Cognitive Deficits in Rats after Forebrain Cholinergic Depletion are Reversed by a Novel NO Mimetic Nitrate Ester

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Many conditions adversely affecting learning, memory, and cognition are associated with reductions in forebrain acetylcholine (ACh), most notably aging and Alzheimer's disease. In the current study, we demonstrate that bilateral depletion of neocortical and hippocampal ACh in rats produces deficits in a spatial learning task and in a recently described, delayed visual matching-to-sample task. Oral administration of the novel nitrate, GT1061 (4-methyl-5-(2-nitroxyethyl) thiazole HCl), and the acetylcholinesterase inhibitor, donepezil, reversed the cognitive deficits in both memory tasks in a dose-dependent manner. GT1061 was superior in the delayed matching-to-sample task. GT1061 was absorbed rapidly after oral administration, crossed the blood brain barrier, and achieved brain concentrations that were slightly higher than those found in plasma. The activity of GT1061 was NO mimetic: soluble guanylyl cyclase (sGC) was activated, but selectivity was observed for sGC in the hippocampus relative to the vasculature; and hippocampal levels of phosphorylated ERK1/2, which is a postulated intermediary in the formation of long-term memory, were increased. The beneficial effect on visual and spatial memory task performance supports the concept that stimulating the NO/sGC/cGMP signal transduction system can provide new, effective treatments for cognitive disorders. This approach may be superior to that of current drugs that attempt only to salvage the residual function of damaged cholinergic neurons.

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INTRODUCTION

Alzheimer's disease (AD) is characterized by disruption of both excitatory amino acid and cholinergic neurotransmission. Loss of cholinergic neurons, and subsequent deficits in cholinergic neurotransmission in the hippocampus and cerebral cortex, is strongly correlated with clinical signs of cognitive impairment and dementia in AD patients (Mesulam, 2004). The use of acetylcholinesterase inhibitor drugs, such as donepezil (Aricept), is based upon the cholinergic hypothesis of memory loss and cognitive deficit (Bartus *et al*, 1982). These drugs have represented the sole therapeutic response to AD until the recent introduction of the *N*-methyl-D-aspartate (NMDA) receptor low affinity antagonist, memantine. The clinical efficacy of acetylcholinesterase inhibitors has been questioned (Courtney *et al*,

2004), and memantine is not approved for mild to moderate forms of AD (Areosa *et al*, 2005), thus the imperative for new therapeutic agents to treat AD and other dementias remains.

Signal transduction by the NO/soluble guanylyl cyclase (sGC)/cGMP system modulates synaptic transmission and plasticity in the CNS, and *in vivo* studies suggest an important role for NO/cGMP in the acquisition of new learning and memory (Bernabeu *et al*, 1997; Prickaerts *et al*, 2002; Prickaerts *et al*, 1997). This is supported by numerous *in vitro* studies indicating a role in long-term potentiation (LTP), clustering of presynaptic and postsynaptic proteins (suggesting a role in synaptogenesis), and enhancement of presynaptic neurotransmitter release. Aberrant NO signaling in the brains of AD patients has also been reported (Bonkale *et al*, 1995). Thus, sGC activation and cGMP formation in the brain may be an effective strategy for mitigating the cognitive dysfunction that occurs as a consequence of cholinergic deficits in the CNS. In contrast to the acetylcholinesterase inhibitors that attempt to augment the residual function of a degenerating cholinergic system, NO mimetics are postulated to bypass this system and modulate the function of signaling pathways downstream from cholinergic receptor activation.

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Most recently, two studies have directly implicated a role for dysfunction of NO/cGMP signaling in AD. In hippocampal slices, disruption of LTP by amyloid- β peptide (A β) was shown to result from a malfunction of NO/cGMP signaling (Puzzo *et al*, 2005). Further, Snyder *et al* (2005) reported that A β produced a depression of NMDA-evoked currents via endocytosis of NMDA receptors, mediated via dissociation of the synaptic protein, PSD95, from the NMDA receptor. As both sGC and neuronal nitric oxide synthase (NOS) are known to be associated with PSD95 in the brain (Russwurm *et al*, 2001), this work suggests that A β specifically disrupts hippocampal NO/cGMP signaling. It was concluded that intervention with NO donors or drugs enhancing NO/cGMP signaling might provide a novel approach to treatment of AD (Puzzo *et al*, 2005).

Classical nitrates, of which glyceryl trinitrate (GTN) is the archetype, are NO mimetics, in that they manifest many aspects of the biological activity of NO *in vivo* (Bennett *et al*, 1994; Thatcher *et al*, 2004). Although the hypotensive effect of GTN itself precludes its use in CNS-directed therapy, we have previously reported a novel nitrate that demonstrates: (a) attenuated hypotensive activity relative to GTN; (b) neuroprotection in the middle cerebral artery occlusion model of ischemic stroke; and (c), reversal of scopolamine-induced cognition deficits in rats tested in the Morris water task (MWT) (Reynolds *et al*, 2002; Smith *et al*, 2000).

Prusky *et al* (2004) recently demonstrated long-delay trial-unique visual matching-to-sample (DMTS) performance in rats, implicating essential roles for both the hippocampus and perirhinal cortex. Thus, the hippocampus is important not only for spatial memory, but also for visual recognition memory, a cognitive process that is disrupted in AD (Lindeboom *et al*, 2002). In this paper, we use this DMTS task to examine the reversal of a cholinergic deficit in recognition memory by a novel nitrate, GT1061 (4-methyl-5-(2-nitroxyethyl) thiazole HCl), and compare both to effects on spatial working memory and to the effects of donepezil in the same tasks. A potential mechanism of action of this novel compound in the hippocampus also is described.

MATERIALS AND METHODS

All procedures for animal experimentation were undertaken in accordance with the principles and guidelines of Canadian Council on Animal Care.

Behavioral Studies

Moving hidden platform Morris water task: surgery. In these studies, 90 of 100 young adult female Long-Evans rats received bilateral injections of 192 IgG-saporin into the nucleus basalis magnocellularis and medial septum to deplete the cholinergic innervation of the cerebral cortex and hippocampus, respectively (Pizzo *et al*, 1999). Rats were anaesthetized with isoflurane and under stereotaxic guidance (coordinates: medial septum; AP +0.6, L 1.2, D -7.5, -6.8 from bregma; nucleus basalis; AP -1, L $\pm$ 3, D -7.5), 192 IgG-saporin (0.1 μ g in 1.0 μ l of phosphate-buffered saline, pH 7.4, over 15 s) was infused into each site. The remaining 10 animals received sham surgery. All rats were allowed to recover for 14 days.

MWT testing. In the moving hidden platform version of the MWT, the platform was moved to a new location within the pool every other day according to a random sequence of locations. Thus, for two consecutive days the platform remained in the same location. On the third day, the platform was moved to a new location for another pair of days, and this was continued until six locations had been completed. Rats received eight trials each day. Data were collected using a video camera and microcomputer automated system (HVS 2020 Plus system). On every trial, time and distance to find the platform was automatically recorded and analyzed. Following recovery from surgery, the rats were divided into 10 groups; sham operated vehicle control ($n=10$), lesion vehicle control ($n=10$), four groups, each receiving one dose of donepezil (0.05, 0.5, 0.1, and 1 mg/kg) ($n=10$), and four groups, each receiving one dose of GT1061 (0.5, 1, 5, and 10 mg/kg) ($n=10$). Each rat received the appropriate drug by gavage 60 min before testing each day. Each day the rats were tested in the MWT. Following the last day of testing, the animals were perfused with 4% paraformaldehyde and their brains removed, cryoprotected, and sectioned for acetylcholinesterase staining (Karnovsky and Roots, 1964).

Delayed matching-to-sample task. The testing apparatus and DMTS training and testing procedures are detailed in a previous publication (Prusky *et al*, 2004). Briefly, adult female Long-Evans rats ($n=10$) were first trained on a simple visual discrimination task, and then in four stages, were gradually shifted to the trial-unique, one-trial learning task. Throughout the testing, rats were rewarded for identifying and swimming toward a correct picture where they could escape from water to a submerged platform. The black-and-white pictures were displayed on computer monitors to eliminate nonvisual cues. The pictures were drawn from a set of more than 100 and on any given day no picture appeared on more than one trial, such that, relative familiarity of pictures is irrelevant. Similarly, the probability of any specific picture being used as a sample or a foil was equal. In the final stage, during each trial, rats viewed a picture in a sample phase, and then in a choice phase, discriminated the sample picture from a novel picture with high accuracy in a different pool located in the same room. The left/right position of the correct picture varied randomly to make the task nonspatial. The delay dependence of memory for the sample picture was measured systematically by varying the delay between the sample and choice phases. All animals received approximately 6 weeks of presurgical training. At 1-week after surgery, DMTS performance was remeasured. On the basis of the presurgical memory performance we selected a 4-min interval for postsurgical testing of drug effects. Presurgical performance at this delay interval between sample and choice was well-above chance for each rat, but was less than maximum accuracy (Figure 1). Thus, it afforded us the opportunity to measure deficits, drug effects, and dose response, at a moderate level of memory difficulty (thereby avoiding floor or ceiling effects). Performance in the DMTS task before and after surgery was compared at the 4-min interval. Drug or drug vehicle was administered by gavage 30-min before testing. Each rat was tested with vehicle solution and three doses each of GT1061 and donepezil, thus, the experimental

Figure 1 Mean preoperative visual matching-to-sample performance at each training delay interval. Choice accuracy was assessed in each animal during 24 trials at each interval, and declined systematically with increasing delays. Data are expressed as the mean $\pm$ SEM ($n = 7$).

design was fully 'within subjects'. The doses of GT1061 were 1, 5, and 10 mg/kg and for donepezil were 0.1, 0.5, and 1.0 mg/kg. There was at least a 48-h time interval between successive tests of drugs or vehicle. The order of testing was always vehicle, GT1061 5, 10, and 1 mg/kg and then donepezil 0.5, 1.0, and 0.1 mg/kg. Between GT1061 and donepezil testing we retested vehicle administration, and found that performance was not different from that observed with the first vehicle test ($55.0 \pm 2.2\%$ correct *vs* $57.3 \pm 1.9\%$ correct, $p = 0.35$, Student's *t*-test for paired data). On each day of testing, each rat received 12 memory trials and the percentage correct choices was calculated. In a separate experiment, we assessed whether GT1061 enhanced cognitive performance in nonlesioned animals. Rats ($n = 4$) were trained as described above and DMTS performance evaluated after oral administration of 10 mg/kg GT1061. This was compared to DMTS performance assessed on the day before and the day subsequent to drug administration.

Surgery involving DMTS rats. After initial training, the cholinergic innervation of the hippocampus and cerebral cortex was depleted by intraventricular microinjection of 192-IgG saporin (1.8 μ g in 5 μ l of phosphate-buffered saline, pH 7.4, over 15 s) into the lateral ventricle of each cerebral hemisphere (coordinates: AP -1.1 , L 1.5 , D -4.2 from bregma).

Pharmacokinetic studies. Male Sprague-Dawley rats (250–275 g) were fasted overnight, anaesthetized for surgery with a combination of Hypnorm/Versed and then catheters (30-gauge Teflon fused to 0.02-inch inside diameter Tygon[®]) were inserted into the abdominal aorta and abdominal vena cava for blood sampling and drug administration, respectively. Animals were administered 50 μ mol/kg (11.2 mg/kg) GT1061 (prepared in saline) by the i.v., i.p., or oral routes and blood samples (250 μ l) obtained from the aortic catheter at various times. Blood was immediately centrifuged and 100 μ l aliquots of plasma were assayed for GT1061 by gas chromatography with electron capture detection. The data were fitted to a 2-compartment open model using PKSolutions software (Summit Research Services, Montrose, CO) to derive pharmacokinetic

parameters. To assess the distribution of GT1061 into the CNS, rats were fasted overnight and then administered 25 μ mol/kg (5.6 mg/kg) GT1061 by gavage. Five minutes later the animals were killed and a blood sample was obtained. The brain was removed and one hippocampus dissected out and frozen in liquid nitrogen, procedures lasting approximately 5 min to complete. The hippocampus was weighed, pulverized, and then both plasma and hippocampus samples assayed for GT1061 by gas chromatography. To assess potential sex and strain differences in the pharmacokinetic properties of GT1061, female Long-Evans rats were instrumented as described above, and blood levels of GT1061 determined after oral administration of 50 μ mol/kg GT1061. The area under the plasma concentration-time curve from 0 to 120 min was 82.1 ± 36.2 μ mol-min/l (SEM, $n = 5$) for female Long-Evans rats, and 125 ± 28.1 μ mol-min/l (SEM, $n = 6$) for male Sprague-Dawley rats. These values were not significantly different ($p = 0.354$, Student's *t*-test for unpaired data).

Hippocampal and aortic sGC activity. Cholinergic innervation of the hippocampus and cerebral cortex of female Long-Evans rats was depleted by intraventricular microinjection of 192-IgG saporin, as described above, and 2 weeks later the aortae and hippocampi from sham-operated and lesioned animals were removed. The preparation of the 100 000 g supernatant fractions of aorta and hippocampus, and enzyme assay conditions for assessment of sGC activity were as described (Bennett *et al*, 1992). Samples (0.2 ml) contained 2 mM L-cysteine and GT1061 (10 μ M–10 mM). Reactions were initiated by the addition of 44–79 μ g hippocampal supernatant protein or 20–31 μ g aortic supernatant protein and terminated by the addition of 0.8 ml of 50 mM sodium acetate (pH 4.0) followed by heating at 90°C for 3 min. The cGMP was quantitated by radioimmunoassay and the protein content determined using bovine serum albumin as the standard.

Phosphorylation of ERK1/2. To test for *in vivo* activation of ERK1/2, separate groups of nonlesioned animals were administered GT1061 (1 mg/kg i.p. (4.5 μ mol/kg)) or phosphate-buffered saline. At 60 min after drug administration, the brain of each animal was rapidly removed, dissected into regions (hippocampus, frontal cortex, and cerebellum), and homogenized in lysis buffer containing 50 mM TRIS, pH 7.4, 50 mM sodium pyrophosphate, 1 mM EDTA, 1% Triton X-100, 1 mM dithiothreitol, 1 mM PMSF, 1 mM NaF, 1 mM NaVO₄ and protease inhibitors (Roche Applied Science). Phosphorylated ERK1/2 (pERK) was determined by ELISA using a commercially available kit (Assay Designs Inc., Ann Arbor, MI). In some experiments, rats were administered the NMDA receptor antagonist, MK-801 (1 mg/kg, i.p.), 20 min before GT1061. For *in vitro* brain slice experiments, rats were anesthetized with halothane, decapitated, and the brain rapidly excised into ice-cold sucrose-substituted Krebs' buffer (342 mM sucrose, 4.8 mM KCl, 1.2 mM KH₂PO₄, 1.3 mM CaCl₂, 1.2 mM MgSO₄, 25 mM NaHCO₃, and 10 mM glucose, pH 7.4, saturated with 95% O₂/5% CO₂). The hippocampus was isolated, 400 μ m slices were cut with a McIlwain tissue chopper, and a small scalpel blade was used to isolate the CA1 region of each slice. Hippocampal mini-slices containing the CA1 region were

equilibrated for 3 h in Krebs' buffer (118 mM NaCl, 4.8 mM KCl, 1.2 mM KH_2PO_4 , 1.3 mM CaCl_2 , 1.2 mM MgSO_4 , 25 mM NaHCO_3 , and 10 mM glucose, saturated with 95% O_2 /5% CO_2) at room temperature, and then exposed to drug or vehicle for 30 min. At the end of the incubation, the slices were homogenized in lysis buffer, and the content of pERK determined by ELISA.

Synthesis of GT1061. GT1061, prepared under GMP conditions, was obtained from GB Therapeutics Ltd (Mississauga, ON, Canada). GT1061 was synthesized from 4-methyl-5-(2-hydroxyethyl)-1,3-thiazole by adaptation of standard acidic nitration procedures, followed by salt formation with HCl and crystallization. Purity as assessed by chromatography and spectroscopy was greater than 98%. GT1061 is a crystalline white solid: mp 107–108°C (d).

Statistical analyses. In the MWT and the DMTS task, data were evaluated with a repeated measures ANOVA procedure (SPSS 11.0) as previously recommended (Kirk, 1982; Ringo, 1991). *Post hoc* tests were conducted using Fisher's LSD method. Data from other experiments were analyzed by the appropriate statistical test, as indicated. Differences were considered to be statistically significant if $p < 0.05$.

RESULTS

Histology

Examination of the acetylcholinesterase stained sections using NIH Image densitometry showed that after bilateral injections of 192 IgG-saporin into the nucleus basalis magnocellularis and medial septum, animals exhibited significantly less acetylcholinesterase staining in both the neocortex and hippocampus as compared to sections obtained from control animals ($t = -12.9$, $p < 0.0001$, and $t = -7.38$, $p < 0.0001$, respectively, Figure 2). Compared to the focal injections of 192 IgG-saporin, we found a similar degree of depletion of acetylcholinesterase in the neocortex and hippocampus after intraventricular administration of 192 IgG-saporin (both p 's > 0.6).

Moving Hidden Platform Morris Water Task

We conducted separate ANOVAs for donepezil-treated and GT1061-treated groups on latency to find the platform on days when it was in novel locations. There were two within subject factors, day and trial. In the case of donepezil, the ANOVA shows that the groups did differ significantly from one another ($F(5,46) = 2.55$, $p < 0.04$). The animals improved significantly over days ($F(5,230) = 6.07$, $p < 0.0001$) and trials ($F(7,322) = 184.67$, $p < 0.0001$). In the case of GT1061, the ANOVA shows that the groups did differ significantly from one another ($F(5,48) = 3.33$, $p < 0.01$). The animals improved significantly over days ($F(5,240) = 7.52$, $p < 0.0001$) and trials ($F(7,336) = 90.3$, $p < 0.0001$). In both cases, *post hoc* analysis shows that the sham control group exhibited shorter latencies compared to all other groups (all p 's < 0.0001).

For further illustration of the main trend, Figure 3a shows the average latency to find the hidden platform on trial 2 on days when the platform was in novel locations. Previous

Figure 2 Acetylcholinesterase staining in rat brain after either bilateral injections of 192-IgG saporin into the nucleus basalis magnocellularis and medial septum (middle panels), or intraventricular injections of 192 IgG-saporin (bottom panels). Stained sections from vehicle control (top panels), or 192 IgG-saporin-treated animals were analyzed using NIH image densitometry. Animals treated with 192 IgG-saporin exhibited significantly less staining in both the cerebral cortex ($t = -12.9$, $p < 0.0001$) and hippocampus ($t = -7.38$, $p < 0.0001$) compared to sections from the control animals.

Figure 3 Effect of oral administration of GT1061 (0.5–10 mg/kg) and donepezil (0.05–1 mg/kg) in the moving hidden platform Morris water task. (a) The data represents the average time to find the hidden platform on trial 2 on days when the platform was in novel locations. Data are expressed as the mean $\pm$ SEM ($n = 10$). The groups did not differ significantly from one another ($F(9,78) = 1.66$, $p = 0.11$, repeated measures ANOVA). (b) The data represents the average time to find the hidden platform on trial 2 on days when the platform was in the same location. Data are expressed as the mean $\pm$ SEM ($n = 10$). * $p < 0.05$ vs lesion, repeated measures ANOVA ($F(9,78) = 2.09$, $p = 0.04$).

work in our laboratory has shown that this trial provides a good index of new learning. Trial 1 performance was not different between any of the groups and all of the groups improved by approximately 16 s between trial 1 and 2. An ANOVA on the trial 2 data revealed that the treatment groups were not different from one another ($F(9,78) = 1.66$, $p = 0.11$), indicating that in lesioned animals, neither donepezil nor GT1061 altered the animal's capacity for new learning.

In similar fashion, we analyzed the latency to find the platform on days when the hidden platform remained in the same location as the previous day for groups receiving donepezil. The ANOVA shows that the groups did not differ significantly from one another ($F(5,46) = 0.42$, $p = 0.83$). The groups did improve significantly over days ($F(5,230) = 4.4$, $p < 0.0007$), and trials ($F(7,322) = 59.4$, $p < 0.0001$). For the groups receiving GT1061 there was a significant difference between the groups ($F(5,48) = 2.39$, $p = 0.05$). The groups did improve significantly over days ($F(5,25) = 3.62$, $p < 0.004$). Again, for simplicity of illustration, Figure 3b shows the group data for performance on trial 2 for all days when the platform was in the same location as the previous day. Earlier work has shown that this is a sensitive index of memory performance, which should reveal effects from the previous day's learning and the first trial reminder. The trial 1 performance showed a similar trend to that of trial 2, but we selected trial 2 because of its lower within group variance and hence greater sensitivity. A repeated measures ANOVA examining the group differences on trial 2 only when the platform was in the same locations as the previous day shows that the groups differed significantly from one another ($F(9,78) = 2.09$, $p = 0.04$). With donepezil, doses of 0.1 mg/kg or higher produced significant benefit in cholinergic-depleted rats (p 's < 0.01). GT1061, at an oral dose of 10 mg/kg, produced the same beneficial effect on memory performance as did donepezil ($p = 0.54$) and produced performance that was not different from sham lesion rats ($p = 0.60$) but superior to depleted control rats ($p < 0.002$). There were no significant differences in swim speeds associated with cholinergic depletions or either drug (all p 's > 0.2).

Delayed Matching to Sample Task

The overall repeated measures ANOVA on percent correct showed statistically significant differences between conditions ($F(7,63) = 25.0$, $p < 0.001$). Before surgery, rats made 75.0% correct choices at a 4-min delay between sample and choice phases and this was significantly decreased to approximately 55.0% in the same animals after cholinergic depletion ($p < 0.0006$). This deficit in the visual memory task was large and sustained throughout the experiment. All doses of GT1061 significantly improved performance. Indeed, the highest dose of GT1061 restored performance in the DMTS task to a level that was not significantly different from the presurgery level (Figure 4). The two higher doses of donepezil improved performance, but the lowest dose (0.1 mg/kg) had no effect. In this assay, GT1061 was equipotent (on a per mg basis) with donepezil at the 1 mg/kg dose, and at 10 mg/kg, the improvement in performance was significantly greater than that of any of the

Figure 4 Effect of oral administration of GT1061 (1–10 mg/kg) and donepezil (0.1–1 mg/kg) in the DMTS task, which measures recognition memory performance. The decreased accuracy in the task after cholinergic depletion was reversed at all doses of GT1061 and at higher doses of donepezil. Data are expressed as the mean $\pm$ SEM ($n = 10$). * $p < 0.01$ vs postsurgery, repeated measures ANOVA ($F(7,63) = 25.0$, $p < 0.001$).

doses of donepezil (p 's < 0.01). It should be noted that the oral bioavailability of donepezil approaches 100%, whereas that of GT1061 is about 20% (see below). In the study that assessed whether GT1061 improved performance in nonlesioned animals, rats made $77 \pm 2\%$ (SEM) correct choices on the trial day before GT1061, and $83.5 \pm 4.9\%$ correct choices on the day after. Oral administration of 10 mg/kg GT1061 improved performance to $87.8 \pm 4.3\%$ correct choices, but the increase was not statistically significant ($F(2,6) = 3.4$, $p = 0.161$).

Pharmacokinetic Studies

GT1061 was rapidly absorbed after i.p. or oral administration with peak plasma concentrations occurring 3 min after dosing (Figure 5). Calculation of the AUCs indicated an oral bioavailability of $20.5 \pm 11.7\%$ (range of 9–41.3%), and an i.p. bioavailability of $52.8 \pm 10.4\%$ (range of 36.6 to 64.7%). After i.v. administration, values for V_d (l/kg), Cl (l/min/kg) and elimination half-life (min) (derived from V_d and Cl) were 2.5, 0.079, and 21.6, respectively. For oral administration the values were: 3.4, 0.072, and 32.6, and for i.p. administration the values were: 2.2, 0.08, and 19.0. Calculations for oral and i.p. V_d and Cl values used the derived bioavailability values for the fractional dose absorbed. In the brain bioavailability study, the mean plasma level of GT1061 5 min after oral dosing was 0.52 ± 0.25 nmol/ml (SD, $n = 6$), whereas the mean tissue concentration of GT1061 in hippocampus (expressed as nmol/g wet weight, roughly equivalent to nmol/ml assuming that 1 g of tissue has a volume of 1 ml) was 0.67 ± 0.42 nmol/g wet weight (SD, $n = 6$) (Figure 6). Plasma and hippocampal levels of GT1061 were not significantly different from each other (Student's t -test for paired data), and in every animal the brain levels of GT1061 were higher than the plasma levels (Figure 6). The ratio of brain to blood GT1061 was 1.3 ± 0.28 , a value that may underestimate the true brain penetration of GT1061, since during the time between killing and completion of the dissection of the hippocampus (ca. 5 min) further metabolic biotransformation of GT1061 in the brain tissue may have occurred.

Figure 5 Time course of plasma concentrations of GT1061 after i.v., i.p., and oral administration of 50 $\mu\text{mol/kg}$ GT1061. Data are expressed as the mean $\pm$ SEM ($n=5-6$). The oral bioavailability of GT1061 was $20.5 \pm 11.7\%$.

Figure 6 Plasma and hippocampal concentrations of GT1061 (25 $\mu\text{mol/kg}$) 5 min after oral administration. The mean plasma concentration of GT1061 was 0.52 ± 0.25 nmol/mL (SD, $n=6$) and the mean tissue concentration of GT1061 in hippocampus was 0.67 ± 0.42 nmol/g wet weight (SD, $n=6$). These values were not significantly different to each other (Student's *t*-test for paired data). The ratio of brain to plasma GT1061 was 1.3 ± 0.28 .

GT1061 exhibited pharmacokinetic properties similar to classic nitrates such as GTN and isosorbide dinitrate. Absorption was rapid, and bioavailability after oral administration was somewhat variable (9–41%). The large volume of distribution (2.2–3.4 l/kg, depending on route of administration) indicated that the drug was distributed extensively in tissues. This was clearly evident in the brain, where drug concentrations exceeded those in plasma. Extrapolation of the plasma levels of drug obtained in the pharmacokinetic study to the doses of drug that demonstrated efficacy in the behavioral tests suggest that therapeutic concentrations of GT1061 are in the submicromolar range.

Hippocampal and Aortic sGC Activity

GT1061 caused a concentration-dependent increase in sGC activity in both aortic and hippocampal homogenates (Figure 7), although the degree of activation in the hippocampus was markedly greater. Enzyme activation by GT1061 in hippocampal homogenates from control and

Figure 7 Comparison of activation of sGC by GT1061 in the hippocampus and aorta after depletion of cholinergic innervation of the hippocampus and cerebral cortex by intraventricular microinjection of 192 IgG-saporin. The basal levels of sGC activity in hippocampal homogenates from control and lesioned animals were 7.7 ± 1.8 and 8.4 ± 2.2 pmol/min/mg protein, respectively, and in aortic homogenates from control and lesioned animals were 86.1 ± 9.4 and 82.3 ± 9.1 pmol/min/mg protein, respectively. Data are expressed as the fold increase in activity over basal (mean $\pm$ SEM, $n=7$). Enzyme activation was not altered in homogenates from lesioned animals. Enzyme activation was significantly greater ($p < 0.05$) in homogenates from hippocampus compared to aorta at concentrations of GT1061 of 0.3 mM and above (one-way ANOVA (($F(3,27) = 6.02-9.59$, $p < 0.05$) and Newman-Keuls *post hoc* test).

lesioned animals was virtually identical, indicating that cyclic GMP signaling remained intact in cholinergic-deficient animals. Enzyme activity in hippocampal homogenates increased 16- to 20-fold at the highest concentration of GT1061, whereas in the aorta, the maximal increase in activity was only 2- to 3-fold. In broken cell preparations, with very rare exceptions, nitrates do not activate sGC above basal levels in the absence of certain added thiols (Artz *et al*, 2001; Zavorin *et al*, 2001); GT1061 is not an exception.

Phosphorylation of ERK1/2

In the *in vivo* assessment of phosphorylated ERK1/2 (pERK), GT1061 increased the levels of pERK in the cerebral cortex, but had no effect in the cerebellum (Figure 8a). In the hippocampus, pERK levels were increased by GT1061, but the increase did not reach statistical significance. However, when GT1061 was administered to animals that had been pretreated with the NMDA receptor antagonist, MK-801, the increase in hippocampal pERK caused by GT1061 was significant (Figure 8a, inset), suggesting that in the hippocampus, the effects of GT1061 are more evident in circumstances where input from upstream effectors has been compromised. Qualitatively, this is consistent with the lack of a significant effect of GT1061 on cognition in unlesioned animals. We also compared the effect of GT1061 to that of the cholinergic muscarinic receptor agonist, carbachol, in hippocampal mini-slices containing the CA1 region (Figure 8b). Both drugs significantly increased the amount of pERK in this *in vitro* preparation.

Figure 8 Effect of GT1061 on ERK phosphorylation. (a) Levels of pERK1/2 in different brain regions 60 min after a single i.p. injection of GT1061 (1 mg/kg). Data are expressed as a percentage of the levels of pERK measured in saline-injected animals (mean $\pm$ SEM, $n = 6$). * $p < 0.05$ vs control, Student's *t*-test for unpaired data. Inset: Levels of pERK in the hippocampus 60 min after a single i.p. injection of GT1061 (1 mg/kg) with or without prior treatment of animals with the NMDA receptor antagonist, MK-801 (1 mg/kg, i.p.). Data are expressed as a percentage of the levels of pERK measured in saline-injected animals (mean $\pm$ SEM, $n = 6$). * $p < 0.05$ vs control, Student's *t*-test for unpaired data. (b) Levels of pERK1/2 in the CA1 region of hippocampal mini-slices *in vitro* after a 30 min incubation with GT1061 (50 μ M) or the cholinergic muscarinic receptor agonist, carbachol (50 μ M). Both compounds induced an equivalent increase in the content of pERK. Data are expressed as the mean $\pm$ SEM ($n = 5$). * $p < 0.05$ vs control, repeated measures ANOVA ($F(2,8) = 9.62$, $p = 0.0074$). Pair wise comparisons were performed using Newman-Keuls *post hoc* test.

DISCUSSION

Development of pharmacologically effective treatments that alleviate cognitive disorders is a central objective in neuroscience. The high incidence of dementia, in particular AD, in the aging population provides strong impetus to search for new compounds with the potential to reverse the deficits in short-term memory. The neuropathological changes underlying age-related cognitive decline and AD are complex and poorly understood. However, the loss of forebrain cholinergic neurons that project to the cerebral cortex and hippocampus is a hallmark of this disease. In addition, the deposition of A β in both the brain parenchyma and cerebral blood vessels, and the formation of intracellular neurofibrillary tangles containing tau-protein are also observed. Studies in Alzheimer's patients have revealed that the extent of cholinergic depletion correlates with the severity of memory deficits (Mesulam, 2004). Thus, animal models of cortical and hippocampal cholinergic depletion have been widely used to induce relevant

cognitive deficits for the purpose of identifying potential new therapeutic compounds for the treatment of AD.

It is well established that dysfunction in the hippocampus causes spatial learning deficits in tasks such as the MWT (Redish and Touretzky, 1998; Silva *et al*, 1998a). However, the role of the hippocampus in visual memory and recognition tasks has been controversial. It was recently confirmed that the hippocampus is essential for nonspatial visual recognition memory in the rat (Prusky *et al*, 2004). The observation of similarities between cognitive processing in the rat and human is important, but the development of a reliable rat model of recognition memory deficit also provides a powerful tool for development of drugs for AD and neurodegenerative diseases. It is noteworthy that visual association tests are reported to detect, with high specificity, a substantial proportion of AD patients up to a year before diagnosis (Lindeboom *et al*, 2002). This paper demonstrates that novel nitrates are a new class of cognition enhancing agents, and also validates the use of the rat DMTS model by assay of the AD drug donepezil. In the original DMTS work (Prusky *et al*, 2004), the perirhinal cortex was lesioned by aspiration and the hippocampus lesioned by direct injection of excitotoxic doses of NMDA. We show herein that intraventricular infusion of 192 IgG-saporin induces a reliable cholinergic lesion resulting in visual recognition memory impairment, which is reversed by treatment with GT1061 and donepezil. We note here that with the present results we cannot determine the extent to which damage to noncholinergic cells may contribute to the memory deficit.

Our observation that a novel nitrate, NO mimetic compound exerts beneficial effects in multiple learning paradigms, supports the idea that activation of the sGC/cGMP signal transduction cascade enhances synaptic efficacy in brain regions responsible for the acquisition, storage, and retention of new learning. Activation of NO/cGMP signaling represents a new drug target for AD, a disease characterized by generalized, progressive decline in cognitive function due to diffuse neuronal loss and synaptic failure in the hippocampus and cerebral cortex (Scheff and Price, 2003; Selkoe, 2002).

Acetylcholine (ACh), a major modulatory neurotransmitter in the brain, is critically involved in learning and memory. In the cortex and hippocampus, the procognitive actions of ACh are mediated via muscarinic receptors whose activity is primarily excitatory, and involves multiple different ionic conductances. In the hippocampus, cholinergic muscarinic receptor activation leads to increased tissue levels of cGMP (Tonnaer *et al*, 1991). Importantly, the inhibition of the slow afterhyperpolarizing current (a calcium-activated potassium conductance that underlies spike-frequency adaptation) induced by muscarinic receptor activation in hippocampal CA1 pyramidal neurons can be blocked by inhibitors of sGC and PKG (Krause and Pedarzani, 2000). These data suggest that the neuromodulatory effects of ACh in the brain involve, at least in part, the NO/sGC/cGMP signal transduction cascade.

Behavioral studies have demonstrated that NMDA receptors also play an important role in both spatial working memory and long-term memory processes. Blockade of NMDA receptors, or inhibition of NOS activity, impairs performance in the Y-maze test, a model of spatial

working memory (Yamada *et al*, 1996). The impairment induced by NMDA receptor blockade could be reversed by intracerebroventricular administration of L-arginine, a nitrosothiol NO donor, or a cGMP analog. Taken together, these studies suggest that both ACh and glutamate activate receptor systems coupled to the NO/sGC/cGMP signal transduction cascade and that this biochemical cascade is important for synaptic plasticity and the formation of memory.

In previous studies, we found that the novel nitrate, GT715, was less potent than GTN for relaxation of vascular smooth muscle and for decreasing mean arterial pressure, and more effective than GTN for activating sGC in the hippocampus. In simile with these data, the activation of sGC in broken cell preparations of aorta and hippocampus revealed that GT1061 selectively increased enzyme activity in the hippocampus (16- to 20-fold in hippocampus vs 2- to 3-fold in aorta), confirming that the structure of the nitrate NO mimetic influences tissue selectivity for sGC activation. Additionally, sGC activity was not altered in tissue homogenates from lesioned animals, indicating that cGMP signaling remained intact after depletion of cholinergic innervation of the hippocampus. Soluble GC is a heterodimer of α and β subunits, and recent studies of the distribution of α_1 and α_2 subunits of sGC have shown that the α_2 subunit is expressed in relatively high abundance in brain, especially in the hippocampus, where the $\alpha_2\beta_1$ rather than the $\alpha_1\beta_1$ heterodimer of sGC is the major isoform present (Mergia *et al*, 2003). Furthermore, the α_2 subunit interacts with a PDZ domain of PSD95 (Russwurm *et al*, 2001). As this scaffolding protein couples the NMDA receptor with neuronal NOS, this suggests an important role for the $\alpha_2\beta_1$ heterodimer isoform in synaptic transmission. The enhanced activation of hippocampal sGC by GT1061 suggests that there is selectivity for targeting the $\alpha_2\beta_1$ isoform of sGC.

A number of kinase signaling cascades may be involved in signal transduction by NO/cGMP resulting in enhanced cognition and memory; chief among these being the ERK cascade (Lu *et al*, 1999). Activation of ERK1/2 is believed to play a critical role in cognition and memory mediated through phosphorylation and activation of CREB, and may serve as a point of convergence for a number of upstream signaling molecules, including glutamate acting via NMDA receptors and ACh acting via cholinergic muscarinic and nicotinic receptors (Sweatt, 2004). Activation of transcription regulating factors, such as CREB, leading to protein synthesis is a requirement for the consolidation of new learning into long-term memory (Silva *et al*, 1998b). ERK1/2 activation can be mediated via the protein kinases, PKA or PKC, whereas CREB can be activated and phosphorylated via CaMK IV, PKA, and RSK2, the last mediating the activation of CREB by the ERK cascade (Xing *et al*, 1996). Studies suggest that PKG provides a parallel pathway to PKA-signaling in both phases of LTP, with PKG and PKA pathways performing complementary roles (Lu *et al*, 1999). NO/cGMP and PKG contribute to CREB phosphorylation, mediated by the ERK cascade, and also possibly in part via the CaMK pathway (Lu *et al*, 1999). Interestingly, experiments with YC-1, an agent that augments sGC activation, produced an enhancement of LTP in rat hippocampus and amygdala via an NO/cGMP/PKG/ERK pathway culminating

in CREB phosphorylation (Chien *et al*, 2003). In *in vitro* brain slice experiments, we found that GT1061 increased the levels of pERK in the CA1 region of the hippocampus. In addition, the finding that *in vivo* administration of GT1061, at doses associated with cognitive improvement, increased ERK1/2 phosphorylation in the hippocampus and cerebral cortex, provides evidence for a mechanism of GT1061 cognition enhancement via sGC/cGMP signaling mediated at least in part by the ERK cascade.

Arancio and co-workers reported recently that activation of sGC or addition of a cGMP analog were able to protect against the A β -induced impairment of hippocampal CA1 region LTP (Puzzo *et al*, 2005). In mouse hippocampal slices treated with agents in the presence of A β_{1-42} before induction of LTP, A β inhibited LTP and blocked elevation of pCREB in response to tetanus; the activation of sGC had no effect on basal LTP and pCREB levels in the absence of the tetanus stimulation. A further study provided a pathway for A β -induced disruption of NO/cGMP signaling initiated by binding of A β to the α_7 nicotinic ACh receptor (Snyder *et al*, 2005). Treatment of neurons with A β was reported to produce a depression of NMDA-evoked currents and reduced CREB phosphorylation via a mechanism involving dissociation of the synaptic protein, PSD95, from the NMDA receptor. It is reasonable to assume that this event will seriously disrupt NO/cGMP signaling, since both sGC and nNOS are known to be bound to NMDA receptor-associated PSD95 (Brenman and Bredt, 1997; Mergia *et al*, 2003; Russwurm *et al*, 2001). Thus, evidence is growing justifying intervention with NO mimetic drugs in treatment of AD.

In summary, we have shown for the first time that the visual recognition memory deficit induced by cholinergic depletion in a rat model can be fully reversed by drug administration. All current clinical agents approved for use in mild to moderate AD attempt to salvage the residual functionality of damaged and degenerating cholinergic neurons, whereas we demonstrate herein an alternative approach whereby the function of the signaling apparatus is modulated to compensate for damage to the cholinergic circuitry. The drug that supports this postulate is a novel nitrate NO mimetic, GT1061. Importantly, this compound displays significant oral bioavailability and brain penetration, and efficacy that, in the visual memory recognition DMTS task, was superior to the clinical AD drug, donepezil. Moreover, GT1061 reverses the cognitive deficits induced by depletion of forebrain ACh in two different learning paradigms, reinforcing both the previous observations with the novel nitrate, GT715, and the potential for development of nitrate NO mimetics as new therapeutic agents. GT1061 induces a selective activation of sGC in the hippocampus compared to the vasculature, predicting that the beneficial CNS actions of the drug will occur at lower plasma concentrations than will any potential undesirable hypotensive actions. GT1061 was observed to activate a pathway in the hippocampus, the ERK cascade, (a) known to be modulated by sGC/cGMP signaling, and (b) which represents a necessary pathway in the formation of long-term memory. Novel nitrates, as represented by GT1061, are a class of compounds that can be exploited in the discovery and development of new therapeutic agents for the treatment of neurodegenerative diseases such as Alzheimer's disease.

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