ADHD in adolescence and adulthood, with a special focus on the dopamine transporter and nicotine

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The persistence of attention deficit hyperactivity disorder (ADHD) into adolescence and adulthood has now been accepted as a clinical entity. The rate of prevalence among adults is assumed to be from 2% to 4%. With increasing age, a symptom change has to be considered; disturbance of attention becomes more prominent, whereas hyperactivity often diminishes or changes to inactivity. Neuroimaging studies show a high striatal dopamine transporter (DAT) availability in most adults with ADHD; this can be reduced by stimulants. Nicotine seems to have a stimulant-like action on the DAT. In most adults with ADHD, therapy has to be multimodal, combining psychotherapy and medication. Methylphenidate is the first-line drug in adult ADHD; further options are amphetamine and noradrenaline reuptake inhibitors. Nonresponders to methylphenidate seem to have no elevated DAT availability prior to therapy. Combination with other psychiatric disorders occurs frequently in adults with ADHD; in these patients additional pharmacological treatment with special regard to the comorbid disease is recommended.

Keywords: attention deficit hyperactivity disorder; adolescence; adulthood; brain imaging techniques; single photon emission computed tomography; dopamine transporter; nicotine; diagnosis; therapy; methylphenidate

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Symptomatology and diagnosis

The core symptoms of ADHD are inattention, hyperactivity, and impulsivity. In most descriptions of ADHD in the 1980s and the early 1990s, it seemed that hyperactivity had to be present in every case as a striking symptom, but with growing knowledge of ADHD it became evident that not all patients—in particular girls—present hyperactivity. Since 1994, with the fourth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM IV) three types of ADHD have been differentiated:

- Combined type (6 or more symptoms of hyperactivity/impulsivity as well as of inattention);
- Inattentive type (6 or more symptoms of inattention);
- Hyperactive/impulsive type (6 or more symptoms of hyperactivity/impulsivity).

A change in symptoms with increasing age is characteristic of ADHD. After puberty, hyperactivity often...
changes to inactivity; therefore, ADHD often is not accepted as a diagnosis in adults. Impulsivity normally lessens with age. Adults with ADHD seem to have a problem with willpower: activity can be overexerted, but if a task is not attractive to the person, this results in a lack of effort. Or, inattention can change to hyperfocussing, when the person is attracted by a task.

With adults, differing patterns of comorbidity and symptom heterogeneity pose new conceptual, diagnostic, and treatment challenges. While core symptoms are often overt problems in children, in adults subtler executive dysfunction appears. Even though the growing consensus is that ADHD is a disorder of executive functions (EF), the details of the EF/ADHD connection remain unclear and may be far more complex in adults.4

In Table I examples are given for the changes of the 18 DSM-IV symptoms from childhood to adulthood. The 6-question Adult Self-Report Scale – V 1.1 (ASRS - V 1.1) Screener (http://www.hcp.med.harvard.edu/ncs/fpdir/adhd) is a subset of the WHO’s 18-question Adult Self-Report Scale – V 1.1 (ASRS - V 1.1) Symptom Checklist. The patient should fill in checkmarks. Four or more checkmarks in the darkly shaded areas may indicate that the symptoms are consistent with adult ADHD (Figure 1). Wender developed a set of characteristics to specify both childhood criteria and current ADHD symptoms.5 He pointed out affective lability, which is not mentioned in DSM-IV, as a frequent symptom in adult ADHD.

Prevalence of ADHD in adulthood

The prevalence of ADHD in children according to DSM-IV criteria varies from 2.4% to 19.8%.6 Concerning persistence into adulthood, most authors describe a rate of about 50%. The largest follow-up study, which investigated 197 Chinese children after 15 years, showed a rate of persistence of 70%.7 Generally, the degree of prevalence (1% to 6% in adults) depends on the view of the reporter in the initial assessment. Most instruments consist of some form of self-report, and in adulthood it is often not possible to ask information of parents or persons with a close relationship to the patient. Patients with ADHD are often not aware of their symptoms, or do not report the severity of symptoms.

Neurobiological basis of ADHD

Current interest in the neurobiological basis of ADHD originally commenced in the 1970s. Neurochemical, neurophysiological, and radiological attributes were noted, proving, in particular, abnormalities in the dopaminergic and noradrenergic system. Genetic investigations showed increased evidence that genetic components were present in most cases of ADHD, which is now seen as the psychiatric disease with the highest heritability.9 Neuroimaging studies with magnetic resonance imaging (MRI), photon-emission tomography (PET), and single photon-emission computed tomography (SPECT) suggest involvement of striatal structures in ADHD.10-13 Molecular genetic studies described an involvement of a polymorphism of the dopamine transporter (DAT) 1 gene in ADHD (eg, a higher rate of homozygosity of the 10-repeat allele in the 3’untranslated region of exon 15 of the DAT gene on chromosome 5p15.3).14-19 With regard to the suspected striatal impairment in ADHD and the known effect of methylphenidate on the

Clinical research

Selected abbreviations and acronyms

| Abbreviation | Definition |
|--------------|------------|
| ADHD         | attention deficit hyperactivity disorder |
| DAT          | dopamine transporter |
| MRI          | magnetic resonance imaging |
| PET          | photon-emission tomography |
| SPECT        | single photon-emission computed tomography |

ASRS 6 Questions

1. How often do you have trouble wrapping up the final details of a project, once the challenging parts have been done? [Never, Rarely, Sometimes, Often, Very often]
2. How often do you have difficulty getting things in order when you have to do a task that requires organization? [Never, Rarely, Sometimes, Often, Very often]
3. How often do you have problems remembering appointments or obligations? [Never, Rarely, Sometimes, Often, Very often]
4. When you have a task that requires a lot of thought, how often do you avoid or delay getting started? [Never, Rarely, Sometimes, Often, Very often]
5. How often do you fidget or squirm with your hands or your feet when you have to sit down for a long time? [Never, Rarely, Sometimes, Often, Very often]
6. How often do you feel overly active and compelled to do things, like you were driven by a motor? [Never, Rarely, Sometimes, Often, Very often]

Figure 1. Adult Self-Report Scale (ASRS) Screener: 4 or more checkmarks in the shaded areas may indicate symptoms of adult ADHD. ADHD, attention deficit hyperactivity disorder.
DAT mainly localized in this section of the brain, since the end of the 1990s DAT has been investigated in patients with ADHD. This is carried out by using radioactive marked ligands which are known to bind effectively with the DAT system. A group of investigators from Boston used altropane marked with iodine-123; our group from Munich and Philadelphia used the cocaine derivative TRODAT-1 marked with technetium 99m. Both studies showed a clearly higher availability of DAT in the striatum of adult patients with ADHD compared with the normal healthy controls. While DAT availability was found to be raised to 17% in the investigations using TRODAT-1 compared with the control collective (Figure 2), the group using altropane demonstrated a rise of 70% in 6 patients;

| 1. How often do you have trouble wrapping up the final details of a project, once the challenging parts have been done? | Often does not follow through on instructions and fails to finish schoolwork, chores, or duties in the workplace (and due to oppositional behavior or failure to understand instructions) |
| 2. How often do you have difficulty getting things in order when you have to do a task that requires organization? | Often has difficulty organizing tasks and activities |
| 3. How often do you have problems remembering appointments or obligations? | Is often forgetful in daily activities |
| 4. When you have a task that requires a lot of thought, how often do you avoid or delay getting started? | Often avoids, dislikes, or is reluctant to engage in tasks that require sustained mental effort (such as schoolwork or homework) |
| 5. How often do you fidget or squirm with your hands or feet when you have to sit down for a long time? | Often fidgets with hands or feet or squirms in seat |
| 6. How often do you feel overly active and compelled to do things, like you were driven by a motor? | Is often “on the go” or often acts as if “driven by a motor” |
| 7. How often do you make careless mistakes when you have to work on a boring or difficult project? | Often fails to give close attention to details or makes careless mistakes in schoolwork, work, or other activities |
| 8. How often do you have difficulty keeping your attention when you are doing boring or repetitive work? | Often has difficulty sustaining attention in tasks or play activities |
| 9. How often do you have difficulty concentrating on what people say to you, even when they are speaking to you directly? | Often does not seem to listen when spoken to directly |
| 10. How often do you misplace or have difficulty finding things at home or at work? | Often loses things necessary for tasks or activities (eg, toys, school assignments, pencils, books, or tools) |
| 11. How often are you distracted by activity or noise around you? | Is often easily distracted by extraneous stimuli |
| 12. How often do you leave your seat in meetings or other situations in which you are expected to remain seated? | Often leaves seat in classroom or in other situations in which remaining seated is expected |
| 13. How often do you feel restless or fidgety? | Often runs about or climbs excessively in situations in which it is inappropriate (in adolescents or adults, may be limited to subjective feelings of restlessness) |
| 14. How often do you have difficulty unwinding and relaxing when you have time to yourself? | Often has difficulty playing or engaging in leisure activities quietly |
| 15. How often do you find yourself talking too much when you are in social situations? | Often talks excessively |
| 16. When you’re in a conversation, how often do you find yourself finishing the sentences of the people you are talking to, before they can finish them themselves? | Often blurts out answers before questions have been completed |
| 17. How often do you have difficulty waiting your turn in situations when turn taking is required? | Often has difficulty awaiting turn |
| 18. How often do you interrupt others when they are busy? | Often interrupts or intrudes on others (eg, butts into conversations or games) |

Table I. Comparison of ADHD symptoms in adulthood according to ASRS (http://www.med.nyu.edu/psych/assets/adhdscreen18.pdf.) in the left column and in childhood according to DSM-IV in the right column. ADHD, attention deficit hyperactivity disorder.
this percentage was reduced to around 30% in a larger group of 19 patients on continuation of the altropane study (Spencer T, personal communication). Meanwhile, a rise in DAT availability was also detected in the basal ganglia of children with ADHD by means of an $[{}^{123}I]$ N-(3-iodopropen-2-yl)-2-carbo-methoxy-3β-(4-chlorphenyl) tropane (IPT)-SPECT investigation, following initial results with N-$ω$-fluoropropyl-2 β-carboxymethoxy-3 β-[4-iodophenyl] tropane (FP-CIT). Interestingly, a rise in the DAT could not be detected in a SPECT investigation with β-CIT; this could be due to the extremely slow kinetics of β-CIT (measurement 1 day after application) or another specificity of this radiotracer, which also binds with serotonin transporters and possibly holds other substructures of DAT compared with altropane, FP-CIT, IPT, or TRODAT-1. Our own initial results revealed that the DAT was raised not only in the hyperactive-impulsive and combined type, but also in the inattentive type of ADHD (Figure 3). It may be deduced from a neurochemical view that with an increase in DAT, which transports dopamine back from the synaptic cleft, less dopamine is available in the synaptic cleft of dopamine-dependent neurons. The results obtained by our group showed not only the impairment of DAT but also reported for the first time in vivo and intraindividually in patients with ADHD, that impaired metabolism can be improved with methylphenidate: on administration of 5 mg 3 times daily it was seen that there was a clear reduction in DAT availability in all patients after a period of 4 weeks (Figure 2). With this very low dose, most of these patients demonstrated a lower concentration of DAT than did the control group. In a study involving six children with ADHD, an IPT-SPECT investigation confirmed a marked reduction in DAT density under administration of methylphenidate. Regarding normal healthy individuals, other authors were able to show evidence of a similar reduction in DAT under methylphenidate in a PET investigation with $[^{11}C]$ cocaine. In conclusion, the SPECT investigations on DAT confirmed the supposition that with ADHD an impairment is present mainly in the dopaminergic system. This can be improved with the introduction of stimulants. Our initial results show that nonresponders to methylphenidate do not reveal raised DAT in striatum prior to therapy, whereas the responders seem to have a high DAT availability (Figure 4).
It should be noted, for possible diagnostic applications of SPECT, that the DAT concentration decreases with increase of age and that nicotine may influence DAT availability. The lowering of DAT with increasing age may be an explanation for our observation, that most adults need lower doses of methylphenidate compared with children and adolescents. Nicotine seems to have a similar effect on the DAT as do stimulants (Figure 3).

Investigating potential reasons for the lower elevation in the TRODAT-1 study compared with the altropane study, we found a subgroup of patients with relatively little increase in DAT despite severe clinical impairments; further questioning of these patients revealed that all of them were cigarette smokers. Comparison of 11 smoking nonmedicated patients with ADHD with sex- and age-matched nonsmoking drug-naive adults with ADHD showed significantly higher DAT density in the nonsmoking patients, despite higher ADHD scores for the smokers. This finding suggests that nicotine may act directly on DAT in the same manner as stimulants. In a self-trial, DAT availability was reduced by over 50% 5 hours after intake of 20 mg methylphenidate in a slow-release formulation; wearing a nicotine skin patch, equaling 10 to 20 cigarettes daily, for 5 hours. In the same subject 3 months later, an effect similar to that of methylphenidate was seen (Figure 5), confirming the view that the lowering of DAT after use of nicotine is not a sign of a neurochemical adaptation associated with chronic exposure to nicotine. Further evidence comes from the reevaluation of a woman with ADHD, who was smoking 15 cigarettes per day at the time.

Figure 4. Specific binding of striatal DAT, measured with \([99mTc]TRODAT-1\) SPECT in 18 adults with ADHD (percentage of deviation from values of controls of the same age) in relation to CGI values after 10 weeks of intake of methylphenidate. DAT, dopamine transporter; SPECT, single photon emission computed tomography; ADHD, attention deficit hyperactivity disorder.

Figure 5. DAT availability in the striatum of a 53-year-old nonsmoker before (A), after 5 hours of intake of 20 mg methylphenidate (Ritalin SR®) (B), and 3 months later after 5 hours of wearing a 17.5 mg/24 h nicotine skin patch (C), shown by specific accumulation of \([Tc-99m]TRODAT-1\) in SPECT scans. DAT, dopamine transporter; SPECT, single photon emission computed tomography; STR, specific binding in striatum; BKG, specific binding in the cerebellum as background.

Figure 6. DAT availability in the striatum of a 29-year-old female smoker with ADHD before (A), after 4 weeks of intake of 3x5 mg methylphenidate (Ritalin®) per day (B), and 2 years later after 1 year of cessation of nicotine abuse and intake of methylphenidate, shown by specific accumulation of \([Tc-99m]TRODAT-1\) in SPECT scans. DAT, dopamine transporter; ADHD, attention deficit hyperactivity disorder; SPECT, single photon emission computed tomography; STR, specific binding in striatum; BKG, specific binding in the cerebellum as background.

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of first investigation with TRODAT-1, showing no remark-
able elevation of DAT; 4 weeks of intake of 3x5 mg
methylphenidate led to a marked reduction similar to the
other patients in this group. One year later, the patient
stopped the nicotine and methylphenidate intake; 2 years
after the first investigation she returned with increased
complaints of ADHD. At this time she underwent another
TRODAT-1 SPECT scan, presenting with a 19% elevation
compared with the first scan with nicotine and a 61% ele-
vation compared with the second scan with nicotine and
methylphenidate (Figure 6). This finding is in accordance
with the opinion that the effect of nicotine on DAT does
not result in a persistent loss of DAT. In this context, it is
of interest that an investigation with $[^{123}\text{I}]\beta$-CIT SPECT
showed no altered striatal uptake in smokers versus con-
trols, a potential parallel to the above mentioned findings
of no DAT elevation in ADHD with this method.

**Treatment of adults**

Most patients need multimodal therapy: psychoeducation
concerning the special aspects of ADHD is essential.
Many patients have problems with their self-esteem due
to a misunderstanding of the symptoms and a childhood
with parents suffering from ADHD—especially impulsiv-
ity—and they need additional psychotherapy.

Treatment with stimulants is the drug therapy of first
choice and is approved in the USA for adults (d-
methylphenidate and amphetamines); stimulants for
adult ADHD are as effective as in children. An investi-
gation performed in the year 2000 in Germany showed
that only 0.024% of the adults registered in the public
health system—that is 90% of the adult population—was
treated with methylphenidate in Germany at this time
(Figure 7). That is about one out of 40,000 and includes
all kinds of treatments, including for narcolepsy and
other psychiatric diseases.

In the treatment of adults with ADHD we start with a
daily dose of 5 mg, reaching 15 mg methylphenidate after
14 days of treatment. Titration up to 40 mg follows, if the
patient needs more than the initial end point of the 15-
mg dose. But more than 70% of our patients did not tol-
erate doses above 30 mg without having marked side
effects. Frequently, they felt an increased inner tension.
With higher doses, they also described a state resembling
the impairments they experienced before the treatment.
Most of them had a good stabilization with a combina-
tion of methylphenidate in a dose up to 20 mg and addi-
tional antidepressants.

With all patients we try to start monotherapy with stimu-
lants. When methylphenidate has no positive effect on
depressive or other comorbid symptoms, we start an addi-
tional medication after 4 weeks of treatment. Some
patients do not tolerate methylphenidate well; they feel
more depressed after starting the methylphenidate med-
ication, but they are not as inattentive as before the begin-
ing of therapy; in this situation we change from
methylphenidate to amphetamines; some patients have
more benefits from this kind of stimulants acting in dual
ways. When selecting the additional medication it is impor-
tant to regard the comorbid disorder. We experienced
good clinical response with the following medication:

- Depression: venlafaxine 18.75-150 mg/d
- Depression combined with severe distraction: amisul-
pride 25 – 100 mg/d
- OCD: sertraline 50-100 mg/d
- Borderline syndrome: venlafaxine 37.5 mg -150 mg/d
- Slight autistic symptoms: fluoxetine 10-20 mg/d

The use of tricyclic antidepressants is problematic, as the
long-term effect is not as good as that of treatment with
new antidepressants. The change seems to take place after
3 months. The side effects often lead to treatment dropouts.

In recent years the norepinephrine reuptake inhibitor
atomoxetine was introduced as an additional possibility
to treat ADHD. This substance does not influence the
dopamine in the striatum, but in the prefrontal cortex,
where dopamine transport is mediated by norepineph-
rine transporter.
Trastorno de hiperactividad con déficit de atención de adolescentes y adultos, con especial énfasis en el transportador de la dopamina y en la nicotina

Hoy, se acepta que el trastorno de hiperactividad con déficit de atención (THDA) puede persistir como entidad clínica hasta la adolescencia y vida adulta. La tasa de prevalencia entre los adultos varía entre el 2% y el 4%. Con el envejecimiento se produce un cambio sintomático; el trastorno de atención se agudiza, mientras que la hiperactividad suele disminuir o se torna en inactividad. Los estudios de neuroimagen revelan una gran disponibilidad del transportador de dopamina del estriado (DAT) entre la mayoría de los adultos con THDA; los estimulantes reducen su concentración. Al parecer, la nicotina estimula el DAT. El tratamiento de la mayoría de los adultos con THDA es multimodal — combinación de psicoterapia y medicación. El metilfenidato es el fármaco preferido para el THDA del adulto; otras posibilidades son la anfetamina y los inhibidores de la captación de noradrenalina. Al parecer, las personas que no responden al metilfenidato no muestran una disponibilidad elevada de DAT antes del tratamiento. Los adultos con THDA suelen padecer otros trastornos psiquiátricos; se recomienda un tratamiento farmacológico adicional de estos pacientes, prestando especial atención a la enfermedad asociada.

Hyperactivité et déficit de l’attention (HADA) à l’adolescence et à l’âge adulte: mise au point sur le transporteur de la dopamine et la nicotine

La persistencia de un trastorno de hiperactividad y de un déficit de la atención a la adolescencia y a la edad adulta (HADA) está mantenida aceptada en tant que entidad clínica. La prevalencia en los adultos es estimada entre 2 y 4 %. Se ha tenido cuenta de los síntomas que se modifican con la edad: el trastorno de la atención se agrava cuando la hiperactividad disminuye o se muda en inactividad. Las etudes de neuro-imagerie montan una presencia striata elevada del transportador de la dopamina (TD) en la gran parte de los adultos diagnosticados con HADA, que puede ser disminuida por los estímulos. La nicotina parece una misma acción sobre el TD que los estimulantes. Chez la gran parte de los adultos sufriendo de HADA, el tratamiento debe ser multimodal, abarcando la psicoterapia y los medicamentos. La méthylphénidate es el tratamiento de primera intención en el adulto sufriendo de HADA; las otras opciones son la amphetamina y los inhibidores de la recaptación de noradrenalina. El taux de TD antes de la misma no parece ser elevado entre los no respuesta en el méthylphénidate. D'autres troubles psychiatriques son souvent associés au HADA chez l'adulte; chez ces patients il est recommandé de traiter le trouble comorbide.

REFERENCES

1. Spencer T, Biederman J, Wilens T, et al. A large, double-blind, randomized clinical trial of methylphenidate in the treatment of adults with attention-deficit/hyperactivity disorder. Biol Psychiatry. 2005;57:456-463.
2. Swanson JM, Sergeant JA, Taylor E, Sonuga-Barke EJ, Jensen PS, Cantwell DP. Attention-deficit hyperactivity disorder and hyperkinetic disorder. Lancet. 1998;351:429-433.
3. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders. 4th Edition. Washington, DC: American Psychiatric Association; 1994.
4. Wolf LE, Wasserstein J. Adult ADHD. Concluding thoughts. Ann N Y Acad Sci. 2001;931:396-408.
5. Wender PH. Attention-Deficit Hyperactivity Disorder in Adults. New York, NY, Oxford, UK: Oxford University Press; 1995.
6. Faraone SV, Sergeant J, Gillberg C, Biederman J. The worldwide prevalence of ADHD: is it an American condition? World Psychiatry. 2003;2:104-113.
7. Yan W. An investigation of adult outcome of hyperactive children in Shanghai. Chin Med J. 1996;109:877-880.
8. Yan W. An investigation of adult outcome of hyperactive children in Shanghai. Psychiatry Clin Neurosci. 1998;52(suppl):S303-S305.
9. Faraone SV. Genetics of childhood disorders: XX. ADHD, Part 4: is ADHD genetically heterogeneous? J Am Acad Child Adolesc Psychiatry. 2000;39:1455-1457.
10. Bush G, Valera EM, Seidman LJ. Functional neuroimaging of attention-deficit/hyperactivity disorder: a review and suggested future directions. Biol Psychiatry. 2005;57:1273-1284.
11. Castellanos FX, Tannock R. Neuroscience of attention-deficit/hyperactivity disorder: the search for endophenotypes. Nature Rev Neurosci. 2002;3:617-628.
12. Krause KH, Dresel SK, Krause J, la Fougeré C, Ackenheil M. The dopamine transporter and neuroimaging in attention deficit hyperactivity disorder. Neurosci Biobehav Rev. 2003;27:605-613.
13. Spencer TJ, Biederman J, Madras BK, et al. In vivo neuroreceptor imaging in attention-deficit/hyperactivity disorder: a focus on the dopamine transporter. Biol Psychiatry. 2005;57:1293-1300.
14. Chen CK, Chen SI, Mill J, et al. The dopamine transporter gene is associated with attention deficit hyperactivity disorder in a Taiwanese sample. Mol Psychiatry. 2005;8:393-396.
15. Cook EH Jr, Stein MA, Krasowski MD, et al. Association of attention-deficit disorder and the dopamine transporter gene. Am J Hum Genet. 1995;56:993-998.
16. Cornish KM, Manly T, Savage R, et al. Association of the dopamine transporter (DAT1) 10/10-repeat genotype with ADHD symptoms and response inhibition in a general population sample. Mol Psychiatry. 2005;10:686-698.
17. Faraone SV, Perlis RH, Doyle AE, et al. Molecular genetics of attention-deficit/hyperactivity disorder. Biol Psychiatry. 2005;57:1313-1323.
18. Gill M, Daly G, Heron S, Hawi Z, Fitzgerald M. Confirmation of association between attention deficit hyperactivity disorder and a dopamine transporter polymorphism. Mol Psychiatry. 1997;2:311-313.
19. Waldman ID, Rowe DC, Abramowitz A, et al. Association and linkage of the dopamine transporter gene and attention-deficit hyperactivity disorder in children: Heterogeneity owing to diagnostic subtype and severity. Am J Hum Genet. 1998;63:1767-1776.
20. Dougherty DD, Bonab AA, Spencer T, Rauch SL, Madras BK, Fischman AJ. Dopamine transporter availability in patients with attention deficit hyperactivity disorder. Lancet. 1999;354:2132-2133.
21. Dresel SH, Kung MP, Plossl K, Meegalla SK, Kung HF. Pharmacological effects of dopaminergic drugs on in vivo binding of [99mTc]TRODAT-1 to the central dopamine transporters in rats. Eur J Nucl Med. 1998;25:31-39.
22. Dresel S, Krause J, Krause KH, et al. Attention deficit hyperactivity disorder: binding of [99mTc]TRODAT-1 to the dopamine transporter before and after methylphenidate treatment. Eur J Nucl Med. 2000;27:1518-1524.
23. Krause KH, Dresel SH, Krause J, Kung HF, Tatsch K. Increased striatal dopamine transporter in adult patients with attention deficit hyperactivity disorder as measured by single photon emission computed tomography. Neurosci Lett. 2000;285:107-110.
24. Cheon KA, Ryu YH, Kim JW, Cho DY. The homozygosity for 10-repeat allele at dopamine transporter gene and dopamine transporter density in Korean children with attention deficit hyperactivity disorder: relating to treatment response to methylphenidate. Eur Neuropsychopharmacol. 2005;15:95-101.
25. Al Younis ICH. Attention deficit hyperactivity disorder: neuroimaging before and after treatment with methylphenidate in children. J Nucl Med. 2002;43(suppl 5):347P.
26. van Dyck CH, Quinlan DM, Cretella LM, et al. Unaltered dopamine transporter availability in adult attention deficit hyperactivity disorder. Am J Psychiatry. 2002;159:309-312.
27. Vles JS, Feron FJ, Hendriksen JG, Jolles J, van Kroonenburgh MJ, Weber WE. Methylphenidate down-regulates the dopamine receptor and transporter system in children with attention deficit hyperkinetic disorder (ADHD). Neuropediatrics. 2003;34:77-80.
28. Volkow ND, Wang GJ, Fowler JS, et al. Dopamine transporter occupancies in the human brain induced by therapeutic doses of oral methylphenidate. Am J Psychiatry. 1998;155:1325-1331.
29. Krause J, la Fougere C, Krause KH, Ackenhel M, Dresel SH. Influence of striatal dopamine transporter availability on the response to methylphenidate in adult patients with ADHD. Eur Arch Psychiatry Clin Neurosci. 2005;255:428-431.
30. Volkow ND, Ding YS, Fowler JS, et al. Dopamine transporters decrease with age. J Nucl Med. 1996;37:554-559.
31. Krause KH, Dresel SH, Krause J, Kung HF, Tatsch K, Ackenhel M. Stimulant-like action of nicotine on striatal dopamine transporter in the brain of adults with attention deficit hyperactivity disorder. Int J Neuropsychopharmacol. 2002;5:111-113.
32. Court JA, Lloyd S, Thomas N, et al. Dopamine and nicotinic receptor binding and the levels of dopamine and homovanillic acid in human brain related to tobacco use. Neuroscience. 1998;87:63-78.
33. Staley JK, Krishnan-Sarin S, Zoghbi S, et al. Sex differences in [123I]beta-CIT SPECT measures of dopamine and serotonin transporter availability in healthy smokers and nonsmokers. Synapse. 2001;41:275-284.
34. Faraone SV, Spencer T, Aleardi M, Pagano C, Biederman J. Meta-analysis of the efficacy of methylphenidate for treating adult attention-deficit/hyperactivity disorder. J Clin Psychopharmacol. 2004;24:24-29.
35. Ferber L v, Lehmkuhl G, Köster I, et al. Methylphenidatgebrauch in Deutschland: Versichertenbezogene epidemiologische Studie über die Entwicklung von 1998 bis 2000. Dtsch Arztebl. 2003;100:A41-46.
36. Michelson D, Adler L, Spencer T, et al. Atomoxetine in adults with ADHD: two randomized, placebo-controlled studies. Biol Psychiatry. 2003;53:112-120.
37. Stahl SM. Neurotransmission of cognition, part 1. Dopamine is a hitchhiker in frontal cortex: noradrenaline transporters regulate dopamine. J Clin Psychiatry. 2003;64:4-5.