

Evidence That Methylphenidate Enhances the Saliency of a Mathematical Task by Increasing Dopamine in the Human Brain

Nora D. Volkow, M.D.

Gene-Jack Wang, M.D.

Joanna S. Fowler, Ph.D.

Frank Telang, M.D.

Laurence Maynard, M.D.

Jean Logan, Ph.D.

Samuel J. Gatley, Ph.D.

Naomi Pappas, M.S.

Christopher Wong, M.S.

Paul Vaska, Ph.D.

Wei Zhu, Ph.D.

James M. Swanson, Ph.D.

Objective: Methylphenidate is the most commonly prescribed drug for attention deficit hyperactivity disorder (ADHD), yet its therapeutic mechanisms are poorly understood. The objective of this study was to assess if methylphenidate, by increasing dopamine (neurotransmitter involved in motivation) in brain, would enhance the saliency of an academic task, making it more interesting.

Method: Healthy subjects (N=16) underwent positron emission tomography with [¹¹C]raclopride (dopamine D₂ receptor radioligand that competes with endogenous dopamine for binding) to assess the effects of oral methylphenidate (20 mg) on extracellular dopamine in the striatum. The authors compared the effects of methylphenidate during an academic task (solving mathematical problems with monetary reinforcement) and a neutral task (passively viewing cards with no remuneration). In parallel, the effects of methylphenidate on the interest that the academic task elicited were also evaluated.

Results: Methylphenidate, when coupled with the mathematical task, significantly increased extracellular dopamine, but this did not occur when coupled with the neutral task. The mathematical task did not increase dopamine when coupled with placebo. Subjective reports about interest and motivation in the mathematical task were greater with methylphenidate than with placebo and were associated with dopamine increases.

Conclusions: The significant association between methylphenidate-induced dopamine increases and the interest and motivation for the task confirms the prediction that methylphenidate enhances the saliency of an event by increasing dopamine. The enhanced interest for the task could increase attention and improve performance and could be one of the mechanisms underlying methylphenidate's therapeutic effects. These findings support educational strategies that make schoolwork more interesting as nonpharmacological interventions to treat ADHD.

(*Am J Psychiatry* 2004; 161:1173–1180)

Attention deficit hyperactivity disorder (ADHD) is the most commonly diagnosed behavioral disorder of childhood (1). Methylphenidate and amphetamine are the most frequently used treatments for ADHD (1). Despite methylphenidate's widespread use, its mechanisms of action are poorly understood. Methylphenidate blocks the dopamine transporters, and this indirect dopamine agonist effect may be critical for its therapeutic effects (2). Specifically, stimulant-induced dopamine increases are believed to decrease background firing rates and increase signal-to-noise ratio of striatal cells (3), which we postulate as a mechanism for improving attention by enhancing task-related neuronal cell firing. Since dopamine also modulates salience and motivation (4), we postulate that small and brief task-induced dopamine increases would be amplified by methylphenidate, which would increase the saliency of the task and thus improve performance (5).

Using positron emission tomography (PET), we have shown that a relatively large therapeutic dose of oral methylphenidate (60 mg) significantly blocked dopamine transporters in all subjects but did not increase extracellu-

lar dopamine in all of them (6). We reasoned that methylphenidate-induced increases in dopamine were due not only to dopamine transporter blockade but also to the rate of dopamine release, which is mostly a function of dopamine cell firing (7). Since dopamine cells fire in response to salient stimuli, we postulated that methylphenidate-induced dopamine transporter blockade (the main mechanism for removal of extracellular dopamine [8]) would amplify dopamine-induced increases to salient stimuli and thus be context dependent. In the present study we tested whether methylphenidate would enhance increases in extracellular dopamine in response to an academic task (solving mathematical problems with monetary reinforcement), relative to a neutral task (passively viewing scenery cards with no remuneration). We used a mathematical task because this type of task is frequently used as a measure to assess treatment efficacy in ADHD. Although in a classroom setting performance is not monetarily remunerated, it is often reinforced by grading (9). We chose as the neutral task passive viewing of scenery pictures rather than a mathematical task with no remuneration because

TABLE 1. Plasma Methylphenidate Concentration Over Time in 16 Healthy Subjects During Performance of a Math Task With a Monetary Reward and a Neutral Noncompensated Task^a

Time (minutes)	Plasma Methylphenidate Concentration (ng/ml)			
	During Math Task		During Neutral Task	
	Mean	SD	Mean	SD
60	9.0	6.2	8.8	6.0
90	9.4	4.6	8.0	3.8
120	8.9	3.5	8.2	3.6

^a Following oral methylphenidate administration (20 mg), subjects underwent positron emission tomography scans while solving mathematical problems with monetary reinforcement (math task) or passively viewing pictures of scenery with no remuneration (neutral task). The concentration of methylphenidate in plasma did not differ between conditions.

we wanted to optimize the contrast between a task that might maximize dopamine increases (mathematical task) and a task that would minimize dopamine increases (viewing neutral pictures).

We used PET imaging with [¹¹C]raclopride, a dopamine D₂ receptor ligand sensitive to competition with endogenous dopamine (10), to measure the relative changes in extracellular dopamine induced by methylphenidate when administered before the mathematical task and before the neutral task. Because [¹¹C]raclopride binding is highly reproducible (11), differences in binding between placebo and intervention predominantly reflect intervention-induced changes in extracellular dopamine (12). Our a priori hypothesis was that methylphenidate, by blocking dopamine transporters, would amplify the dopamine increases induced by the task, thus enhancing its saliency.

Method

Subjects

Sixteen healthy subjects (14 men and two women; mean age=35 years, SD=8) selected from a pool of subjects who responded to an advertisement were studied. Subjects were initially screened by phone and then evaluated at Brookhaven National Laboratory by a physician (G.-J.W.) to ensure that there was 1) no current or past psychiatric or neurological disease, drug abuse, or significant medical illness; 2) no medications being taken (including over-the-counter medications); and 3) no pregnancy. Physical examination and laboratory test results were also obtained. Magnetic resonance imaging scans were not performed, so we could not corroborate a lack of structural brain abnormalities. On average, out of every five subjects initially screened one was recruited, and all of the 16 subjects recruited completed the study. Prescan urine tests ensured the absence of any psychoactive drug, including sedative drugs, and the absence of pregnancy in the female subjects. Subjects were monetarily compensated for their participation in the study. Written informed consent was obtained from all subjects after complete description of the study and following the guidelines set by the Institutional Review Board at Brookhaven National Laboratory.

Scans

PET scans were run on a whole-body, high-resolution positron emission tomograph (Siemens/CTI ECAT HR+, with 4.6×4.6×4.2 mm National Electrical Manufacturers Association resolution at center of field of view and 63 slices) in three-dimensional dy-

namic acquisition mode using [¹¹C]raclopride as a dopamine D₂ receptor ligand (13). Methods for positioning of subjects, catheterizations, transmission scans, and blood sampling and analysis have been published (11). Briefly, emission scans were started immediately after injection of 4–8 mCi of [¹¹C]raclopride (specific activity: 0.5–1.5 Ci/μmol at the end of bombardment). A series of 20 emission scans were obtained from time of injection up to 54 minutes. Arterial sampling was used to measure total carbon-11 and unchanged [¹¹C]raclopride in plasma.

Subjects were scanned four times with [¹¹C]raclopride under four conditions defined by the drug/task combinations: 1) neutral task preceded by placebo administration (placebo/neutral task); 2) neutral task preceded by methylphenidate administration (methylphenidate/neutral task); 3) mathematical task preceded by placebo administration (placebo/math task); and 4) mathematical task preceded by methylphenidate administration (methylphenidate/math task). Subjects were scanned on 2 days and were blinded as to whether they received methylphenidate or placebo. On one day the first scanning condition was placebo/neutral task and the second condition 2 hours later was methylphenidate/math task. On the other day, the first scanning condition was placebo/math task and the second condition 2 hours later was methylphenidate/neutral task. The order of these two sessions was balanced across subjects. The placebo (calcium carbonate) or methylphenidate (20 mg) was given orally 60 minutes prior to the mathematical and the neutral tasks, which were started 15 minutes prior to the injection of [¹¹C]raclopride. Venous blood was drawn for quantification of plasma concentrations of methylphenidate prior to and 60, 90, and 120 minutes after oral methylphenidate using capillary gas chromatography/mass spectrometry (14).

Stimulation

For the mathematical task, the subjects were asked to answer a series of mathematical problems that were preselected for each subject so that they would respond correctly to 80% of them. This also required that the task difficulty be adjusted depending on performance during the study. Problems were presented in a series of colored cards at a rate of one card per minute. Correct responses were remunerated by 25 cents to one dollar depending on the difficulty of the question, which were chosen so that subjects would earn approximately 50 dollars per session. For the neutral task, subjects were presented with cards that showed images of scenery and were not asked to provide responses nor were they remunerated. The mathematical and the neutral tasks began 15 minutes prior to radiotracer injection and continued for 45 minutes.

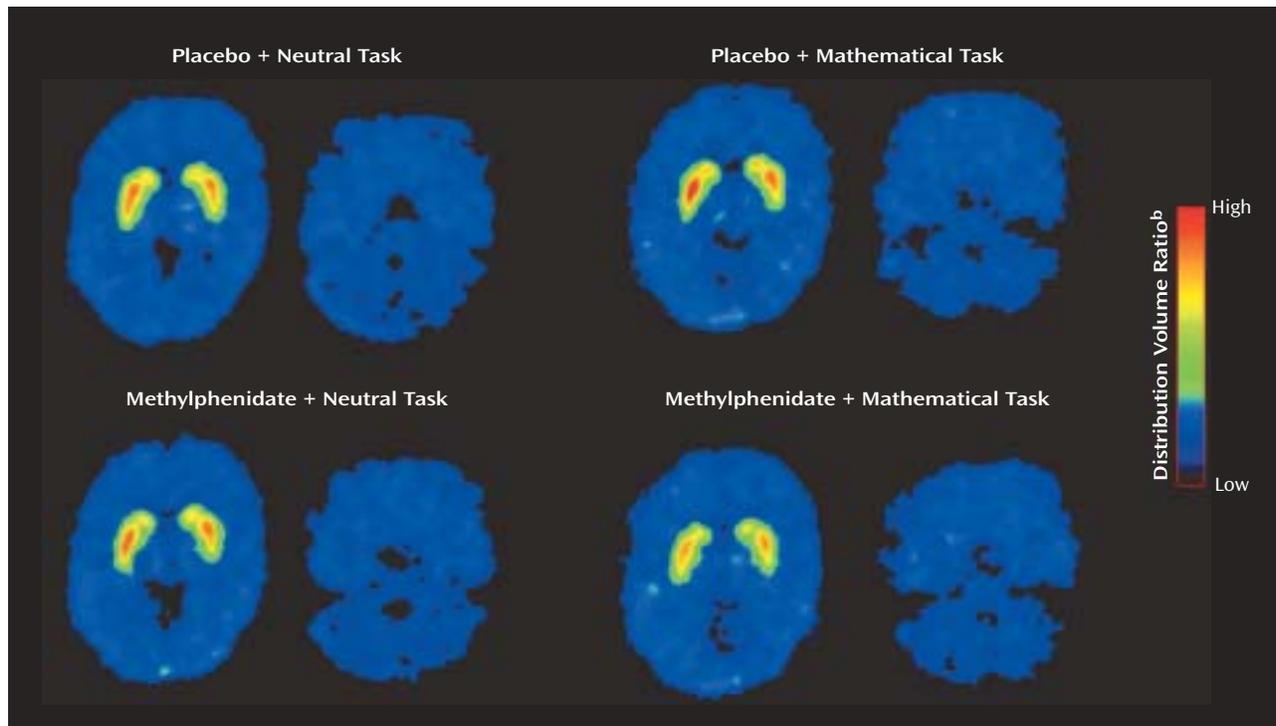
Behavioral Measures

Subjects were asked to provide subjective ratings (from 1 to 10 for which 1=low and 10=high) of the mathematical task for the following descriptors: interesting, exciting, motivating, boring, and tiresome. These ratings were obtained just prior to [¹¹C]raclopride injection and 15 minutes after initiation of the mathematical task. Heart rate and blood pressure were monitored continuously. The behavioral measures were lost for one of the subjects.

Image Analysis

Regions of interest were outlined for the striatum and cerebellum as described previously (11). Briefly, regions of interest were initially outlined on the individual's summed baseline [¹¹C]raclopride image (images obtained between 15 to 54 minutes) and were then projected into the dynamic [¹¹C]raclopride images to generate time activity curves for the striatum and cerebellum. These time activity curves for tissue concentration along with the time activity curves for unchanged tracer in plasma were used to calculate [¹¹C]raclopride's transfer constant from plasma to brain (K₁) and the distribution volume, which corresponds to the equi-

FIGURE 1. Distribution Volume of [¹¹C]Raclopride at the Level of the Striatum and Cerebellum During PET Scans of a Healthy Subject Performing a Math Task With a Monetary Reward or a Noncompensated Neutral Task After Receiving Placebo or Methylphenidate^a



^a Following oral administration of methylphenidate (20 mg) or placebo, subjects underwent positron emission tomography scans while solving mathematical problems with monetary reinforcement (math task) or passively viewing pictures of scenery with no remuneration (neutral task). Binding of [¹¹C]raclopride in the striatum was lowest for the math task in the methylphenidate condition.

^b Equilibrium measurement of the ratio of tissue concentration to plasma concentration in the striatum relative to the cerebellum.

librium measurement of the ratio of tissue concentration to plasma concentration, in the striatum and cerebellum using a graphical analysis technique for reversible systems (15). The ratio of distribution volume in the striatum to that in the cerebellum corresponds to $(B_{\max}/K_d)+1$ and is insensitive to changes in cerebral blood flow (16). Differences between conditions were quantified as percent change in B_{\max}/K_d with respect to the placebo/neutral task condition, which was used as the baseline condition. The effects of methylphenidate on the mathematical task were quantified as percent change in B_{\max}/K_d with respect to the placebo/math task condition.

Data Analysis

Differences in K_1 , distribution volume, and B_{\max}/K_d between conditions were tested using repeated analysis of variance (ANOVA), and post hoc *t* tests were then used to determine in which conditions the effects differed from the baseline condition (placebo/neutral task). The effects of methylphenidate on the descriptor ratings of the mathematical task were tested using paired *t* tests between the placebo/math task condition and the methylphenidate/math task condition. Pearson product-moment correlations were used to assess the relationship between the changes in dopamine between the placebo/math task condition and the methylphenidate/math task condition and the changes in the descriptor ratings for the two conditions.

Results

The plasma methylphenidate concentrations were the same in the methylphenidate/neutral task and the meth-

ylphenidate/math task conditions, and they were in the typical therapeutic range of about 8–9 ng/ml at the time when the PET [¹¹C]raclopride measures were obtained (Table 1). The distribution volume images for one of the subjects are shown in Figure 1 for the four testing conditions: placebo/neutral task (condition used as baseline), placebo/math task (assessing the effects of the mathematical task by itself), methylphenidate/neutral task (assessing the effects of methylphenidate by itself) and methylphenidate/math task (assessing the combined effects of methylphenidate and the mathematical task).

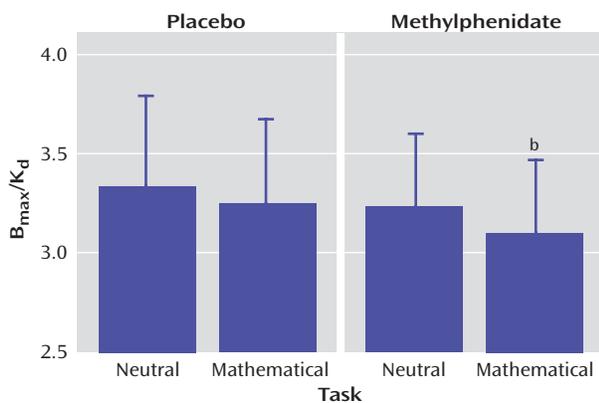
There were no differences across conditions in K_1 in the striatum or cerebellum, indicating that there were no significant condition-related differences in radiotracer delivery (Table 2). There were no differences in the distribution volume in cerebellum. The repeated-measures ANOVA showed significant differences across the conditions for distribution volume in the striatum ($F=8.2$, $df=3$, 63 , $p<0.001$). Post hoc *t* tests showed that when compared with the baseline condition, distribution volume measures in the striatum were significantly lower for the methylphenidate/math task condition ($p<0.0001$) but did not differ for the placebo/math task and the methylphenidate/neutral task conditions (Table 2). Similarly, the repeated-measures ANOVA showed significant differences across the condi-

TABLE 2. Measures of [¹¹C]Raclopride in the Striatum and Cerebellum During PET Scans of 16 Healthy Subjects Performing a Math Task With a Monetary Reward or a Noncompensated Neutral Task After Receiving Placebo or Methylphenidate

Scanning Condition ^a	Transfer Constant From Plasma to Brain (K ₁)				Distribution Volume			
	Striatum		Cerebellum		Striatum		Cerebellum	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
Placebo/neutral task	0.09	0.01	0.10	0.01	0.40	0.07	1.74	0.22
Placebo/math task	0.09	0.01	0.10	0.01	0.40	0.05	1.67	0.22
Methylphenidate/neutral task	0.09	0.01	0.10	0.01	0.40	0.05	1.67	0.18
Methylphenidate/math task	0.09	0.01	0.10	0.01	0.38	0.05	1.54 ^b	0.19

^a Following oral administration of methylphenidate (20 mg) or placebo, subjects underwent positron emission tomography scans while solving mathematical problems with monetary reinforcement (math task) or passively viewing pictures of scenery with no remuneration (neutral task).

^b Significantly different from the placebo/neutral task condition (which was considered the baseline), per post hoc paired t test ($p < 0.0001$).

FIGURE 2. Methylphenidate-Induced Changes in Dopamine Transporter Binding Potential (B_{max}/K_d) in the Striatum of 16 Healthy Subjects Performing a Math Task With a Monetary Reward or a Noncompensated Neutral Task^a

^a Following oral administration of methylphenidate (20 mg) or placebo, subjects underwent positron emission tomography scans while solving mathematical problems with monetary reinforcement (math task) or passively viewing pictures of scenery with no remuneration (neutral task). Bars represent standard deviations of the mean values.

^b Significantly different from the placebo/neutral task condition (which was considered the baseline), per post hoc paired t test ($p \leq 0.05$).

tions for the striatal B_{max}/K_d measures ($F=2.75$, $df=3, 63$, $p < 0.05$). Post hoc t tests showed that when compared with the baseline condition they were significantly lower for the methylphenidate/math task condition but did not differ for the placebo/math task and the methylphenidate/neutral task conditions (Figure 2). This indicates that the only condition for which the dopamine changes were large enough to be detected by the [¹¹C]raclopride method was when methylphenidate was administered with the mathematical task.

Analysis of the behavioral measures showed a significant effect of methylphenidate in the perception of the mathematical task by the subjects. Methylphenidate significantly increased the ratings of the task as being interesting, exciting, motivating, and less tiresome (Table 3).

The correlation analyses showed a significant association between the changes in extracellular dopamine between the methylphenidate/math task and the placebo/math task conditions and the changes in the rating of inter-

esting ($r=0.67$, $df=14$, $p=0.006$), exciting ($r=0.52$, $df=14$, $p < 0.05$), and motivating ($r=0.67$, $df=14$, $p=0.006$) (Figure 3).

The changes in B_{max}/K_d during the methylphenidate/math task condition were not correlated with the plasma methylphenidate concentration (60 minutes: $r=0.26$, $df=14$, $p=0.34$; 90 minutes: $r=0.33$, $df=14$, $p=0.21$), indicating that differences in methylphenidate metabolism were not responsible for the variability in the dopamine increases. Correlations between the dopamine changes during the methylphenidate/math task conditions and age revealed a significant negative correlation: the younger the subjects, the larger the increases in dopamine ($r=-0.68$, $df=15$, $p=0.004$).

Discussion

Here we have shown that methylphenidate-induced increases in dopamine are context dependent. Methylphenidate increased extracellular dopamine in the striatum when administered before subjects were required to perform a remunerated mathematical task but not when administered before subjects passively viewed scenery cards. Since dopamine cell activity is sensitive to environmental stimulation (salient and novel stimuli activate dopamine cells [17]), this suggests a mechanism to explain why methylphenidate-induced increase in dopamine depends on the conditions of its administration.

The context dependency of stimulant-induced dopamine increases has been demonstrated in laboratory animals for cocaine, which like methylphenidate increases dopamine by blocking the dopamine transporters (18). For example, cocaine-induced increases in dopamine in the nucleus accumbens were larger when animals were given cocaine in an environment where they had previously received cocaine than when they received it in a novel environment (19) or when animals self-administered cocaine than when cocaine administration was involuntary (20). Also methylphenidate-induced increases in dopamine in the prefrontal cortex were significantly greater when rats were restrained at the time of administration than when they were not (21).

In clinical studies, the context dependency of stimulant medication was one of the first effects noted in early studies of "hyperactivity" (22) and has been replicated in sub-

TABLE 3. Descriptor Ratings for a Math Task With a Monetary Reward Performed by 16 Healthy Subjects Following Administration of Placebo and Methylphenidate^a

Descriptor	Rating by Subjects				Analysis	
	Placebo		Methylphenidate			
	Mean	SD	Mean	SD	t (df=15)	p
Interesting	6.9	1.9	8.9	1.6	3.6	0.003
Exciting	6.9	2.1	8.6	1.8	3.8	0.004
Motivating	7.5	1.8	8.6	1.8	2.1	0.05
Tiresome	3.2	2.1	2.3	2.0	2.3	0.05
Boring	2.8	2.2	1.9	2.4	1.2	0.29

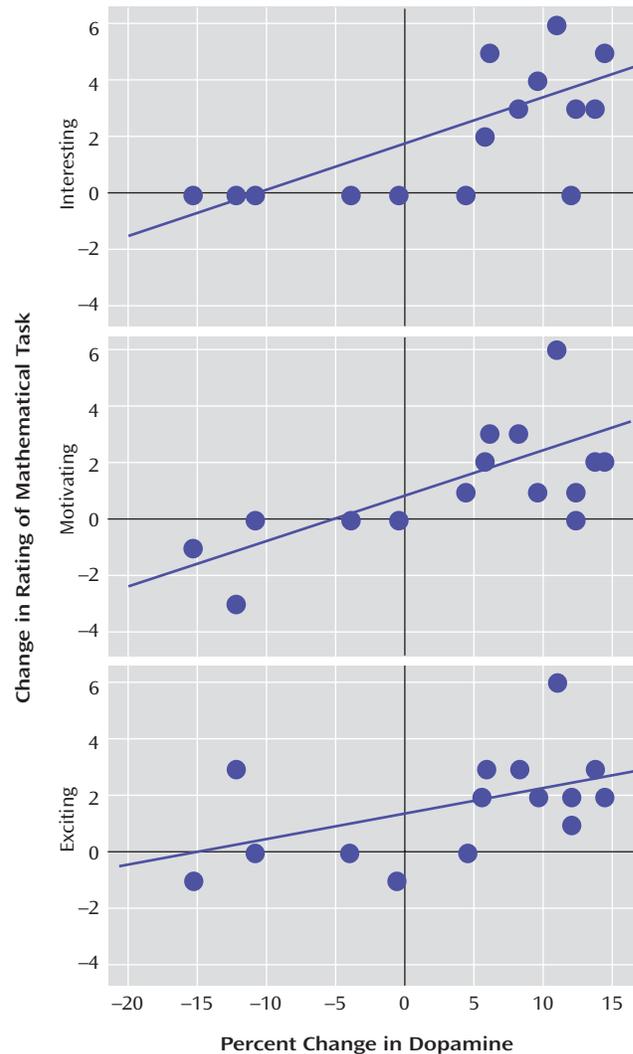
^a Following oral administration of methylphenidate (20 mg) or placebo, subjects underwent positron emission tomography scans while solving mathematical problems with monetary reinforcement. For each descriptor, subjects gave a rating that ranged from 1 to 10, with 1 being low and 10 being high.

sequent studies in children with ADHD (23). Recently, context dependency was documented in a laboratory school setting in which children with ADHD treated with methylphenidate showed larger reductions in placebo-adjusted activity level in the classroom than in the playground setting (24). Similarly, in a previous PET study we showed that methylphenidate-induced increases in extracellular dopamine were larger when it was administered with a salient stimulus (visual display of food in food-deprived subjects) than with a neutral stimulus (recall of family genealogy) (25).

In this study, the pairing of 20 mg of methylphenidate with the neutral task did not increase extracellular dopamine. This differs from a study in which we showed dopamine increases with 60 mg of methylphenidate when given with a neutral stimulus and is likely to reflect the much larger dose used for that study. However, the current findings also differ from a previous study in which we showed dopamine increases with 20 mg of methylphenidate when given with a neutral stimulus in food-deprived subjects (25). This discrepancy is likely to reflect food deprivation, which has been shown to affect extracellular dopamine (26). Further studies are required to assess if methylphenidate's effects in the human brain are sensitive to acute food deprivation in ways other than that of the effects of food on drug absorption.

Methylphenidate significantly increased the descriptor ratings of interesting, exciting, and motivating for the mathematical task and showed a tendency to decrease the rating for the task descriptor tiresome. The increases in these descriptors were associated with methylphenidate-induced dopamine increases when given with the mathematical task. This suggests that the amplification of small dopamine increases due to the task by methylphenidate may be the mechanism underlying this drug's ability to make tasks more salient to the subject. Although most of methylphenidate's therapeutic effects have emphasized its effects on attention, here we document an effect of methylphenidate on motivation as reflected by the subjective report of how the subject experiences the task in terms of descriptors of interesting, exciting, and motivating, which

FIGURE 3. Relationship Between the Changes in Extracellular Brain Dopamine During a Math Task With a Monetary Reward and Changes in Descriptor Ratings of the Math Task Following Administration of Methylphenidate in 16 Healthy Subjects^a



^a Changes were determined by comparing values after oral methylphenidate (20 mg) with values after placebo.

we consider to be characteristics of salience in this setting. We postulate that methylphenidate's ability to increase the saliency of the task will increase the motivation it elicits and that this will result in improved performance. Indeed, others have pointed to the potential relevance that methylphenidate's effects on saliency may have on its therapeutic effects in ADHD (27). Also, methylphenidate when used for other neuropsychiatric conditions has been shown to be effective in the treatment of apathy, which is an extreme state of lack of motivation (28).

The levels of extracellular dopamine in the striatum are set by two processes: tonic dopamine cell firing (maintains baseline steady-state levels and sets the overall responsiveness of the dopamine system) and phasic dopamine cell firing (leads to fast dopamine changes that highlight the

saliency of stimuli) (29). It had been previously hypothesized that therapeutic effects of methylphenidate are due to increased tonic dopamine levels, which stimulate dopamine autoreceptors and attenuate phasic dopamine increases (30). However, we now postulate that the long-lasting blockade of dopamine transporters by methylphenidate actually amplifies stimuli-induced dopamine increases (their magnitude and duration). Both the current results and those from a prior PET study in which we showed that methylphenidate amplified dopamine increases induced by display of food in food-deprived subjects (25) support the notion of an amplification of dopamine increases by oral methylphenidate rather than attenuation. Thus we postulate that methylphenidate will operate to amplify the saliency value of stimuli to which the subject may be exposed during everyday routines and which by themselves may have been insufficient to elicit dopamine responses strong enough and lasting long enough to signal saliency and drive and maintain interest and attention for the period required to complete the relevant task.

In this study the mathematical task, even with monetary remuneration, was insufficient by itself to elicit a measurable response. This most likely reflects the limited sensitivity of the methodology. Although some studies have shown decreases in [¹¹C]raclopride binding in striatum with behavioral interventions (31, 32), others have failed to show an effect after exercise (33) or after food stimulation (25). These differences most likely reflect the magnitude and duration of the dopamine changes elicited by the various behavioral interventions.

In this study methylphenidate, when given with the neutral task, was insufficient to elicit a measurable response. This most likely reflects in addition to the limited sensitivity of the methodology the relatively small dose used (20 mg for a group whose mean weight was 70 kg) and the lack of saliency of the neutral task. Note that we have reported increases with dopamine after 60 mg of methylphenidate when given in a neutral condition (5) but have also reported increases with 20 mg when given with salient stimuli other than money (25). These results offer a new hypothesis of why combining pharmacological and psychosocial interventions is more effective than medication alone. The traditional explanation is that a given dose of medication produces a specific effect by blocking dopamine transporters and increasing dopamine, which operates to focus attention and that this in turn enhances the impact of behavioral techniques (such as tokens earned for attending to an academic task in the classroom). However, if token reinforcers in behavioral interventions operate like money in the present study, then the behavioral intervention may operate to magnify the stimulant-induced increase in dopamine for a given dose of medication. This may explain why in the Multimodal Treatment Study of Children With ADHD (34), the combination of behavior modification with medication produced the same or

slightly superior reduction of symptoms at a lower dose of methylphenidate (35).

Methylphenidate-induced increases in dopamine when co-administered with the mathematical task declined as a function of age. An age-related blunting in stimulant-induced dopamine increases has also been observed after intravenous (36) and oral methylphenidate (5) and after intravenous amphetamine in healthy subjects (37). This could reflect a decrease in dopamine release with aging. Indeed, microdialysis studies have shown that stimulant-induced increases in extracellular dopamine are significantly lower in older than in younger animals (38, 39).

The following are study limitations. First, although changes in [¹¹C]raclopride binding are linearly related to extracellular dopamine, the precise relationship with synaptic dopamine is not understood (10). Second, to avoid the need for four arterial cannulations, we conducted the studies over 2 rather than 4 days. This required us to always give methylphenidate during the second study of the day. However, the order effect cannot explain the differences observed with methylphenidate when given with the neutral versus the mathematical task, since for both conditions methylphenidate was given at the same time of day. Third, the experimental design did not allow us to evaluate the effects of methylphenidate on performance, since the difficulty of the problems was individually adjusted so that each subject would achieve 80% correct performance. Last, because the mathematical test conditions were always paired with monetary reward, we cannot exclude the possibility that methylphenidate enhanced the saliency of the monetary reward rather than of the mathematical test itself. A study that assesses the effects of methylphenidate on a mathematical task that is not remunerated would enable research to address this issue. This is relevant because most of the behavioral strategies in ADHD use interventions that reinforce good behaviors (such as token reinforcers that can be exchanged for computer games, individual time with staff, special outings, and time in the playground, which in a way are similar to the use of money in this study).

In summary, these results provide evidence that methylphenidate's effects on dopamine are sensitive to the conditions of its administration. Dopamine increases were larger when methylphenidate was given with a task that required cognitive performance that was remunerated than when it was given with a task that did not require performance and was not remunerated. They also show that methylphenidate enhanced the motivational saliency of the cognitive task, as evidenced by the perception of the task as more interesting, exciting, and motivating. The significant association between methylphenidate-induced dopamine increases and the perception of the task as interesting and motivating lead us to postulate that methylphenidate's therapeutic effects may be secondary to its ability to enhance stimuli-induced dopamine increases, thus making them more motivationally salient and

thereby improving performance. This brings to light the possibility that methylphenidate's improvement of performance may be secondary to its motivational effects, which would explain why stimulants improve performance of a boring task in normal healthy individuals and why unmedicated ADHD children perform properly when the task is salient (40). These findings support educational strategies that make schoolwork more interesting as non-pharmacological interventions to treat ADHD.

Received Oct. 1, 2003; revision received Dec. 2, 2003; accepted Dec. 4, 2003. From the Medical and Chemistry Departments, Brookhaven National Laboratory, Upton, N.Y.; the Department of Psychiatry and the Department of Applied Mathematics, State University of New York at Stony Brook, Stony Brook, N.Y.; and the University of California at Irvine Child Development Center, Irvine, Calif. Address reprint requests to Dr. Volkow, National Institute on Drug Abuse, 6001 Executive Blvd., Rockville, MD 20853; nvolkow@nida.nih.gov (e-mail).

This research was carried out at Brookhaven National Laboratory under support by the U.S. Department of Energy (Office of Biological and Environmental Research) (DE-AC02-76 CH00016) and by NIDA grants DA-09490-01 and DA-06278.

The authors thank D. Schlyer and M. Schueller for Cyclotron operations; D. Warner for PET operations; R. Ferrieri, V. Garza, P. King, C. Shea, and Y. Xu for radiotracer preparation and analysis; N. Netusil, M. Jayne, and P. Carter for patient care; and T. Cooper for plasma methylphenidate analyses.

References

- Greenhill LL, Pliszka S, Dulcan MK, Bernet W, Arnold V, Beitchman J, Benson RS, Bukstein O, Kinlan J, McClellan J, Rue D, Shaw JA, Stock S (American Academy of Child and Adolescent Psychiatry): Practice parameter for the use of stimulant medications in the treatment of children, adolescents, and adults. *J Am Acad Child Adolesc Psychiatry* 2002; 41(2 suppl):265–495
- Volkow ND, Wang G-J, Fowler JS, Gatley SJ, Logan J, Ding Y-S, Hitzemann R, Pappas N: Dopamine transporter occupancies in the human brain induced by therapeutic doses of oral methylphenidate. *Am J Psychiatry* 1998; 155:1325–1331
- Kiyatkin EA, Rebec GV: Dopaminergic modulation of glutamate-induced excitations of neurons in the neostriatum and nucleus accumbens of awake, unrestrained rats. *J Neurophysiol* 1996; 75:142–153
- Berridge KC, Robinson TE: What is the role of dopamine in reward: hedonic impact, reward learning, or incentive salience? *Brain Res Rev* 1997; 28:309–369
- Volkow ND, Wang G, Fowler JS, Logan J, Gerasimov M, Maynard L, Ding Y, Gatley SJ, Gifford A, Franceschi D: Therapeutic doses of oral methylphenidate significantly increase extracellular dopamine in the human brain. *J Neurosci* 2001; 21:RC121
- Volkow ND, Wang GJ, Fowler JS, Logan J, Franceschi D, Maynard L, Ding YS, Gatley SJ, Gifford W, Swanson JM: Relationship between blockade of dopamine transporters by oral methylphenidate and the increases in extracellular dopamine: therapeutic implications. *Synapse* 2002; 43:181–187
- Wightman RM, Robinson DL: Transient changes in mesolimbic dopamine and their association with "reward." *J Neurochem* 2002; 82:721–735
- Giros B, Jaber M, Jones SR, Wightman RM, Caron MG: Hyperlocomotion and indifference to cocaine and amphetamine in mice lacking the dopamine transporter. *Nature* 1996; 379:606–612
- Swanson J, Wigal S, Greenhill L, Browne R, Waslick B, Lerner M, Williams L, Flynn D, Agler D, Crowley KL, Fineberg E, Regino R, Baren M, Cantwell D: Objective and subjective measures of the pharmacodynamic effects of Adderall in the treatment of children with ADHD in a controlled laboratory classroom setting. *Psychopharmacol Bull* 1998; 34:55–60
- Breier A, Su TP, Saunders R, Carson RE, Kolachana BS, de Bartolomeis A, Weinberger DR, Weisenfeld N, Malhotra AK, Eckelman WC, Pickar D: Schizophrenia is associated with elevated amphetamine-induced synaptic dopamine concentrations: evidence from a novel positron emission tomography method. *Proc Natl Acad Sci USA* 1997; 94:2569–2574
- Volkow ND, Fowler JS, Wang GJ, Dewey SL, Schlyer D, MacGregor R, Logan J, Alexoff D, Shea C, Hitzemann R, Wolf AP: Reproducibility of repeated measures of ^{11}C raclopride binding in the human brain. *J Nucl Med* 1993; 34:609–613
- Dewey SL, Smith GS, Logan J, Brodie JD, Fowler JS, Wolf AP: Striatal binding of the PET ligand ^{11}C -raclopride is altered by drugs that modify synaptic dopamine levels. *Synapse* 1993; 13:350–356
- Hall H, Kohler C, Gawell L, Farde L, Sedvall G: Raclopride, a new selective ligand for the dopamine-D2 receptors. *Prog Neuropsychopharmacol Biol Psychiatry* 1988; 12:559–568
- Srinivas NR, Hubbard JW, Quinn D, Korchinski ED, Midha K: Extensive and enantioselective presynaptic metabolism of dl-threo-methylphenidate in humans. *Prog Neuropsychopharmacol Biol Psychiatry* 1991; 15:213–220
- Logan J, Fowler JS, Volkow ND, Wolf AP, Dewey SL, Schlyer DJ, MacGregor RR, Hitzemann R, Bendriem B, Gatley SJ, Wolf AP: Graphical analysis of reversible radioligand binding from time-activity measurements applied to [^{11}C -methyl]-(-)-cocaine PET studies in human subjects. *J Cereb Blood Flow Metab* 1990; 10:740–747
- Logan J, Volkow ND, Fowler JS, Wang GJ, Dewey SL, MacGregor R, Schlyer D, Gatley SJ, Pappas N, King P, Vitkun S, Wolf AP: Effects of blood flow on [^{11}C]raclopride binding in the brain: model simulations and kinetic analysis of PET data. *J Cereb Blood Flow Metab* 1994; 14:995–1010
- Hollerman JR, Schultz W: Dopamine neurons report an error in the temporal prediction of reward during learning. *Nat Neurosci* 1998; 1:304–309
- Ritz MC, Lamb RJ, Goldberg SR, Kuhar MJ: Cocaine receptors on dopamine transporters are related to self-administration of cocaine. *Science* 1987; 237:1219–1223
- Duvauchelle CL, Ikegami A, Asami S, Robens J, Kressin K, Castaneda E: Effects of cocaine context on NAcc dopamine and behavioral activity after repeated intravenous cocaine administration. *Brain Res* 2000; 862:49–58
- Hemby SE, Co C, Koves TR, Smith JE, Dworkin SI: Differences in extracellular dopamine concentrations in the nucleus accumbens during response-dependent and response-independent cocaine administration in the rat. *Psychopharmacology (Berl)* 1997; 133:7–16
- Marsteller DA, Gerasimov MR, Schiffer WK, Geiger JM, Barnett CR, Borg JS, Scott S, Ceccarelli J, Volkow ND, Molina PE, Alexoff DL, Dewey SL: Acute handling stress modulates methylphenidate-induced catecholamine overflow in the medial prefrontal cortex. *Neuropsychopharmacology* 2002; 27:163–170
- Conners KC: The effect of dexedrine on rapid discrimination and motor control of hyperkinetic children under mild stress. *J Nerv Ment Dis* 1966; 142:429–433
- Porrino LJ, Rapoport JL, Behar D, Ismond DR, Bunney WE Jr: A naturalistic assessment of the motor activity of hyperactive boys, II: stimulant drug effects. *Arch Gen Psychiatry* 1983; 40:688–693
- Swanson JM, Gupta S, Williams L, Agler D, Lerner M, Wigal S: Efficacy of a new pattern of delivery of methylphenidate for the treatment of ADHD: effects on activity level in the classroom

- and on the playground. *J Am Acad Child Adolesc Psychiatry* 2002; 41:1306–1314
25. Volkow ND, Wang G-J, Fowler JS, Logan J, Jayne B, Franceschi D, Wong C, Gattley SJ, Gifford A, Ding Y-S, Pappa N: "Non-hedonic" food motivation in humans involves dopamine in the dorsal striatum. *Synapse* 2002; 44:175–180
 26. Acworth IN, Ressler K, Wurtman RJ: Feeding-associated alterations in striatal neurotransmitter release. *Ann NY Acad Sci* 1989; 575:596–598
 27. Wilkison PC, Kircher JC, McMahon WM, Sloane HN: Effects of methylphenidate on reward strength in boys with attention-deficit hyperactivity disorder. *J Am Acad Child Adolesc Psychiatry* 1995; 34:897–901
 28. Marin RS, Fogel BS, Hawkins J, Duffy J, Krupp B: Apathy: a treatable syndrome. *J Neuropsychiatry Clin Neurosci* 1995; 7:23–30
 29. Grace AA: The tonic/phasic model of dopamine system regulation and its implications for understanding alcohol and psychostimulant craving. *Addiction* 2000; 95(suppl 2):S119–S128
 30. Seeman P, Madras BK: Anti-hyperactivity medication: methylphenidate and amphetamine. *Mol Psychiatry* 1998; 3:386–396
 31. Koeppe MJ, Gunn RN, Lawrence AD, Cunningham VJ, Dagher A, Jones T, Brooks DJ, Bench CJ, Grasby PM: Evidence for striatal dopamine release during a video game. *Nature* 1998; 393:266–268
 32. Pappata S, Dehaene S, Poline JB, Gregoire MC, Jobert A, Delforge J, Frouin V, Bottlaender M, Dolle F, Di Giamberardino L, Syrota A: In vivo detection of striatal dopamine release during reward: a PET study with [¹¹C]raclopride and a single dynamic scan approach. *Neuroimage* 2002; 16:1015–1027
 33. Wang GJ, Volkow ND, Fowler JS, Franceschi D, Logan J, Pappas NR, Wong CT, Netusil N: PET studies of the effects of aerobic exercise on human striatal dopamine release. *J Nucl Med* 2000; 41:1352–1356
 34. Multimodal Treatment Study of Children With ADHD (MTA) Cooperative Group: A 14-month randomized clinical trial of treatment strategies for attention deficit/hyperactivity disorder. *Arch Gen Psychiatry* 1999; 56:1073–1086
 35. Swanson JM, Kraemer HC, Hinshaw SP, Arnold LE, Conners CK, Abikoff HB, Clevenger W, Davies M, Elliott, G, Greenhill LL, Hechtman L, Hoza B, Jensen PS, March JS, Newcorn JH, Owens L, Pelham WE, Schiller E, Severe J, Simpson S, Vitiello B, Wells CK, Wigal T, Wu M: Clinical relevance of the primary findings of the MTA: success rates based on severity of ADHD and ODD symptoms at the end of treatment. *J Am Acad Child Adolesc Psychiatry* 2001; 40:168–179
 36. Volkow ND, Wang GJ, Fowler JS, Logan J, Schlyer D, Hitzemann R, Lieberman J, Angrist B, Pappas N, MacGregor R, Wolf AP: Imaging endogenous dopamine competition with [¹¹C]raclopride in the human brain. *Synapse* 1994; 16:255–262
 37. Laruelle M, Abi-Dargham A, van Dyck CH, Rosenblatt W, Zea-Ponce Y, Zoghbi SS, Baldwin RM, Charney DS, Hoffer PB, Kung HF, Innis RB: SPECT imaging of striatal dopamine release after amphetamine challenge. *J Nucl Med* 1995; 36:1182–1190
 38. Gerhardt GA, Maloney RE Jr: Microdialysis studies of basal levels and stimulus-evoked overflow of dopamine and metabolites in the striatum of young and aged Fischer 344 rats. *Brain Res* 1999; 816:68–77
 39. Harada N, Nishiyama S, Satoh K, Fukumoto D, Kakiuchi T, Tsukada H: Age-related changes in the striatal dopaminergic system in the living brain: a multiparametric PET study in conscious monkeys. *Synapse* 2002; 45:38–45
 40. Grinspoon L, Hedblom P: *The Speed Culture: Amphetamine Use and Abuse in America*. Cambridge, Mass, Harvard University Press, 1975