

The Safest and Most Effective Solution for the Candidiasis and Mercury Cycle

By Lyn Hanshew, M.D.

One of the most difficult organisms to diagnose and successfully treat is *Candida albicans*. *C. albicans* is a yeast that is considered part of the normal flora of the alimentary canal and is also, commonly found in the genitourinary system. By six months of age, 90% of infants test positive for *Candida* and almost all adults test positive. *C. albicans* is an opportunistic organism and exists in a competitive environment with other organisms to control its growth. This equilibrium in the gut and genitourinary tract can be disrupted by changes in the pH, high levels of simple sugars, hormone imbalance, antibiotic exposure that alters the make-up and quantity of the flora constituents and suppression of the immune system by toxins and drugs such as steroids. If this happens, overgrowth of *Candida* can occur and this organism can burrow through the walls of the intestine, enter the blood stream and make its way throughout the body.

Continued on page 2

In This Issue

The Safest and Most Effective Solution for the Candidiasis and Mercury Cycle

..... 2

Results RNA
1 888 823 3869
www.resultsrna.com

The Safest and Most Effective Solution for the Candidiasis and Mercury Cycle

by Lyn Hanshew, M.D.



One of the most difficult organisms to diagnose and successfully treat is *Candida albicans*. *C. albicans* is a yeast that is considered part of the normal flora of

the alimentary canal and is also, commonly found in the genitourinary system. By six months of age, 90% of infants test positive for *Candida* and almost all adults test positive. *C. albicans* is an opportunistic organism and exists in a competitive environment with other organisms to control its growth. This equilibrium in the gut and genitourinary tract can be disrupted by changes in the pH, high levels of simple sugars, hormone imbalance, antibiotic exposure that alters the makeup and quantity of the flora constituents and suppression of the immune system by toxins and drugs such as steroids. If this happens, overgrowth of *Candida* can occur and this organism can burrow through the walls of the intestine, enter the blood stream and make its way throughout the body.

Candidemia and disseminated candidiasis are the most common causes of morbidity and mortality in hospitalized patients, especially in the ICU. A predisposed host is one who is on broad spectrum antibiotics, immunosuppressive drugs, parenteral nutrition and central

venous catheters. Data was extracted from the Prospective Antifungal Therapy Alliance database and demonstrated the incidence of candidemia caused by *C. albicans* was 45.6% and 54.4% was caused by non-*C. albicans* species. The overall crude 12 week mortality was 35.2%. There are no pathognomonic signs or symptoms (Horn DL et al. 2009). The clinical clues are fever of unknown origin or signs of severe sepsis while on antibiotics, and multiple, non-tender, nodular erythematous cutaneous lesions. The evaluation of new antifungal agents and the precise role of prophylactic therapy are needed. (Singhi S. 2009, Perlthoth J, et al 2007)

Due to the ubiquitous presence of *Candida*, diagnostic testing is of little help. There are a variety of tests that practitioners use to diagnose yeast overgrowth, that may include stool tests, blood tests, live blood cell tests, etc., but none of these tests are reliable. They may or may not detect an infection of Candidiasis. (Pappas PG 2004)

Therefore, practitioners often diagnose and individuals often self-diagnose based upon ambulatory versus hospitalized status and a list of common symptoms associated with the gastrointestinal and genitourinary systems, immune, endocrine and the neurological system, including mental and emotional symptoms. This list includes: thrush, indigestion,

acid reflux, abdominal gas and bloating, diarrhea, constipation, rectal itching, vaginitis headaches, migraines, excessive fatigue, inability to think clearly or concentrate, poor memory, hyperactivity, mood swings cravings for alcohol or sweets, anxiety, depression, irritability, dizziness, itching, acne, eczema, athlete's foot, sinus inflammation, persistent cough, sore throat, earache, pre-menstrual syndrome, low sex drive, muscle weakness, sensitivity to fragrances and/or other chemicals and chronic pain. (Crook W 1983, 2005) This wide-ranging list is significant and indicative of the toxicity associated with *Candida*.

A successful treatment strategy includes killing the organism and correcting the underlying causes for *Candida* overgrowth and dissemination. Pharmaceutical anti-fungals agents can be grouped into three classes based on their site of action: azoles which inhibit the synthesis of ergosterol, a major constituent of the fungal cell membrane, are considered fungistatic; polyenes also interact with ergosterol and cause membrane leakage resulting in cell death and 5-fluorocytosine inhibits protein synthesis.

Nystatin is a polyene, oral anti-fungal used primarily for the treatment of oral candidiasis. It is not absorbed out of the gut and has few mild gastrointestinal side effects. Amphotericin B is a fungicidal polyene administered intravenously,

which commonly can cause serious multi-organ damage and result in death. It is not frequently used since the development of the azole class of agents. This class has less side effects, ranging from mild nausea, vomiting and diarrhea, headache, dizziness and skin rash to severe blistering, bruising, jaundice and seizures. Side-effects for 5-fluorocytosine include bone marrow suppression, gastrointestinal, liver, kidney and central nervous system toxicity. Efficacy and safety issues and development of resistance limits the usefulness of all of these agents.

Historically, non-pharmaceutical approaches to killing or limiting the growth of *Candida* have not been well-researched. Thai and Ignacio (2005) demonstrated that grape seed extract had comparable fungistatic activity to Miconazole in suppressing *C. albicans* growth. They found that tea tree oil, probiotic supernatant and garlic showed no decrease in growth rates of *C. albicans*.

The good news is there is now a safe and effective solution for the serious issue of candidiasis. The average person in the US has between 400-800 potentially toxic, immunosuppressive, carcinogenic, endocrine-disrupting, and gene-damaging chemicals stored in his cells and is therefore at risk for candidiasis. Mercury, lead, cadmium, arsenic, pesticides, insecticides, dioxins, furans, phthalates, VOCs, and PCBs are just some of the foreign substances that have created an excessive toxic body burden of harmful chemicals.

Of all the toxins, mercury is the most destructive to the neurological, immune and endocrine systems. It is a deadly mutagen causing DNA damage. Mercury contributes to or causes many

illnesses including autism, autoimmune diseases, Alzheimer's disease, cancers, heart disease, endocrine problems, and neurological and behavioral disorders. Murray and Kidby (1975) demonstrated that Mercury is incorporated into the cell wall of yeast. This is the key to the toxic effects of candidiasis and the basis of the Herxheimer reaction. Mercury being released from dying cells is extremely toxic to surrounding tissues. The list of symptoms associated with candidiasis is the same list of symptoms associated with mercury toxicity. It is impossible to cure candida overgrowth without removing the toxic burden of mercury.

Prior to the discovery of antibiotics, health practitioners used colloidal silver to safely and effectively treat infections. In 1914 the medical journal *Lancet* reported phenomenal results from silver use stating it to be absolutely harmless, non-toxic to humans, and highly germicidal. In 1929, over 5 million prescriptions for silver-based products were issued in the United States alone. In fact, colloidal silver has been proven useful against all species of fungi, parasites, bacteria, protozoa, and viruses. Even to this day, properly formulated colloidal silver is still one of the most effective, safe, antimicrobials known to man with no risk of resistance developing.

To effectively reduce toxic body burden of harmful toxins and infections, Results RNA® has created Total Body Detox® composed of two revolutionary intra-oral spray formulas, Advanced Cellular Silver (ACS) 200® Extra Strength and Advanced Cellular Zeolite (ACZ) nano® Extra Strength to effectively and safely kill pathogens and remove toxic body burden. ACZ nano® Extra Strength and ACS 200®

Extra Strength have extraordinary characteristics.

ACS 200® Extra Strength represents a major advancement in medical-use silver technology and excels in both safety and efficacy. ACS 200® Extra Strength demonstrates a much broader pathogen kill spectrum than traditional prescription antibiotics, antifungal, or antiviral preparations. Far more advanced in both safety and efficacy than traditional colloidal silver, ACS 200® Extra Strength is a 200 ppm (parts per million) silver that has been proven capable of rapidly killing an enormous array of disease-causing organisms, literally oxidizing the cell wall of Gram-positive and Gram-negative bacteria, as well as viruses, fungi, parasites and spirochetes. Independent studies of ACS 200® demonstrate "complete kill" against *Candida* in less than 3 minutes. An FDA protocol oral toxicity independent study was conducted by Pacific BioLabs in Hercules, CA. Using mega doses of ACS 200®, there were no toxic signs observed throughout the study.

Now that the *Candida* organisms are dead, how do we address the mercury that has been released? ACZ nano® Extra Strength is a zeolite product that has many significant qualities which make it a superior choice over other detoxification or chelation methods, including other zeolite-based products. In urine challenge studies, ACZ nano® Extra Strength has been independently proven to increase urinary output of mercury, lead and other toxic metals by several thousand percent. It is interesting to note that extremely toxic mercury levels were recorded in the urine of patients while taking ACZ nano® Extra Strength who had undetectable mercury levels in their baseline urine.

These results show just how difficult it is for the body to remove mercury and other toxins without an effective chelator present. Traditional chelating agents have significant limitations in safely removing mercury, lead, cadmium and arsenic. One drawback is that agents such as EDTA have high affinity for essential nutrient minerals such as calcium and remove them simultaneously with toxins. If not carefully monitored, this removal of calcium can be quite dangerous and can cause rapid muscle weakness and potentially heart damage. With a weak bond, these chelating agents can “pull” out mercury from the tissues and then “drop” mercury into the bloodstream where it can redeposit in the brain or other vital organs. If this happens the patient’s condition is likely to worsen. ACZ nano® Extra Strength irreversibly binds toxins and safely removes them through the urinary tract. This prevents Herxheimer reactions and prevents mercury from being recycled into other Candida cells.

Total Body Detox® comprised of ACS 200® Extra Strength to safely and effectively kill Candida and ACZ nano® Extra Strength to irreversibly bind and excrete the mercury and other toxins that are released, is the safest and most effective solution to the serious health threat of candidiasis.

About the Author

Dr. Hanshew practiced medicine in the Seattle area for 15 years. She achieved Board-Certified in Family Medicine and Bariatric Medicine. She also has specialized training in Anti-Aging Medicine, Natural Hormone Replacement and Environmental Toxicity issues relating to the exponential rise in the incidence and successful treatment of Lyme disease,

Autism, Fibromyalgia, ADD, Chronic Fatigue, Multiple Sclerosis, Obesity, Anxiety, Depression and Cancer.

References

Crook W. The Yeast Connection. 1983

Crook W, Dean C, Cass H, Crook E. The Yeast Connection and Women’s Health. 2005

Horn DL et al. Epidemiology and outcomes of candidemia in 2019 patients: data from the prospective antifungal therapy alliance registry.. Clin Infect Dis. 2009 Jun 15;48(12):1695-703

Ignacio,C and Thai,D. Comparative analysis of antifungal activity of natural remedies versus miconazole nitrate salt against Candida albicans. Biological Sciences Dept. California Polytechnical State Institute. 2005

Murray and Kidby. Sub-cellular Location of Mercury in Yeast Grown in the Presence of Mercuric Chloride” 1975

Pappas,PG Rex JH, Sobel,JD et al. Guidelines for Treatment of Candidiasis. Clinical Infectious Diseases 2004;38:161-89

Perkins,Cynthia. The Candida Solution.

PerltrothJ, Choi B, Spellberg B. Nosocomial fungal infections:epidemiology, diagnosis and treatment. Med Mycol. 2007 Jun;45(4);321-46

Singhi S and Deep A. Invasive candidiasis in pediatric intensive care units. Indian J Pediatrics 2009 Oct;76(10): 1033-44 Epub 2009 Nov12

Truss CO. “Tissue injury induced by

Candida albicans: Mental and neurologic manifestations.” Journal of Orthomolecular Psychiatry 1978 7:1, 17:37