L}\), respectively, if plasma concentrations of all other amino acid concentrations would remain the same.

Discussion

This study investigated (1) plasma LNAA concentrations and presumptive brain influx of individual LNAA in HT1 patients in comparison to control participants, (2) the effect of phenylalanine supplementation in HT1 patients on plasma LNAA biochemistry as well as on presumptive brain LNAA influx, and (3) minimal plasma phenylalanine concentrations necessary to increase the presumptive brain phenylalanine influx. The main findings of this study are threefold. Firstly, although plasma phenylalanine concentrations were low to normal, the presumptive brain influx

<table>
<thead>
<tr>
<th>Table 2. The effect of phenylalanine supplementation.</th>
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<tbody>
<tr>
<td><strong>Patient 1</strong></td>
</tr>
<tr>
<td>Number of samples</td>
</tr>
<tr>
<td>Phe supplementation (mg/kg/day)</td>
</tr>
<tr>
<td>Plasma phenylalanine (µmol/L)</td>
</tr>
<tr>
<td>Plasma tyrosine (µmol/L)</td>
</tr>
<tr>
<td>Plasma phe/tyr ratio</td>
</tr>
<tr>
<td>Brain phenylalanine influx (nmol/(min* g))</td>
</tr>
<tr>
<td>Brain tyrosine influx (nmol/(min* g))</td>
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</table>

Results on plasma biochemistry and presumptive brain phenylalanine and tyrosine influx in four HT1 patients before and during phenylalanine supplementation, expressed as means.

https://doi.org/10.1371/journal.pone.0185342.t002
of phenylalanine was largely impaired in HT1 patients compared to control participants. Secondly, phenylalanine supplementation tended to improve the presumptive brain phenylalanine influx in HT1 patients, having only a small negative effect on plasma tyrosine concentrations and presumptive brain tyrosine influx. Thirdly, to ensure sufficient brain phenylalanine influx in HT1 patients, minimal plasma phenylalanine concentrations may need to be higher than considered thus far.

Before discussing these results in more detail, some methodological issues are addressed. With regard to the plasma amino acid measurements, unfortunately, plasma tryptophan concentrations could not be studied and therefore possible disturbances in presumptive brain tryptophan influx with implications for cerebral serotonin metabolism could not be investigated. In addition, the timing of blood sampling and fasting status is unfortunately unknown. It is known that there is a large variation of plasma phenylalanine concentrations in HT1 patients during the day, with low concentrations especially occurring in the afternoon [5,7].

With regard to calculations for presumed brain influx, the \( K_m \) and \( V_{max} \)-values used in this study have been determined in rats. However, although absolute values differ between humans and rats, LNAA transport characteristics, as determined in brain microvascular endothelial cells of humans, have shown a strong correlation to those in rats [27]. Thus, we assume that inter-species differences are not a major issue. We acknowledge that the presumptive brain influx of LNAA, based on LNAA transport characteristics and plasma concentrations, do not directly reflect brain LNAA availability. However, as lumbar puncture to obtain cerebral spinal

Fig 2. Presumptive brain influx of phenylalanine comparing HT1 patients and controls. Presumptive brain influx of phenylalanine of both HT1 patients and controls plotted against age. The straight line represents the mean presumptive brain phenylalanine influx of controls. The dashed lines represent the -1SD and -2SD of the presumptive brain influx of controls.

https://doi.org/10.1371/journal.pone.0185342.g002

![Graph showing presumptive brain influx of phenylalanine](https://doi.org/10.1371/journal.pone.0185342.g002)
fluid is rather invasive and brain phenylalanine concentrations could not be measured by
normal magnetic resonance spectroscopy techniques, this theoretical method is used. In addition,
the calculated presumptive brain amino acid influx has already been shown useful in the
design of arginine-fortified and LNAA-optimized amino acid supplements in glutaric aciduria
type 1 and MSUD [24,28].

Our results clearly suggest a non-optimal brain LNAA homeostasis in HT1 patients. With
regard to our first research question, except for tyrosine and phenylalanine, no strong differ-
ences in plasma LNAA concentrations between HT1 patients and controls were observed.
Plasma tyrosine concentrations were largely increased, although still within the recommended
range for HT1 patients [1]. Plasma phenylalanine concentrations were decreased as also previ-
ously reported by various study groups [5,7,8]. All other LNAA were within the normal range.
Presumptive brain influx of individual LNAA, however, showed a different profile. While pre-
sumptive brain tyrosine influx was similarly increased in HT1 patients compared to controls
as were plasma tyrosine concentrations, this was not true for other LNAA, especially phenyl-
alanine. Presumptive brain phenylalanine influx in HT1 patients compared to controls was
much more decreased than could be expected only based on plasma phenylalanine values.
Moreover, presumptive brain influx of all other LNAA was slightly reduced in HT1 patients
compared to controls (although mostly within the normal range with z-scores around -1),
de spite normal plasma concentrations. Although the clinical significance of this disturbed
brain LNAA biochemistry requires further investigation, the observed neurocognitive impair-
ments in HT1 patients suggest a possible relationship.

Regarding the second research question, previous studies have shown dietary interven-
tions in inborn errors of amino acid metabolism to be very effective in increasing brain
LNAA concentrations and thereby possibly improving the neurocognitive outcome [22,25,
26]. Phenylalanine supplementation in HT1 patients has not only been related to increased
plasma phenylalanine concentrations and disappearance of eczema as a peripheral conse-
quence of low plasma phenylalanine concentrations, but has also been suggested to be
associated with improved brain functioning with better psychomotor development and
disappearance of cortical myoclonus [8]. In the present study, plasma phenylalanine as
well as tyrosine concentrations seem to increase after phenylalanine supplementation.
Despite higher plasma tyrosine concentrations and resulting competition for transport
across the BBB, presumptive brain phenylalanine influx was increased, although it should
be noticed that values of control participants are never reached. Only one patient (patient
2, Table 2) seemed not to respond to the phenylalanine supplementation. Plasma phenyl-
alanine nor tyrosine concentrations were increased in this patient, possibly caused by an
amount of supplementation that was too low or problems with dietary compliance.

With regard to the third research question, based on the results of the present study, plasma
phenylalanine concentrations in HT1 patients would need to be increased more than consid-
ered thus far to ensure adequate brain phenylalanine influx. In previous studies in HT1
patients, hypophenylalaninemia was defined by plasma phenylalanine concentrations below
40 µmol/L [7] or below 30 µmol/L [8]. Based on our study (without the inclusion of trypto-
phan), plasma phenylalanine concentrations need to be increased even further to maintain
adequate presumptive brain influx of phenylalanine in HT1 patients. Unfortunately, this esti-
mated brain influx is based on a theoretical model not knowing real brain phenylalanine con-
centrations in HT1 patients. Thimm et al. showed increased tyrosine concentrations and
decreased concentrations of serotonin metabolites in the cerebral spinal fluid in HT1 patients,
but did not report on concentrations of other LNAA such as phenylalanine [14]. However,
low concentrations of serotonin metabolites could indicate that tryptophan, the precursor of
serotonin, is outcompeted by tyrosine at the BBB, like we suggest with this model for phenylalanine.

**Conclusion**

The pathophysiological mechanisms of neuropsychological impairment in HT1 are not fully understood and need further investigation. This theoretical experiment suggests a role for disturbed brain LNAA transport. While plasma tyrosine concentrations and presumptive brain tyrosine influx were similarly markedly elevated in HT1 patients, presumptive brain phenylalanine influx was much more affected than could be expected only based on its plasma values. To improve presumptive brain phenylalanine influx in HT1 patients, additional plasma phenylalanine supplementation has shown to be effective, but it seems that plasma phenylalanine concentrations need to be increased even further than was previously suggested.

**Author Contributions**

**Conceptualization:** Willem G. van Ginkel, Danique van Vliet, Francjan J. van Spronsen.

**Data curation:** Willem G. van Ginkel, Danique van Vliet, M. Rebecca Heiner-Fokkema, Francjan J. van Spronsen.

**Formal analysis:** Willem G. van Ginkel, Danique van Vliet, Johannes G. M. Burgerhof, Francjan J. van Spronsen.

**Investigation:** Willem G. van Ginkel, Danique van Vliet, Francjan J. van Spronsen.

**Methodology:** Willem G. van Ginkel, Danique van Vliet, Johannes G. M. Burgerhof, Francjan J. van Spronsen.

**Project administration:** Willem G. van Ginkel, Danique van Vliet, Francjan J. van Spronsen.

**Resources:** Johannes G. M. Burgerhof, Pim de Blaauw, M. Estela Rubio Gozalbo, M. Rebecca Heiner-Fokkema, Francjan J. van Spronsen.

**Supervision:** M. Rebecca Heiner-Fokkema, Francjan J. van Spronsen.

**Validation:** Willem G. van Ginkel, Danique van Vliet, M. Rebecca Heiner-Fokkema, Francjan J. van Spronsen.

**Visualization:** Willem G. van Ginkel, Danique van Vliet.

**Writing – original draft:** Willem G. van Ginkel, Danique van Vliet, Francjan J. van Spronsen.

**Writing – review & editing:** Willem G. van Ginkel, Danique van Vliet, Johannes G. M. Burgerhof, Pim de Blaauw, M. Estela Rubio Gozalbo, M. Rebecca Heiner-Fokkema, Francjan J. van Spronsen.

**References**


