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Genetically determined interaction between the dopamine transporter and the D2 receptor on prefronto-striatal activity and volume in humans

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Abstract

Dopamine modulation of neuronal activity during memory tasks identifies a non-linear inverted-U shaped function. Both the dopamine transporter (DAT) and dopamine D2 receptors (encoded by *DRD2*) critically regulate dopamine signaling in the striatum and in prefrontal cortex during memory. Moreover, *in vitro* studies have demonstrated that DAT and D2 proteins reciprocally regulate each other presynaptically. Therefore, we have evaluated the genetic interaction between a *DRD2* polymorphism (rs1076560) causing reduced presynaptic D2 receptor expression and the *DAT* 3'-VNTR variant (affecting DAT expression) in a large sample of healthy subjects undergoing BOLD - fMRI during memory tasks and structural MRI. Results indicated a significant *DRD2/DAT* interaction in prefrontal cortex and striatum BOLD activity during both working memory and encoding of recognition memory. The differential effect on BOLD activity of the *DAT* variant was mostly manifest in the context of the *DRD2* allele associated with lower presynaptic expression. Similar results were also evident for gray matter volume in caudate. These interactions describe a non-linear relationship between compound genotypes and brain activity or gray matter volume. Complementary data from striatal protein extracts from wild-type and D2 knock-out animals (*D2R* $-/-$) indicate that DAT and D2 proteins interact *in vivo*. Taken together, our results demonstrate that the interaction between genetic variants in *DRD2* and *DAT* critically modulates the non-linear relationship between dopamine and neuronal activity during memory processing.

Keywords

working memory; Recognition Memory; FMRI; Dopamine; Transport; D2; Receptor

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Introduction

The relationship between dopamine levels and neuronal activity in prefrontal cortex and in striatum (Alexander et al., 1986; Goldman-Rakic, 1996) describes a non-linear function (inverted-U) (Seamans and Yang, 2004). In other words, there is a critical range of dopamine stimulation within which neuronal activity is more focused, i.e. less neuronal activity for better behavioral performance (Mattay et al., 2003). Above or below this critical range of dopamine stimulation, neuronal activity becomes more diffuse and behavioral performance deteriorates (Mattay et al., 2003). Earlier studies have demonstrated that this relationship is strongly modulated by dopamine D1 receptors (Seamans and Yang, 2004). However, several additional mechanisms regulate dopamine levels, including D2 receptor signaling as well as reuptake by the dopamine transporter (DAT). Studies in animals and in humans strongly implicate D2 receptors and the DAT in regulating working memory performance and prefronto-striatal neuronal activity (Arnsten et al., 1995; Kimberg et al., 2001; Mozley et al., 2001; Glickstein et al., 2002; Cropley et al., 2006; Kellendonk et al., 2006; Mehta and Riedel, 2006; Chou et al., 2007). Recent studies focusing on the relationship between D2 and DAT have also shown that DAT activity is regulated by D2 receptors (Meiergerd et al., 1993; Parsons et al., 1993; Dickinson et al., 1999; Mortensen and Amara, 2003). Mice genetically engineered to lack the D2 receptor display reduced activity of DAT (Dickinson et al., 1999). D2 agonist and antagonist agents modify the kinetics of dopamine reuptake by DAT (Kimmel et al., 2001). Moreover, DAT is also regulated by dopamine D2 receptors through a direct protein-protein interaction which facilitates the recruitment of intracellular DAT to the cell surface, thereby enhancing dopamine clearance (Lee et al., 2007). Finally, this physical interaction appears to be more specific for presynaptic D2 receptors (Bolan et al., 2007), which is supported by results showing that mice lacking DAT have complete loss of function of presynaptic D2 receptors (Jones et al., 1999).

By a mechanism of alternative splicing, the D2 receptor gene (*DRD2*) encodes two molecularly distinct isoforms, D2S and D2L. D2L acts mainly at postsynaptic sites and D2S serves presynaptic autoreceptor functions (Khan et al., 1998; Usiello et al., 2000). The D2S isoform appears to be involved in regulation of prefronto-striatal synaptic plasticity associated with long-term potentiation, centrally implicated in the physiology of memory (Centonze et al., 2004). Consistently, we have demonstrated that a novel intronic *DRD2* polymorphism (rs1076560, G>T) of the D2 receptor affects both relative expression of D2S in prefrontal cortex/striatum and activity of the striato-thalamic-prefrontal pathway during working memory in healthy subjects (Zhang et al., 2007) and in patients with schizophrenia (Bertolino et al., 2008a).

A functional variable number of tandem repeat (VNTR) polymorphism in the 3' untranslated region of the DAT gene has been described (Vandenberg et al., 1992). Alleles of this polymorphism range from 3 to 11 repeats, with the 9- and 10-repeat alleles by far the most common (Vandenberg et al., 1992). As compared to the 9-repeat allele, the 10-repeat allele has been associated with increased gene expression both *in vitro* (Mill et al., 2002; VanNess et al., 2005) and *in vivo* (Heinz et al., 2000). Consistently, several studies have reported that the 10-repeat allele is associated with more focused cortical activity during memory and attention in healthy subjects as well as in patients with ADHD (Fossella et al., 2002; Cornish et al., 2005; Bertolino et al., 2006a; Johanson et al., 2006; Schott et al., 2006; Caldu et al., 2007).

Based on this prior evidence, we used fMRI in healthy subjects to explore the relationship of *DRD2* and *DAT* functional polymorphisms with brain activity during working memory. To confirm the working memory results in another cognitive setting known to involve the

prefronto-striatalthalamic-prefrontal circuit, we also evaluated the interaction of these two genotypes with brain activity during encoding of recognition memory. Our hypothesis was based on the physiological role of dopamine in regulating neuronal signal-to-noise ratios in this pathway, on earlier BOLD fMRI studies (Bertolino et al., 2006a; Zhang et al., 2007; Bertolino et al., 2008b; Bertolino et al., 2008a), on known trophic action of dopamine on neurons and synapses (Nieoullon, 2002), and on earlier *in vitro* experiments demonstrating direct interaction between D2 and DAT. Thus, we hypothesized that these two genes interact in modulating the non-linearity of cortical and subcortical activity in humans during memory performance and in modulating gray matter volume (assessed respectively with BOLD fMRI and Voxel-Based-Morphometry). Moreover, to further substantiate the possibility that the effects in humans might depend on a direct interaction between these two proteins, we analyzed this possibility by co-immunoprecipitation. Importantly, using mouse striata from wild-type (WT) and *DRD2* knock-out mice, we were able to show a physical interaction between DAT and D2 proteins *in vivo*.

Materials and Methods

Subjects

We studied one hundred forty two healthy subjects (57 males, mean age 27.4 ± 7.3). The present study was approved by the local IRB at the University of Bari. Moreover, after complete description of the study to the subjects, written informed consent was obtained. For additional demographics, see Table 1.

Genotype determination

SNP rs1076560 of *DRD2* and the 3' *VNTR DAT* genotypes were determined as in (Bertolino et al., 2006a; Zhang et al., 2007; Bertolino et al., 2008b).

3' VNTR DAT—Genotyping of the DAT1 40-bp repeat (VNTR) polymorphism in the 3' untranslated region (rs# 28363170) was determined using forward 5'-TGTGGTGTAGGGAACGGCCTGAG-3' and reverse 5'-CTTCCTGGAGGTCACGGCTCAAGG-3' primers. DNA amplification by polymerase chain reaction (PCR) of the 40-base pair repeat alleles was performed as described elsewhere (Szekeres et al., 2004). PCR products were separated by 4% agarose gel electrophoresis, visualized by ultraviolet transillumination and fragment sized by comparison with invitrogen 100bp DNA ladder. To confirm the results obtained with agarose gel electrophoresis, genomic DNA fragments were PCR-amplified using fluorescent labeled forward primer, resolved on an ABI Prism 3100 DNA sequencer (Applied Biosystems, Foster City, CA, USA) and analyzed with Genotyper software.

DRD2—SNP rs1076560 was analyzed with allele-specific PCR primers as described (Papp et al., 2003) or SNaPshot (Applied Biosciences (ABI), Foster City CA) (Zhang et al., 2005).

As in several earlier studies (Heinz et al., 2000; Jacobsen et al., 2000; Cheon et al., 2003; Durston et al., 2005; van Dyck et al., 2005; Gilbert et al., 2006; Schott et al., 2006; Laucht et al., 2007), since the number of subjects homozygous for the 9-repeat allele of the *DAT* was very small, we grouped all subjects carrying at least one 9-repeat allele in one group, 9-repeat carriers. After genotype determination, the groups were divided based on *DRD2*, *DAT*, and *DRD2-DAT* genotypes. The Ns were as follows: *DRD2* (GG=110, GT=32); *DAT* (9-repeat carriers =91, 10/10-repeat=51); *DRD2-DAT* (GT 9-repeat carriers=22, GG 9-repeat carriers=69, GT 10/10-repeat=10, GG 10/10-repeat carriers=41). No *DRD2 TT* subjects were observed in this sample. The allelic distribution of both genes was in Hardy Weinberg equilibrium (*DRD2* df 1, $\chi^2 > 2.1$, $p > 0.1$, *DAT* df 1, $\chi^2 = 1.2$, $p > 0.2$).

Functional imaging tasks and acquisition parameters

N-Back Working Memory paradigm—The genotype groups in this sample were matched for all demographic variables except gender ($p < 0.05$). During fMRI, all subjects completed a blocked paradigm of the N-Back task with a 2-Back working memory condition and a non-memory guided control condition 0-Back (Bertolino et al., 2004). This paradigm has been extensively used to evaluate activity of prefrontal cortex and striatum. “N-back” refers to how far back in the sequence of stimuli the subject had to recall. The stimuli consisted of numbers (1–4) shown in random sequence and displayed at the points of a diamond-shaped box. There was a visually paced motor task which also served as a non-memory guided control condition (0-Back) that simply required subjects to identify the stimulus currently seen. In the working memory condition, the task required recollection of a stimulus seen two stimuli (2-Back) previously while continuing to encode additionally incoming stimuli. As in earlier studies in which we used this paradigm, both numerical and spatial information together were used to guide subjects response to the task. Performance data were recorded as the number of correct responses (accuracy) and as reaction time.

Each subject was scanned using the same MR scanner with a gradient-echo echo planar imaging (EPI) sequence using the following parameters: 20 contiguous slices echo time=30 msec, repetition time= 2000 msec; field of view 24 cm; matrix 64×64 (Bertolino et al., 2004a; Bertolino et al., 2004b). We used a simple block design in which each block consisted of eight alternating 0-Back and 2-Back conditions (each lasting 30 seconds), obtained in 4 min and 8 sec, 120 whole-brain fMRI volumes. The first four scans at the beginning of each time series were acquired to allow the signal to reach a steady state and were not included in the final analysis.

Recognition memory paradigm—102 subjects of the original sample performed this additional paradigm. The genotype groups in this sample were matched for all demographic variables, including sex (all $p > 0.1$). The fMRI paradigm consisted of the encoding (“indoor” or “outdoor”) and subsequent retrieval (“new” or “old”) of novel, complex scenes, a task that has consistently been shown to produce activation of the prefrontal cortex and of the striatum in human neuroimaging experiments (Hariri et al., 2003; Bertolino et al., 2006c). Stimuli of neutral valence were presented in a blocked paradigm which provides robust power and sensitivity for BOLD signal change in the hippocampal region. Four encoding blocks were followed by four retrieval blocks in an interleaved design with a passive rest condition, resulting in a total of 16 blocks. Each block was 20 sec. long, producing a total scan time of 5.33 minutes. During encoding blocks, subjects viewed six images, presented serially for 3 sec each, and determined whether each image represented an “indoor” or “outdoor” scene (Hariri et al., 2003). An equal number of “indoor” and “outdoor” scenes were presented in each encoding block. All scenes were of neutral emotional valence and were derived from the International Affective Picture System (Lang et al., 1997). During subsequent retrieval blocks, subjects again viewed six images, presented serially for 3 sec each, and determined whether each scene was “new” or “old.” In each retrieval block, half the scenes were “old” (i.e., presented during the encoding blocks) and half were “new” (i.e., not presented during the encoding blocks). The order of “indoor” and “outdoor” scenes as well as “new” and “old” scenes were randomly distributed throughout the encoding and retrieval blocks, respectively. During the interleaved rest blocks, subjects were instructed to fixate on a centrally presented cross-hair. Before the beginning of each block, subjects viewed a brief (2 sec) instruction: “Indoor or Outdoor?”, “New or Old?”, or “Rest”. However, because of the blocked paradigm, the retrieval phase is actually a mixture between encoding and retrieval: subjects in this phase view an equal number of new and old stimuli. The new stimuli are likely to engage encoding mechanisms in the hippocampus, and this activity would be mixed in with any retrieval related activity. Therefore, because of this limitation, we decided not to analyze the retrieval data (for review, see (Schacter

and Wagner, 1999; Squire et al., 2004)). During scanning, all subjects responded by button presses with their right hand, allowing for determination of behavioral accuracy and reaction time.

Each subject was scanned using a GE Signa 3T scanner (General Electric, Milwaukee, WI). BOLD functional images were acquired with a gradient-echo echo planar imaging (EPI) sequence and covered 24 axial slices (4 mm thick, 1mm gap) that began at the cerebral vertex and encompassed the entire cerebrum and the majority of the cerebellum (repetition time/echo time, 2000/28 msec; field of view, 24 cm; matrix, 64 × 64).

BOLD-fMRI image analysis

Preprocessing and statistical analyses of the fMRI data: Analysis of the fMRI data was completed using statistical parametric mapping (SPM5; <http://www.fil.ion.ucl.ac.uk/spm>). Images for each subject were realigned to the first volume in the time series to correct for head motion (< 2.5 mm of translation, <1.5° rotation), spatially normalized into a standard stereotactic space (Montreal Neurological Institute, MNI, template) using a 12 parameter affine model and smoothed to minimize noise and residual differences in gyral anatomy with a Gaussian filter, set at 10 mm full-width at half-maximum. Voxel-wise signal intensities were ratio normalized to the whole-brain global mean. For each experimental condition, a box car model convolved with the hemodynamic response function (HRF) at each voxel was modeled. Predetermined condition effects at each voxel were calculated using a t-statistic, producing a statistical image for the contrasts of 2-Back versus 0-Back (N-Back) and of encoding versus rest (recognition memory). All these individual contrast images were then used in second-level random effects models to determine task-specific regional responses at the group level for the entire sample.

To detect the association of *DRD2* and *DAT* genotypes and fMRI activation in the working memory and recognition memory cortical and subcortical networks, the contrast images of all subjects were included in whole-brain image analyses for all predetermined condition effects using second-level random effects models for ANCOVA analyses within SPM5 covarying for gender. For recognition memory data, ANOVA was used instead because the genotype groups were matched for all demographic variables. Because of earlier fMRI studies of working memory and recognition memory (Bertolino et al., 2006a; Bertolino et al., 2006c; Zhang et al., 2007; Bertolino et al., 2008b; Bertolino et al., 2008a) demonstrating main effects of these two genes, a statistical threshold of $p < 0.005$, $k=3$, uncorrected, was used to evaluate main effects of *DRD2* and *DAT*. On the other hand, the same statistical threshold with a further False Discovery Rate (FDR) small volume correction for multiple comparisons (Meyer-Lindenberg et al., 2008) (using a 10mm radius sphere centered around the coordinates in caudate, putamen, and prefrontal cortex published in previous studies $p=0.05$ (Hofer et al., 2007; Koch et al., 2007; Tan et al., 2007)), was used to identify significant responses for the interaction between the two genes in these anatomical regions. These coordinates included: $x -36, y 9, z 59$; $x -36, y 30, z 23$; $x 20, y -3, z 17$; $x -18, y 3, z 15$; $x 8, y 12, z 8$ (Tan et al., 2007); $x -18, y 3, z 15$ (Hofer et al., 2007); $x 42, y 12, z 36$ (Koch et al., 2007). Because we did not have *a priori* hypotheses regarding the activity of brain regions outside of the prefrontal cortex and striatum, we used a statistical threshold of $p=0.05$, corrected for multiple comparisons across all voxels, for these whole-brain comparisons. All fMRI results are reported in Montreal Neurological Institutes (MNI) coordinates system.

Voxel-based morphometry acquisition and processing—Eighty six subjects of the original sample were included in these analyses. Again, genotype groups in this sample were matched for all demographic variables except gender ($p < 0.05$). 3D structural MRI images were obtained on a 3T GE MR scanner using a T1-weighted SPGR sequence (TR/TE/NEX=25/3/1;

flip angle, 6°; matrix size, 256 × 256; field of view, 25 × 25 cm) with 124 sagittal slices (slice thickness=1.3 mm, in-plane resolution of 0.94 × 0.94). VBM analysis was performed using a unified segmentation protocol (Ashburner and Friston, 2005) implemented in SPM5 (<http://www.fil.ion.ucl.ac.uk/spm>). Briefly, this protocol involves a number of fully automated preprocessing steps, including: extraction of brain, spatial normalization into stereotactic MNI space, segmentation into gray and white matter and CSF compartments, correction for volume changes induced by spatial normalization (modulation), and smoothing with a 8 mm full width at half maximum (FWHM) isotropic gaussian kernel.

Global effects of *DRD2* and *DAT* genotypes on gray matter volume were evaluated using an ANCOVA in SPM5. We controlled for potential confounds in our statistical model including linear and quadratic expansions of age (Buchel et al., 1996), gender, and total brain volume. Because of our strong *a priori* hypothesis, we used a region of interest (ROI) approach using the “Human aal atlas” within the Wake Forest University PickAtlas 1.04 (<http://www.fmri.wfubmc.edu/cms/>). Statistical non-stationary inference (Hayasaka et al., 2004) was performed at the cluster level at $p < 0.05$ corrected within the caudate ROI by using the ns toolbox implemented in SPM5 to avoid increased false positive rate due to non-stationary nature of structural images (<http://fmri.wfubmc.edu/cms/NS-General>). VBM data are reported with reference to the MNI standard space within SPM5.

Membrane preparations, Western blotting and immunoprecipitation analyses—

Mouse striatal extracts from D2R^{-/-} (Baik et al., 1995) and wild-type (WT) littermates were rapidly dissected and proteins were extracted as previously described (Tirota et al., 2008), with minor modifications. Briefly, striata were homogenized on ice with a polyethylene pestle, in a Membrane Extraction Buffer, (MEB 50 mM TRIS pH 7.5, 150 mM NaCl, 5mM EDTA pH 8.0), with protease and phosphatase inhibitors cocktail (Sigma). Samples were sedimented by centrifugation (40 minutes, 70000 × g, 4°C), and the pellets were resuspended in MEB supplemented with 10 mM 3-[(3-cholamidopropyl) dimethylammonio]-1-propanesulfonate (CHAPS) and gently rotated at 4°C for 40 minutes. The solubilized fractions were finally centrifuged, to remove the insoluble material (10 minutes, 18000×g at 4°C). Supernatants represent solubilized membrane extracts.

For Immunoprecipitation (IP) assays, 500 µg of membrane extracts were incubated for 4 h with 20 µl protein-A/G Sepharose (GE Lifesciences) at 4° C, in 500 µl of PBS supplemented by protease and phosphatase inhibitors cocktail and centrifuged at 3,000 × g for 5 min. The supernatants were incubated overnight at 4°C with 5 µg of the appropriate antibody (anti-DAT # sc-14002, from Santa Cruz Biotechnology; or mouse monoclonal anti-D2R raised against a peptide corresponding to the mouse D2R sequence from aa 309 to 322 (Doi et al. 2006). Sepharose beads were washed four times by centrifugation in MEB buffer, then resuspended in Laemmli sample buffer and resolved on SDS 12% PAGE. As negative control, immunoprecipitations were performed using either normal mouse or rabbit IgG. For western blots, 50 µg of membrane extracts were loaded per lane and separated on 12% SDS-PAGE. Proteins were transferred to polyvinylidene fluoride (PVDF) membrane (Millipore). Non-specific binding was blocked in 5% non-fat dry milk in 0.05% Tween 20 in PBS for 1h, at room temperature. Immuno-complexes were revealed by using appropriate peroxidase-conjugated secondary antibodies (Jackson Immuno-Research) along with a chemiluminescent reagent (SuperSignal West-Pico, Pierce).

RESULTS

Demographics and Behavioral Memory Performance

ANOVAs and χ^2 tests indicated that there were no significant differences between genotype groups in any demographic variable (all $p > 0.1$), with the exception of gender which was

unequally distributed for *DRD2* genotype and for *DRD2-DAT* compound genotypes (respectively, $\chi^2=6.5$, df 1, $p=0.01$, $\chi^2=10.1$, df 3, $p=0.01$).

Separate ANOVAs on behavioral working memory and encoding of recognition memory performance did not demonstrate any statistically significant main effect or interaction of the two polymorphisms on either accuracy or reaction time during working memory or encoding of recognition memory (all $F_{1,137}<2$, all $p>0.15$) thus allowing us to examine the effect of complex genotypes on brain activity independent of behavioral variation. Covarying these analyses for gender did not modify the results. For additional results, see also Table 1.

Neuroimaging results of the N-Back Working Memory Paradigm

Main effect of task—Consistent with prior reports (Callicott et al., 1999; Callicott et al., 2000; Bertolino et al., 2004; Bertolino et al., 2006a), a pattern of brain regions was activated during working memory. These anatomical regions included bilateral prefrontal cortex, bilateral parietal cortex, the anterior cingulate, the head of the caudate, the putamen and bilateral cerebellum in both groups.

Genotype main effects and interaction during working memory—ANOVA of the main effect of *DRD2* genotype revealed several clusters in which *GT* subjects had greater activity than *GG* subjects. These areas included the middle and inferior frontal gyri, the anterior cingulate and the right putamen (Table 2). The inverse contrast did not reveal any significant difference. ANOVA of the main effect of *DAT* genotype revealed several clusters in left middle and inferior frontal gyri in which *9-repeat* carriers had greater activity than *10/10-repeat* subjects (Table 2). The inverse contrast did not reveal any statistically significant difference. These main effects of *DRD2* and *DAT* are internally consistent as both *GT DRD2* genotype and the *9-repeat DAT* genotype are expected to lead to enhanced dopamine release.

We next addressed the question whether *DRD2* and *DAT* interact with each other. We expected that the genotypes potentially associated with putative intermediate levels of dopamine would show more focused activity for a similar level of behavioral performance (Williams and Castner, 2006). ANOVA revealed several locales with a statistically significant interaction between the two genotypes, including the caudate bilaterally and the left middle frontal gyrus (Table 2, Figure 1). Analysis of the BOLD signal change outside of SPM revealed an interaction in the right and left head of the caudate (Figure 1A-B; left caudate $F_{1,137}=9.1$, $p=0.002$; post hoc with Fisher LSD: *GT 9-carrier repeat*>*GT 10/10 repeat* $p=0.02$; *GG 10/10 repeat*>*GG 9-carrier repeat* $p=0.01$; right caudate $F_{1,137}=9.2$, $p=0.002$; post hoc: *GT 9-carrier repeat*>*GT 10/10 repeat* $p=0.03$; *GG 10/10 repeat*>*GG 9-carrier repeat* $p=0.009$) such that subjects with *9-repeat genotype* of the *DAT* have greater activity in the context of *DRD2 GT* genotype, whereas the opposite is true in the context of the *GG DRD2* genotype.

A polynomial regression also revealed a non-linear relationship between compound genotype and BOLD signal in right and left head of the caudate (respectively, $R^2=0.09$, $F=7.46$, $p=0.0008$; $R^2=0.09$, $F=6.99$, $p=0.001$, Figure 2A). Similarly, in the left middle frontal gyrus subjects with *9-carrier repeat* genotype had greater activity based on *GT DRD2* genotype (Figure 1C-D; $F_{1,137}=7.3$, $p=0.007$; post hoc: *GT 9-carrier repeat*>*GT 10/10 repeat* $p=0.005$; *GG 10/10 repeat*>*GG 9-carrier repeat* $p=0.5$). A polynomial regression also revealed a non-linear relationship between compound genotype and BOLD signal in left DLPFC ($R^2=0.054$, $F=4.0$, $p=0.01$, Figure 2B).

Additional results are reported in Table 2.

Neuroimaging results of the recognition memory paradigm

Main effect of task—Consistent with prior reports (Hariri et al., 2003; Bertolino et al., 2006c; Bertolino et al., 2008b), we found significant bilateral activation of the prefrontal cortex (including dorsolateral prefrontal and ventrolateral prefrontal cortex), and the striatum (including the head of the caudate and putamen).

Genotype main effects and interaction during recognition memory—ANOVA of the main effect of *DRD2* genotype revealed that *GT* subjects had statistically significant greater activity than *GG* in the left putamen and right inferior frontal gyrus (Table 3). The inverse contrast did not reveal any significant difference. ANOVA of the main effect of *DAT* genotype revealed that carriers of the *9-repeat* genotype had statistically significant greater activity than *10/10-repeat* subjects bilaterally in the caudate, and in right inferior and middle frontal gyri as well as in left middle frontal gyrus (Table 3). The inverse contrast did not reveal any statistically significant difference. ANOVA also revealed several locales with a statistically significant interaction between the two genotypes bilaterally in the caudate and in right middle frontal gyrus. Analysis of the BOLD signal change in the left head of the caudate revealed an interaction (Figure 3A-B, $F_{1,103}=6.1$, $p=0.01$; post hoc with Fisher LSD: *GT 9-carrier repeat* > *GT 10/10 repeat* $p=0.004$; *GG 10/10 repeat* vs. *GG 9-carrier repeat* $p=0.5$). Analysis of the BOLD signal change in middle frontal gyrus revealed another interaction (Figure 3C-D, $F_{1,103}=4.3$, $p=0.03$; post hoc: *GT 9-carrier repeat* > *GT 10/10 repeat* $p=0.03$; *GG 10/10 repeat* vs *GG 9-carrier repeat* $p=0.7$).

A polynomial regression also revealed a non-linear relationship between compound genotype and BOLD signal in the head of the caudate and in middle frontal gyrus (respectively, $R^2=0.072$, $F=4.05$, $p=0.02$, $R^2=0.048$, $F=2.6$, $p=0.07$, Figure 4A and B).

Voxel-based morphometry—ANOVA demonstrated a significant effect of *DRD2* genotype on caudate gray matter volume. *GG* subjects had greater gray matter volume in right caudate compared with *GT* subjects ($x\ 11$, $y\ 1$, $z\ 17$, $k=377$, $Z=2.84$, $p=0.04$). No statistically significant main effect of *DAT* genotype was found. However, ANOVA demonstrated a statistically significant interaction between *DRD2* and *DAT* genotypes in the left and right caudate (respectively, $x\ -18$, $y\ 5$, $z\ 19$, $k=567$, $Z=3.1$, $p=0.01$; $x\ 12$ $y\ 9$ $z\ 14$, $k=37$, $Z=2.63$, $p=0.06$; Figure 5A and B). Extraction of gray matter content from the left caudate cluster confirmed the interaction ($F_{1,82}=16.2$, $p=0.0001$; post hoc with Fisher LSD: *GT 9-carrier repeat* > *GT 10/10 repeat* $p=0.001$; *GG 10/10 repeat* > *GG 9-carrier repeat* $p=0.01$). Polynomial regression also revealed a non-linear relationship between compound genotype and gray matter in the caudate (respectively, $R^2=0.18$, $F=9.46$, $p=0.0001$, Figure 5C).

Direct interaction between Dopamine D2 Receptors and DAT proteins *in vivo*

Previous data indicate that reduction of D2 receptors or DAT strongly influence neuronal activity as well as the presence of an interaction between the different genotypes. Importantly, a direct protein-protein interaction between DAT and D2 has been recently demonstrated. This interaction promotes the recruitment of DAT to the plasma membrane and regulates DAT function in transfected cells as well as in primary culture of rat midbrain neurons (Lee et al., 2007). Disruption of DAT/D2 interaction has been postulated to decrease dopamine uptake (Lee et al., 2007). We thus assessed whether these two proteins might physically interact *in vivo*. To do this we used newly generated anti-D2 receptor antibodies (Doi et al., 2006; Tirotta et al., 2008) to perform immunoprecipitation analyses from striatal mouse extracts. As control of our experiments, we used D2 receptor knockout striatal extracts. Importantly, we found that D2 receptors interact with DAT *in vivo* (Fig. 6A). Indeed, using either DAT antibodies (IP DAT, Fig. 6A) or D2 receptor antibodies (IP D2; Fig. 6B), we were able to respectively immunoprecipitate either D2 (Fig. 6A) or DAT (Fig. 6B) from striatal extracts of wild-type

(WT) animals. This interaction is abolished in the brain of knockout animals ($D2R^{-/-}$) in which immunoprecipitation analyses, using either the D2 or DAT specific antibodies, did not detect the presence of a band, as expected (Fig. 6), further confirming the specificity of the antibodies.

DISCUSSION

Consistent with earlier functional imaging studies, with the known distribution of dopamine projections and of D2S and DAT in the synapse, our results suggest a robust interaction between *DRD2* rs1076560 genotype and the 3' *VNTR* polymorphism of the *DAT*. More specifically, *DRD2* and *DAT* genotypes have independent effects on activity of brain areas involved in the working memory network with *DRD2* heterozygotes and *DAT 9-repeat* carriers having greater activity. Moreover, these two genotypes interact in modulating response of the cortical and subcortical working memory network, including the striatum and the dorsolateral prefrontal cortex. The data obtained strongly suggest that the effect of *DAT* alleles is especially manifest in the context of *DRD2* GT genotype. Also, these two genotypes identify similar main effects and interactions in the striatum and dorsolateral prefrontal cortex during encoding of recognition memory, providing replication of the effects in the context of another cognitive paradigm. Consistent with earlier literature, these interactions identify non-linear relationships between putative dopamine levels and brain activity. As a further demonstration of the functional effects of these polymorphisms, these two genotypes also interacted in determining gray matter volume of the caudate. Once again, the effect of *DAT* is mostly evident in the context of *DRD2* GT genotype. Moreover, to strengthen these results in humans and the *in vitro* earlier results of a physical D2-DAT interaction, we also demonstrate that these two proteins indeed interact *in vivo* as indicated by the immunoprecipitation analyses from striatal mouse extracts providing a possible molecular mechanism of the effects in humans.

The present results are consistent with earlier studies from our group and others indicating that *DRD2* and *DAT 3' VNTR* genotypes have independent effects on activity of the striatum and of the prefrontal cortex during different cognitive challenges (Smolka et al., 2005; Bertolino et al., 2006a; Yacubian et al., 2007; Zhang et al., 2007; Bertolino et al., 2008b). *DRD2* rs1076590 heterozygotes and *DAT 9-repeat* carriers have greater activity in these brain areas compared with the other genotypes. Moreover, this is the first study addressing the interaction between these two genotypes on *in vivo* measures of brain activity. The molecular mechanisms responsible for the effects we have measured *in vivo* with BOLD fMRI in striatum and in prefrontal cortex may be several. D2 receptors are abundantly expressed in the striatum, where the D2S isoform is mostly presynaptic and inhibits dopamine release. Thus, *DRD2 GT* genotype associated with less D2S may increase dopamine levels in the striatum thus increasing its activity. DATs in the striatum are also found presynaptically and are responsible for dopamine reuptake. Similar to *DRD2*, the effect of *DAT 9-carrier* genotype may be associated with less dopamine reuptake, greater dopamine levels and, eventually, greater striatal activity. As for the interaction between the two genes, recent evidence has clearly demonstrated that D2 receptors and DATs are often coupled in presynaptic terminals and they reciprocally regulate each other. Therefore, the interaction we demonstrate in the striatum can be easily understood in that the *DRD2 GT* genotype is associated with reduced expression of presynaptic D2 receptors which might induce less presynaptic inhibition and greater release of dopamine. Similarly, the *9-repeat* allele of the *DAT*, associated with lower expression of DATs, very likely induces diminished reuptake of dopamine and thereby greater striatal activity. Our data further suggest that the effect of *DRD2 GT* genotype might be compensated by the *10-repeat* allele of the *DAT*, or, conversely, that the effect of *DRD2 GG* genotype is compensatory of the *9-repeat* allele of the *DAT*. These results are also consistent with the VBM data in striatum suggesting an interaction of these two genes on gray matter volume. This effect can be explained by the known trophic action of dopamine on neurons and synapses (Nieoullon, 2002). Interestingly, a physical interaction between DAT and D2 has been reported and the

responsible regions have been identified in *in vitro* studies (Lee et al., 2007). These findings are nicely complemented by our immunoprecipitation results which provide *in vivo* evidence that, in the mouse brain, D2 receptors interact with DAT as demonstrated by the reciprocal ability of D2 and DAT antibodies to immunoprecipitate either DAT or D2, respectively in WT striatal extracts. The specificity of this interaction is indicated by the total absence of DAT or D2 in immunoprecipitates from D2 receptor knockout mice. Disruption of this interaction has been postulated to decrease dopamine uptake (Lee et al., 2007). Thus, absence or reduction of D2 might prevent/impair the formation of this complex, leading to aberrant DA release and reuptake. In agreement with this contention, blockade of DAT activity by cocaine stimulates an outstanding elevation of the extracellular dopamine levels, in the striatum of D2R^{-/-} mice, which is well above that evoked in WT mice (Rouge-Pont et al., 2002).

Explanation of the interaction of these two genetic variants in prefrontal cortex may be more complex. D2 receptors in prefrontal cortex are also found presynaptically on dopamine terminals (Pickel et al., 2002) modulating dopamine release and D2S seem to be expressed relatively more abundantly in prefrontal cortex (Zhang et al., 2007). In the cortex, DATs tend to be found on nonvaricose axon segments of small diameter which make symmetric synapses with GABA neurons (Sesack et al., 1998; Lewis et al., 2001), with a distribution similar to that of some D₂ receptors (Haber et al., 1995; Pickel et al., 2002; Negyessy and Goldman-Rakic, 2005). Thus, both D2 receptors and DATs may act through GABA neurons to indirectly modulate prefrontal pyramidal neuron firing (Goldman-Rakic, 1996). An alternative explanation of this interaction in prefrontal cortex is that the cortico-striato-thalamic-cortical is critical for cognition and memory (Alexander et al., 1986). An important modulator of this circuit is dopamine. Specifically, greater release of dopamine in the striatum increases activity of the whole network (Tisch et al., 2004). Thus, it is possible that the effect of *DRD2* and *DAT* genotypes as well as their interaction might be associated with modulation of dopamine release in the striatum which would increase activity in the whole cortico-striato-thalamic-cortical circuit (Tisch et al., 2004).

Like in earlier studies in healthy humans which used other genes to study dopamine signaling (Yacubian et al., 2007; Bertolino et al., 2008b), in the present study the interaction between *DRD2* and *DAT* genotypes identifies non-linear relationships of dopamine signaling with neuronal activity in the caudate and in prefrontal cortex during working memory. This interaction between the two genes is consistent with a large series of experiments suggesting a non-linear inverted-U shaped relationship between prefrontal dopamine levels and prefrontal neuronal activity (Williams and Castner, 2006; Vijayraghavan et al., 2007). Our data are consistent with these earlier studies investigating dopamine modulation of prefrontal neuronal activity. Moreover, together with previous studies (Yacubian et al., 2007; Bertolino et al., 2008b), the present results suggest that dopamine modulation of neuronal activity may follow a non-linear dose-response curve also in other brain areas, including the striatum. Our data also extend earlier studies suggesting that this non-linear inverted-U relationship is not exclusively determined by D1 receptor signaling but also includes D2 receptors and DATs. In this regard, we note that the profile of the curve may change if changing the order of compound genotypes on the y-axis. However, we had preselected the genotype order to reflect the hypothesis that genotypes putatively associated with intermediate levels of dopamine (those in which the effect of *DRD2* would be compensated by the *DAT* effect or vice versa) would be more efficient (less BOLD activity for a given level of behavioral performance). This hypothesis is consistent with several other studies in the literature (reviewed in (Seamans and Yang, 2004; Williams and Castner, 2006)) indicating that too much or too little dopamine activity is detrimental for memory and it is associated with reduced efficiency.

Some potential limitations of the present data have to be discussed. The working memory and VBM samples were not matched for gender, and we covaried the relative analyses for this

variable to control for the possibility that gender modulates the interaction between genotypes on brain activity or gray matter volume. Therefore, we cannot fully discount the possibility of a gender effect on the variables we have measured. However, this was not the main purpose of our study and exploring a three-way interaction would have probably required an even larger number of subjects to achieve sufficient statistical power.

Some of the imaging findings are lateralized. One reason for these results may be the nature of cognitive requests of task paradigms. For example, in several of our earlier papers with the N-Back paradigm findings were either lateralized or they were statistically more robust in left DLPFC (Bertolino et al., 2004; Bertolino et al., 2006a; Bertolino et al., 2006b; Bertolino et al., 2008a). Therefore, a likely reason for lateralization in DLPFC is the verbal vs. non-verbal nature of the two memory tasks.

Another limitation of the present study is that we have not measured dopamine receptors and transporters directly in humans. Therefore, the effects demonstrated with BOLD fMRI are not necessarily related to dopamine levels or release. Our results might be related to plasticity mechanisms associated with different molecular pathways present at the same time but not necessarily related to these two genotypes. However, the immunoprecipitation results in the animal experiments at the very least suggest that these interactions are possible *in vivo*.

In conclusion, the data of the present study suggest a dopamine-related mechanism modulating activity of the prefronto-striatal network during memory performance. This genetically modulated mechanism may be important to determine brain phenotypes found in several disorders implicating dopamine dysregulation of prefronto-striatal networks, including Schizophrenia, drug abuse, and Parkinson's Disease.

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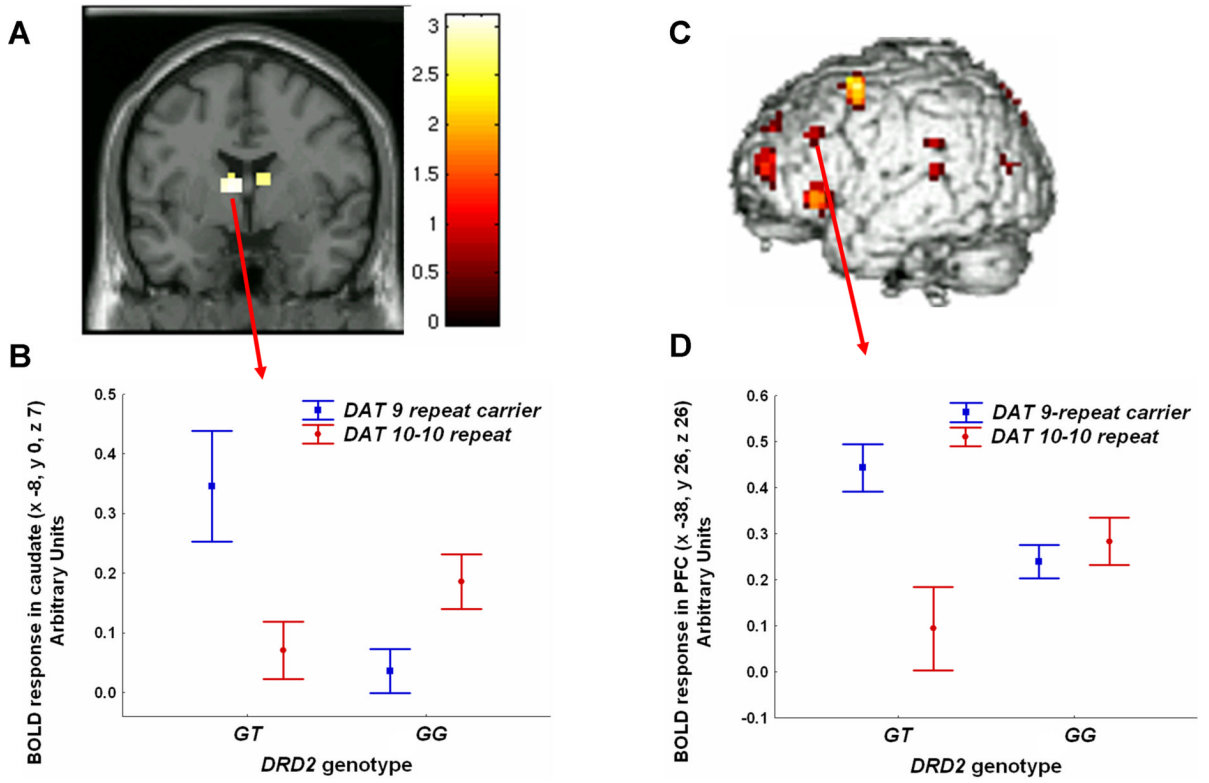


Figure 1.
A: Coronal MRI section through the caudate nuclei indicating locales with *DRD2-DAT* genotype interaction on BOLD response during working memory (image thresholded at $p < 0.005$, uncorrected); **B:** mean \pm SE of BOLD response in caudate of the interaction between *DRD2* and *DAT* genotypes. **C:** 3D rendering indicating the interaction between *DRD2-DAT* genotypes on cortical working memory; **D:** mean \pm SE confidence intervals of BOLD response of the interaction between *DRD2-DAT* genotype in left middle frontal gyrus activity during working memory.

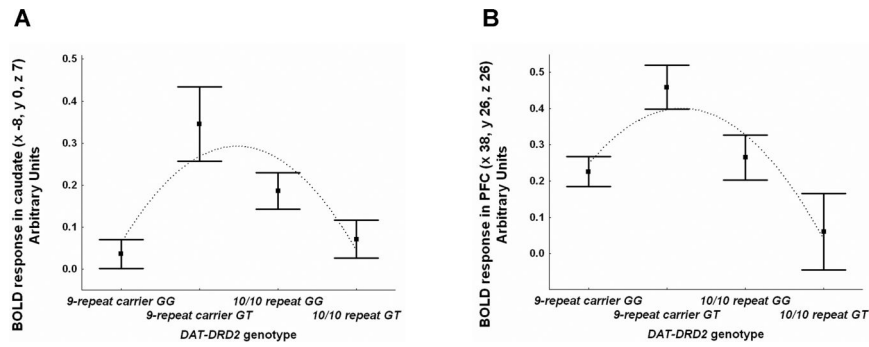


Figure 2. mean \pm SE of BOLD response in caudate (A) and in middle frontal gyrus (PFC, B) during working memory in compound genotype groups. Since we expected that the genotypes potentially associated with putative intermediate levels of dopamine would show more focused activity for a similar level of behavioral performance, genotype groups are ordered from putative less dopamine reuptake (*DAT 9-carriers* have less DAT expression) and release (*DRD2 GG* subjects have greater presynaptic D2 mRNA) to greater reuptake and release (*DAT 10–10 repeat DRD2 GT* subjects). The relationship between compound genotype and BOLD response is non-linear. Genotypes on the y-axis are ordered to reflect the nature of our hypothesis about the relationship between putative levels of dopamine stimulation determined by compound genotypes and BOLD activity for a given level of performance. Thus, genotypes at the two extremes of the curve are hypothesized to have intermediate levels of dopamine stimulation and more efficient BOLD activity (less activity for a given behavioral performance).

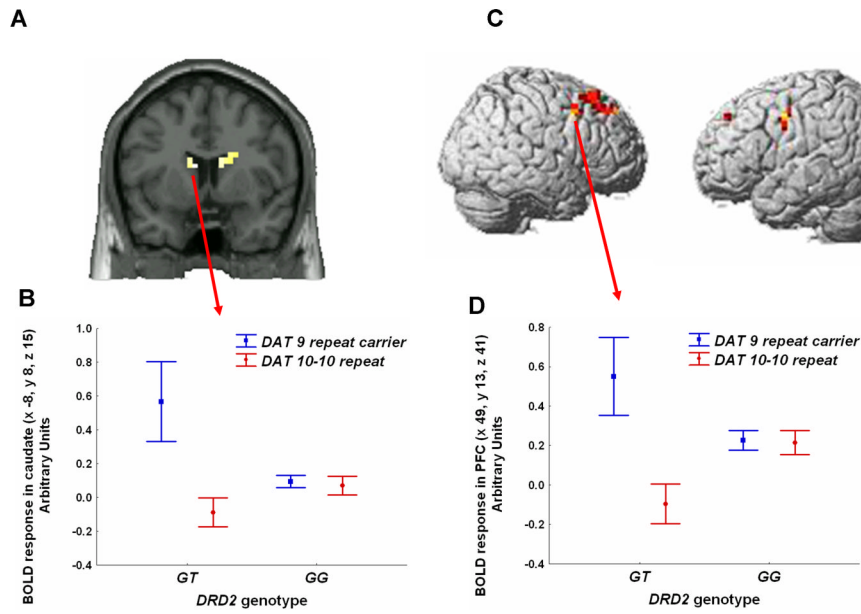


Figure 3.

A: Coronal MRI section through the caudate nuclei indicating locales with *DRD2-DAT* genotype interaction on BOLD response during encoding of recognition memory (image thresholded at $p < 0.005$, uncorrected); **B:** mean \pm SE of BOLD response of the interaction between *DRD2* and *DAT* genotypes. **C:** 3D rendering indicating the interaction between *DRD2-DAT* genotypes on cortical activity during encoding of recognition memory; **D:** mean \pm SE of BOLD response of the interaction between *DRD2-DAT* genotype in right middle frontal gyrus activity during encoding of recognition memory.

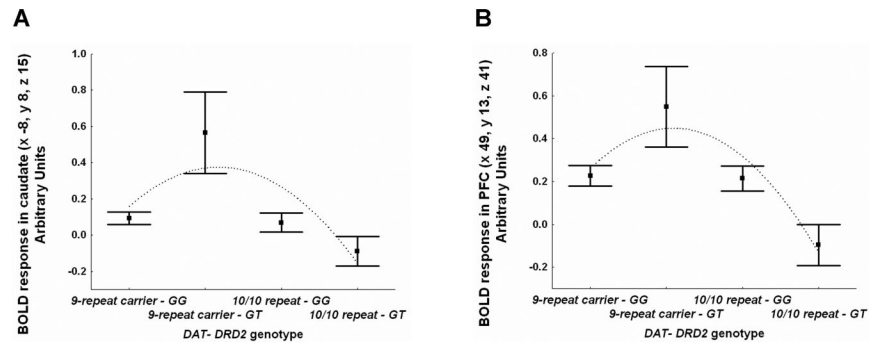


Figure 4. mean \pm SE of BOLD response in caudate (**A**) and in middle frontal gyrus (PFC, **B**) during encoding of recognition memory in compound genotype groups ordered from putative less dopamine reuptake (*DAT 9-carriers* have less DAT expression) and release (*DRD2 GG* subjects have greater presynaptic D2 mRNA) to greater reuptake and release (*DAT 10–10 repeat DRD2 GT* subjects). The relationship between compound genotype and BOLD response is non-linear.

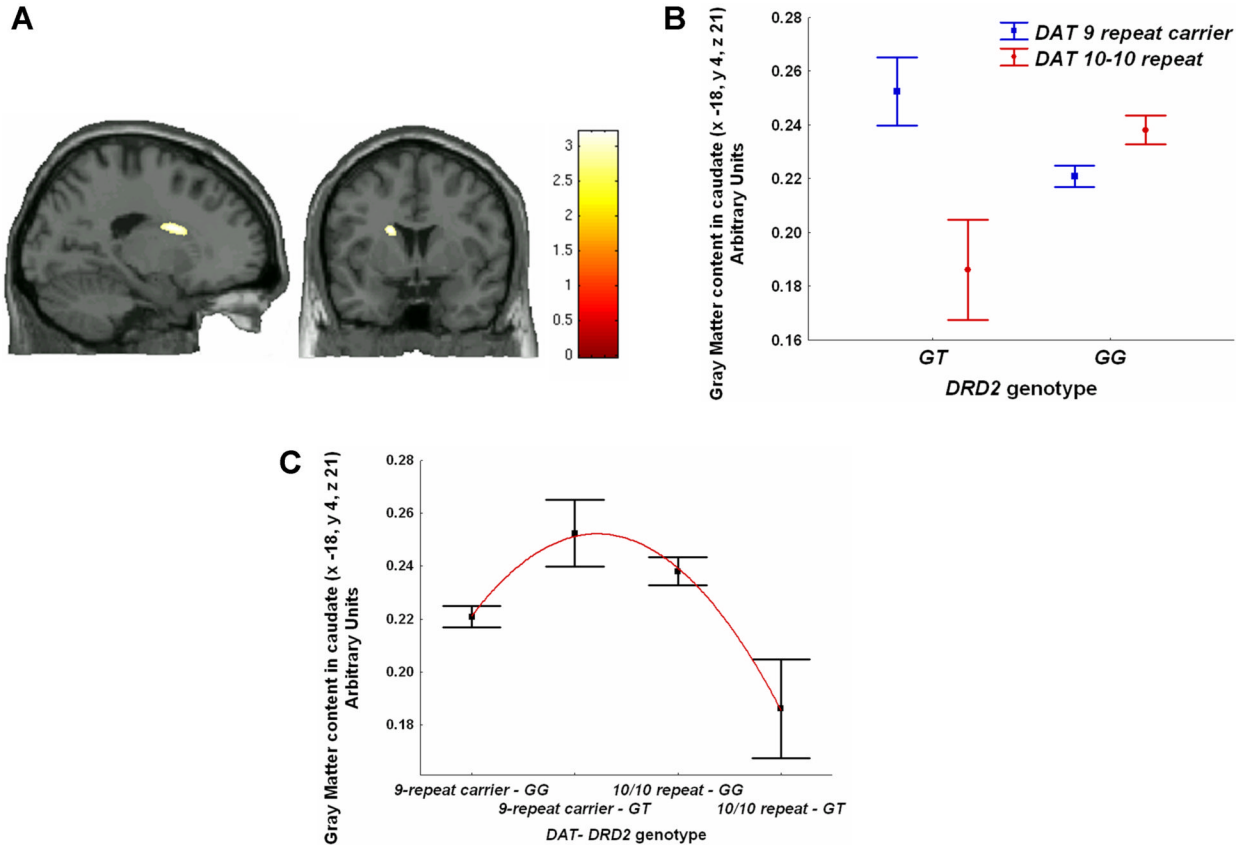


Figure 5.
A: Sagittal and Coronal MRI sections through the caudate nuclei indicating locales with *DRD2*-*DAT* genotype interaction on gray matter content (image thresholded at $p < 0.005$, uncorrected);
B: mean \pm SE of gray matter content of the interaction between *DRD2* and *DAT* genotypes;
C: mean \pm SE of gray matter content in caudate in compound genotype groups ordered from putative less dopamine reuptake (*DAT 9-carriers* have less *DAT* expression) and release (*DRD2 GG* subjects have greater presynaptic *D2* mRNA) to greater reuptake and release (*DAT 10–10 repeat DRD2 GT* subjects). The relationship between compound genotype and gray matter content is non-linear.

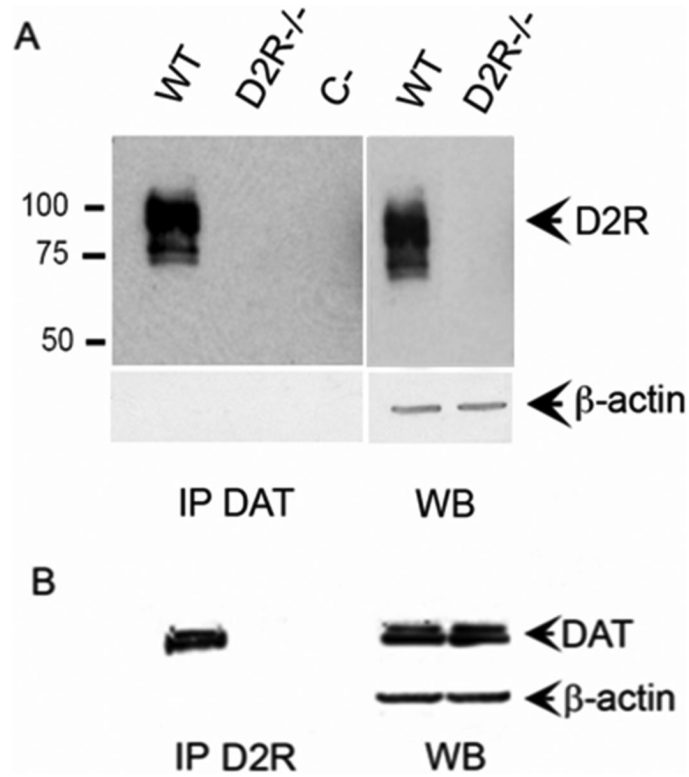


Figure 6. Immunoprecipitation experiments show D2R-DAT interaction *in vivo*
 Striatal membranes were prepared from WT and D2R^{-/-} mice. 500 μ g proteins were used for Immunoprecipitation experiments using either DAT or D2R specific antibodies. **A)** Samples were immunoprecipitated with anti-DAT and revealed with anti-D2R antibodies (first two lanes). Note the presence of immunoprecipitated D2R only in WT striata but not from D2R^{-/-} striatal membranes; C-, control experiments were performed using normal IgG rabbit. Western Blots (WB) were performed using 50 μ g striatal membranes showing the presence of D2R only in WT extracts. **B)** Same as in A, but proteins were precipitated with anti-D2R antibody and revealed with anti-DAT antibody; C-, control experiments were performed using normal IgG mouse. Note that DAT was precipitated only from WT extracts. Western Blots as in A were revealed using anti-DAT antibody showing presence of DAT in WT and D2R^{-/-} extracts. Actin was used as internal control of loaded quantities of striatal extracts in A and B.

Table 1
Demographics, N-back working memory and encoding of recognition memory behavioral performance.

Working Memory		Age	IQ	Handedness	Working Memory Test (2 Back)	Working Memory Test (2 Back)	reaction time (msec)
Total N (males)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	
DRD2							
GG	110(37♂)	27.7 (±7.4)	113.5 (±13.0)	0.75 (±0.43)	67.7 (±16.7)	523.2 (±248.9)	
GT	32(19♂)	28.4 (±7.9)	111.4 (±14.7)	0.73 (±0.32)	69.4 (±15.7)	499.7 (±246.0)	
DAT							
DRD2-DAT							
10/10 repeat	51(21♂)	28.6 (±7.1)	115.7 (±10.8)	0.70 (±0.43)	70.2 (±15.4)	517.0 (±254.8)	
9 repeat carriers	91(35♂)	27.4 (±7.6)	111.6 (±14.4)	0.76 (±0.38)	66.9 (±17.0)	518.1 (±244.7)	
DRD2-DAT							
GG-10/10 repeat	41(13♂)	28.6 (±7.1)	117.9 (±9.6)	0.72 (±0.43)	69.4 (±16.3)	507.3 (±261.5)	
GG-9 repeat carriers	69(24♂)	27.1 (±7.5)	111.0 (±14.0)	0.76 (±0.42)	66.7 (±17.0)	533.0 (±242.4)	
GT-10/10 repeat	10(8♂)	28.4 (±7.6)	107.0 (±11.5)	0.62 (±0.43)	73.2 (±11.3)	553.9 (±235.4)	
GT-9 repeat carriers	22(11♂)	28.4 (±8.2)	114.1 (±15.6)	0.78 (±0.25)	67.6 (±17.3)	473.3 (±252.2)	
Recognition Memory							
Total N (males)	Age	IQ	Handedness	Recognition Memory Test (encoding)	Recognition Memory Test (encoding)	reaction time (msec)	
Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)		
DRD2							
GG	82(23♂)	26.2 (±5.9)	112.6 (±11.4)	0.7 (±0.47)	94.8 (±3.9)	1129.5 (±187.3)	
GT	20(9♂)	26.5 (±5.8)	112.5 (±12.2)	0.7 (±0.36)	92.5 (±5.5)	1170.6 (±221.5)	
DAT							
10/10 repeat	38(10♂)	27.5 (±5.3)	113.7 (±10.4)	0.66 (±0.46)	94.6 (±3.9)	1140.4 (±192.2)	

Recognition Memory		Age	IQ	Handedness	Recognition Memory Test (encoding)	Recognition Memory Test (encoding)
	Total N (males)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)	Mean (sd)
9 repeat carriers	64(22♂)	25.6 (±6.1)	112.1 (±12.0)	0.73 (±0.44)	94.2 (±4.6)	1136.1 (±196.8)
DRD2-DAT						
GG-10/10 repeat	33(7♂)	27.0 (±5.6)	114.5 (±9.7)	0.62 (±0.49)	94.7 (±3.9)	1113.3 (±182.6)
GG-9 repeat carriers	49(16♂)	25.7 (±6.2)	111.7 (±12.1)	0.76 (±0.46)	94.8 (±3.9)	1140.0 (±191.7)
GT-10/10 repeat	5(3♂)	30.2 (±2.2)	109.0 (±5.7)	0.87 (±0.16)	94.2 (±3.7)	1303.3 (±183.1)
GT-9 repeat carriers	15(6♂)	25.5 (±6.0)	113.2 (±11.9)	0.70 (±0.39)	92.0 (±6.0)	1123.2 (±220.0)

Table 2

Statistics and Montreal Neurological Institute coordinates for the effects of DRD2 and DAT genotypes as well as for their interaction on brain activity during working memory.

Region (Brodmann's area)	Main effect of DRD2			
	Talairach coordinates (x y z)	k	Z value	P uncorrected
Right Inferior Frontal Gyrus (BA 45)	59 23 6	25	3.30	0.000
Right Middle Frontal Gyrus (BA 10)	30 52 -3	7	2.74	0.003
Right Putamen	26 7 -6	3	2.61	0.004
Right Anterior Cingulate (BA 32)	15 26 19	8	2.9	0.002
Region (Brodmann's area)	Main effect of DAT			
	Talairach coordinates (x y z)	k	Z value	P uncorrected
Left Middle Frontal Gyrus (BA 6)	-41 6 51	8	3.18	0.001
Left Middle Frontal Gyrus (BA 8/9)	-49 20 37	11	2.83	0.002
Left Inferior Frontal Gyrus (BA 47)	-45 22 -7	7	3.09	0.001
Region (Brodmann's area)	Interaction between DAT and DRD2			
	Talairach coordinates (x y z)	k	Z value	p (FDR-corrected)
Left Middle Frontal Gyrus (BA 6)	-38 6 51	15	4.17	0.019
Left Middle Frontal Gyrus (BA 9)	-38 26 26	19	3.21	0.019
Left Caudate Body	-8 0 7	32	3.04	0.019
Right Caudate Body	11 -6 18	14	2.95	0.019
Region (Brodmann's area)	Interaction between DAT and DRD2			
	Talairach coordinates (x y z)	k	Z value	P uncorrected
Right Caudate Body	26 -72 21	6	3.48	0.000
Left Caudate Body	-34 49 33	7	3.31	0.000
Left Middle Frontal Gyrus (BA 10)	-38 52 11	9	3.23	0.001
Left Cingulate Gyrus (BA 32)	-11 24 34	19	3.21	0.001
Right Superior temporal Gyrus (BA 39)	38 -53 24	4	3.19	0.001
Left Inferior Frontal Gyrus (BA 47)	-45 22 -7	17	3.19	0.001
Left Superior Temporal Gyrus (BA 22)	-63 -35 16	3	3.00	0.001

Table 3

Statistics and Montreal Neurological Institute coordinates for the effects of DRD2 and DAT genotypes as well as for their interaction on brain activity during encoding of recognition memory.

Main effect of DRD2				
Region (Brodmann's area)	Talairach coordinates (x y z)	k	Z value	P uncorrected
Left Putamen	-22 5 14	6	3.10	0.001
Right Inferior Frontal Gyrus (BA47)	48 22 -1	6	2.88	0.002

Main effect of DAT				
Region (Brodmann's area)	Talairach coordinates (x y z)	k	Z value	P uncorrected
Thalamus, Medial Dorsal Nucleus	-8 -18 11	54	3.57	0.000
Left Precentral Gyrus (BA 6)	-55 -2 42	21	3.38	0.000
Inferior Frontal Gyrus (BA 47)	-19 10 -16	4	3.20	0.001
Supramarginal Gyrus (BA 40)	41 -46 34	3	3.10	0.001
Thalamus	4 -4 8	17	2.92	0.002
Clastrum	30 18 -1	8	2.84	0.002

Interaction between DAT and DRD2				
Region (Brodmann's area)	Talairach coordinates (x y z)	k	Z value	p (FDR-corrected)
Right Caudate Body	- 8 8 15	15	3.06	0.047
Left Caudate Body	-11 7 17	2	3.00	0.047
Right Middle Frontal Gyrus (BA 8)	49 13 41	10	2.83	0.047

Interaction between DAT and DRD2				
Region (Brodmann's area)	Talairach coordinates (x y z)	k	Z value	P uncorrected
Inferior Frontal Gyrus (BA 47)	-22 10 -16	11	3.24	0.001
Precuneus (BA 19)	26 -78 35	4	3.19	0.001
Precentral Gyrus (BA 6)	-41 -5 42	13	3.10	0.001
Precentral Gyrus (BA 4)	59 -19 42	4	3.08	0.001
Thalamus, Medial Dorsal Nucleus	4 -21 8	5	3.00	0.001
Caudate Body	19 -9 24	4	2.88	0.002
Caudate Tail	26 -30 -17	3	3.11	0.001
Inferior Frontal Gyrus (BA 45)	49 19 9	5	2.84	0.002
Inferior Frontal Gyrus (BA 13)	30 14 -10	8	2.76	0.003