Importance  Growth hormone–releasing hormone (GHRH) has been previously shown to have cognition-enhancing effects. The role of neurotransmitter changes, measured by proton magnetic resonance spectroscopy, may inform the mechanisms for this response.

Objective  To examine the neurochemical effects of GHRH in a subset of participants from the parent trial.

Design  Randomized, double-blind, placebo-controlled substudy of a larger trial.

Setting  Clinical research unit at the University of Washington School of Medicine.

Participants  Thirty adults (17 with mild cognitive impairment [MCI]), ranging in age from 55 to 87 years, were enrolled and successfully completed the study.

Interventions  Participants self-administered daily subcutaneous injections of tesamorelin (Theratechnologies Inc), a stabilized analogue of human GHRH (1 mg/d), or placebo 30 minutes before bedtime for 20 weeks. At baseline and weeks 10 and 20, participants underwent brain magnetic resonance imaging and spectroscopy protocols and cognitive testing and provided blood samples after fasting. Participants also underwent glucose tolerance tests before and after intervention.

Main Outcomes and Measures  Brain levels of glutamate, inhibitory transmitters γ-aminobutyric acid (GABA) and N-acetylaspartylglutamate (NAAG), and myo-inositol (MI), an osmolyte linked to Alzheimer disease in humans, were measured in three 2 × 2 × 2-cm³ left-sided brain regions (dorsolateral frontal, posterior cingulate, and posterior parietal). Glutamate, GABA, and MI levels were expressed as ratios to creatine plus phosphocreatine, and NAAG was expressed as a ratio to N-acetylaspartate.

Results  After 20 weeks of GHRH administration, GABA levels were increased in all brain regions (P < .04), NAAG levels were increased (P = .03) in the dorsolateral frontal cortex, and MI levels were decreased in the posterior cingulate (P = .002). These effects were similar in adults with MCI and older adults with normal cognitive function. No changes in the brain levels of glutamate were observed. In the posterior cingulate, treatment-related changes in serum insulin-like growth factor 1 were positively correlated with changes in GABA (r = 0.47; P = .001) and tended to be negatively correlated with MI (r = −0.34; P = .06). Consistent with the results of the parent trial, a favorable treatment effect on cognition was observed in substudy participants (P = .03). No significant associations were observed between treatment-related changes in neurochemical and cognitive outcomes. Glucose homeostasis in the periphery was not reliably affected by GHRH administration and did not account for treatment neurochemical effects.

Conclusions  Twenty weeks of GHRH administration increased GABA levels in all 3 brain regions, increased NAAG levels in the frontal cortex, and decreased MI levels in the posterior cingulate. To our knowledge, this is the first evidence that 20 weeks of somatotropic supplementation modulates inhibitory neurotransmitter and brain metabolite levels in a clinical trial, and it provides preliminary support for one possible mechanism to explain favorable GHRH effects on cognition in adults with MCI and in healthy older adults.
Circulating levels of growth hormone (GH) and insulin-like growth factor 1 (IGF-1) decline across the life span, a condition referred to as somatopause. Most of the GH measured in plasma is produced in the pituitary, with some central nervous system tissues, such as the hippocampus, also producing GH. Several different peptides act as regulators for the GH system: 1 that is inhibitory, somatostatin, and 2 that are excitatory, ghrelin and GH-releasing hormone (GHRH). As with GH, GHRH levels decrease with age. When given as an injectable supplement, GHRH acts on its own receptors and directly stimulates GH release and IGF-1 synthesis by the liver. Unlike extrinsic GH supplementation, GHRH administration modulates intrinsic regulatory systems and evokes a normal pulsatile pattern of GH release.

The behavioral effects of somatotropic supplementation have been examined in numerous animal studies and in 6 double-blind placebo-controlled clinical studies to date. Supplementation with IGF-1 improves spatial and reference memory in aged rats and, if given to younger animals for 21 months, halts the behavioral decline in spatial memory that occurs with aging. In humans, 3 of 4 GH supplementation studies demonstrated improved cognitive function for 6 to 24 months, and 1 did not. We have conducted 2 randomized clinical trials of GHRH administration. In the first study, 6 months of treatment with GHRH (1-29 NH2) (sermorelin acetate; Serono Laboratories) improved performance on tests of fluid intelligence (working memory, planning and organization, selective attention, and processing speed) in healthy older adults. Our subsequent trial replicated and extended this work to demonstrate positive cognitive effects not only in healthy older adults but also in adults with amnestic mild cognitive impairment (MCI) who are at increased risk of Alzheimer disease (AD) dementia.

Although the favorable effects of somatotropic supplementation on neuronal activity, including synaptic transmission and receptor density, are robust in animal studies, research on the neurobiological effects of these hormones in humans is quite limited. In 1 study, the cerebral glucose metabolic rate in the frontal cortex, as measured with positron emission tomography, was positively correlated with serum IGF-1 levels in older adults. In another imaging study, serum IGF-1 levels were positively correlated with brain levels of N-acetylaspartate (NAA), a putative marker of neuronal activity, in adults with persistent childhood GH deficiency. Imaging studies using magnetic resonance spectroscopy (MRS) survey the common brain metabolites of choline-containing compounds, creatine plus phosphocreatine (Cr), myo-inositol (MI), NAA, and the sum of glutamate plus glutamine, spectra that often include contributions from γ-aminobutyric acid (GABA). New techniques are increasingly used, such as J-resolved point-resolved spectroscopy (J-PRESS), which permits separation of spectral composites into constituent neurotransmitters (glutamate, glutamine, N-acetylaspartylglutamate [NAAG], and GABA), along with established approaches (eg, Mescher-Garwood [MEGA] PRESS) to target single neurotransmitters with high specificity. Herein, we describe the results of a brain imaging study using MRS to examine the effects of GHRH on inhibitory and excitatory neurotransmitters in a subgroup of participants enrolled in a large randomized clinical 20-week trial. In light of results from studies demonstrating brain areas vulnerable to aging- or AD-related processes, we evaluated dorsolateral frontal, posterior cingulate, and posterior parietal regions for changes.

Methods

The 20-week parent study was registered on ClinicalTrials.gov (NCT00257712) and was approved by the University of Washington and Veterans Affairs Puget Sound Health Care System institutional review boards and by the Veterans Affairs Research and Development Committee. The Consolidated Standards of Reporting Trials (CONSORT) flowchart describing participant flow through the parent trial is published elsewhere. At baseline and at weeks 10 and 20, participants completed magnetic resonance imaging (MRI) and MRS imaging protocols and cognitive testing. At baseline and other specified time points during the study, blood samples were collected for measurements of IGF-1, insulin, and glucose after participants had been fasting. Participants also completed oral glucose tolerance tests before and after treatment to further assess GHRH effects on glucose homeostasis.

Participants

Written informed consent was obtained for all MRS substudy participants. From 152 older adults with normal cognitive function (86 adults) or amnestic MCI (66 adults) enrolled in the parent trial, 30 participants (18 men) were enrolled in the MRS substudy (9 with MCI and 7 with normal cognitive function received placebo; 8 and 6, respectively, received GHRH). All participants successfully completed the imaging protocols.

Intervention

In the parent study, participants were randomized in a 1:1 ratio in blocks of 4 to receive either placebo or 1.0 mg/d of tesamorelin (henceforth, GHRH), a human GHRH analogue (acetate salt of N-[trans-3-hexenoyl]-human GHRH [1-44] amide; Theratechnologies Inc), which results in a pulsatile GH response and elevated serum IGF-1 to young adult normal levels. Injected subcutaneously 30 minutes before bedtime for 20 weeks. The GHRH dose was reduced because of an adverse event (eg, arthralgia and fluid retention) within the first 4 weeks of treatment in 4 substudy participants (2 women with normal cognitive function and 1 woman and 1 man with MCI), and the GHRH dose was increased for 1 substudy participant (a man with MCI) when his serum concentration of IGF-1 failed to increase by at least 15% over baseline. Each unblinded dose adjustment was yoked with a similar adjustment for a placebo-treated participant. Other details regarding blinding, compliance, safety monitoring, and dose adjustments in the parent trial are published elsewhere.

Cognition

Four parallel versions of the cognitive protocol were administered at baseline, at weeks 10 and 20, and after a 10-week
washout (no MRS data were collected at week 30) in counter-balanced order. The protocol included tests of executive function (Stroop Color-Word Interference, Task Switching, Self-ordered Pointing Test, word fluency) and episodic memory (story recall, Hopkins Verbal Learning Test, Visual-Spatial Learning Test, delayed match-to-sample task). Composite scores were constructed by summing change z scores within cognitive domain before analysis. Additional details regarding cognitive testing and outcomes are published elsewhere.15

Assays
Serum levels of IGF-1 were initially measured with a 2-site immunoradiometric assay, which included an extraction step to remove IGF-1–binding proteins (DSL/Beckman Coulter; 2.6% intra-assay and 4.5% interassay coefficients of variation). When the manufacturer discontinued this assay, IGF-1 was measured using radioimmunoassay (Mediagnost/IBL America; 3.5% intra-assay and 5.2% interassay coefficients of variation). The 2 assays correlated linearly and all radioimmunoassay values were normalized to the immunoradiometric assay standard with a linear correction algorithm.

Insulin and glucose were measured under fasting and challenge conditions in light of well-established GH effects on energy metabolism. For the oral glucose tolerance test, blood was first collected after a 12-hour fast for glucose and insulin measurements; participants then consumed a 75-g dextrose solution, and blood was sampled 1 and 2 hours later. At screening, participants meeting American Diabetes Association standard glycemic criteria for diabetes, indicated by fasting plasma glucose levels above 125 mg/dL (to convert to millimoles per liter, multiply by 0.0555) or 2-hour oral glucose tolerance test levels above 199 pg/mL, were excluded from enrollment.

Imaging Acquisition and Processing
Before MRI, participants were given instructions to hydrate for 24 hours by drinking eight 8-oz glasses of water and to avoid caffeine for 2 hours before the procedure. They were positioned comfortably on the 3T Philips scanner bed. Supplemental padding was used to minimize head movement in an 8-channel head coil. Localizers in all 3 planes were obtained with a 3-dimensional fast low-angle shot sequence, followed by a high-resolution magnetization-prepared rapid gradient echo (MPRAGE) volume acquisition in the sagittal plane (field of view, 24 cm; 256 × 256 matrix; isotropic 1 × 1 × 1-mm3 voxels; 10 minutes). The MPRAGE sequence was then reformatted to generate coronal and axial image volumes.

Next, 2 × 2 × 2-cm3 voxels were localized in the left-sided dorsolateral frontal, posterior cingulate, and posterior parietal regions (Figure 1). Three-plane voxel coordinates and anatomic landmarks were tagged and screen saved to ensure accurate voxel relocation for follow-up MRI at weeks 10 and 20. At each voxel location, higher-order shimming was performed, followed by a 24-echo water-suppressed PRESS acquisition (echo time [TE], 35–380 milliseconds in 15-millisecond steps; repetition time, 2 seconds; number of signals acquired, 16; 2048 points; bandwidth, 2000; approximately 14 minutes with shimming and setup), with water-suppressed acquisitions used to evaluate for eddy current evaluation. Data were saved to a Linux workstation and processed by using custom MATLAB software (MathWorks) with C-scripts for TE-averaged (sum of 24 echoes) and J-PRESS data sets (spectra reformatted as described elsewhere).20 Metabolites measured included GABA, NAAG, glutamate, glutamine, and MI, for J-PRESS data sets and choline-containing compounds, Cr, and NAA for TE-averaged data sets. Metabolites were referenced to Cr and NAAG to NAA by convention.30 Gray-white (G/W) matter voxel fractions were obtained by using SPM software (http://www.fil.ion.ucl.ac.uk) with custom MATLAB scripts for voxel extraction (modified from the original source code by Mikael Montelius, MS).

Statistical Analysis
The analytic plan was similar to that used in the parent trial.15 The primary outcomes for the MRS substudy included the in-
hibitory transmitters GABA and NAAG and the excitatory transmitter glutamate. Although GHRH-related changes from baseline to week 20 were predicted in light of the behavioral results, changes at week 10 were examined in post hoc exploratory analyses. Before analysis, multiple regression and correlation procedures were used to create residualized change scores at weeks 10 and 20 (ie, residuals from the regression of longitudinal data on the baseline value). Residuals that serve as change scores are adjusted for baseline differences and are inherently more stable than delta values.31 Age, sex, and G/W matter voxel fractions were used as covariates in all models.

The GABA/Cr, NAAG/NAA, and glutamate/Cr ratios were subjected to a single multivariate analysis of variance (MANOVA), with treatment group (GHRH vs placebo), diagnosis (MCI vs normal cognitive function), and brain region (dorsolateral frontal, posterior cingulate, or posterior parietal) as independent variables. When the MANOVA results proved significant, subsequent univariate analyses of variances (ANOVAs) were conducted on the constituent outcomes. When appropriate, pairwise comparisons by group, diagnosis, or brain region were performed with use of ANOVAs rather than t tests to permit covariates in the model. Similarly structured exploratory ANOVAs were performed on secondary neurochemical outcomes (all referenced relative to Cr including MI, choline-containing compounds, NAA, and glutamine). The effects of GHRH on cognition and glucose homeostasis in MRS substudy participants were examined in post hoc analyses. Correlational analyses explored relationships between baseline and treatment-related changes in neurochemical outcomes, serum IGF-1 levels, glucose homeostasis, and cognitive composites of executive function and verbal episodic memory scores (composites that were sensitive to GHRH effects in the parent trial). Baseline neurochemical differences by diagnosis (MCI vs normal cognitive function) were also examined by using ANOVA, with adjustment for age, sex, and G/W matter voxel fraction. All analyses were performed with Stata software (StataCorp).32

### Results

At baseline, characteristics of the participants were comparable across treatment groups for diagnosis, cognitive status, age, and educational level; body composition; and serum IGF-1, fasting insulin, and glucose levels (Table 1); no differences in metabolites were observed by treatment group assignment or diagnosis (data not shown).

The omnibus MANOVA on GABA/Cr, NAAG/NAA, and glutamate/Cr indicated a significant effect of GHRH (main effect of treatment, F2,19 = 6.17; P = .003), which was comparable in adults with MCI and in healthy older adults (no treatment × diagnosis interaction; P = .48). The results of subsequent ANOVAs on constituent outcomes indicated that for GABA/Cr, ratios increased with 20 weeks of GHRH administration (main effect of treatment, F2,24 = 13.62; P = .001; Figure 2) in all brain regions sampled (no treatment × region interaction; P = .98) for adults with or without MCI (no treatment × diagnosis interaction; P = .90) (eFigure, A).

Separate exploratory analyses by brain region confirmed widespread effects of GHRH on GABA/Cr ratios (dorsolateral frontal region, F2,22 = 4.57 [P = .04]; posterior cingulate, F2,22 = 7.37 [P = .01]; posterior parietal, F2,23 = 5.58 [P = .03]). (The posterior parietal region sampled included relatively more white matter, and J-PRESS tends to overestimate GABA in white matter [J.E.J., written communication, July 2, 2012].) There were regional differences in NAAG/NAA ratios in response to GHRH administration (treatment × region interaction, F2,22 = 4.06; P = .02) (Figure 3); the results of post hoc analyses indicated treatment-related increases in regions with relatively more gray (frontal or cingulate region) than white matter (parietal region) for both diagnostic groups (no treatment × diagnosis interaction; P = .77) (eFigure, B), most notably in the frontal cortex (P = .03). The univariate analysis of glutamate/Cr ratios indicated no significant main or interaction effects involving treatment (P > .35).

### Table 1. Baseline Characteristics by Treatment Group and Diagnosis

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>GHRH Group Mean (SD)</th>
<th>Placebo Group Mean (SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Normal Cognitive Function</td>
<td>MCI</td>
</tr>
<tr>
<td>Participants, No. (No. female)</td>
<td>6 (3)</td>
<td>8 (2)</td>
</tr>
<tr>
<td>Age, y</td>
<td>66.8 (9.0)</td>
<td>69.4 (8.3)</td>
</tr>
<tr>
<td>Educational level, y</td>
<td>17.5 (2.6)</td>
<td>17.8 (2.4)</td>
</tr>
<tr>
<td>MMSE</td>
<td>29.3 (1.0)</td>
<td>29.4 (1.4)</td>
</tr>
<tr>
<td>Story recall scoreb</td>
<td>60.3 (14.3)</td>
<td>40.1 (16.4)</td>
</tr>
<tr>
<td>BMI</td>
<td>25.7 (3.2)</td>
<td>28.1 (3.9)</td>
</tr>
<tr>
<td>IGF-1, ng/mL</td>
<td>158 (92)</td>
<td>231 (78)</td>
</tr>
<tr>
<td>Insulin, μIU/mL</td>
<td>4.2 (0.8)</td>
<td>8.0 (5.6)</td>
</tr>
<tr>
<td>Glucose, mg/dL</td>
<td>97 (7.2)</td>
<td>104 (11.0)</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index (calculated as weight in kilograms divided by height in meters squared); GHRH, growth hormone–releasing hormone; glucose, fasting plasma glucose concentration; IGF-1, fasting serum concentration of insulinlike growth factor 1; insulin, fasting plasma insulin concentration; MMSE, Mini-Mental State Examination score (maximum score, 30).

* Data represent mean (SD) values unless otherwise noted.

** Significant baseline difference by diagnosis (P < .05).
The result of secondary analyses indicated that MI/Cr ratios decreased with 20 weeks of GHRH administration in the posterior cingulate ($F_{23} = 11.57; \ P = .002$) in adults with or without MCI (no treatment × diagnosis interaction; $P = .21$) (eFigure, C), but not in the dorsolateral frontal ($P = .61$) or parietal ($P = .53$) regions. This difference was supported by a treatment × region interaction in the omnibus analysis (ANOVA, $F_{2,50} = 3.88; \ P = .03$) (Figure 4). There were no other treatment effects on neurochemical measurements, and GABA, NAAG, and MI levels were unchanged from baseline to 10 weeks of treatment (all $P > .14$).

Post hoc analysis of GHRH effects on executive function and verbal memory composite scores indicated that the benefits observed for the larger group in the parent trial were also observed in the smaller group of MRS participants (MANOVA for main effect of treatment, with adjustment for age, educational level, and Mini-Mental State Examination score, $F_{2,20} = 4.02; \ P = .03$) (Table 2). Cognitive response to GHRH did not differ between participants who opted in and those who opted out of the MRS substudy ($P = .53$). There were no reliable effects of treatment on peripheral measures of glucose homoeostasis. Although fasting insulin levels increased for GHRH-treated adults with MCI in the parent trial ($P = .005$), this effect failed to reach significance in the substudy ($P = .12$).

Exploratory correlational analyses indicated that in the posterior cingulate region, the baseline GABA/Cr ratio was positively correlated with the serum IGF-1 concentration ($r = 0.40; \ P = .03$), as was the change from baseline to week 20 in these measures for the entire MRS substudy cohort ($r = 0.47; \ P = .001$). Although changes in GABA/Cr ratios and fasting insulin concentrations were associated for the substudy cohort ($r = 0.41; \ P = .03$), when the fasting insulin concentration was included as an additional covariate in the structured ANOVA for GABA/Cr, the initial findings remained unchanged. Treatment-related changes in MI/Cr ratios tended to be negatively correlated with IGF-1 ($r = -0.34; \ P = .06$). In the dorsolateral frontal cortex, although the serum IGF-1 concentration and NAAG/NAA ratio were positively correlated at baseline ($r = 0.41; \ P = .02$), treatment-related changes in these measures from baseline to week 20 were not ($r = -0.07; \ P = .71$). There were no other associations involving neurochemical outcomes, serum IGF-1 and fasting insulin levels, and cognitive composite scores.
scores (all \( P > .05 \)). A summary of the correlational analyses is provided online (eTable).

### Discussion

In adults with MCI and healthy older adults, 20 weeks of GHRH administration increased GABA/Cr ratios in the left dorsolateral frontal, left posterior cingulate, and left posterior parietal regions and NAA/Glu ratios in the left dorsolateral frontal region. Treatment with GHRH did not affect glutamate, the predominant excitatory neurotransmitter in the brain. Myo-inositol, an osmolyte and precursor of phospholipids that has been found to be elevated with increasing neuropathological severity associated with AD, was decreased in the left posterior cingulate region after treatment. Although the implications of M1 changes have been the source of much speculation, this and our other biochemical findings suggest that GHRH may have beneficial effects on neurotransmitter and osmotic changes linked to aging and AD.

The role of GABA as the major inhibitory neurotransmitter, ubiquitous in the central nervous system, is well characterized (reviewed by Rossignol). Although the animal literature suggests that GABA receptor changes may underlie the beneficial memory effects of somatotropic supplementation, to our knowledge, our findings provide the first evidence that GHRH administration increases inhibitory neurotransmitter levels in the brain. The diverse functional role of NAAG as a peptide neurotransmitter has only recently been characterized (reviewed by Neale et al.). In a few postmortem studies of AD, NAAG levels are reduced (levels in the occipital lobe are less than levels in the temporal lobe, which are less than levels in the frontal lobe). Unlike neurotransmitters that act on ionotropic receptors, NAAG acts on metabotropic glutamate receptor 3, present on both neurons and glia. A secondary form of NAAG, NAAG2, was recently identified, and its function and distribution are still undetermined. In the absence of clear evidence to indicate that NAAG has effects other than inhibiting neurotransmission, perhaps the most parsimonious conclusion from our data at this point is that 20 weeks of GHRH treatment enhances inhibitory features of brain network activity (GABA and NAAG).

A range of studies demonstrates aging-related changes in GABA and an important role of efficient inhibitory processing for successful cognition. In the auditory cortex, the synthesis enzyme for GABA, glutamic acid decarboxylase, is reduced with aging within most cortical layers (II-V). Although this effect is not as widespread in the parietal cortex, enzyme reductions in the hippocampus support focused involvement in that region. Aging- or disease-related changes in the balance of inhibitory to excitatory activity in the brain may represent an important marker of brain function efficiency. Majidi and colleagues reported that the expression of proteins associated with excitatory and inhibitory postsynaptic sites in the prefrontal and parietal cortical regions were altered in older cognitively impaired mice compared with older unimpaired or young mice, implicating a potential important role of excitatory-inhibitory tone in aging-related cognitive impairment.

In a chemically induced model of behavioral deficit using the noncompetitive \( N \)-methyl-\( \alpha \)-aspartate receptor antagonist phencyclidine, transplantation of embryonic medial ganglionic eminence into the frontal lobe either before or after phencyclidine treatment selectively causes growth of GABAergic interneurons and a reversal of cognitive impairment. In theoretical models, it has been suggested that deficits in GABA result in poorer performance because of an increase in background noise relative to stable stimulus strength, whereas supplementation overcomes this condition. Consistent with this interpretation, reduced levels of dorsolateral prefrontal GABA in humans are related to greater impulsivity and increased perseveration in schizophrenia.

Our study has several limitations. The sample size was small and only 3 brain regions were analyzed with MRS. Although the method used (J-PRESS) has been shown to be sufficient and reproducible for measuring GABA, further studies using more sensitive editing approaches (eg, MEGA-PRESS) would be helpful to improve sensitivity. The anatomic coordinates selected for examination in the brain may have precluded our ability to detect associations between GHRH effects on neurochemical outcomes and cognition in our study. For example, in studies in which the anatomic target for MRS is highly localized to the behavior being surveyed, close concordance between GABA and function is observed; a high correlation was demonstrated between visual discrimination tasks.
and magnetoencephalographic measures of gamma band activity (probably mediated by GABA\(^2\))33, functional MRI findings, and GABA in the visual cortex.\(^2\)47 Similar findings are reported for motor control and GABA in the supplementary motor area\(^4\) and for tactile discrimination and GABA in the sensorimotor cortex\(^5\); control regions selected to assess cortical GABA levels remote from the site of interest demonstrated no concordance with performance.\(^4\)\(^9\)\(^5\)\(^0\)

Future work evaluating metabolite-behavior changes induced by GHRH in targeted task-specific regions will be useful to evaluate whether GHRH-related effects on inhibitory neurotranschemistry tone might account for GHRH-related improvements in cognitive function. Our data do not permit us to resolve whether brain neurochemical treatment effects were due to indirect GHRH effects in the periphery or direct GHRH effects in the brain. Although the role of GH on glucose homeostasis is well established, we did not observe consistent treatment effects on glucose or insulin under fasting or challenge conditions, and treatment-related change in neurochemical outcomes were not correlated with these measures. The effects of GHRH on brain glucose metabolism have yet to be explored in clinical trials.

In summary, 20 weeks of GHRH administration increased brain levels of the inhibitory transmitters GABA and NAAG and decreased MI levels both in adults with MCI and in healthy older adults. These treatment-related changes are consistent with amelioration of aging- or disease-related biochemical processes in the brain. Although larger and longer-duration treatment trials are needed to confirm these preliminary findings, our pilot study provides additional evidence to support positive GHRH effects on brain function in normal and pathological aging.

**REFERENCES**


3. Donahue CP, Kosik KS, Shors TJ. Growth hormone is produced within the hippocampus where it responds to age, sex, and stress. Proc Natl Acad Sci U S A. 2006;103(15):6031-6036.


17. Arwert LI, Veltman DJ, Deijen JB, Lammersma AA, Jonker C, Drent AA, Jonker C, Drent ML. Memory performance and...


