## **MODERN INDICATIONS FOR THE USE OF OPIPRAMOL**

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#### **SUMMARY**

Opipramol is considered as a pharmacological agent that does not fit the classification taking into account the division of antidepressants, antipsychotics and anxiolytics. It has a structure related to tricyclic antidepressants but it has a different mechanism of action, i.e. binding to sigma1 and to sigma2 sites. It has been regarded as an effective drug in general anxiety disorders together with other agents like SSRI's, SNRI's, buspirone and pregabalin for many years. It can however also be indicated in other conditions, e.g. it may be used as a premedication in the evening prior to surgery, positive results are also observed in psychopharmacological treatment with opipramol in somatoform disorders, symptoms of depression can be significantly reduced in the climacteric syndrome. The latest data from literature present also certain dangers and side effects, which may result due to opipramol administration. Mania may be induced not only in bipolar patients treated with opipramol, but it can be an adverse drug reaction in generalized anxiety disorder. This analysis shows however that opipramol is an important drug still very useful in different clinical conditions.

Key words: opipramol – depression - anxiety disorders - somatoform disorders

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### **INTRODUCTION**

Depressive and anxiety disorders are nowadays treated in different ways. The most important methods in this field remain pharmacology, psychotherapy or a combination of those two methods (Krysta 2014, Klasik 2012). Antidepressants are the most frequently prescribed drugs in general anxiety disorder (GAD) (Reinhold 2015), or in panic disorder (PD). In that condition sometimes, for a limited period of time, benzodiapines are used as an additional drug (Locke 2015). Agents like mirtazapine and SSRI's are effective in the treatment of PTSD (Schneier 2015). Antidepressants in combination with other medication are also beneficial in somatoform disorders (Kleinstauber 2014). A thymoleptic drug which has an anti-anxiety and weak antidepressive action, used mainly in the pharmacological treatment of general anxiety and somatoform disorders is opipramol (Krupka-Matuszczyk 2012).

## INDICATIONS FOR THE USE OF OPIPRAMOL

Opipramol was introduced to the treatment in the 60s. It was synthesised as one of the tricyclics similar to previously enteredimipramine, however, clinical trials have not allowed to qualify opipramol in the group of antidepressants. Nevertheless, in view of these dativeaction, opipramol had been extensively used, especially in ambulatory practice (Jaracz 2003). It does not inhibit

the neuronal uptake of norepinephrine and/or serotonin (Müller 2004). It has a different mechanism of action. It binds to sites sigma1 and sigma2, and no reuptakeinhibiting properties are observed. This drug is widely prescribed in many European countries, like Germany and it has a strong anxiolytic efficacy the treatment of GAD (Möller 2001). It is suggested that effects at sigma2 sites are involved in the anxiolytic properties of opipramol (Volz 2004). According to Murawiec (2012) opipramol does not fit the classification taking into account the division of antidepressants, antipsychotics and anxiolytics. Therefore it requires a separate discussion as it is relatively poorly explored. It has been considered as a potent drug in general anxiety disorders for many years along with SSRI's, SNRI's, buspirone and pregabalin, as it has anxiolytic efficacy superior to placebo (Boerner 2007a, Boerner 2007b, Gale 2007). The duration of treatment depends on the indication and the clinical condition of the patient. The dose Proposed by Prusiński (2003a) is150 mg for several months (of course if there are no major side effects). According to Indian authors (Mohapatra 2013) the recommended dose for adults is 50 mg in the morning, 50mg in the afternoon, and 100 mg in the evening. It is proposed that after achieving an improvement, the dose is reduced to reach the maintenance dose of 50 mg 2 times/day. But if necessary, the dose can be also raised to 100 mg 3 times daily. It should not be discontinued quickly, because its sudden withdrawal, especially if higher doses, it can cause nausea, vomiting, anorexia, headache and insom-

nia. In the elderly it is advisable to use lower dosesabout1/3 oreven1/2. The most common mistake made during pharmacotherapy in the outpatient is that opipramol is used in very small doses and the drug is takien over too short a period (Prusiński 2003b). Except from the traditional form of the drug introduced in the 60s, a search for new pharmacological options has been made. The appearance of sustained release opipramol dihydrochloride matrix tablets turned out to be a new attitude in the treatment of depression and anxiety (Gönüllü 2006). Opipramol can also be indicated in other conditions, e.g. it may be used as a premedication in the evening prior to surgery (Hueppe 2011, Gerlach 2002). On the other hand in case of surgical interventions, a modification of opipramol therapy can be necessary in a perioperative period in order to avoid interactions (Redel 2013). Positive results are also observed in the treatment with opipramol in somatoform disorders (Freyberger 1998, Volz 2004) and the specific anxiolytic effect of opipramol include in particular somatic symptoms of anxiety (Krupka-Matuszczyk 2013, Święcicki 2013). Symptoms of depression can be significantly reduced in the climacteric syndrome (van Lith & Motke 1983). Opipramol, is not a hypnotic drug, but it promotes sleep, so it can be used widely in the treatment of some forms of insomnia (Müller 1998). Attempts are also made to use opipramol in posttraumatic stress syndrome (PTSD) and gastrointestinal dyskinesias (Vademecumleków 1997). It can have a positive effect in a so called "drug rebound headache", which refers to headaches occurring every day in patients with migraine and tension headaches as a consequence of taking analgesics or ergotamine every day. Opipramol turned out to be effective, when taken by increments from half tablet up to three in 6 days, and the treatment was then continued for 4-6 weeks (Prusinski 1999). Some possible indications for opipramol are still the matter of research. Animal studies suggest that there could be a possible clinical application in peptic ulcer patients with a comorbid depression (Dursun 2009). Opipramol, as potent sigma ligand, could theoretically be an anti-ischemic agent helpful in neurological conditions (Rao 1990).

## **TOLERANCE AND SIDE EFFECTS**

As already mentioned above, opipramol has found wide applications, which resulted not only from its effectiveness, but also from the tolerance by patients and relatively few side effects (Prusiński 2003b). Fatal poisonings with this agent are very rare (Skopp 1996). Anyway the data from literature draw the attention to few, but possible side effects. Mania may be induced in some cases not only in bipolar patients treated with opipramol (Firoz 2015), but it can be an adverse drug reaction in generalized anxiety disorder (Kar 2015). Although opipramol is a drug prescribed orally, some rare attempts to its parenteral use have ben described. Barańska-Rybak et al. (2014) published a case report of recurrent abscesses with a coexisting fever following the injection of a mixture of tramadol, opipramol, and clonazepam. Another interesting case report describes a patient with unipolar depressive treated with maprotiline and opipramol with bioptically proven severe acute hepatitis caused by those two antidepressants (Braun 1998). In spite of those rare adverse events reported in the literature, opipramol is generally regarded as a safe pharmacological agent useful in different clinical conditions.

# CONCLUSIONS

Opipramol is an efficient drug in general anxiety disorders along with SSRI's, SNRI's, buspirone and pregabalin having an efficacy better than placebo. It can also be indicated in other conditions, e.g. it may be used as a premedication in the evening prior to surgery, positive results are also observed in psychopharmacological treatment with opipramol in somatoform disorders, symptoms of depression can be significantly reduced in the climacteric syndrome. It is a safe drug and has few side effects. A few reports show that e.g. mania may be induced not only in bipolar patients treated with opipramol, but it can have an adverse drug reaction in generalized anxiety disorder. The above analysis shows that opipramol is an important drug with a wide range of indications.

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# References

- 1. Barańska-Rybak W, Błażewicz I, Kąkol M, Roter M & Nowicki R: Cutaneous manifestations of injectable drug use: hidden secrets. Cutis 2014; 93:185-187.
- 2. Boerner RJ: Generalized anxiety disorder (GAD) diagnosis and therapy. Med Monatsschr Pharm 2007a; 30:401-408.
- 3. Boerner RJ: Pathologic worry. MMW Fortschrit Med 2007b; 14:46-49.
- Braun JS, Geiger R, Wehner H, Schaffer S & Berger M: Hepatitis caused by antidepressive therapy with maprotiline and opipramol. Pharmacopsychiatry 1998; 31:152-155.
- Dursun H, Albayrak F, Bilici M, Koc F, Alp HH, Candar T et al.: Gastroprotective and antioxidant effects of opipramol on indomethacin-induced ulcers in rats. Yakugaku Zasshi 2009; 129:861-869.
- Firoz K, Khaleel A, Rajmohan V, Kumar M & Raghuram T: Mania induced by opipramol. Indian J Psychol Med 2015; 37:96-98.
- Freyberger HJ, Grabe HJ, Maier W, Janca A & Stoll KD: Opipramol (Insidon) in the treatment of somatoform disorders. Fortschr Neurol Psychiatr 1998; 66 Suppl 1:S25-30.

- 8. Gale CK & Millichamp J: Generalised anxiety disorder. BMJ Clin Evid 2007; 20: pii: 1002.
- 9. Gerlach K, Uhlig T, Plathof J, Klassen A, Stoll KD, Schmucker P et al.: Effects of opipramol as an evening anaesthesiologic premedication. Neuropsychobiology 2002; 46:161-166.
- 10. Gönüllü U, Uner M, Yener G & Altinkurt T: Introduction of sustained release opipramoldihydrochloride matrix tablets as a new approach in the treatment of depressive disorders. Int J Biomed Sci 2006; 2:337-343.
- 11. Hueppe M, Hartge D, Stoll KD, Ros A, Schmucker P & Gerlach K: Opipramol improves subjective quality of sleep the night prior to surgery: confirmatory testing of a double-blind, randomized clinical trial. Neuropsychobiology 2011; 64:24-31.
- 12. Jaracz J & Rybakowski J: Pharmacological and clinical properties of opipramol in the light o new studies. Farmakoter PsychiatNeurol 2003; 4:115-121.
- 13. Kar SK & Garg K: Mania, a probable adverse drug reaction of opipramol in a patient with generalized anxiety disorder. Aust N Z J Psychiatry 2015; In press, pii: 0004867415583337.
- 14. Klasik A, Krysta K & Krzystanek M: Impact of psychotherapy and antidepressive treatment on cognitive functions in patients treated for depression. Psychiatr Danub 2012; 24(Suppl 1):S130-134.
- 15. Kleinstauber M, Witthoft M, Steffanowski A, van Marwijk H, Hiller W & Lambert MJ: Pharmacological interventions for somatoform disorders in adults. Cochrane Database Syst Rev 2014. In press, 11:Cd010628. doi:10.1002/14651858.CD010628.pub2.
- 16. Krupka-Matuszczyk I: Opipramol: drug traditionally used in the treatment of anxiety and depression. Case studies. Rev Clin Neuropsychiatr 2012; 4:131-134.
- 17. Krysta K, Krzystanek M, Janas-Kozik M, Klasik A, Krupka-Matuszczyk I: Impact of pharmacological and psychological treatment methods of depressive and anxiety disorders on cognitive functioning. J Neural Transm 2014; In press, doi:10.1007/s00702-014-1282-3.
- 18. Krysta K: Safe and effective use of benzodiazepines in psychiatric disorders and other comorbid conditions. Psychiatria 2014; 11:9-14.
- 19. Leutgeb U: Opipramol as nighttime medication. Dtsch Arztebl Int 2009; 106:321; author reply 321-322.
- 20. Locke AB, Kirst N & Shultz CG: Diagnosis and management of generalized anxiety disorder and panic disorder in adults. Am Family Physician 2015; 91:617-624.
- 21. Mohapatra S, Rath NM, Agrawal A & Verma J: Opipramol: a novel drug. Delhi Psychiatry Journal 2013; 16:409–411.

- 22. Möller HJ, Volz HP, Reimann IW & Stoll KD: Opipramol for the treatment of generalized anxiety disorder: a placebo-controlled trial including an alprazolam-treated group. J Clin Psychopharmacol 2001; 21:59-65.
- 23. Müller WE & Siebert B: Opipramol im Vergleich zu anderen Therapeutika - Neuepharmakologische Data Fortschr Neur Psych 1998; 66(Suppl 1):9–12, 15.
- 24. Müller WE, Siebert B, Holoubek G & Gentsch C: Neuropharmacology of the anxiolytic drug opipramol, a sigma site ligand. Pharmacopsychiatry 2004; 37(Suppl 3):S189-197.
- 25. Murawiec S: Outsider's hard fate. The case of opipramol. Rev Clin Neuropsychiatr 2012; 1:40–45.
- 26. Prusiński A: Opipramol (Pramolan): renesans leku i jego aktualne zastosowanie. Przew Lek 2003a, 6:46-52.
- Prusiński A: Opipramol (Pramolan®) in neuroses therapy. Psychiatria w Praktyce Ogólnolekarskiej 2003b; 3:57–62.
- 28. Prusiński A: Drug rebound headaches. Neurol Neurochir Pol 1999; 32(Suppl 6):31-37.
- 29. Rao TS, Cler JA, Mick SJ, Ragan DM, Lanthorn TH, Contreras PC et al.: Opipramol, a potent sigma ligand, is an anti-ischemic agent: neurochemical evidence for an interaction with the N-methyl-D-aspartate receptor complex in vivo by cerebellar cGMP measurements. Neuropharmacology 1990; 29:1199-1204.
- 30. Redel A, Hommers LG, Kranke P, Schwemmer U & Prasser C: Modification of perioperative psychiatric drug therapy. Anasthesiol Intensivmed Notfallmed Schmerzther 2013; 48:10-17.
- Reinhold JA & Rickels K: Pharmacological treatment for generalized anxiety disorder in adults: an update. Expert Opin Pharmacother 2015; 16:1669-1681.
- 32. Schneier FR, Campeas R, Carcamo J, Glass A, Lewis-Fernandez R, Neria Y et al.: Combined mirtazapine and SSRI treatment of PTSD: A placebo-controlled trial. Depress Anxiety 2015; 32:570-579.
- 33. Skopp G, Miltner E & Aderjan R: Fatal poisoning with the antidepressive agent opipramol. Forensic Sci Int 1996; 77:45-51.
- 34. Święcicki Ł: Practical aspects of pharmacotherapy of anxiety is there a place for opipramol. Psychiatria 2013; 10:63-66.
- 35. Vademecum leków: Ośrodek Informacji Naukowej Polfa, Warszawa, 1997.
- 36. van Lith ND, Motke JC: Opipramol in the climacteric syndrome. A double-blind, placebo-controlled trial. Maturitas 1983; 5:17-23.
- 37. Volz HP: Somatoform disorders what must the general practitioner know? MMW Fortschr Med 2004; 146:27-28, 30.

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