



Ten Things Physicians and Patients Should Question

1 Don't use homeopathic medications, non-vitamin dietary supplements or herbal supplements as treatments for disease or preventive health measures.

Alternative therapies are often assumed safe and effective just because they are “natural.” There is a lack of stringent quality control of the ingredients present in many herbal and dietary supplements. Reliable evidence that these products are effective is often lacking, but substantial evidence exists that they may produce harm. Indirect health risks also occur when these products delay or replace more effective forms of treatment or when they compromise the efficacy of conventional medicines.

2 Don't administer a chelating agent prior to testing urine for metals, a practice referred to as “provoked” urine testing.

Metals are ubiquitous in the environment and all individuals are exposed to and store some quantity of metals in the body. These do not necessarily result in illness. Scientific studies demonstrate that administration of a chelating agent leads to increased excretion of various metals into the urine, even in healthy individuals without metal-related disease. These “provoked” or “challenge” tests of urine are not reliable means to diagnose metal poisoning and have been associated with harm.

3 Don't order heavy metal screening tests to assess non-specific symptoms in the absence of excessive exposure to metals.

Individuals are constantly exposed to metals in the environment and often have detectable levels without being poisoned. Indiscriminant testing leads to needless concern when a test returns outside of a “normal” range. Diagnosis of any metal poisoning requires an appropriate exposure history and clinical findings consistent with poisoning by that metal. A patient should only undergo specific metal testing if there is concern for a specific poisoning based on history and physical examination findings.

4 Don't recommend chelation except for documented metal intoxication which has been diagnosed using validated tests in appropriate biological samples.

Chelation does not improve objective outcomes in autism, cardiovascular disease or neurodegenerative conditions like Alzheimer's disease. Edetate disodium is not FDA-approved for any condition. Even when used for appropriately diagnosed metal intoxication, chelating drugs may have significant side effects, including dehydration, hypocalcemia, kidney injury, liver enzyme elevations, hypotension, allergic reactions and essential mineral deficiencies. Inappropriate chelation, which may cost hundreds to thousands of dollars, risks these harms, as well as neurodevelopmental toxicity, teratogenicity and death.

5 Don't remove mercury-containing dental amalgams.

Mercury-containing dental amalgams release small amounts of mercury. Randomized clinical trials demonstrate that the mercury present in amalgams does not produce illness. Removal of such amalgams is unnecessary, expensive and subjects the individual to absorption of greater doses of mercury than if left in place.



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6

Don't use phenytoin or fosphenytoin to treat seizures caused by drug toxicity or drug withdrawal.

With rare exceptions, phenytoin is ineffective for convulsions caused by drug or medication toxicity. Phenytoin has been demonstrated to be ineffective for the treatment of isoniazid-induced seizures and withdrawal seizures and may potentially be harmful when used to treat seizures induced by theophylline or cyclic antidepressants. First-line treatment of toxin-induced seizures and withdrawal seizures is benzodiazepines, followed by additional medications that act through agonism at the GABA A receptor, such as barbiturates.

7

Don't recommend "detoxification" through colon cleansing or promoting sweating for disease treatment or prevention.

No objective scientific evidence supports a role for colonic irrigation for "detoxification." No US FDA-approved colonic hydrotherapy systems exist for nonmedical purposes like colon cleansing. Colonic cleansing through hydrotherapy, laxatives or cathartics may result in cramping, pain, dehydration, electrolyte imbalances, infections and bowel perforation. Promoting sweating doesn't produce clinically relevant toxin elimination. Methods to promote sweating may cause heat stroke, dehydration, burns, myocardial injury, carbon monoxide poisoning and liver or kidney damage, which might compromise toxin elimination.

8

Don't order tests to evaluate for or diagnose "idiopathic environmental intolerances," "electromagnetic hypersensitivity" or "mold toxicosis."

These diagnoses reflect labels to indicate that patients have adverse non-allergic reactions to normal environmental stimuli. These diagnoses are made on the bases of self-reported symptoms or non-validated testing procedures. Although these conditions have been widely promoted, evidence-based assessments fail to support these diagnoses as disease entities. Labeling a patient with these diagnoses may adversely affect the patient's lifestyle, obscure ascertainment of the etiology of their symptoms and promote unnecessary testing.

9

Don't perform hair or nail testing for "metal poisoning" screening in patients with nonspecific symptoms.

The proper clinical assessment for potential exposure to metals must consider the precise exposure, symptoms, signs, route of exposure and dose. Hair and nail testing are rarely required, frequently unreliable and provide limited utility after metal exposures. A patient should undergo tailored testing for a specific metal exposure based on an appropriate evaluation. Non-specific hair and nail testing for multiple metals subjects patients to potentially harmful diagnostic mislabeling and subsequent detrimental therapy.

10

Don't perform fasciotomy in patients with snake envenomation absent direct measurement of elevated intracompartmental pressures.

Crotalinae snakebites produce findings mimicking compartment syndrome that are rarely indicative of actual compartment syndrome. Myonecrosis results from venom toxicity rather than elevated compartment pressures. Fasciotomy does not prevent, and may worsen, necrosis. In some cases with elevated compartment pressures, treatment with antivenom and without fasciotomy was successful. No available evidence indicates when fasciotomy should be performed in the management of snakebites. If considered, fasciotomy should not be performed without first documenting elevated compartment pressure.

How This List Was Created

The American College of Medical Toxicology's (ACMT's) Board of Directors established a *Choosing Wisely*[®] work group in 2013 to develop a list of items for the *Choosing Wisely*[®] campaign. Members of the work group were chosen to represent various practice settings within the field of medical toxicology, including ambulatory, acute and population-based practice. Work group members included the President of the College, the Chair of the Practice Committee, the Chair of the Positions and Guidelines committee and other academic leaders within the medical toxicology community. All work group members also represented the American Academy of Clinical Toxicology (AACT). The first list was released by the work group in 2013 and in 2014, the work group reconvened to develop a second list of items for the campaign. A second preliminary list was disseminated to all members of ACMT and AACT for review, commentary and potential additions. Additional feedback was solicited from leaders within the field of medical toxicology. The work group reviewed all responses, and narrowed the list to the final five items based on a review of scientific evidence, relevance to the specialty and greatest opportunity to improve care, reduce cost and reduce harm to patients. The final list was approved by the ACMT Board of Directors and the AACT Board of Trustees.

The ACMT and AACT disclosure and conflict of interest policies can be found at www.acmt.net and www.clintox.org respectively.

Sources

- Woodward KN. The potential impact of the use of the homeopathic and herbal medicines on monitoring the safety of prescription products. *Hum Exp Toxicol*. 2005;24:219–33.
Thompson E, Barron S, Spence D. A preliminary audit investigating remedy reactions including adverse events in routine homeopathic practice. *Homeopathy*. 2004;93:203–9.
De Smet PA. Health risks of herbal remedies. *Drug Saf*. 1995;13:81–93.
Farah MH, Edwards R, Lindquist M, Leon C, Shaw D. International monitoring of adverse health effects associated with herbal medicines. *Pharmacoepidemiol Drug Saf*. 2000;9(2):105–12.
Drew AK, Myers SP. Safety issues in herbal medicine: implications for the health professions. *Med J Aust*. 1997;166:538–41.
- Charlton N, Wallace KL. American College of Medical Toxicology position statement on post-chelator challenge urinary metal testing. *American College of Medical Toxicology*; 2009 Jun [cited 2013 Apr 23]. Available from: http://www.acmt.net/cgi/page.cgi/zine_service.html?aid=2999&zine=show.
Risher JF, Amler SN. Mercury exposure: evaluation and intervention the inappropriate use of chelating agents in the diagnosis and treatment of putative mercury poisoning. *Neurotoxicology*. 2005 Aug;26(4):691–9.
- McKay C, Holland M, Nelson L. A call to arms for medical toxicologists: the dose, not the detection, makes the poison. *Internet J Med Toxicol*. 2003;6(1):1.
Schober SE, Sinks TH, Jones RL, Bolger PM, McDowell M, Osterloh J, Garrett ES, Canady RA, Dillon CF, Sun Y, Joseph CB, Mahaffey KR. Blood mercury levels in US children and women of childbearing age, 1999–2000. *JAMA*. 2003;289(13):1667–74.
- Nonstandard uses of chelation therapy. *Med Lett Drugs Ther*. 2010 Sep 20;52(1347):75–6.
Kosnett MJ. Chelation for heavy metals (arsenic, lead, and mercury): protective or perilous? *Clin Pharmacol Ther*. 2010 Sep;88(3):412–5.
Nissen SE. Concerns about reliability in the Trial to Assess Chelation Therapy (TACT). *JAMA*. 2013 Mar 27;309(12):1293–4.
Risher JF, Amler SN. Mercury exposure: evaluation and intervention the inappropriate use of chelating agents in the diagnosis and treatment of putative mercury poisoning. *Neurotoxicology*. 2005 Aug;26(4):691–9.
U.S. Food and Drug Administration. FDA warns marketers of unapproved 'chelation' drugs. *FDA Consumer Health Information*. 2010 October;1.
- Bellinger DC, Trachtenberg F, Barregard L, Tavares M, Cernichiari E, Daniel D, McKinlay S. Neuropsychological and renal effects of dental amalgam in children. A randomized clinical trial. *JAMA*. 2006 Apr 19;295(15):1775–83.
Factor-Litvak P, Hasselgren G, Jacobs D, Begg M, Kline J, Geier J, Mervish N, Schoenholtz S, Graziano J. Mercury derived from dental amalgams and neuropsychologic function. *Environ Health Persp*. 2003 May;111(5):719–23.
- Goldberg MJ, Spector R, Miller G. Phenobarbital improves survival in theophylline-intoxicated rabbits. *J Toxicol Clin Toxicol*. 1986;24(3):203–11.
Blake KV, Massey KL, Hendeles L, Nickerson D, Neims A. Relative efficacy of phenytoin and phenobarbital for the prevention of theophylline-induced seizures in mice. *Ann Emerg Med*. 1988 Oct;17(10):1024–8.
Miller J, Robinson A, Percy AK. Acute isoniazid poisoning in childhood. *Am J Dis Child*. 1980 Mar;134(3):290–2.
Saad SF, el-Masry AM, Scott PM. Influence of certain anticonvulsants on the concentration of gamma-aminobutyric acid in the cerebral hemispheres of mice. *Eur J Pharmacol* 1972 Mar;17(3):386–92.
Okamoto M, Rosenberg HC, Boisse NR. Evaluation of anticonvulsants in barbiturate withdrawal. *J Pharmacol Exp Ther*. 1977 Aug;202(2):479–89.
Chance JF. Emergency department treatment of alcohol withdrawal seizures with phenytoin. *Ann Emerg Med*. 1991 May;20:520–2.
Sharma AN, Hoffman RJ. Toxin-related seizures. *Emerg Med Clin North Am*. 2011 Feb;29(1):125–39.
Hung OL, Shih RD. Antiepileptic drugs: the old and the new. *Emerg Med Clin North Am*. 2011 Feb;29(1):141–50.
- Colon cleansing. *Med Lett Drugs Ther*. 2009 May 18;51(1312):39–40.
Acosta RD, Cash BD. Clinical effects of colonic cleansing for general health promotion: a systematic review. *Am J Gastroenterol*. 2009 Nov;104(11):2830–6.
Kenttämies A, Karkola K. Death in sauna. *J Forensic Sci*. 2008 May;53(3):724–9.
Mishori R, Otubu A, Jones AA. The dangers of colon cleansing. *J Fam Pract*. 2011 Aug;60(8):454–7.
Rodhe A, Eriksson A. Sauna deaths in Sweden, 1992–2003. *Am J Forensic Med Pathol*. 2008 Mar;29(1):27–31.

8

Baliatsas C, VanKamp I, Lebre E, Rubin GJ. Idiopathic environmental intolerance attributed to electromagnetic fields (IEI-EMF): a systematic review of identifying criteria. *BMC Public Health*. 2012 Aug 11;12:643.

Boyd I, Rubin G, Wessely S. Taking refuge from modernity: 21st century hermits. *J R Soc Med*. 2012 Dec;105:523-9.

Hausteiner C, Bornschein S, Zilker T, Henningsen P, Förstl H. Dysfunctional cognitions in idiopathic environmental intolerances (IEI) - an integrative psychiatric perspective. *Toxicol Lett*. 2007 Jun 15;171(1-2):1-9.

Rubin GJ, Hillert L, Nieto-Hernandez R, van Rongen E, Oftedal G. Do people with idiopathic environmental intolerance attributed to electromagnetic fields display physiological effects when exposed to electromagnetic fields? A systematic review of provocation studies. *Bioelectromagnetics*. 2011 Dec;32(8):593-609.

Staudenmayer H, Binkley KE, Leznoff A, Phillips S. Idiopathic environmental intolerance: Part 2: causation analysis applying Bradford Hill's criteria to the psychogenic theory. *Toxicol Rev*. 2003;22:247-61.

Staudenmayer H, Binkley KE, Leznoff A, Phillips S. Idiopathic environmental intolerance: Part 1: a causation analysis applying Bradford Hill's criteria to the toxicogenic theory. *Toxicol Rev*. 2003;22:235-46.

9

Calabrese EJ. Hormesis is central to toxicology, pharmacology and risk assessment. *Hum Exp Toxicol*. 2010 Apr;29(4):249-61.

Frisch M, Schwartz BS. The pitfalls of hair analysis for toxicants in clinical practice: three case reports. *Environ Health Perspect*. 2002 Apr;110(4):433-6.

Seidel S, Kreutzer R, Smith D, McNeel S, Gilliss D. Assessment of commercial laboratories performing hair mineral analysis. *JAMA*. 2001 Jan;285:67-72.

Steindel SJ, Howanitz PJ. The uncertainty of hair analysis for trace metals. *JAMA*. 2001 Jan 3;285(1):83-5.

10

Cumpston KL. Is there a role for fasciotomy in Crotalinae envenomations in North America? *Clin Toxicol (Phila)*. 2011 Jun;49(5):351-65.

Hall EL. Role of surgical intervention in the management of crotaline snake envenomation. *Ann Emerg Med*. 2001 Feb;37:175-80.

Tanen DA, Danish DC, Grice GA, Riffenburgh RH, Clark RF. Fasciotomy worsens the amount of myonecrosis in a porcine model of crotaline envenomation. *Ann Emerg Med*. 2004 Aug;44(2):99-104.

Bucarechi F, De Capitani EM, Hyslop S, Mello SM, Fernandes CB, Bergo F, Nascimento FB. Compartment syndrome after South American rattlesnake (*Crotalus durissus terrificus*) envenomation. *Clin Toxicol (Phila)*. 2014 Jul;52(6):639-41.

Mazer-Amirshahi M, Boutsikaris A, Clancy C. Elevated compartment pressures from copperhead envenomation successfully treated with antivenin. *J Emerg Med*. 2014 Jan;46(1):34-7.

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About the American College of Medical Toxicology and the American Academy of Clinical Toxicology

The American College of Medical Toxicology (ACMT) is an association of physicians with recognized expertise in the diagnosis, management and prevention of human poisoning and other adverse health effects due to medications, occupational and environmental toxins and biological agents. ACMT's mission is to advance quality care of poisoned patients and public health through physicians who specialize in consultative, emergency, environmental, forensic and occupational toxicology. ACMT values the importance of research and evidence based practice in combating human poisoning.

The American Academy of Clinical Toxicology (AACT) is a multidisciplinary organization uniting scientists and clinicians in the advancement of research, education, prevention and treatment of diseases caused by chemicals, drugs and toxins. AACT's mission is to promote the study of health effects of poisons, encourage the development of new therapies and treatment in clinical toxicology, and define the position of clinical toxicologists on toxicology-related issues.

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