

New data on Metaxalone (Skelaxin) and serotonin toxicity (serotonin syndrome): warning of potential for fatalities if combined with serotonin re-uptake inhibitors

Gillman, P. K. PsychoTropical Research, Bucasia, Queensland, Australia

Abstract

Evidence has emerged that **Metaxalone (Skelaxin)** is a weak monoamine oxidase inhibitor (MAOI) which when used in large doses may be sufficiently potent to induce **serotonin toxicity (ST)** (aka Serotonin Syndrome), but only if and when it is combined with a serotonin reuptake inhibitor (SRI). Metaxalone was introduced just over 50 years ago and is used widely for the treatment of musculoskeletal pain.

It was approved in the days when less extensive pharmacological data concerning drugs existed. Now, theoretical computational (*in silico*) techniques allow a degree of useful prediction of the likely receptor and enzyme targets of drugs (this *in silico* modelling has indicated metaxalone is likely to be a weak MAOI — see below). Also, it is an ‘oxazolidinone’, structurally and pharmacologically related to antibiotics like linezolid, which definitely do exhibit both MAOI potency, and also the clinical effect of inducing ST. That strongly suggests the wisdom of vigilance concerning the probability of ST with metaxalone.

As of June of 2018 several case reports have described what is probably severe ST in patients who have received metaxalone in combination with an SRI. The fact that such reports have only recently emerged may, at least partly, reflect the increasing knowledge and understanding of ST. The importance of such understanding has been highlighted recently by various incorrect and misleading warnings emanating from several drug regulatory agencies (WHO, FDA etc.) concerning ST, as well as continuing publication of ‘reviews’ which do not reflect current expert knowledge of those drug interactions that are capable of precipitating ST: a recent egregious example being the review by Werneke et al. {Werneke, 2016 #19895}. These misleading reviews and ‘official’ warnings include triptans and antiemetic drugs like ondansetron. The explication of why it is logical to dismiss ondansetron and triptans as suspects, but conclude that Metaxalone is ‘guilty’, contains valuable lessons in understanding pharmacology, toxicology and serotonin toxicity (1-3). The same applies to methylthionium (methylene blue). These lessons have, evidently, not yet gained general recognition despite all the computers in use, nor have the appropriate interaction warnings been successfully incorporated into clinical management protocols.

Keywords: Metaxalone, oxazolidinone antibiotics, linezolid, MAOIs, methylthioninium (methylene blue), moclobemide, SSRIs, SRIs, SNRIs, cyclobenzaprine, drug interactions, Serotonin Toxicity, Serotonin Syndrome, severe, fatal, warning.

Introduction and background

This revision (2018) updates my original warning from 2014. Perhaps we should not be too surprised that an understanding of the spectrum concept of **serotonin toxicity (ST)** mechanisms is continuing to provide useful elucidation of the probable mode of action of various drugs (2). The utility of such an understanding has been clearly illustrated by the story of methylene blue, dealt with extensively both on my website and in my peer reviewed publications (3). More recently its utility has been exemplified in the rebuttal of the warnings issued about triptans and ST (3-5) and about ondansetron and ST (6-10).

The above review (3) explained how an understanding of the potencies and mechanisms of ST, and their important interactions, leads to precise and reliable predictions. Almost the only possible explanation of **severe** ST involving usual therapeutic doses of drugs is the classic and potentially fatal interaction of an MAOI with an SRI.

This chain of events was started in Nov 2014 when Dr Pete Bentham alerted me to recent case reports suggesting severe ST involving metaxalone (11, 12).

The fact that metaxalone has been used for such a long time (more than 50 years) with no apparent reports of ST would at first seem to make ST unlikely: after all one might suppose it would surely have occurred and been reported somewhere? Maybe it has been and no-one has yet unearthed the reports? (which seems to be the case, see below) I have attempted to ascertain this, with limited success, it is not as easy as one might think. Such considerations notwithstanding, all this suggest that it is a weak MAOI, probably reversible, like moclobemide. Therefore, it is only when it is ingested in large doses and in combination with a serotonin reuptake inhibitor (SRI) that ST is exhibited.

That is precisely the same picture that we do indeed see with the reversible MAO-A selective drug moclobemide (13, 14). Some people tried using combinations of moclobemide and SSRIs, but it proved to be a rather delicate and brave balancing act which could (and did) easily go seriously amiss (15-19): there were deaths, some published, some only reported to me privately. Most people stopped doing it fairly quickly. I summed it up with this comment: 'The above data indicate clearly that such combinations represent a predictably risky strategy. Moclobemide, if combined with any drug exhibiting significant SRI potency, produces a risk of severe ST and also the possibility of fatalities even with 'therapeutic' doses.' (16).

I recap on this story because there are close parallels and lessons between moclobemide, linezolid and metaxalone whereby small

increases in dosage (or increases in blood levels due to pharmacokinetic interactions) may rapidly lead to catastrophic and fatal toxicity.

The current evidence about metaxalone provides a warning signal that is sufficiently strong that it mandates further investigation. One arm of this investigation should be the effort to ascertain whether other cases of toxicity with this drug involve typical ST manifestations and whether these have occurred only in the presence of (S)SRIs (which is what I would confidently predict). So far that would appear to be the case. It would be surprising if there were not a few more cases of severe or fatal ST from metaxalone overdoses in subjects on SRIs out there somewhere.

Case Reports: updated June 2018

These initial cases (11, 12) were brought to my attention by Dr Pete Bentham. My research since then has revealed further previously reported, but unrecognised, cases where metaxalone appears to be implicated in ST (20, 21). There is a further report from 2015, of 2 cases of probable ST involving ODs of metaxalone and therapeutic doses of an SRI (22), exactly as I predicted in my original warning a year ago. And since then (23).

There are also one or two post-mortem toxicology reports of suicides or deaths that involve metaxalone & SRIs. In such cases there is usually no information indicating symptoms before death, so establishing that such cases represent ST is impossible. Nevertheless, these do constitute another suspicious signal (24-26).

The Huska and Martini cases (20, 22) fall into that minuscule group of case reports about ST in the last decade which are actually useful. They both contain elements which immediately make one think it is likely to be severe ST involving an MAOI and an SRI. The patients presented acutely complaining of feverishness and shaking, temperatures rose rapidly, and they exhibited severe tremulousness, shivering, hyperreflexia, clonus, and muscle rigidity.

This is very much the classic picture of ST with dramatic and rapid onset dominated by neuro-muscular symptoms and resolving rapidly following cessation of medication (and the use of cyproheptadine). It is instructive to compare these cases with the cyclobenzaprine cases (see below) and the poor cases put forward by the FDA which were adduced to justify the idea of ST with ondansetron: those cases, quite apart from not meeting formal diagnostic criteria, just do not look like ST.

The other metaxalone cases described are all 'probable' or 'definite' ST and involved an SRI, except for one of the cases reported by Bosak. The investigation of this case involved steps to ascertain what drugs had been taken as part of the overdose, but as is so often the case, sources of information were limited and of uncertain reliability. Although drug screening failed to find SRI drugs, it remains entirely possible the patient

had ingested an SRI prior to the overdose (viz. I am sure the screening did not include chlorpheniramine, a potent SRI available ‘OTC’).

These cases provide a very strong indication that metaxalone is an MAOI. The detailed reasoning that justifies this statement is contained in the other extensive information about ST, both in my peer-reviewed papers, and on my website. In short, it is the same as the picture we see with moclobemide. Overdoses of moclobemide alone do not cause serotonergic symptoms or ST, but when it is mixed with even small doses of SRIs serious cases of ST are frequent, and fatal cases are well-documented. Thus, we would expect no signs of ST with metaxalone by itself, but we would expect it if large doses of metaxalone are combined with therapeutic doses of an SRI.

That is precisely what the above cases illustrate.

New data on MAOI potency of Metaxalone

Another avenue of investigation is to establish more precisely the likelihood of it being a significant MAO inhibitor. This can be done with computer modelling techniques, and there is good reason for pursuing this because many similar compounds have been studied in the search for antibiotics (27) and one of them, that has now been on the market for some time, linezolid, definitely has significant MAOI activity at high, but sometimes clinically achieved, blood levels and has caused definite ST (28). The other main possibility is to actually assay its MAO activity *in vitro* using similar techniques to those we (Ramsay et al.) used for MB (29). Its ability to modify the tyramine pressor response would also give additional information as would testing in a rat model of ST e.g. see (30). However, that is all time and money. It would be worth doing if there is someone out there with the time and money.

Colleagues that I contacted (in 2014) about this responded promptly and calculations using theoretical modelling on computers (*in silico* modelling) indicated that the drug is indeed a weak monoamine oxidase inhibitor. We have Prof Yelecki in Turkey to thank for this (see acknowledgements below). Prof Yelecki’s computed value is 4.3 μmol for the inhibition of MAO-A and 3.5 μmol for MAO-B: this puts it in the same ballpark as moclobemide and linezolid. Other variables such as metabolism and the degree of brain penetrance mean that prediction of the potency in the human brain is not presently possible with any accuracy. Nevertheless, that is a strong indication that metaxalone has clinically relevant MAOI activity in high doses.

It is precisely because accurate estimation of the potency of drug effects at the relevant sites/receptors in the human brain is so difficult to ascertain that the occurrence or absence of ST with drug combinations is a useful tool for revealing whether the concentrations are actually reaching clinically relevant levels and altering central serotonin neurotransmission.

More and higher quality data from toxicology databases of overdoses in humans would therefore be of value, because it would establish the frequency and severity of ST with different doses and combinations of drugs. Unfortunately, despite recommendations made nearly 20 years ago (31), such data is not routinely collected.

An aside about Cyclobenzaprine

It is also worth noting in passing that one of the other drugs used in the same context as metaxalone is cyclobenzaprine. This drug has also been implicated in possible cases of ST, although there do not appear to be any clear cases of severe toxicity like the ones above with metaxalone (32-34). Previous discussion on this question prompted a similar chain of events to be set in motion as is related above. *In silico* modelling by Tudor Oprea's group has suggested that cyclobenzaprine is a weak SRI, comparable to amitriptyline (35, 36).

I intend to discuss these useful and interesting results about cyclobenzaprine in a separate commentary when I have time. In this immediate context the current data on cyclobenzaprine serves to illustrate this interesting and important point about the drug interactions the precipitate ST. If cyclobenzaprine is a significant SRI in humans then we would expect it to cause serotonergic symptoms, or even ST, when it is taken in overdose (which, like amitriptyline, it does **not** seem to do), but we would not expect it to cause major ST when combined with other (S)SRIs. This is because both drugs are working by the same mechanism of action and our current understanding indicates clearly that this does not lead to any greater elevation of serotonin than would be expected by simply increasing the dose of one or other of the SRI drugs. However, if cyclobenzaprine were to be mixed with an MAOI then the story would be dramatically different and severe life-threatening ST would be likely. Because MAOIs are not used very frequently the result is that cases of ST emerging with cyclobenzaprine would be expected to be rather less common than cases with metaxalone (other things being equal). This is because a lot of people in the general community are taking (S)SRI drugs (like 5-10%) so you do not have to have a very big series of metaxalone overdoses before it becomes inevitable that a proportion of them will have ingested both drugs.

Conclusion

The subtle twist appears to be that metaxalone is just sufficiently weak as an MAOI that in general circumstances it does not cause problems when combined with SRIs. Perhaps it is only in patients who ingest large, supra-therapeutic doses, or develop particularly high blood levels {Surmaitis, 2016 #21005}, that have a risk of ST with SRIs. And thus, metaxalone has managed to stay under the radar for half a century. In some ways it most closely resembles linezolid which definitely increases

serotonin-mediated side effects, but rarely provokes severe toxicity (28, 37-39).

Acknowledgements

The following colleagues have generously assisted with data interpretation and advice:

Prof. Dr. Kemal Yelekci, Kadir Has University Faculty of Engineering and Natural Sciences Department of Bioinformatics and Genetics (Head) Cibali Campus TR-34083 Fatih Istanbul TURKIYE

Rona Ramsay, Centre for Biomolecular Sciences, University of St Andrews, Fife, Scotland, UK

Tudor Oprea, Division of Biocomputing, Department of Biochemistry and Molecular Biology, University of New Mexico School of Medicine, Albuquerque, New Mexico, USA

References

1. Buckley, NA, Dawson, AH, and Isbister, GK, Serotonin syndrome. *BMJ*, **2014**. 348: p. 10.1136/bmj.g1626.
<http://www.ncbi.nlm.nih.gov/pubmed/24554467>
2. Gillman, PK, A review of serotonin toxicity data: implications for the mechanisms of antidepressant drug action. *Biol Psychiatry*, **2006**. 59(11): p. 1046-51.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=16460699
3. Gillman, PK, CNS toxicity involving methylene blue: the exemplar for understanding and predicting drug interactions that precipitate serotonin toxicity. *J Psychopharmacol (Oxf)*, **2011**. 25(3): p. 429-3.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=20142303
4. Evans, RW, Tepper, SJ, Shapiro, RE, Sun-Edelstein, C, et al., The FDA alert on serotonin syndrome with use of triptans combined with selective serotonin reuptake inhibitors or selective serotonin-norepinephrine reuptake inhibitors: American Headache Society position paper. *Headache*, **2010**. 50(6): p. 1089-99.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=20618823
5. Tepper, S, Allen, C, Sanders, D, Greene, A, et al., Coprescription of triptans with potentially interacting medications: a cohort study involving 240,268 patients. *Headache*, **2003**. 43(1): p. 44-8.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=12864757

6. Rojas-Fernandes, C, Can 5-HT₃ antagonists really contribute to serotonin toxicity? A call for clarity and pharmacological law and order. *Drugs - Real World Outcomes*, **2014**: p. 10.1007/s40801-014-0004-3.
<https://link.springer.com/article/10.1007/s40801-014-0004-3>
7. Anon, Summary safety review-serotonin blocking drugs (serotonin antagonists) Aloxi (palonosetron), Anzemet (dolasetron), Kytril (granisetron) and generics, and Zofran (ondansetron) and generics-Serotonin Syndrome. Health Canada, **2014**. May 14: p. <http://www.hc-sc.gc.ca/dhp-mps/medeff/advisories-avis/review-examen/serotonin-eng.php> (accessed Oct 2014).
8. Anon, Can 5-HT₃ antagonists (e.g., ondansetron etc.) really contribute to serotonin toxicity? Hunter Drug Information Service. HDIS Fact Sheet. Hunter New England Health District., **2014**.
9. Gillman, PK, The TGA and Aesop's 'Boy Who Cried Wolf'. *ANZCA Bull*, **2014**. Dec: p. 7.
10. Rosenbaum, H and Gillman, PK, Patient Safety and Methylene Blue-Associated Severe Serotonin Toxicity. *A&A Case Reports*, **2016**. 7(1): p. http://journals.lww.com/aacr/Citation/2016/07010/Patient_Safety_and_Methylene_Blue_Associated.1.aspx.
11. Bosak, AR and Skolnik, AB, Serotonin Syndrome Associated with Metaxalone Overdose. *J Med Toxicol*, **2014**. 10(4): p. 402-5.
<http://www.ncbi.nlm.nih.gov/pubmed/24805103>
12. Ignneri, L, Shaw, C, and Solomon, B, Life-threatening serotonin syndrome after administration of metaxalone in a patient on citalopram. *Crit Care Med*, **2013**. 41(12): p. A285.
13. Isbister, GK, McGettigan, P, and Dawson, A, A fatal case of moclobemide-citalopram intoxication. *J. Anal. Toxicol.*, **2001**. 25(8): p. 716-7.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=11765030
14. Isbister, GK, Hackett, LP, Dawson, AH, Whyte, IM, et al., Moclobemide poisoning: toxicokinetics and occurrence of serotonin toxicity. *Brit J of Clin Pharmacol*, **2003**. 56: p. 441-450.
15. Gillman, PK, Serotonin syndrome: clomipramine too soon after moclobemide. *Int Clin Psychopharmacol*, **1997**. 12: p. 339-342.

http://journals.lww.com/intclinpsychopharm/Abstract/1997/11000/Serotonin_syndrome__clomipramine_too_soon_after.7.aspx

16. Gillman, PK, Moclobemide and the risk of serotonin toxicity (or serotonin syndrome). *CNS Drug Rev*, **2004**. 10: p. 83-85.

<http://onlinelibrary.wiley.com/store/10.1111/j.1527-3458.2004.tb00005.x/asset/j.1527-3458.2004.tb00005.x.pdf?v=1&t=ih5v9pa8&s=967b77c9e3938d24bfb26f8a8118da5b710cfe9a>

17. Pilgrim, JL, Gerostamoulos, D, Woodford, N, and Drummer, OH, Serotonin toxicity involving MDMA (ecstasy) and moclobemide. *Forensic Sci Int*, **2011**.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=21570786

18. Pilgrim, JL, Gerostamoulos, D, and Drummer, OH, Deaths Involving MDMA and the Concomitant Use of Pharmaceutical Drugs. *J. Anal. Toxicol.*, **2011**. 35(4): p. 219-26.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=21513615

19. Pilgrim, JL, Gerostamoulos, D, and Drummer, OH, Deaths involving contraindicated and inappropriate combinations of serotonergic drugs. *Int. J. Legal Med.*, **2010**.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=21120513

20. Huska, MT, Catalano, G, and Catalano, MC, Serotonin syndrome associated with the use of escitalopram. *CNS Spectr*, **2007**. 12(4): p. 270-274 <http://www.cnsspectrums.com/asp/articleDetail.aspx?articleid=1038>.

<http://www.cnsspectrums.com/asp/articleDetail.aspx?articleid=1038>

21. Rivera, A, Gaudino, W, and Weiss, L, Serotonin syndrome in a patient taking tramadol and nefazodone concomitantly: A case report. *Arch. Phys. Med. Rehabil.*, **2003**. 84(9): p. E23.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=13680712

22. Martini, DI, Nacca, N, Haswell, D, Cobb, T, et al., Serotonin syndrome following metaxalone overdose and therapeutic use of a selective serotonin reuptake inhibitor. *Clin Toxicol (Phila)*, **2015**. 53(3): p. 185-7.

<http://www.ncbi.nlm.nih.gov/pubmed/25671244>

23. Surmaitis, RM, Nappe, TM, and Cook, MD, Serotonin syndrome associated with therapeutic metaxalone in a patient with cirrhosis. *Am. J. Emerg. Med.*, **2016**. 34(2): p. 346 e5-6.

<https://www.ncbi.nlm.nih.gov/pubmed/26159819>

24. Moore, KA, Levine, B, and Fowler, D, A fatality involving metaxalone. *Forensic Sci Int*, **2005**. 149(2-3): p. 249-51.

<http://www.ncbi.nlm.nih.gov/pubmed/15749367>

25. Poklis, JL, Ropero-Miller, JD, Garside, D, and Winecker, RE, Metaxalone (Skelaxin)-related death. *J. Anal. Toxicol.*, **2004**. 28(6): p. 537-41.
<http://www.ncbi.nlm.nih.gov/pubmed/15516312>
26. Gruszecki, AC, Kloda, S, Simmons, GT, Daly, TM, et al., Polydrug fatality involving metaxalone. *J. Forensic Sci.*, **2003**. 48(2): p. 432-4.
<http://www.ncbi.nlm.nih.gov/pubmed/12665007>
27. Ramsay, RR and Gravestock, MB, Monoamine oxidases: to inhibit or not to inhibit. *Mini Rev Med Chem*, **2003**. 3(2): p. 129-36.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=12570845
28. Kulkarni, RR and Kulkarni, PR, Linezolid-induced near-fatal serotonin syndrome during escitalopram therapy: case report and review of literature. *Indian J Psychol Med*, **2013**. 35(4): p. 413-6.
<http://www.ncbi.nlm.nih.gov/pubmed/24379509>
29. Ramsay, RR, Dunford, C, and Gillman, PK, Methylene blue and serotonin toxicity: inhibition of monoamine oxidase A (MAO A) confirms a theoretical prediction. *Br J Pharmacol*, **2007**. 152(6): p. 946-51.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=17721552
30. Shioda, K, Nisijima, K, Yoshino, T, and Kato, S, Mirtazapine abolishes hyperthermia in an animal model of serotonin syndrome. *Neurosci. Lett.*, **2010**. 482(3): p. 216-9.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=20655983
31. Gillman, PK, Serotonin Syndrome: History and Risk. *Fundam. Clin. Pharmacol.*, **1998**. 12(5): p. 482-491.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=9794145
32. Day, LT and Jeanmonod, RK, Serotonin syndrome in a patient taking Lexapro and Flexeril: a case report. *Am. J. Emerg. Med.*, **2008**. 26(9): p. 1069 e1-3.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=19091289
33. Keegan, MT, Brown, DR, and Rabinstein, AA, Serotonin syndrome from the interaction of cyclobenzaprine with other serotonergic drugs. *Anesth. Analg.*, **2006**. 103(6): p. 1466-8.
http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=17122225
34. Gillman, PK, Is there sufficient evidence to suggest cyclobenzaprine might be implicated in causing serotonin toxicity? *Am. J. Emerg. Med.*, **2009**. 27(4): p. 509-510.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=19555629

[http://www.ajemjournal.com/article/S0735-6757\(09\)00128-4/abstract](http://www.ajemjournal.com/article/S0735-6757(09)00128-4/abstract)

35. Gregori-Puigjane, E, Setola, V, Hert, J, Crews, BA, et al., Identifying mechanism-of-action targets for drugs and probes. *Proc Natl Acad Sci USA*, **2012**.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=22711801

36. Mestres, J, Seifert, SA, and Oprea, TI, Linking pharmacology to clinical reports: cyclobenzaprine and its possible association with serotonin syndrome. *Clin Pharmacol Ther*, **2011**. 90(5): p. 662-5.

<http://www.ncbi.nlm.nih.gov/pubmed/21975349>

37. Lawrence, KR, Adra, M, and Gillman, PK, Serotonin Toxicity Associated with the Use of Linezolid: A Review of Postmarketing Data. *Clin Infect Dis*, **2006**. 42: p. 1578-83.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=16652315

38. Lodise, TP, Patel, N, Rivera, A, Tristani, L, et al., Comparative evaluation of serotonin toxicity among veterans affairs patients receiving linezolid and vancomycin. *Antimicrob Agents Chemother*, **2013**. 57(12): p. 5901-11.

<http://www.ncbi.nlm.nih.gov/pubmed/24041888>

39. Butterfield, JM, Lawrence, KR, Reisman, A, Huang, DB, et al., Comparison of serotonin toxicity with concomitant use of either linezolid or comparators and serotonergic agents: an analysis of Phase III and IV randomized clinical trial data. *J. Antimicrob. Chemother.*, **2012**. 67(2): p. 494-502.

http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?cmd=Retrieve&db=PubMed&dopt=Citation&list_uids=22139199