
CNS SPECTRUMS

CME Review Article

A neuroscientific update on monoamine oxidase
and its inhibitors

This activity is sponsored by the Neuroscience Education Institute



CME Information

Accreditation and credit designation statements

The Neuroscience Education Institute is accredited by the Accreditation Council for Continuing Medical Education to provide continuing medical education for physicians.

The Neuroscience Education Institute designates this enduring material for a maximum of 1.5 *AMA PRA Category 1 Credits*[™]. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

Target audience

This activity has been developed for prescribers specializing in psychiatry. There are no prerequisites. All other health care providers interested in psychopharmacology are welcome for advanced study, especially primary care physicians, nurse practitioners, psychologists, and pharmacists.

Statement of need

Most practicing mental health clinicians are familiar with the various first-line treatments and can select among them to recommend an appropriate initial treatment. However, there are documented gaps between established best practices and actual practice for addressing residual symptoms, side effects, and treatment-resistant depression.

To help address these professional practice gaps, quality improvement efforts need to provide education regarding (1) monitoring patients over time for residual symptoms, side effects, and nonadherence; (2) adjusting treatment to address residual symptoms, side effects, and other issues that may affect response or adherence; and (3) applying strategies to address treatment-resistance.

Learning objectives

After completing this activity, participants should be better able to:

- Explain the pathophysiology and epidemiology of major depressive disorder (MDD), atypical depression, and treatment-resistant depression review
- Modify treatment strategies in order to address challenges of treating depression including safety concerns (eg, adverse effects, drug/food interactions), noncompliance, and relapse
- Integrate counseling strategies for treatment-resistance into clinical practice to improve patient outcomes

Date of release/expiration

Released: December, 2013

CME credit expires: November, 2016

Sponsor

This activity is sponsored by the Neuroscience Education Institute.

Acknowledgment of Financial Support

This activity is supported by an educational grant from Mylan Specialty L.P.

Activity instructions

This CME activity is in the form of a printed article and incorporates instructional design to enhance your retention of the information and pharmacologic concepts that are being presented. You are advised to review this activity from beginning to end, and then complete the posttest and activity evaluation. The estimated time for completion of this activity is 90 minutes.

NEI disclosure policy

It is the policy of the Neuroscience Education Institute to ensure balance, independence, objectivity, and scientific rigor in all its educational activities. Therefore, all individuals in a position to influence or control content development are required by NEI to disclose any financial relationships or apparent conflicts of interest. Although potential conflicts of interest are identified and resolved prior to the activity being presented, it remains for the participant to determine whether outside interests reflect a possible bias in either the exposition or the conclusions presented.

These materials have been peer reviewed to ensure the scientific accuracy and medical relevance of information presented and its independence from commercial bias. NEI takes responsibility for the content, quality, and scientific integrity of this CME activity.

Disclosure Statements

Author

Thomas L. Schwartz, MD, is an associate professor in the department of psychiatry at SUNY Upstate

Medical University in Syracuse, NY. Dr. Schwartz receives research support from Bristol-Myers Squibb, Cephalon, and Cyberonics.

No writing assistance was utilized in the production of this article.

Content editor

Debbi Ann Morrissette, PhD, is an adjunct professor of biological sciences at California State University in San Marcos and at Palomar Community College in San Marcos, CA, and senior medical writer at the Neuroscience Education Institute in Carlsbad, CA. Dr. Morrissette has no financial relationships to disclose.

CNS Spectrums peer review

All CME articles are peer reviewed in accordance with the strict standards of *CNS Spectrums* and in accordance with requirements and recommendations of the International Committee of Medical Journal Editors. The Editorial policies of the journal *CNS Spectrums* and peer review of all articles that appear in the journal are managed independently by Cambridge University Press, and no financial relationship exists between the CME provider and Cambridge for this service.

Additional peer reviewer

Mark D. Williams, MD, is an assistant professor in the department of psychiatry and psychology at the Mayo Clinic in Rochester, MN. Dr. Williams has no financial relationships to disclose.

Design staff

Nancy Muntner is the director of medical illustrations at the Neuroscience Education Institute in Carlsbad, CA. She has no financial relationships to disclose.

Program development

Sheri Mills is the director of program development at the Neuroscience Education Institute in Carlsbad, CA. She has no financial relationships to disclose.

Steve Smith is the president and chief operating officer at the Neuroscience Education Institute in Carlsbad, CA. He has no financial relationships to disclose.

Disclosed financial relationships with conflicts of interest have been reviewed by the Neuroscience Education Institute CME Advisory Board Chair and resolved. All faculty and planning committee members have attested that their financial relationships do not affect their ability to present well-balanced, evidence-based content for this activity.

Disclosure of off-label use

This educational activity may include discussion of unlabeled and/or investigational uses of agents that are not currently labeled for such use by the FDA. Please consult the product prescribing information for full disclosure of labeled uses.

Disclaimer

Participants have an implied responsibility to use the newly acquired information from this activity to enhance patient outcomes and their own professional development. The information presented in this educational activity is not meant to serve as a guideline for patient management. Any procedures, medications, or other courses of diagnosis or treatment discussed or suggested in this educational activity should not be used by clinicians without evaluation of their patients' conditions and possible contraindications or dangers in use, review of any applicable manufacturer's product information, and comparison with recommendations of other authorities. Primary references and full prescribing information should be consulted.

A neuroscientific update on monoamine oxidase and its inhibitors

Thomas L. Schwartz*

Psychiatry Department, SUNY Upstate Medical University, Syracuse, New York, USA

The goal of this brief review is to explain the role of monoamine oxidase enzymes in the neurobiology, etiology, and presentation of psychiatric illnesses, primarily major depressive disorder. This article will initially focus on the basic science and function of the monoamine oxidase system and some proposed neuropsychiatric symptoms that may arise if this enzyme system is altered by genetic predisposition. These findings and theories will next be translationally discussed in regard to clinical application pertaining to enzyme inhibition and the treatment of major depressive and other psychiatric disorders.

Received 23 August 2013; Accepted 9 September 2013; First published online 19 November 2013

Key words: Antidepressant treatment, major depressive disorder, monoamine oxidase inhibition.

Introduction

The overly simplistic monoamine deficiency hypothesis that states that major depressive disorder (MDD) arises out of low concentrations of serotonin, dopamine, and/or norepinephrine is often combated and debated.¹ Newer theories such as monoamine receptor excesses, brain-derived neurotrophic factor deficiencies, hypofrontality, or hyperactive amygdala activity are potentially valid in theory now as well.^{2,3} However, in practice, most antidepressant treatments (ADTs) that are indicated and approved function to facilitate increases in monoamine levels, ie, selective serotonin reuptake inhibitors (SSRI), selective serotonin-norepinephrine reuptake inhibitors (SNRI), selective norepinephrine-dopamine reuptake inhibitors (NDRI), and tricyclic antidepressants (TCA). Other ADTs manipulate monoamine receptors, ie, serotonin antagonist-reuptake inhibitors (SARI), serotonin partial agonist-reuptake inhibitors (SPARI), and norepinephrine agonist-serotonin antagonists (NASA) as a way to change neurotransmission rates and effect an antidepressant response through the monoamine systems as well.^{4,5}

These ADTs also have in common the fact that they will raise only 1 or 2 monoamines' central nervous

system (CNS) concentrations at a time. The monoamine oxidase inhibitor (MAOI) ADT functions uniquely to increase all 3 monoamines simultaneously.³ This addresses the monoamine hypothesis of depression directly by inhibiting the degradation of all 3 monoamines by lowering monoamine oxidase-A (MOA-A) (preferentially) and monoamine oxidase-B (MOA-B) enzyme activity within CNS neurons.

Despite the MAOI class's ability to robustly increase all 3 monoamines, these ADTs have largely fallen out of use due to the risk of drug-drug interactions and dietary reactions. These may cause increased risk (hypertensive crisis/serotonin syndrome) to the patient versus more conventional ADTs, and also, the MAOIs require quite a bit of time educating and informing the patient prior to treatment initiation.⁶ There likely is also increased fear of use among younger clinicians due to little training and experience with MAOIs in psychiatry residency clinics, as well as increased patient fear if patients should self-research the MAOI class of drugs online. Clinicians also underutilize the MAOI, likely because there are more facts about interactions, washout periods, titration schedules, and combination-augmentation limitations that they are forced to memorize or reference at the point of service, making competent MAOI use time-insensitive for the busy practitioner.

Despite many ADTs being available for use, MDD still remains a difficult-to-treat psychiatric disorder with a 16% lifetime risk, 60% recurrence rate, and a 30%

*Address for correspondence: Thomas L. Schwartz.

(Email: schwartzt@upstate.edu)

This activity is supported by an educational grant from Mylan Specialty L.P.

chance of becoming chronic in nature.⁷⁻¹² With very few ADTs noted to be in the research pipeline in general, and especially very few, if any, that will work outside of the monoamine hypothesis,¹³ psychopharmacologists likely have to become more comfortable and adept at using available treatments from each pharmacological family while awaiting the next, novel breakthrough ADT to be developed and marketed.¹⁴

The remainder of this article is dedicated to better describing and understanding the MAOI class of ADT. From basic science to MDD etiological theory, information will be presented in order to give the reader a well-rounded sense of why the MAOI class of antidepressants should continue to have a clear place in the treatment of treatment-resistant MDD, especially where risk-benefit analysis is more meaningful and acceptable.

MAO Functioning in the CNS

MAO-A and MAO-B enzymes' main function is to lower CNS concentration of monoamines. The brain seeks to maintain homeostasis in most ways, and governing the amount and activity of available monoamines is no exception. If dopamine levels are too high, MAO levels are noted to increase to compensate. If serotonin levels become too low, then MAO activity should lower as well to leave adequate serotonin supplies left for the CNS to function optimally.¹⁵ An example might make use of the dopamine hypothesis of schizophrenia,¹⁶ where too much dopamine is felt to promote positive symptoms of hallucinations or delusions. The opposite may happen in MDD, or even attention deficit hyperactivity disorder (ADHD), where more deficient dopamine activity levels may produce cognitive symptoms of inattention, inability to make decisions, poor concentration, and loss of vigilance.³ In either case, monoamine levels and receptor activity are partially maintained by the MAO enzyme system, and when optimal, no symptoms likely occur.

It is likely that in a certain subset of MDD patients, the monoamine deficiency hypothesis is clearly true. For example, some patients may have elevated MAO concentrations or activity that systematically lower monoamines, causing brain functioning to change, which yields a myriad of MDD symptoms. For example, prefrontal cortex (PFC) and anterior cingulate (ACC) neuroanatomic areas have been found to have excess MAO activity (34%) in MDD studies, and suicide victims show hypothalamic elevations as well.¹⁷⁻²¹ Particularly, MAO-A increases in the PFC would lower dopamine and norepinephrine; this could theoretically predispose patients to executive dysfunction and to incorrect negative emotional valences being assigned to social situations. In the ACC, this might promote inattention,

poor concentration, and, more likely, provide for a loss of vigilance or an inability to stay on task.³

As a teaching example, consider the following 3 cases. In the first case, imagine a patient who is depressed and has inherited genes for a defective serotonin transporter (SERT), also known as a reuptake system. If this transporter system is overly aggressive, then serotonin is removed from synapses too quickly and MDD symptoms may emerge. Knowing this, the choice of an SSRI would make clinical and neuroscientific sense, as the reuptake pumps that are causing the depression are too active, and should be inhibited and shut down. In this way, the drug mechanism of action perfectly corrects the underlying brain pathology that is likely causing this individual patient's particular depressive symptoms. Here, the drug corrects the defective protein's excessive activity (SERT system). (See Figures 1A-1C.)

The second MDD case involves a patient who has inherited genes for MAO-A, whereby the enzymes produced are overly active and aggressively destroy and deplete brain monoamines, thus causing clinical depressive symptoms. Choosing an MAOI treatment, even initially, would be worth the clinical risk perhaps, given that the drug's mechanism of action may actually directly reverse the cause of the depressive symptoms at hand. Again, the drug corrects the defective and overly aggressive catabolic protein (enzyme) in this case too (see Figure 1D).

In the final case, imagine that our second patient with the aggressive MAO-A enzyme activity comes to be seen in a primary care or psychiatric care practice in a first episode of MDD. Generally this patient is placed on a front-line treatment, likely an SSRI. Sequenced Treatment Alternatives to Relieve Depression (STAR*D) trial results²² would suggest that only 1/3 of these patients will remit on the SSRI. The next guideline and practical step would be to switch to an SNRI or NDRI, which would allow another batch of patients to remit.²³ Still others will remain fully depressed. For these, a third step of using sedating antidepressants, antidepressant combinations, or TCA monotherapies is often required. After this, a reasonable minority of patients will remain unremitted, and those who gain a remission are very likely to relapse within a year. In these cases, use of novel antidepressants is often fraught with treatment response and loss of response, or "poop out." In some of these cases, it is possible that the ADT provides a temporary increase in CNS monoamine levels, only to be systematically thwarted by an inherited and aggressive MAO-A activity the patient's brain possesses. It is possible that brain homeostasis prevails in these cases where the depressed brain detects the extra monoamines provided by the SSRI being used and adjusts by increasing its levels of MAO availability or

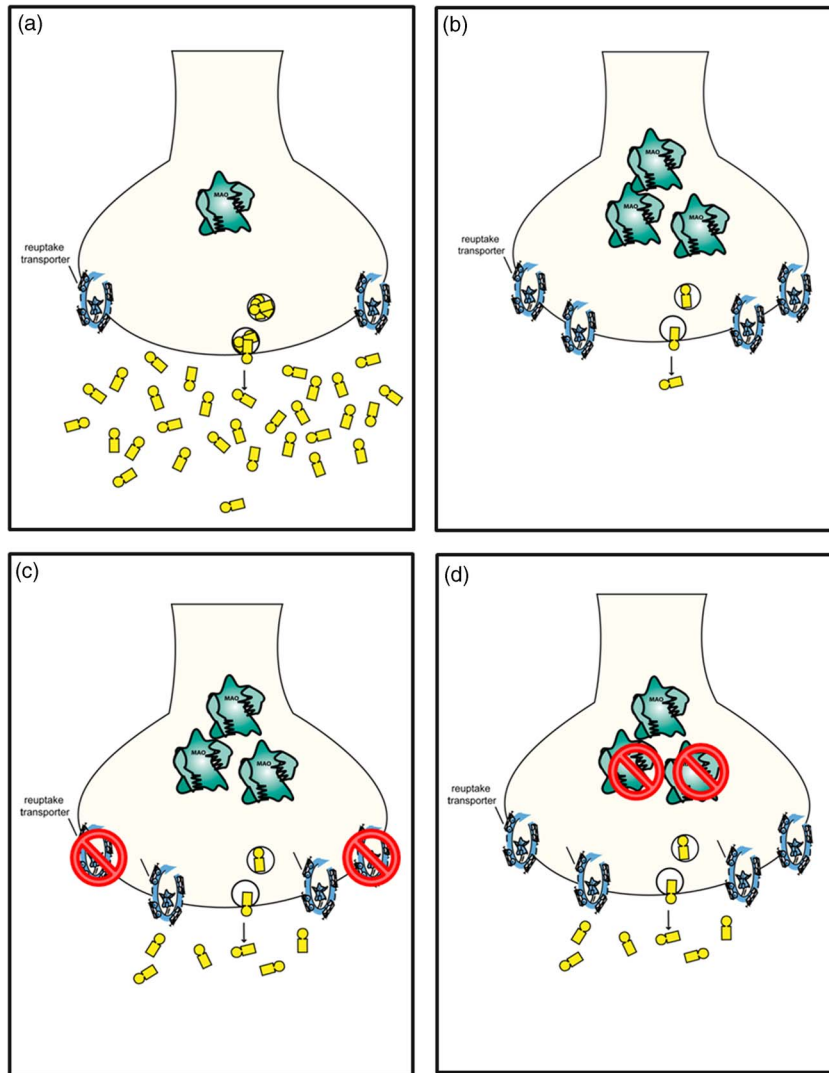


FIGURE 1. Monoamine deficiency and treatment in depression. (A) In the healthy brain, levels of both reuptake transporters and monoamine oxidase-A (MAO-A) allow for adequate monoamine levels. (B) According to the monoamine hypothesis of depression, the depressed brain experiences too much reuptake and/or MAO-A activity, leading to reduced levels of monoamines in the synapse. (C) Treatment with a first-line antidepressant, such as an SSRI, leads to increased levels of monoamines in the synapse by blocking reuptake. (D) Treatment with an MAOI leads to increased synaptic monoamine levels by blocking the enzymatic activity of MAO-A. Copyright 2013. Neuroscience Education Institute. Used with permission.

activity.¹⁵ Assuming that each ADT therapeutic trial here would take 3 months to titrate and await effect, this patient's depression would have been treated for at least 9 months with little hope of sustained remission, as the root etiologic cause of the MDD in this case was not addressed. If the practitioner knew of the inherent MOA-A abnormality at treatment initiation, this could have been explained to the patient, and the greater inherent risk of MAOI use could be justified and possibly accepted by patient and prescriber alike. In this case, lowering MAO-A activity with an MAOI may have directly addressed the cause of excessive monoamine depletion instead of adding the SSRI, SNRI, NDRI, etc as temporary fixes.

MAO-A as a Genetic Risk Factor in Major Depressive Disorder

Genetically, there are some findings that may support the overactive MAO-A corollary of the monoamine deficiency hypothesis of MDD. First, a single nucleotide polymorphism mutation called T941G has a G allele, which, when homozygously inherited from both parents, yields a more robust MAO-A enzyme to be created that is 75% more efficient at monoamine degradation and is more often associated with MDD than its less active T allele.²⁴ There is also a variable tandem repeat mutation (MAOA-uVNTR) where the long allele is associated with greater risk to develop MDD, worse outcomes in MDD

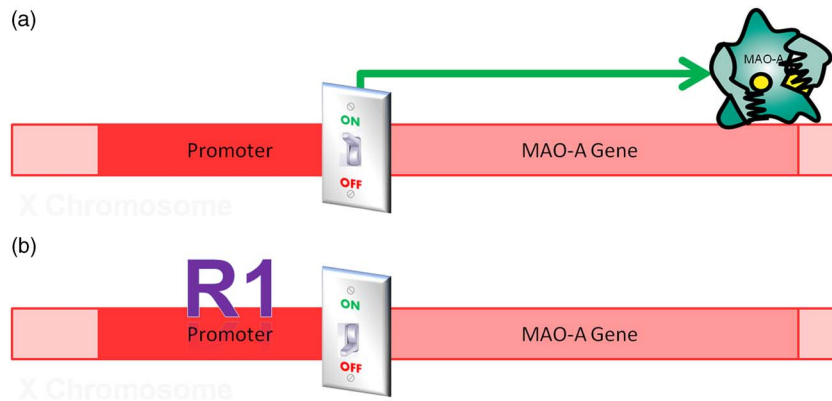


FIGURE 2. The R1 repressor protein modulates MAO-A gene expression. R1 is an upstream transcriptional repressor of the MAO A gene. (A) When R1 is not bound to the promoter region of the MAO-A gene (located on the X chromosome), MAO-A is expressed, leading to degradation of monoamines. (B) Binding of R1 to the promoter region of the MAO-A gene shuts off expression of MAO-A, leading to reduced degradation of monoamines. If levels of R1 are low (due to genetic and/or environmental factors), excessive expression of MAO-A would be expected to cause deficiency in monoamine levels. Copyright 2013 Neuroscience Education Institute. Used with permission.

patients who experienced childhood trauma, and greater suicide risks in male MDD patients.^{17,24,25-30} Outside of MDD, this gene may also dictate a patient's temperament or personality style. For example, a cortico-limbic neuroanatomic circuit was detailed in a functional neuroimaging study by Buckholtz et al.³¹ These authors determined that a network, composed of area BA 10 of the ventromedial prefrontal cortex (vmPFC), the ACC, and the amygdala, exists and functions under significant genetic control based largely on the MAO-A gene. Here, increased MAO-A functioning (long repeat alleles) in the vmPFC allows for a loss of top-down control in the amygdala with a resultant hyperactivity here. In these cases, patients often have a loss of harm avoidance, more impulsivity, and a loss of prosocial reward-seeking, all predominantly in male subjects.³¹ This supports the growing stress-diathesis-laden theory that MDD patients inherit genes for aberrant proteins. When there is enough inheritance of risk combined with environmentally stressful interactions, these dysfunctional proteins (enzymes, receptors, transporters, transcription factors, etc) change brain neurocircuitry functioning and symptoms can develop. Sometimes, symptoms may represent as personality traits that predispose patients to greater interpersonal and social stress that may lead secondarily to MDD development, or perhaps these inherited abnormal proteins may lead directly to depressive symptoms, ie, poor concentration, suicidality, etc.³² Regardless, the above MAO genetic findings and translational correlations suggest a potential etiology for depressive symptoms in subsets of MDD patients.

As noted above, aggressive MAO-A activity may deplete monoamines and facilitate the development of MDD symptoms in some patients. Another depressogenic mechanism may be related to a potential loss of

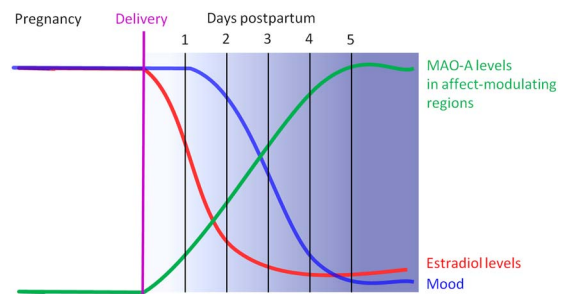


FIGURE 3. MAO-A activity, estrogen, and postpartum depression. In the weeks following delivery, estradiol levels are drastically reduced. As estrogen is an inhibitor of monoamine oxidase A (MAO-A), it is not surprising that MAO-A levels increase during the postpartum period, leading to increased degradation of monoamines and a concomitant increase in the risk for depressive symptoms. Copyright 2013 Neuroscience Education Institute. Used with permission.

CNS neuroprotection when MAO-A is too active, and brain atrophy may occur. In fact, several studies have shown that patients with more chronic MDD appear to have a loss in brain tissue volume.^{33,34} It also appears that a byproduct of MAO activity is the generation of chemicals that are toxic and detrimental to neuronal health.³⁵ For example, increases in reactive oxygen species (ROS), ie, oxygen ions and peroxides, have been noted in depressed patients, which allows for decreased mitochondrial ATP production, altered cerebral energy metabolism, and abnormal increases in cell death and damage to occur. Additionally, R1 has been associated with increased MDD findings as well. R1 is an upstream transcription factor protein whose function is to actually lower the production of functioning MAO-A proteins. R1 binds to the MAO-A gene's promoter region and inhibits it (Figure 2). Post-mortem studies in MDD patients suggest that R1 is 37% deficient in treated and

untreated MDD patients and also in suicide victims.^{19,36} It suggests in these cases that excessive MAO-A activity was likely increased, and monoamine levels and neurotransmission were decreased by default.

There may also be a gender difference in the role that MAO enzymes play in the development of MDD symptoms. Women typically have a higher prevalence

of MDD diagnosis and treatment compared to men. One possible explanation may involve MAO-A levels again. First, sex steroidal hormones (estrogen/testosterone) inhibit MAO activity and promote creation of greater serotonin CNS monoamine concentrations. One clinical example revolves around postpartum depression, where estrogen levels drop quickly after delivery. Postpartum

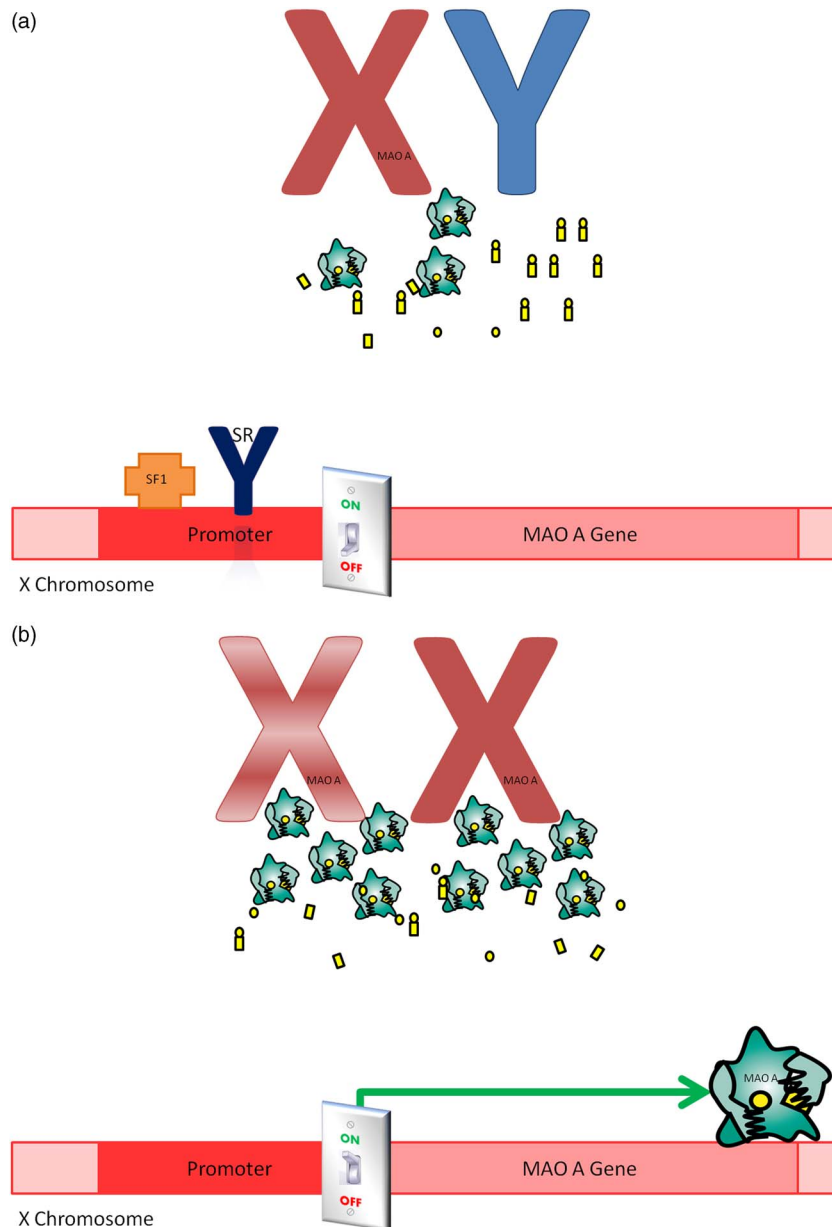


FIGURE 4. Sex differences in MAO-A gene expression. (A) In males, the SRY protein produced by the sex-determining region of the Y sex chromosome, which is consequently present only in males, regulates the expression of the MAO-A gene, which is located on the X chromosome. When SRY, along with a protein called steroidogenic factor 1 (SF1), binds to the promoter region of the MAO-A gene, expression of MAO-A is inhibited and monoamine levels are elevated. (B) Unlike males, females have 2 copies of the X-chromosome; 1 X chromosome is usually inactive (silenced) in every cell of the female body. However, the silencing of the X chromosome is not always absolute, and this incomplete X inactivation can lead to an excess expression of MAO-A in females. Differential expression of MAO-A from incomplete X activation and the absence of the SRY protein may contribute to the increased prevalence and heritability of MDD in females. Copyright 2013 Neuroscience Education Institute. Used with permission.

TABLE 1. MAOI dietary guidelines

Foods to Avoid*	Foods Allowed
Dried, aged, smoked, fermented, spoiled, or improperly stored meat, poultry, and fish	Fresh or processed meat, poultry, and fish; properly stored pickled or smoked fish
Broad bean pods	All other vegetables
Aged cheeses	Processed cheese slices, cottage cheese, ricotta cheese, yogurt, cream cheese
Tap and unpasteurized beer	Canned or bottled beer and alcohol
Marmite	Brewer's and baker's yeast
Soy product/tofu	Peanuts
Sauerkraut, kimchee	
Banana peel	Bananas, avocados, raspberries
Tyramine-containing nutritional supplement	

*Not necessary for 6 mg transdermal or low-dose oral selegiline
 Copyright 2013 Neuroscience Education Institute. Used with permission.

depression is common in this time frame, and it has been found that the abrupt loss of estrogen compounds postpartum will allow for a 43% increase in MAO-A activity and subsequent losses in synaptic monoamines (Figure 3). Here, Sacher et al³⁷ found and measured these remarkable increases in MAO-A binding activity postpartum using functional neuroimaging techniques. Interestingly, the MAO-A gene resides on both female X chromosomes. Generally, only 1 gene on 1 X-chromosome is active, but in some MDD women, X-inactivation may occur, allowing both X chromosomes to produce MAO-A simultaneously, leading to greater monoamine degradation and MDD symptoms to potentially develop. Also, in males, the Y-chromosome carries a gene whose protein's (SRY) function is to inhibit or limit MAO-A enzyme production. Women do not carry a Y-chromosome and so are missing this check and balance capability again leading to more MAO-A development and activity³⁸⁻⁴⁰ (Figure 4). In summary, certain genetic findings may be used to explain the etiology of MDD symptoms in certain patients. Especially in women, excesses in MAO-A concentrations or activity may be a main etiologic factor.

Conclusions

Evidence appears to be mounting, and it makes intuitive sense neuroscientifically that MAO functioning is related to psychosocial functioning. In this article, the discussion focused largely on normal MAO-A functioning and monoamine homeostasis initially and transitioned to a larger discussion regarding the way abnormalities in MAO-A functioning could lead to temperament changes or the genesis of MDD symptoms directly. One caveat would be to indicate that MDD is likely more complicated than having just the 1 or 2 gene mutations mentioned here.^{41,42} There are likely hundreds of genes that would have to be aberrant, making many dysfunctional proteins to create abnormal

hyper- or hypofunctioning neurocircuits. These would have to interact and coalesce in the face of key environmental factors to create enough MDD symptoms in a patient to make said patient fully syndromal in order to be diagnosed and treated with pharmacotherapy or psychotherapy.

The author does wish the reader to consider that, in a certain subset of MDD patients, that MAO-A dysfunction, especially hyperactivity, may be a key factor among many that may lend to recurring, resistant, or refractory MDD. The use of MAOI agents should not be a treatment of last resort after 10-12 failed antidepressant trials occur or when the patient has become chronic with 2-3 years of unmitigated MDD. Clinically, female patients with recurring MDD and frequent loss of therapeutic antidepressant treatment response should be considered earlier than this for MAOI utilization given the neuroscientific basis presented in this article. There are no randomized or controlled trials in this area, of course. These scenarios rely on the clinician's previous experience in treating this patient population and the willingness to embrace a theoretical biologically oriented formulation. Additionally, the risk-benefit analysis of MAOI use must be considered. Key factors include the knowledge of drug-drug interactions (DDI), whereby a clinician should not add drugs with serotonergic reuptake inhibition to MAOI due to risk of serotonin syndrome. Outside of the customarily avoided SSRI, SNRI, and TCA, the reader should be aware that other drugs may have these properties hidden in their mechanisms of action. For example, the opiates tramadol, meperidine, fentanyl, methadone, and tapentadol all have serotonin reuptake inhibition (SRI) properties,⁴³ as do the antihistamines brompheniramine and chlorpheniramine.⁴⁴ Factually, clinicians often also forget that both cyclobenzaprine (for muscle pain) and carbamazepine (for epilepsy and bipolar disorder) are also TCA in structure. They are fairly devoid of SRI properties, but are still relatively contraindicated with

MAOI use. These DDI can be intimidating to the prescriber of the MAOI, and the author often combines 2 basic rules as part of informed consent when prescribing MAOI to patients: (1) use 1 pharmacy and (2) carry and use a diet restriction card. The first rule is to alert the patient to use 1 single large chain pharmacy store routinely to fill his or her MAOI prescription. This single-pharmacy approach allows 1 agency to monitor all prescription drugs the individual patient receives, and large chain pharmacies deploy likely the most up to date computer tracking systems for DDI given their liability in dispensing conflicting medications.⁴⁵ This provides a double-checking system between the prescriber and the dispenser of the MAOI. Patients should also be instructed to bring all over the counter (OTC) vitamins, minerals, pain medications, cold, and allergy medicines to the pharmacist's counter to seek counseling from the pharmacist *every time*, as even OTC drugs can create serotonin syndrome complications. The second rule deals with the better-known tyramine hypertensive crisis, whereby ingested food containing tyramine (aged cheese, fava beans, banana peel, tap beer, marmite, sauerkraut, soy products, herring, or old or inappropriately stored meat, poultry, or fish, etc) may cause a robust release of norepinephrine, causing extreme hypertension, stroke, or heart attack in some cases. The rule here requires the patient to be aware of, and even carry a card delineating the foods that cannot be ingested while taking an MAOI (Table 1). Unfortunately, there is no double-check system in place, as the prescriber cannot ask the dispenser to monitor and counsel on diet in the same way it can counsel on DDI. Often, patients in practice seem the most worried that they will need to follow a hypertension diet (low salt), a cardiac diet (low fat), or a diabetic diet (low sugar), but in practice once they are told of the seemingly random list of about 20 foods to avoid, most patients seem compliant and become knowledgeable quickly regarding dietary restrictions. Using these 2 safety rules makes prescribing MAOI less intimidating than memorizing and navigating the many different MAOI interaction facts for both patient and clinician. The author hopes that these clinical pearls will allow for the appropriate and timely use of the MAOI class of antidepressants when warranted and after appropriate informed consent is given to the recurrent or treatment-resistant MDD patient.

Disclosures

Thomas L. Schwartz receives research support from Bristol-Myers Squibb, Cephalon, and Cyberonics.

No writing assistance was utilized in the production of this article.

REFERENCES:

- Lacasse JR, Leo J. Serotonin and depression: a disconnect between the advertisements and the scientific literature. *PLoS Med.* 2005; **2**: e392.
- Belmaker RH, Agam G. Major depressive disorder. *N Engl J Med.* 2008; **358**: 55-68.
- Stahl SM. *Stahl's Essential Psychopharmacology: Neuroscientific Basis and Practical Application*, 4th ed. Cambridge, UK: Cambridge University Press; 2013.
- Sadock BJ, Sadock VA, Sussman N. *Kaplan & Sadock's Pocket Handbook of Psychiatric Drug Treatment*, 5th ed. Philadelphia: Lippincott, Williams & Wilkins; 2011.
- Stahl SM. *Stahl's Essential Psychopharmacology: The Prescriber's Guide*, 4th ed. Cambridge, UK: Cambridge University Press; 2011.
- Berlim MT, Fleck MP, Turecki G. Current trends in the assessment and somatic treatment of resistant/refractory major depression: an overview. *Ann Med.* 2008; **40**(2): 149-159.
- Greden JF. The burden of recurrent depression: causes, consequences, and future prospects. *J Clin Psychiatry.* 2001; **62**(Suppl 22): 5-9.
- Kessler RC, Berglund P, Demler O, et al. The epidemiology of major depressive disorder: results from the National Comorbidity Survey Replication (NCS-R). *JAMA.* 2003; **289**(23): 3095-3105.
- Keller MB, Boland RJ. Implications of failing to achieve successful long-term maintenance treatment of recurrent unipolar major depression. *Biol Psychiatry.* 1998; **44**(5): 348-360.
- Keller MB, Shapiro RW. "Double depression": superimposition of acute depressive episodes on chronic depressive disorders. *Am J Psychiatry.* 1982; **139**(4): 438-442.
- Mueller TI, Leon AC. Recovery, chronicity, and levels of psychopathology in major depression. *Psychiatr Clin North Am.* 1996; **19**(1): 85-102.
- Fava M, Rush AJ, Trivedi MH, et al. Background and rationale for the Sequenced Treatment Alternatives to Relieve Depression (STAR*D) study. *Psychiatr Clin North Am.* 2003; **26**(2): 457-494; x.
- Murrough JW, Charney DS. Is there anything really novel on the antidepressant horizon? *Curr Psychiatry Rep.* 2012; **14**(6): 643-649.
- Schwartz TL, Stahl SM. Optimizing antidepressant management of depression: current status and future perspectives. In Cryan JF, Leonard BE, eds. *Depression: From Psychopathology to Pharmacotherapy*. Vol. 27. Basel, Switzerland: Karger; 2010: 54-67.
- Sacher J, Rabiner EA, Clark M, et al. Dynamic, adaptive changes in MAO-A binding after alterations in substrate availability: an in vivo [(11)C]-harmine positron emission tomography study. *J Cereb Blood Flow Metab.* 2012; **32**(3): 443-446.
- Howes OD, Kapur S. The dopamine hypothesis of schizophrenia: version III—the final common pathway. *Schizophr Bull.* 2009; **35**(3): 549-562.
- Du L, Faludi G, Pakovits M, et al. High activity-related allele of MAO-A gene associated with depressed suicide in males. *Neuroreport.* 2002; **13**: 1195-1198.
- De Luca V, Tharmalingam S, Sicard T, Kennedy JL. Gene-gene interaction between MAOA and COMT in suicidal behavior. *Neurosci Lett.* 2005; **383**(1-2): 151-154.
- Johnson S, Stockmeier CA, Meyer JH, et al. The reduction of R1, a novel repressor protein for monoamine oxidase A, in major depressive disorder. *Neuropsychopharmacology.* 2011; **36**(10): 2139-2148.
- Meyer JH. Neuroimaging markers of cellular function in major depressive disorder: implications for therapeutics, personalized medicine, and prevention. *Clin Pharmacol Ther.* 2012; **91**(2): 201-214.

21. Meyer JH, Wilson AA, Sagrati S, *et al.* Brain monoamine oxidase A binding in major depressive disorder: relationship to selective serotonin reuptake inhibitor treatment, recovery, and recurrence. *Arch Gen Psychiatry.* 2009; **66**(12): 1304-1312.
22. Rush AJ, Trivedi MH, Wisniewski SR, *et al.* Acute and longer-term outcomes in depressed outpatients requiring one or several treatment steps: a STAR*D report. *Am J Psychiatry.* 2006; **163**(11): 1905-1917.
23. Zajecka JM, Goldstein C. Combining medications to achieve remission. In Schwartz TL, Petersen TJ, eds. *Depression: Treatment Strategies and Management*, 2nd Ed. New York: Informa; 2010; 54-100.
24. Pitychoutis PM, Zisaki A, Dalla C, Papadopoulou-Daifoti Z. Pharmacogenetic insights into depression and antidepressant response: does sex matter? *Curr Pharm Des.* 2010; **16**(20): 2214-2223.
25. Fan M, Liu B, Jiang T, *et al.* Meta-analysis of the association between the monoamine oxidase-A gene and mood disorders. *Psychiatr Genet.* 2010; **20**(1): 1-7.
26. Gutiérrez B, Arias B, Gastó C, *et al.* Association analysis between a functional polymorphism in the monoamine oxidase A gene promoter and severe mood disorders. *Psychiatr Genet.* 2004; **14**(4): 203-208.
27. Kinnally EL, Huang YY, Haverly R, *et al.* Parental care moderates the influence of MAOA-uVNTR genotype and childhood stressors on trait impulsivity and aggression in adult women. *Psychiatr Genet.* 2009; **19**(3): 126-133.
28. Lung FW, Tzeng DS, Huang MF, Lee MB. Association of the MAOA promoter uVNTR polymorphism with suicide attempts in patients with major depressive disorder. *BMC Med Genet.* 2011; **12**: 74.
29. Schulze TC, Müller DJ, Krauss H, *et al.* Association between a functional polymorphism in the monoamine oxidase A gene promoter and major depressive disorder. *Am J Med Genet.* 2000; **96**(6): 801-803.
30. Xu Z, Zhang Z, Shi Y, *et al.* Influence and interaction of genetic polymorphisms in catecholamine neurotransmitter systems and early life stress on antidepressant drug response. *J Affect Disord.* 2011; **133**(1-2): 165-173.
31. Buckholtz JW, Callicott JH, Kolachana B, *et al.* Genetic variation in MAOA modulates ventromedial prefrontal circuitry mediating individual differences in human personality. *Mol Psychiatry.* 2008; **13**(3): 313-324.
32. Schwartz TL. Psychopharmacological practice: the DSM versus the brain. *Mens Sana Monogr.* 2013; **11**(1): 25-41.
33. Bremner JD, Narayan M, Anderson ER, *et al.* Hippocampal volume reduction in major depression. *Am J Psychiatry.* 2000; **157**(1): 115-118.
34. Sheline YI, Sanghavi M, Mintun MA, Gado MH. Depression duration but not age predicts hippocampal volume loss in medically healthy women with recurrent major depression. *J Neurosci.* 1999; **19**(12): 5034-5043.
35. Andrezza AC, Shao L, Wang JF, Young LT. Mitochondrial complex I activity and oxidative damage to mitochondrial proteins in the prefrontal cortex of patients with bipolar disorder. *Arch Gen Psychiatry.* 2010; **67**(4): 360-368.
36. Thalmeier A, Dickmann M, Giegling I, *et al.* Gene expression profiling of post-mortem orbitofrontal cortex in violent suicide victims. *Int J Neuropsychopharmacol.* 2008; **11**(2): 217-228.
37. Sacher J, Wilson AA, Houle S, *et al.* Elevated brain monoamine oxidase A binding in the early postpartum period. *Arch Gen Psychiatry.* 2010; **67**(5): 468-474.
38. Keating C, Tilbrook A, Kulkarni J. Oestrogen: an overlooked mediator in the neuropsychopharmacology of treatment response? *Int J Neuropsychopharmacol.* 2011; **14**(4): 553-566.
39. Meyers B, D'Agostino A, Walker J, Kritzer MF. Gonadectomy and hormone replacement exert region- and enzyme isoform-specific effects on monoamine oxidase and catechol-O-methyltransferase activity in prefrontal cortex and neostriatum of adult male rats. *Neuroscience.* 2010; **165**(3): 850-862.
40. Wu JB, Chen K, Li Y, Lau YF, Shih JC. Regulation of monoamine oxidase A by the SRY gene on the Y chromosome. *FASEB J.* 2009; **23**(11): 4029-4038.
41. Schwartz TL. Introduction to the special issue focused on the future or psychopharmacological practice. *Clinical Neuropsychiatry.* 2011; **8**(1): 3.
42. Garriock HA, Moreno FA. Genetics of depression: implications for clinical practice yet? *Clinical Neuropsychiatry.* 2011; **8**(1): 37-46.
43. Gillman PK. Monoamine oxidase inhibitors, opioid analgesics and serotonin toxicity. *Br J Anaesth.* 2005; **95**(4): 434-441.
44. Stahl SM, Felker A. Monoamine oxidase inhibitors: a modern guide to an unrequited class of antidepressants. *CNS Spectr.* 2008; **13**(10): 855-870.
45. Ansari J. Drug interaction and pharmacist. *J Young Pharm.* 2010; **2**(3): 326-331.

CME Posttest and Certificate

CME Credit Expires: November 30, 2016

CME posttest study guide

NOTE: The posttest can only be submitted online. The below posttest questions have been provided solely as a study tool to prepare for your online submission. **Faxed/mailed copies of the posttest cannot be processed** and will be returned to the sender. If you do not have access to a computer, contact customer service at 888-535-5600.

- Jackson is a 40-year-old male patient with treatment-resistant depression. In addition to depressed mood, he presents with the complaint of inattention and inability to concentrate. Inattention, poor concentration, and loss of vigilance may be due to:
 - Excess MAO activity in the prefrontal cortex
 - Excess MAO activity in the anterior cingulate cortex
 - Deficient MAO activity in the prefrontal cortex
 - Deficient MAO activity in the anterior cingulate cortex
- Sarah is a 31-year-old patient with treatment-resistant depression. She has recently started taking a monoamine oxidase inhibitor (MAOI) and is having a promising therapeutic response. Females may have increased monoamine oxidase activity compared to males due to:
 - Incomplete X-inactivation
 - The presence of the SRY protein
 - Both of the above
 - Neither of the above
- A 37-year-old man with a history of treatment-resistant depression has been successfully treated with a monoamine oxidase inhibitor (MAOI) for the last year. He recently suffered a broken wrist while playing tennis and is in quite a bit of pain. Which of the following would be an acceptable pain management option for this patient?
 - Hydrocodone
 - Meperidine
 - Tramadol
 - The patient cannot take any of these medications
- Marlena is a 29-year-old patient with major depressive disorder. She has had little therapeutic benefit on previous trials of various SSRIs, SNRIs, or TCAs. You would like to start her on an MAOI, but she is reluctant because she is a vegetarian and does not want to have to give up her favorite foods, including cottage cheese, peanuts, and tofu. You inform her that while taking an MAOI, she would need to avoid:
 - Cottage cheese
 - Peanuts
 - Tofu
 - Cottage cheese and peanuts
 - Cottage cheese and tofu
 - Peanuts and tofu
 - All of the above

CME online posttest and certificate

To receive your certificate of CME credit or participation, complete the posttest and activity evaluation, available only online at <http://www.neiglobal.com/CME> under “CNS Spectrums.” If a passing score of 70% or more is attained (required to receive credit), you can immediately print your certificate. There is no posttest fee. Questions? call 888-535-5600 or email customerservice@neiglobal.com.

Posttest & Evaluation