The influence of atypical antipsychotic drugs on sexual function

Marek J Just
Department of General and Endocrine Surgery, Piekary Medical Centre, Piekary Slaskie, Poland

Abstract: Human sexuality is contingent upon many biological and psychological factors. Such factors include sexual drive (libido), physiological arousal (lubrication/erection), orgasm, and ejaculation, as well as maintaining normal menstrual cycle. The assessment of sexual dysfunction can be difficult due to the intimate nature of the problem and patients’ unwillingness to discuss it. Also, the problem of dysfunction is often overlooked by doctors. Atypical antipsychotic treatment is a key component of mental disorders’ treatment algorithms recommended by the National Institute of Health and Clinical Excellence, the American Psychiatric Association, and the British Society for Psychopharmacology. The relationship between atypical antipsychotic drugs and sexual dysfunction is mediated in part by antipsychotic blockade of pituitary dopamine D2 receptors increasing prolactin secretion, although direct correlations have not been established between raised prolactin levels and clinical symptoms. Variety of mechanisms are likely to contribute to antipsychotic-related sexual dysfunction, including hyperprolactinemia, sedation, and antagonism of a number of neurotransmitter receptors (α-adrenergic, dopaminergic, histaminic, and muscarinic). Maintaining normal sexual function in people treated for mental disorders can affect their quality of life, mood, self-esteem, attitude toward taking medication, and compliance during therapy.

Keywords: schizophrenia, galactorrhea, hyperprolactinemia, mood disorders, anorgasmia

Introduction

Sexual dysfunction is a common problem in people with mood disorders,1 schizophrenia, and other psychotic disorders, with reported prevalence rates of 30%–80% in women and 45%–80% in men.2–4 In clinical practice, sexual disorders are found as a complication in people treated with antipsychotics. The prevalence rate of sexual disorders in men treated with antipsychotics is estimated to be 54% (Figure 1). Among them, 38% have problems in achieving erection, 42% with maintaining erection, at least 19% have ejaculation disorders (reduced volume or no ejaculation), and 58% have reduced orgasm intensity. Furthermore, in men, priapism, galactorrhea (3%), and gynecomastia (6%) may also be observed. Sexual disorders are found in about 30% of women treated with antipsychotics. According to the results obtained by different authors, reduced orgasm quality (33%), reduced ability to reach orgasm (22%), and pain during orgasm (7%) were the most common problems in women treated with antipsychotics. However, hyperprolacitnemia, galactorrhea (5%–19%), and breast swelling (3%) are more frequently found in women than in men. Furthermore, irregular (78%) or complete suppression of menstrual cycles (22%–50%) was observed in women.3 In people receiving psychiatric treatment, the possibility of other concurring factors should also be considered (chronic metabolic diseases and other medications) that can affect sexual function. Sexual dysfunction can also be related to thyroid disorders.6 The concurrence of thyroid disease in mentally ill
Patients may significantly delay the onset of pharmacological treatment, including treatment with antipsychotic drugs, and delay the use of slow transcranial magnetic stimulation and electroconvulsive therapy. Thus, a detailed medical history to confirm or exclude sexual disorders before commencing psychiatric treatment is extremely important. The sexual function of the patient should therefore be evaluated in the premorbid period, during the period from illness onset to treatment initiation, after treatment initiation, and also during exacerbation of mental disease and remission.

Atypical antipsychotics (AAPs) are recommended for the treatment of schizophrenia and other psychotic disorders, including those resulting from organic diseases. These drugs have a more extensive influence on the range of schizophrenia symptoms (deficits, affective, and cognitive) than typical antipsychotic drugs. Moreover, AAPs rarely cause side effects, such as neuroleptic malignant syndrome, versus typical antipsychotic drugs. AAPs may be considered as atypical when characterized by Melzert’s constant (their affinity for 5-HT<sub>2A</sub> and dopamine D<sub>2</sub> receptors [D<sub>2</sub>R] should exceed or be equal to 1.12). Another distinct feature of atypical drugs is the D<sub>2</sub>R-blocking time, which is directly correlated to the dissociation constant (K<sub>d</sub>) of the ligand–receptor complex. AAP drugs form a ligand–receptor complex with a shorter lifespan than typical antipsychotics. Furthermore, the values of their dissociation constant are lower than K<sub>d</sub> for dopamine. AAPs are also used in the treatment of bipolar disorder due to their mood stabilizing properties. They also enhance the antidepressant influence of drugs that are used in the treatment of other affective diseases such as atypical depression, drug-resistant depression, and depression resulting from organic disease.

AAPs can affect sexual function in the following ways: (1) through their specific central effect of blocking dopaminergic receptors in the hypothalamus, contributing to hyperprolactinemia, galactorrhea, menstruation, or erection disorders and reduced libido; (2) through their peripheral effect of blocking α-adrenergic receptors, responsible for dilation of the arteries in the penis; (3) through an unspecified central sedative effect leading to reduced sexual activity.

**The influence of AAPs on the central nervous system**

The influence of AAPs on sexual function is related to the activation of neurotransmitters such as dopamine and serotonin in the central nervous system. This process involves the brain’s dopaminergic and serotonergic structures.

**The influence of dopamine on sexual function**

The dopaminergic structures involved in sexual motivation include the nucleus accumbens, which is a part of the...
in the blood are 10–20 ng/mL in men and 10–25 ng/mL in women.\textsuperscript{21} PRL secretion increases during sleep and is at its highest during the rapid eye movement stage sleep. PRL levels increase to 30 ng/mL between 4 am and 6 am. Current research has revealed that galactorrhea, impotence, azospermia, and lack of libido are observed when PRL levels are $>60$ ng/mL.\textsuperscript{22} Hyperprolactinemia can also lead to reduced blood testosterone levels and can indirectly contribute to reduced sexual activity.\textsuperscript{18,20} According to the research, 80\% of men with PRL blood levels $>50$ ng/mL had reduced libido and suffered from impotence.\textsuperscript{18,20} AAPs as a group cause less hyperprolactinemia than conventional antipsychotics; yet, there is considerable variation among specific drugs.\textsuperscript{9} Of the AAPs, risperidone, amisulpiride, and zotepine have the strongest stimulating effect on PRL synthesis. Asmal et al\textsuperscript{23} showed in their study that PRL levels were increased at the beginning of treatment and were significantly higher in patients treated with risperidone. The average increase in PRL levels was higher in women than in men. However, Suzuki et al\textsuperscript{24} showed that the highest PRL levels (higher in the group of studied women vs men) were measured in the blood of patients treated with risperidone or olanzapine as monotherapy, versus patients taking aripiprazole, quetiapine, and perospirone combined. Furthermore, Jeong et al\textsuperscript{25} found that increased PRL levels during treatment with risperidone was accompanied by sexual dysfunction, especially problems with sexual arousal and penile erection, which were measured by using the Arizona Sexual Experience Scale.\textsuperscript{26} The symptoms of sexual dysfunction disappeared after risperidone withdrawal and the commencement of aripiprazole. The influence of risperidone on the synthesis of PRL and sexual function are combined with its strong affinity for the D\textsubscript{2} receptors and its simultaneous effect on the 5-HT\textsubscript{2} receptors.

Peuskens et al\textsuperscript{27} found that among men receiving risperidone at a dose of 1–16 mg/day, 4.2\%–17.7\% had problems with erection and 3.6\%–17.7\% had a problem with ejaculation. The authors suggested that hyperprolactinemia could be the cause of these dysfunctions. Nevertheless, Jummani and Coffey\textsuperscript{27} observed mild hyperprolactinemia (11–21 ng/mL), premature ejaculation, and impotence in patients treated with risperidone at a dose of 3.5 mg/day. Other AAPs such as clozapine and quetiapine do not display an influence on PRL secretion.\textsuperscript{23} However, Serretti and Chiesa\textsuperscript{28} showed that sexual dysfunction (desire, arousal, and orgasm dysfunction) occurred during treatment with quetiapine and aripiprazole, although PRL blood levels were within the normal range. These observations indicate that other factors such as chronic stress and metabolic diseases (hypertension and diabetes) can be responsible for sexual dysfunction.

PRL influence on sexual function

Antipsychotics have a D\textsubscript{2}-blocking effect and can therefore increase PRL secretion. The normal reference values of PRL in the blood are 10–20 ng/mL in men and 10–25 ng/mL in women.\textsuperscript{21} PRL secretion increases during sleep and is at its highest during the rapid eye movement stage sleep. PRL levels increase to 30 ng/mL between 4 am and 6 am. Current research has revealed that galactorrhea, impotence, azospermia, and lack of libido are observed when PRL levels are $>60$ ng/mL.\textsuperscript{22} Hyperprolactinemia can also lead to reduced blood testosterone levels and can indirectly contribute to reduced sexual activity.\textsuperscript{18,20} According to the research, 80\% of men with PRL blood levels $>50$ ng/mL had reduced libido and suffered from impotence.\textsuperscript{18,20} AAPs as a group cause less hyperprolactinemia than conventional antipsychotics; yet, there is considerable variation among specific drugs.\textsuperscript{9} Of the AAPs, risperidone, amisulpiride, and zotepine have the strongest stimulating effect on PRL synthesis. Asmal et al\textsuperscript{23} showed in their study that PRL levels were increased at the beginning of treatment and were significantly higher in patients treated with risperidone. The average increase in PRL levels was higher in women than in men. However, Suzuki et al\textsuperscript{24} showed that the highest PRL levels (higher in the group of studied women vs men) were measured in the blood of patients treated with risperidone or olanzapine as monotherapy, versus patients taking aripiprazole, quetiapine, and perospirone combined. Furthermore, Jeong et al\textsuperscript{25} found that increased PRL levels during treatment with risperidone was accompanied by sexual dysfunction, especially problems with sexual arousal and penile erection, which were measured by using the Arizona Sexual Experience Scale.\textsuperscript{26} The symptoms of sexual dysfunction disappeared after risperidone withdrawal and the commencement of aripiprazole. The influence of risperidone on the synthesis of PRL and sexual function are combined with its strong affinity for the D\textsubscript{2} receptors and its simultaneous effect on the 5-HT\textsubscript{2} receptors.

Peuskens et al\textsuperscript{27} found that among men receiving risperidone at a dose of 1–16 mg/day, 4.2\%–17.7\% had problems with erection and 3.6\%–17.7\% had a problem with ejaculation. The authors suggested that hyperprolactinemia could be the cause of these dysfunctions. Nevertheless, Jummani and Coffey\textsuperscript{27} observed mild hyperprolactinemia (11–21 ng/mL), premature ejaculation, and impotence in patients treated with risperidone at a dose of 3.5 mg/day. Other AAPs such as clozapine and quetiapine do not display an influence on PRL secretion.\textsuperscript{23} However, Serretti and Chiesa\textsuperscript{28} showed that sexual dysfunction (desire, arousal, and orgasm dysfunction) occurred during treatment with quetiapine and aripiprazole, although PRL blood levels were within the normal range. These observations indicate that other factors such as chronic stress and metabolic diseases (hypertension and diabetes) can be responsible for sexual dysfunction.

PRL influence on sexual function

Antipsychotics have a D\textsubscript{2}-blocking effect and can therefore increase PRL secretion. The normal reference values of PRL
Olanzapine is another AAP that influences PRL levels and sexual function. This results in olanzapine binding with a wide range of dopaminergic (D₁, D₂, D₃, D₄) and serotonergic (5-HT₂A, 5-HT₅, 5-HT₆, 5-HT₉) receptors. According to many studies, olanzapine is characterized by having a lower and more selective affinity for dopaminergic receptors (mainly the mesolimbic pathway) than risperidone. Konarzewska et al. in their own research found a temporary increase in PRL levels in patients treated with olanzapine. This increase was lower in the group of patients treated with olanzapine versus patients treated with risperidone. Similar results were obtained by Kim et al. who observed a lower increase in PRL levels and sexual dysfunction following the change in treatment from risperidone to olanzapine. Baggaley noticed that risperidone showed a significantly higher influence on the development of sexual dysfunction versus olanzapine, quetiapine, and aripiprazole. There are also reports concerning sexual dysfunction and treatment with sertindole, in which orgasm without ejaculation was observed, which was associated with the peripheral α₁-adrenergic antagonism of this drug, and also orgasm with ejaculation of reduced ejaculate volume. Sertindole is a second-generation antipsychotic drug with a high affinity for dopaminergic D₂, serotonergic 5-HT₂A, 5-HT₅, and α₁-adrenergic receptors and a low affinity for other receptors. Muscatello et al., Lewis et al., and Jukić et al. in their studies found that sertindole can cause sexual dysfunction independent of increased PRL levels, more commonly in men than in women.

**The influence of serotonin and noradrenaline on sexual function**

After dopamine, serotonin is the second most important neurotransmitter to influence sexual function. Serotonin mainly inhibits sexual function by stimulating postsynaptic 5-HT₂A and 5-HT₅ receptors, while the stimulation of presynaptic 5-HT₁A autoreceptors increases sexual activity, reducing serotonin secretion from the nerve termini. Furthermore, reduction of the negative symptoms of schizophrenia is combined with the serotonergic system, particularly with antagonism of the 5-HT₂A and/or 5-HT₁A receptors. By blocking these receptors located on dopaminergic neurons, dopaminergic transmission in the mesocortical pathway is intensified, and as a result, cognitive (such as focusing attention and working memory) and executive functions are improved. On the other hand, an increase in transmission in the nigrostriatal pathway reduces the risk of neuroleptic malignant syndrome through disinhibition. Noradrenaline, another neurotransmitter, increases the ability for arousal through its influence on central receptors and inhibits erection by binding with peripheral α₁ receptors.

**The peripheral influence of AAP drugs**

The adrenergic system is one of the peripheral mechanisms that have an influence on penile erection. It is accompanied by the relative functional prevalence of the parasympathetic (β₂-adrenergic activation) over the sympathetic (α₁-adrenergic inhibition) system, necessary to dilate penile arteries and for blood to flow into the cavernous bodies of the penis. One related sexual dysfunction is priapism. In this disorder, a prolonged, painful erection of the penis occurs, which is unrelated to sexual arousal or stimulation. Priapism can be caused by the blocking of α₁-adrenergic receptors by AAP drugs. According to numerous studies, most of the AAP drugs, such as olanzapine, risperidone, and amisulpiride can cause priapism. It is worth noting that olanzapine can cause both reversible and irreversible priapism. The symptoms of reversible priapism after 10 days of olanzapine treatment at a dose of 10 mg/day were observed by Doufik et al. Priapism symptoms disappeared when the treatment was changed from olanzapine to amisulpiride at a dose of 400 mg/day. Furthermore, reversible priapism with olanzapine was also described by Penaskovic et al. This sexual dysfunction appeared after 7 days of olanzapine treatment at a dose of 15 mg/day. However, Jagadheesan et al. described irreversible priapism after treatment with olanzapine. The symptoms of priapism appeared on the sixth day of treatment with olanzapine at a dose of 5 mg/day. Reversible priapism was also observed after treatment with other neuroleptics such as risperidone and ziprasidone. Ginory and Nguyen observed priapism after a month’s therapy with risperidone at a dose of 6 mg/day. Furthermore, Cruzado and Vallejos described priapism that developed during monotherapy with risperidone at a dose of 3 mg/day, after 3 years of using the drug. Due to this adverse effect, risperidone was withdrawn and quetiapine at a dose of 450 mg/day was prescribed for the patient and the side effect disappeared. Karamustafalioglu et al. observed signs of priapism caused by ziprasidone at a dose of 40 mg/day on the seventh day of treatment. The patient felt uncomfortable and stated that the condition had lasted for nearly 8 hours and was not associated with sexual stimulation or desire. Recurring priapism during treatment with ziprasidone at a total daily dose of 160 mg (40 mg bid + 80 mg qhs) was described by Kaufman et al. and with a dose of 140 mg/day (ziprasidone with food, 80 mg at bedtime...
of hyperprolactinemia. Also, the blockade of alpha-1 and
achieve erection and ejaculation, directly and as a result
a factor in reducing sexual drive as well as the ability to
indicates the important role of dopaminergic blockade as
adverse effects may have problems with compliance dur
sexual dysfunction. Patients who do not tolerate these
problems with compliance during
reduction of sexual activity must be considered. When introduc-
ing AAP treatment, knowledge of the pharmacokinetics of the
prescribed drug is important. It is recommended to administer
the whole dose of the drug in the evening, yet sufficiently
early enough so that the peak concentration of the drug in the
blood decreases at the moment of falling asleep (medicine
can be administered even 5–6 hours before sleep). By
dosing olanzapine this way, the sedative effect and adverse
influence on sexual function can be minimized. During
treatment with clozapine or quetiapine, these drugs should
be administered in two daily doses due to their short half-life
and in order to avoid sedative effects during the day. Also,
the evening dose of the drug should be higher than that given
in the morning. Aripiprazole, sertindole, amisulpiride, and
ziprasidone do not show sedative effects as they do not have
any antihistaminergic and anticholinergic effects. Therefore,
they do not have a negative effect on sexual function.

The sedative influence of AAP drugs leading to reduced sexual activity
AAP drugs can adversely influence sexual function due to
their sedative effect. This process involves the blocking
of dopaminergic and 5-HT2A serotonergic receptors,
antihistaminergic effects (blocking of H1 receptors), and an
anticholinergic effect (blocking of muscarinic receptors).
During optimization of AAP treatment with regard to
their antipsychotic influence, sedative symptoms leading to
reduced sexual activity must be considered. When introduc-
ing AAP treatment, knowledge of the pharmacokinetics of the
prescribed drug is important. It is recommended to administer
the whole dose of the drug in the evening, yet sufficiently
early enough so that the peak concentration of the drug in the
blood decreases at the moment of falling asleep (medicine
can be administered even 5–6 hours before sleep).53,54 By
dosing olanzapine this way, the sedative effect and adverse
influence on sexual function can be minimized. During
treatment with clozapine or quetiapine, these drugs should
be administered in two daily doses due to their short half-life
and in order to avoid sedative effects during the day. Also,
the evening dose of the drug should be higher than that given
in the morning.55,56 Aripiprazole, sertindole, amisulpiride, and
ziprasidone do not show sedative effects as they do not have
any antihistaminergic and anticholinergic effects. Therefore,
they do not have a negative effect on sexual function.

Conclusion
Problems during therapy with AAP drugs include a lack of
feedback from patients about the adverse effects caused
by the drugs, particularly when they are combined with
sexual dysfunction. Patients who do not tolerate these
adverse effects may have problems with compliance dur-
ing treatment. In conclusion, current knowledge about the
mechanisms of sexual dysfunction caused by AAP drugs
indicates the important role of dopaminergic blockade as
a factor in reducing sexual drive as well as the ability to
achieve erection and ejaculation, directly and as a result of
hyperprolactinemia. Also, the blockade of alpha-1 and
probably \( \text{z2} \)-adrenergic and muscarinic receptors is related
to some disorders (impotence, priapism), although its final
effects seem to depend on the location (ie, central, peripheral)
of the target receptors to a larger extent, as well as on the
overlapping, “modulating” receptor effects of a drug. The
anti-serotonergic properties of AAP drugs can to some extent
prevent sexual dysfunction.

Acknowledgments
The work was carried out at the Department of General
Surgery, Municipal Hospital in Piekar Slaskie, Poland.
Professional language editing has been performed by a native
English speaker – Peter Kosmider-Jones. This research has
received no specific grant from any funding agency in the
public, commercial, or not-for-profit sectors.

Author contribution
MJJ contributed in the conception of the study, interpretation,
preparation of manuscript, editing, revising, and recruitment.

Disclosure
The author reports no conflicts of interest in this work.

References
1. Grover S, Ghosh A, Sarkar S, Chakrabarti S, Avasthi A. Sexual dysfunc-
tion in clinically stable patients with bipolar disorder receiving lithium.
2. Hamilton LD, Meston CM. Chronic stress and sexual function in
3. Agno A. On the intricate relationship between sexual motivation and
4. Boin AC, Nozoe KT, Polesel DN, Andersen ML, Tufik S. The poten-
tial role of sleep in sexual dysfunction in patients with schizophrenia.
5. Kopecek M, Bares M, Svarc J, Dockery C, Horacek J. Hyperprolactine-
419–422.
6. Ogłodęk E, Szota A, Araszkiewicz A. Electroconvulsive therapy in a
patient with drug-resistant depression and thyroid hormone imbalance.
7. Ogłodęk E, Wiczkowski A, Sieron A, Bilska-Urbani A, Moś D. The
effect of extremely low-frequency magnetic fields on the morphology
757–763.
8. Kristensen D, Hageman I, Bauer J, Jorgensen MB, Correll CU. Antipsy-
chotic polypharmacy in a treatment-refractory schizophrenia population
receiving adjunctive treatment with electroconvulsive therapy. J ECT.
associated with side effects in schizophrenia. J Med Econ. 2014;17(12):
853–861.
10. Connolly A, Taylor D. Factors associated with non evidence-based
prescribing of antipsychotics. Ther Adv Psychopharmacol. 2014;4(6):
247–256.
malignant syndrome after 10 years of treatment. Aust NZ J Psychiatry.