Stop Fluoridation Now: New Research on Fluoride's Brain and Thyroid Toxicity

by Gary Null, PhD, and Martin Feldman, MD

Several years have passed since we published a three-part article on water fluoridation entitled "The Fluoride Controversy Continues: An Update" in the *Townsend Letter* (see the December 2002, January 2003 and February/March 2003 issues). Since that time, new research has emerged that offers health-care professionals and patients alike a more thorough understanding of fluoride's adverse effects on the body.

In this article, we will look at the effects of fluoride on two vital aspects of the body's functioning – the central nervous system and brain and the thyroid mechanism – in more depth than we did in our prior article. We will also explore a newly emerging area of research, which shows that xylitol, a natural sugar substitute used in chewing gums and other products, provides consumers with an effective alternative to the fluoridation of public water supply systems in preventing dental cavities.

reported As we previously, approximately 162 million Americans on public water systems were receiving fluoridated water in 2000.1,2 The US government strongly supports water fluoridation on the grounds that it decreases the rate of dental decay.^{3, 4} Yet studies conducted worldwide have failed to validate this claim. In one large study conducted in 1986/87 for example, there was no statistically significant difference in dental decay rates between fluoridated and nonfluoridated cities.⁵ Other research has found a decline in the dental caries rate in countries that do not have fluoridated water.

The research also shows that fluoride can have a wide range of negative health effects. Many of these effects were discussed in our previous article, where readers can find a full investigation of the topic and supporting references from the scientific literature. By way of review, the following list summarizes some of the major effects of fluoride on the body. This accounting should raise serious questions in anyone's mind about the practice of adding fluoride – which, in the case of water fluoridation, is a toxic waste product of the aluminum and fertilizer industries – to community water systems:

• **Dental fluorosis.** The prevalence of dental fluorosis, in which the teeth become permanently stained, brown and mottled due to fluoride exposure, has increased in both fluoridated and nonfluoridated communities in recent decades.⁶

• Skeletal fluorosis. Fluoride that is deposited in the bones and teeth can cause this crippling disorder. Some experts have suggested that while skeletal fluorosis is rarely reported in the US medical literature, cases may go unrecognized due to a lack of knowledge about the disorder among physicians.⁷

• **Bone fractures.** Studies have found an association between fluoride and the rate of hip fractures among the elderly.⁸⁻¹⁰

• **Cancer.** Links to fluoridation have been revealed in numerous studies, with one finding that about two-thirds of 36 cancer sites in the body were associated with fluoridated water.¹¹

• Enzyme toxicity and genetic damage. Even 1 ppm of fluoride – an amount deemed safe for water fluoridation – can interfere with biological functions such as DNA repair enzyme activity and cause genetic and chromosomal damage.^{12, 13}

• **Reproductive effects.** Fluoride may have negative effects on the male and female reproductive systems, according to several studies.¹⁴⁻¹⁶

• **Pineal gland effects.** Fluoride's adverse impact on this gland may interfere with the hormone melatonin, which regulates sleep cycles, the onset of puberty and other functions.¹⁷

• Elevated blood lead levels. Increased levels of lead in the blood of children have been associated with fluoridated drinking water. High levels of lead, in turn, have been linked to certain health disorders and negative behavioral traits.^{18, 19}

Brain effects of fluoride

One growing area of research has examined the effects of fluoride on the central nervous system and brain. Recent studies have found that fluoride is a neurotoxin which can affect cerebral functioning, with some research associating the ingestion of fluoridated water with reduced intelligence. Scientific references in this area from human and animal studies deserve careful consideration from physicians and consumers who want to know how the multiple sources of fluoride we are exposed to – fluoridated water, fluoridated toothpaste, fruit juices and soft drinks, infant foods, and others $-\max$ affect such a critical aspect of the body's functioning.

A number of studies on the effect of fluoridated water on children's intelligence come from China, where investigators have compared the Intelligence Quotient (IQ) of young people living in high- and low-fluoride areas. These studies include:

• The mean IQ of 222 children, aged 8 to 13 years, in a high-fluoride village was significantly lower (92.02 \pm 13.00) than that of 290 children in a low-fluoride village (100.41 \pm 13.21). Higher levels of fluoride in drinking water were significantly associated with higher rates of mental retardation (IQ <70) and borderline intelligence (IQ 70-79). The researchers concluded that "drinking water fluoride levels greater than 1.0 mg/L may adversely affect the development of children's intelligence."²⁰

• The mean IQ of 60 children, aged 10 to 12 years, in an area with a high level of fluoride in drinking water was significantly lower (92.27 \pm 20.45) than that of 58 children in a low-fluoride area (103.05 \pm 13.86). The high-fluoride area also had more children (21.6%) in the retardation or borderline categories of IQ than did the low-fluoride area (3.4%).²¹

• Among 907 children, aged 8 to 13 years, there was a 15 to 19 point decrease in IQ in children living in an area with a high fluoride exposure compared with those in an area with little or no exposure. Exposure to a high level of fluoride may affect intelligence at an early, rapid stage of development in the embryo and infant.²²

In addition to studies on intelligence, other research on humans has associated fluoride with problems in brain functioning. A study of children who grew up in a coal-burning-pattern high-fluoride area in China found that excessive fluoride intake since early childhood would reduce mental work capacity.23 A study in Mexico found that while children's IQ scores were not influenced by fluoride exposure, other effects on neuropsychological development were associated with chronic exposure (the main source being tap water.) Urinary fluoride correlated positively with an increase in reaction time, which could affect the attention process, and inversely with low scores in visuospatial organization, which could have an impact on the children's reading and writing

abilities.²⁴ A study of structural fumigation workers found that while occupational exposure to sulfuryl fluoride did not cause any widespread pattern of cognitive deficits, such exposure "may be associated with subclinical effects on the central nervous system, including effects on olfactory and some cognitive functions."²⁵

Fluoride's effect on the human brain also extends to the fetus. An examination of 15 therapeutically aborted fetuses in the fifth to eighth month of gestation from a high-fluoride area documented a number of changes in the neurons, undifferentiated neuroblasts and mitochondria. The study concluded that chronic high-fluoride exposure during intrauterine life "may produce certain harmful effects on the developing brain of the fetus."²⁶

Another study shows that low concentrations of a variety of salts, including sodium fluoride, significantly reduce the thermodynamic stability of the human prion protein, and thereby may help promote its conversion to a misfolded form of the protein that accumulates in the brain. Prion diseases are fatal neurological disorders that occur in animals and humans. The human versions include Creutzfeld-Jacob disease, Gerstmann-Straussler-Scheinker disease, fatal familial insomnia and kuru.²⁷

A number of animal studies also have linked fluoride to neurotoxic effects, such as impaired learning and memory abilities, suppression of spontaneous motor activity and poor performance in motor coordination and maze tests.²⁸⁻³¹ A study by Dr. Phyllis Mullenix and colleagues at Forsyth Dental Center in Boston, published in 1995, evaluated the effects of fluoride on the developing brains of rats, using a computer pattern recognition system to quantify various aspects of the animals' behavior. Prenatal exposure to fluoride was associated with hyperactivity in offspring, and exposure during weaning and adulthood was associated with "cognitive deficits." The severity of the behavioral effects increased directly with plasma fluoride levels and concentrations in specific regions of the brain following fluoride ingestion. The researchers said of their findings: "Of course behaviors per se do not extrapolate [across species], but a generic behavioral pattern disruption as found in this rat study can be indicative of a potential for motor dysfunction, IQ deficits and/or learning disabilities in humans."32,33

One would expect these findings to spur the US government's interest in the central nervous system effects of fluoride, a substance it promotes as a safe and effective way to prevent dental caries. But that has not been the case, according to Christopher Bryson in *The Fluoride Deception*. Dr. Mullinex presented her results to government scientists and policy makers in a 1990 seminar held at the National Institutes of Health. Fourteen years later, says Bryson, "the NIH still has not funded any examination of fluoride's central-nervous-system effects and, according to one senior official, does not currently regard fluoride and central-nervous-system effects as a research priority."³⁴

In a 2004 submission to the National Research Council on fluoride's effect on the brain, Ellen Connett of the Fluoride Action Network notes that Dr. Mullinex's study "has gained support from at least six other rat studies.... While these studies have employed different methods than Mullinex et al. (1995), they are consistent in that they have also found that fluoride impacts behavior and/or learning."³⁵

For Dr. Mullinex, says Bryson, the fluoride research seems to have marked the end of her academic career. She was fired from her job at Forsyth Dental Center, where she had been chairwoman of the toxicology department for 11 years, within days of her paper's acceptance for publication. Mullinex says that her dismissal was due to conflict over her decision to publish the findings on fluoride neurotoxicity, according to a report on Salon.com. (One superior has said she was dismissed not because of the fluoride work but because of work-quality problems.)^{36,37}

Numerous other animal studies have documented the effects of fluoride toxicity on the brain. To summarize some of this research from the past few years alone, animal studies have found that: fluoride may cross the blood-brain barrier, accumulate in the hippocampus and inhibit the activity of the enzyme cholinesterase³⁸; high fluoride intake during the early developing stages of life can cause significant neurodegenerative changes in the hippocampus, amygdala, motor cortex and cerebellum³⁹; sodium fluoride can induce DNA damage and apoptosis in the brain⁴⁰; chronic fluoride intoxication during the early stages of life can increase oxidative stress in the brain, disturbing the antioxidant defense system⁴¹; fluoride inhibits some enzymes involved in free-radical metabolism and membrane functioning in the brain and in muscle42; chronic fluoride toxicity reduces the number of neuronal nicotinic acetylcholine receptors (nAChRs), which are involved in cognitive processes such as learning and memory⁴³; selective decreases in the number of nAChRs may

Fluoride's Toxicity

play an important role in the mechanism(s) by which fluoride causes central nervous system dysfunction;⁴⁴ and a high concentration of fluoride and fluoride/iodine combined in drinking water can cause significant changes in the fatty acid composition of brain cells, with a significant decrease in the proportion of unsaturated fatty acid and an obvious increase in saturated fatty acid.⁴⁵

In an editorial in Fluoride, Bruce Spittle discusses biochemical studies that have identified mechanisms by which fluoride could affect brain functioning. He summarizes the research as such: Because fluoride can form a strong hydrogen bond with the amide group, the shape of enzymes may be altered and their activity reduced. Aluminofluoride complexes stimulate various guanine nucleotide binding proteins - called G proteins - which may "mimic or potentiate the action of numerous extracellular signals and significantly affect many cellular responses. Fluoride ions in the presence of trace amounts of aluminum are apparently able to act with powerful pharmacological effects."46

In one animal study, rats administered aluminum fluoride or sodium fluoride in water had increased levels of aluminum in the brain, neural injury and increased deposits of B-amyloid protein in the brain. Similar deposits have been associated with Alzheimer's disease, according to a report in Chemical & Engineering News. The researchers stated that while the small amount of fluoroaluminum complex needed to produce neurotoxic effects was surprising, "perhaps even more surprising" was the amount of sodium fluoride needed - 2.1 ppm.47,48 Another set of experiments found that rats given aluminum fluoride or sodium fluoride in drinking water had occlusions in the blood vessels. In the animals administered aluminum fluoride, the occlusions reduced cerebral blood flow and aerobic metabolism. There was a reduced number of cells in two areas of the hippocampus and changes in neurofilaments in the neocortex that are "usually considered to be related to cell dysfunction."49

Effects on the thyroid mechanism

The interaction between fluoride and iodine also has been the subject of research, with studies finding effects on both the thyroid system and the brain. The data in this area, on fluoride alone or the fluoride/iodine relationship, include:

>

Fluoride's Toxicity

>

• In a study of 7- to 14-year-olds in China, those in a high-fluoride, low-iodine area had an average IQ of 71, compared with 77 in a normal-fluoride, low-iodine area and 96 in a normal-fluoride, normaliodine area. Thyroid effects in children in the high-fluoride, low-iodine area included a TSH of 21 mU/ml (compared with 6 in the normal area), a reverse T3 value of 58 ng/dl (21 in the normal area), and a significantly low reverse T3/T3 ratio of 2.91. IQ and TSH were negatively correlated in the study, and 69% of the children with mental retardation had elevated TSH levels.⁵⁰

• An examination of 165 workers in electrolysis shops of aluminum production with signs of fluoride toxicity also had the following thyroid abnormalities: a moderate reduction in the iodineabsorbing function of the thyroid and a low level of T3 hormone with a normal level of T4 hormone.⁵¹

• A study of 200 children from areas with fluoride in the drinking water found an increase in serum parathyroid levels that was well correlated with fluoride ingestion. An increase in serum parathyroid concentration was associated with a greater severity of clinical and skeletal fluorosis.⁵²

• A study of workers continuously exposed to fluorine found that 51% had reduced T3 hormone. An analysis of their immune status showed that the T lymphocyte count rose but that the functional activity of the cells declined, indicating an impaired cooperation of immunocytes due to imperfect control under the low T3 concentrations.⁵³

• Research found a higher prevalence of goiter (enlargement of the thyroid gland) in two towns in South Africa with high levels of fluoride in the water. 54

Among animal studies on this topic, an examination of the long-term effects of various doses of iodine and fluoride on the pathogenesis of goiter in mice found that both an iodine deficiency and an iodine excess could induce goiter and other functional and histopathological changes in the thyroid. Fluorine affected the thyroid changes caused by either iodine status. The researchers concluded that iodine and fluorine have mutually interacting effects on goiter in mice.55 In a study of pregnant and lactating mice who drank fluoridated water, 14-day-old pups had a 75% decrease in plasma free T4 and reductions of 27% and 17%, respectively, in the cerebellar and cerebral protein concentrations.56

Countering fluoride toxicity

Research suggests that a number of natural substances may oppose the adverse effects of fluoride by preventing or treating the toxicity. In his editorial in Fluoride, Bruce Spittle notes that dietary factors such as an adequate intake of iodine may protect against high-fluoride effects on the brain and IQ. He adds that while animal studies have found a partial recovery of all parameters studied when sodium fluoride and aluminum chloride are withdrawn, "the administration of ascorbic acid, calcium, or vitamin E, alone or in combination, resulted in a more complete recovery fromm the toxic effects.57 Recovery was more pronounced with the combination."58

Other animal studies have reported similar findings. In one, the effects of sodium fluoride (including locomotor behavioral and dental toxicities) were prevented significantly when the animals received calcium carbonate with the fluoride.59 In another study, mice receiving a higher concentration of fluoride in drinking water had a remarkable deterioration in learning capability. However, the administration of a proper concentration of selenium with the fluoride could decrease its toxic effects (a concentration of selenium that was too high produced synergistic toxicities). The findings suggested this mineral "might antagonize the neurotoxicity of fluoride on behavior and morphology."60

Xylitol: a safe alternative

As research continues to raise questions about fluoride's toxic effects, physicians, dental professionals and consumers may want to seek out other ways to prevent dental decay than by consuming fluoridated water. One alternative with considerable merit is the use of xylitol-containing products. Studies and reviews conducted in the past 10 years have shown that this safe, natural sugar substitute is an effective way to reduce the rate of tooth decay.

Xylitol is a sugar alcohol found in certain forest materials (such as birch and beech hardwoods) and foods (such as berries, plums, mushroom and lettuce). It is also produced by the body from food sources. Manufacturers use xylitol not only to replace sugars in foods (it has long been used as a sugar substitute in the diabetic diet) but also to develop xylitolcontaining products for dental use.61, 62 The latter include products such as xylitol chewing gums, lozenges, candies, and mouthrinses that deliver the anticariogenic substance directly to the mouth.

An explanation of xylitol's mechanisms of action in reducing dental decay, along with other background information, is available on Xylitol.com (a Web site provided by the Finnish company Leaf, which makes xylitol products). In essence, xylitol's anticariogenic effect is based on a biochemical feature of this sugar alcohol: It has five carbon atoms instead of the six carbon atoms found in sweeteners such as sorbitol, fructose and glucose. As a result, the bacteria in the mouth that cause dental cariesi streptococcus metans is the worst offender cannot ferment xylitol in their metabolism.63

When a person consumes xylitol the bacteria do not receive the energy supplied by ordinary sugar (sucrose) that enables them to grow, produce acids and launch an "acid attack." During this attack - which lasts more than a half hour after sugar has been consumed - the pH of plaque formed by the bacteria falls below 5.5. Consequently, calcium and phosphate salts begin to dissolve from the surface of the tooth enamel and cavities slowly begin to form. With xylitol, the pH of saliva and plaque does not fall, acid does not form, the bacteria do not absorb well on the teeth, and the plaque level decreases.64

A second anticariogenic effect of xylitol is to stimulate a more alkaline saliva than do other sugar products. This mechanism helps the saliva to correct incipient damage to enamel because the plaque pH may rise above 7 after xylitol products are used. Therefore, the calcium and phosphate salts in the saliva can reach parts of the enamel where they are deficient and the sites begin to harden again.⁶⁵

Research supporting xylitol's effectiveness includes the following:

• A randomized controlled trial of 61 children found a greater shift from higher scores for *streptococcus mutans* (S. *mutans*) to lower scores among subjects who chewed xylitol gum than those in the control group.⁶⁶

• A review of 14 clinical studies published between 1966 and 2001 concluded that the "studies demonstrated a consistent decrease in dental caries, ranging from 30 to 60%, among subjects using sugar substitutes" compared with controls. Subjects using xylitol had the highest reductions in caries.⁶⁷

• A five-year randomized controlled study of 740 10-year-old children found that after three years, the groups using xylitol candies on school days had a highly significant reduction of 35% to 60% in the incidence of caries compared with control groups.⁶⁸

• A randomized controlled trial of 151 subjects found that those who chewed xylitol gum for three months after having used a chlorhexidine (CHX) mouthrinse for 14 days had significantly lower salivary MS levels than did the placebo or control groups. MS levels did not differ significantly at baseline or after the CHX therapy.⁶⁹

• A study of 288 children in a twoyear gum-chewing program re-examined them five years later and found that xylitol gum and, to a lesser extent, xylitol/ sorbitol gum had a long-term protective effect against caries. During the five years after the gum-chewing program ended, xylitol gums reduced the caries risk 59% and xylitol/sorbitol gums reduced the risk 44%. However, the gum chewing should start at least one year before permanent teeth erupt to maximize the long-term protective effects.⁷⁰

 A two-year study of 169 motherchild pairs found a significant reduction in the probability of transmitting mutans streptococci (MS) to children when mothers chewed xylitol gum. Mothers in the xylitol group began chewing the gum three months after delivery, while controls received either chlorhexidine or fluoride varnish treatments at 6, 12 and 18 months after delivery. The children did not chew gum. At 2 years of age, 9.7% of children in the xylitol group (whose mothers began chewing gum three months after delivery) had detectable levels of MS, compared with 28.6% of children in the chlorhexidine group and 48.5% in the fluoride varnish group.⁷¹

• The Michigan Xylitol Programme, which studied children, adults and geriatric subjects who used saliva stimulants, tested the validity of the "penitol-hexitol theory," which holds that sugar alcohols with five hydroxyl groups (penitols such as xylitol) may be more effective cariologically than sugar alcohols with six hydroxyl groups (hexitols such as sorbitol). The accumulated clinical, sialochemical and microbiologic evidence suggested that xylitol is more effective in preventing caries than sorbitol and is cariologically safer.⁷²

• A 40-month double blind cohort study of 1,277 subjects compared the caries rate in groups chewing xylitol, sorbitol or combined gums with a group that received no chewing gum. The gums used in the four xylitol groups were most effective in reducing caries rates. A 100% xylitol pellet gum was the most effective agent.⁷³

• Clinical observations from two chewing gum studies found that after 40 months (permanent dentition) or 18 months (primary dentition), an arrest of caries generally occurred more often in children using xylitol or xylitol/sorbitol gums than in subjects who received no gum or used sucrose gum. The findings suggested that "high-xylitol chewing gum usage can retard or arrest even rampant dentine caries in conditions where effective restoration and prevention programmes have not been instituted..."⁷⁴

While some studies have found that xylitol-containing products did not reduce the levels of S. mutans in saliva,⁷⁵⁻⁷⁷ other studies have confirmed the caries-reducing effects of xylitol and offered insight into its mechanisms of action.⁷⁸⁻⁸⁶

In terms of how best to use xylitol, another Web site, Xylitol.org, offers this advice: A person needs to consume 4 to 12 grams a day to prevent tooth decay. Xylitol is safe for everyone at the cariespreventing level of less than 15 grams a day, and higher intakes yield diminishing benefits. Each piece of "all xylitol" mint or gum contains about 1 gram of xylitol, so a person might start by taking one piece four times a day for a total of 4 grams. Xylitol should be used at least three - and preferably five - times a day. In fact, it may not be effective if it is used only occasionally or even once a day. Use the products immediately after eating (swish water in the mouth first), and replace other gums and mints used between meals with xylitol-containing products.87

A call to arms

The compelling evidence of fluoride's brain and thyroid toxicity – and its adverse effects on other aspects of health – demands an end to the harmful process of water fluoridation. The good news is that readers who want to oppose the use of fluoride in public water systems will find it easier than ever to research this topic and join other health-minded people in advocating an end to fluoridation.

A wealth of timely information is available via the Internet from groups that are concerned about fluoridation. These coalitions have led a revolution in bringing data on fluoride and the fluoridation process into mainstream society through their Web sites. They

Fluoride's Toxicity

provide members and other visitors with the latest research on this topic as it is released, serving as a scientifically sound resource for interested parties. The Web sites also offer guidance on steps individuals can take to oppose fluoridation, such as write letters to local papers and congressional representatives.

The following two groups, for example, do an excellent job of educating people on the effects of fluoride on humans and the environment:

• The Fluoride Action Network (FAN) is an international coalition whose main goals are "to educate the public on the toxicity of fluoride compounds and to end the outdated/hazardous practice of water fluoridation." In addition to providing links to scientific studies, the Web site has a news-tracking page with citations and links for more than 1,200 relevant articles published in the US and worldwide in recent years.

Visit FAN's Web site, www.fluoridealert.org, for information on fluoridation, fluoride pollution, fluorine pesticides, fluoride health effects and much more (802-355-0999). You will also find information on how to become a member of FAN (the basic member rate is \$25 a year) and support this group in its opposition of water fluoridation.

• Second Look is a national nonprofit initiative that facilitates public and scientific examinations of controversial public policy issues, including fluoridation. The initiative explores the health and environmental effects of fluoridation and its complexity as a social and ethical issue. Among other data, Second Look's Web site includes an extensive bibliography of scientific literature on fluoride that has contributed greatly to this article. US citizens can make tax-deductible donations to this initiative. Go tovwww.slweb.org (508-755-7352).

Other Web sites of interest include the following:

>

Help Your Patients get Closer to Nature this Spring

For the 50 million people in the United States who encounter seasonal stresses, pollen in the air can cause watery eyes, headaches, coughing, and sneezing. Allerplex®, Antronex®, and Fen-Gre® from Standard Process can help provide defense against seasonal and environmental stresses.

For more information or to order these products, visit us online at www.standardprocess.com or call 800-558-8740.

^{*}These statements have not been evaluated by the Food & Drug Administration. These products are not intended to diagnose, treat, cure, or prevent any disease.

Fluoride's Toxicity

>

 Citizens for Safe Drinking Water (www.nofluoride.com)

Fluoridation.com (www.fluoridation.com)

 Fluoridedebate.com from Health Way House (www.fluoridedebate.com)

 Parents of Fluoride Poisoned Children (www.bruha.com/pfpc/)

New York State Coalition Opposed to Fluoride (www.orgsites.com/ny/nyscof and www.orgsites.com/ny/nyscof2)

- Massachusetts Communities for Pure Water (www.saveourwater.org)
- Pennsylvania Environmental Network (www.penweb.org/fluoride/)

Another valuable resource is Christopher Bryson's new book, The Fluoride Deception (Seven Stories Press, 2004). We recommend this fine scientific book to readers who are intrigued by the puzzle of how a substance as toxic as fluoride ended up in our drinking water and toothpaste in the first place. The Fluoride Deception is a fascinating investigation of the politics of water fluoridation and the fraud that led to its use.88

Correspondence:

Gary Null, PhD P.O. Box 918, Planetarium Station New York, New York 10024 USA 646-505-4660 Fax 212-472-5139 precisemd@aol.com

The Authors

Gary Null, PhD, has authored more than 50 books on health and nutrition and numerous articles published in leading magazines. His latest title is Gary Null's Power Aging (New American Library, 2003). Null holds a PhD in human nutrition and public health science from the Union Graduate School. He maintains a Web site at www.garynull.com that presents research articles on optimizing health through nutrition, lifestyle factors and alternative medicine.

Thinking of Writing a Book, but...

• You have no time • No experience •

- Don't know where to start .
- Need a writer to work with .
- Need a sizzling proposal to attract a major publisher •
- Or need editing help for an ailing/ incomplete manuscript •

Martin Zucker • (818) 874-9742

Co-author: The Miracle of MSM (Putnam). Natural Hormone Balance for Women (Pocket Books), Preventing Arthritis (Putnam)

Martin Feldman, MD, practices complementary medicine. He is an Assistant Clinical Professor of Neurology at the Mount Sinai School of Medicine in New York City.

References

- Fluoridation statistics 2000: status of water 1. fluoridation in the United States. National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, Atlanta, GA. From www.cdc.gov/ OralHealth/factsheets/fl-stats-us2000.htm.
- Populations receiving optimally fluoridated public 2. drinking water - United States, 2000. MMWR Weekly 2002; 51(07):144-7.
- Fluoride Recommendations Work Group 3. Recommendations for using fluoride to prevent and control dental caries in the United States. Centers for Disease Control and Prevention. MMWR 2001; 50(RR14): 1-42.
- Top achievements in health and well being in the 20th 4. century. Department of Health & Human Services Accountability Report, Fiscal Year 1999. From www.hhs.gov/of/reports/account/acct99/misc/ century.html.
- Brunelle JA, Carlos JP. Recent trends in dental caries in US children and the effect of water fluoridation. J Dent Res 1990; 69 Spec No:723-7; discussion 820-3.
- Fomon SJ, Ekstrand J, Ziegler EE. Fluoride intake and prevalence of dental fluorosis: trends in fluoride intake with special attention to infants. J Public Health Dent 2000; 60(3):131-9
- 7. Hileman B. Fluoridation of water. Chemical & Engineering News 1988: 66:36.
- 8. Li Y, Liang C, Slemenda C, Ji R, et al. Effect of longterm exposure to fluoride in drinking water on risks of bone fractures. J Bone Miner Res 2001; 16(5):932-9.
- Danielson C, Lyon JL, Egger M, Goodenough GK. Hip fractures and fluoridation in Utah's elderly opulation. JAMA 1992; 268(6):746-8.
- Cooper C, Wickham CA, Barker DJ, Jacobsen SJ. 10. Water fluoridation and hip fracture. JAMA 1991; 266:513-4. (Letter, a reanalysis of data presented in a 1990 paper.)
- 11. Takahashi K, Akiniwa K, Narita K. Regression analysis of cancer incidence rates and water fluoride in the USA. based on IACR/IARC (WHO) data (1978-1992). International Agency for Research on Cancer. I Epidemiol 2001; 11(4):170-9.
- 12. Klein W, et al. DNA repair and environmental substances. Zeitschrift fur Angewandte Bader-und Klimaheilkunde 1977; 24(3).
- Mohamed A, Chandler ME. Cytological effects of sodium fluoride on mice. Fluoride 1982; 15(3):110-8.
- Susheela AK, Jethanandani P. Circulating testosterone levels in skeletal fluorosis patients. J Toxicol Clin Toxicol 1996; 34(2):183-9.
- Narayana MV, Chinoy NJ. Effects of fluoride on rat testicular steroidogenesis. *Fluoride* 1994; 27(1):7-12.
 Al-Hiyasat AS, Elbetieha AM, Darmani H.
- Reproductive toxic effects of ingestion of sodium fluoride in female rats. Fluoride 2000: 33(2):79-84.
- 17. Luke J. Fluoride deposition in the aged human pineal gland. Caries Res 2001; 35(2):125-8.
- 18. Masters RD. Silicofluorides and higher blood lead: a national problem that particularly harms blacks. Dartmouth College , Hanover, NH. Videoconference Appalachian Environmental Lab, Frostburg State
- University, Frostburg, MD. November 15, 2001. 19. Masters RD, Coplan MJ, Hone BT, Dykes JE. Association of silicofluoride treated water with elevated blood lead. NeuroToxicology 2000; 21(6):1091-100.
- Xiang Q, Liang Y, Chen L, Wang C, et al. Effect of 20. fluoride in drinking water on children's intelligence. Fluoride 2003; 36(2):84-94.
- Lu Y, Sun ZR, Wu LN, Wang X, et al. Effect of high-21. fluoride water on intelligence in children. Fluoride 2000; 33(2):74-8.
- Li XS, Zhi JL, Gao RO. Effect of fluoride exposure on 22. intelligence in children. Fluoride 1995; 28(4):189-192 (as cited by Second Look).
- 23 Li Y. Li X. Wei S. [Effect of excessive fluoride intake on mental work capacity of children and a preliminary study of its mechanism.] Article in Chinese. Hua Xi Yi Ke Da Xue Xue Bao 1994; 25(2):188-91. Calderon J, Blenda M, Marielena N, Leticia C, et al
- Influence of fluoride exposure on reaction time and visuospatial organization in children. (Annual Conference of the International Society of Environmental Epidemiology) Epidemiology 2000; 11(4):S153 (as cited by Second Look).

- 25. Calvert GM, Mueller CA, Faien JM, Chrislin DW, et al. Health effects associated with sulfuryl fluoride and methyl bromide exposure among structural fumigation workers. Am J Public Health 1998; 88(12):1774-80.
- Du L. [The effect of fluorine on the developing human 26. brain.] Article in Chinese. Zhonghua Bing Li Xue Za Zhi 1992; 21(4): 218-20.
- Apetri AC, Surewicz. Atypical effect of salts on the thermodynamic stability of human prion protein. J Biol Chem 2003; 278(25):22187-92. Epub 2003 Apr03.
- Sun ZR, Liu FZ, Wu LN, et al. Effects of high fluoride drinking water on the cerebral functions of mice. Chinese Journal of Endemiology 2000; 19(4):262-3. As cited and abstracted in Fluoride 2001; 34(1):80.
- 29. Zhang Z, Xu X, Shen X, Xu X. [Effect of fluoride exposure on synaptic structure of brain areas related to learning-memory in mice. | Article in Chinese, Wei Sheng Yan Jiu 1999; 28(4):210-2.
- 30. Vanaja P, Ekambaram P, Jayakumar AR. Effects of sodium fluoride on locomotor behavior and a few biochemical parameters in rats. Environmental Toxicology and Pharmacology 1998; 6:187-91.
- Bhatnagar M, Rao P, Sushma J, Bhatnagar R. 31. Neurotoxicity of fluoride: neurodegeneration in hippocampus of female mice. Indian J Exp Biol 2002; 40(5):546-54.
- 32. Mullinex PJ, Denbesten PK, Schunior A, Kernan WJ. Neurotoxicity of sodium fluoride in rats. Neurotoxicol Teratol 1995; 17(2):169-77.
- Burgstahler AW, Colquhoun J. Neurotoxicity of fluoride. Editorial. Fluoride 1996; 29(2):57-8. 33.
- Bryson C. The fluoride deception. Seven Stories Press, New York, NY, 2004, 13, 20-1. 34 35. Connett E. Fluoride's effect on the brain. Fluoride
- Action Network Pesticides Project submission to National Research Council Committee. April 19, 2004. Bryson, op. cit., 22-3.
- Hertsgaard M, Frazer P. Fear of fluoride. Salon.com, February 17, 1999. From http://archive.salon.com/ news/1999/02/17news.html.
- Zhai JX, Guo ZY, Hu CL, Wang QN, Zhu QX. [Studies on fluoride concentration and cholinesterase activity in rat hippocampus.] Article in Chinese. Zhonghua Lao Dong Wei Sheng Zhi Ye Bing Za Zhi 2003; 21(2):102-4.
- 39. Shivarajashankara YM, Shivashankara AR, Gopalakrishna Bhat P, Muddanna Rao S, Hanumanth Rao S. Histological changes in the brain of young fluoride-intoxicated rats. Fluoride 2002; 35(1): 12-21.
- 40. Chen J, Chen X, Yang K, Xia T, Xie H. [Studies on DNA damage and apoptosis in rat brain induced by fluoride.] Article in Chinese. Zhonghua Yu Fang Yi Xue Za Zhi 2002; 36(4):222-4.
- Shivarajashankara YM, Shivashankara P, Gopalakrishna Bhat P, Hanumanth Rao S. Brain lipid 41. peroxidation and antoxidant systems of young rats in chronic fluoride intoxication. Fluoride 2002; 35(3):197-203 (as cited by Second Look).
- 42. Lakshmi Vani M, Pratap Reddy K. Effects of fluoride accumulation on some enzymes of brain and gastrocnemius muscle of mice. Fluoride 2000; 33(1):17-26 (as cited by Second Look).
- Long YG, Wang YN, Chen J, Jiang SF, et al. Chronic fluoride toxicity decreases the number of nicotinic acetylcholine receptors in rat brain. Neurotoxicol Teratol 2002; 24(6):751-7.
- 44. Chen J, Shan KR, Long YG, Wang YN, et al. Selective decreases of nicotinic acetylcholine receptors in PC12 cells exposed to fluoride. Toxicology 2003; 183(1-3): 235-42
- 45. Shen X, Zhang Z, Xu X. [Influence of combined iodine and fluoride on phospholipid and fatty acid composition in brain cells of rats.] Article in Chinese. Wei Sheng Yan Jiu 2004; 33(2):158-61. Spittle B. Fluoride and intelligence. Fluoride 2000;
- 46. 33(2):49-52.
- Varner JA, Jensen KF, Horvath W, Isaacson RL. 47. Chronic administration of aluminum-fluoride or sodium-fluoride to rats in drinking water: alterations in neuronal and cerebrovascular integrity. Brain Res 1998; 784(1-2):284-98.
- 48. Brain damage in rats from fluoridated water. Chemical & Engineering News; April 27, 1998; 29.
- 49. Isaacson RA, Varner JA, Kensen KF. Toxin-induced blood vessel inclusions caused by the chronic administration of aluminum and sodium fluoride and their implications for dementia. Annals of the New York Academy of Sciences 1997; 825:152-66. As abstracted in Fluoride 1998; 31(2):96-9.
- Lin FF, Aihaiti, Zhao HX, Lin J, et al. The relationship of a low-iodine and high-fluoride environment to subclinical cretinism in Xinjiang. Iodine Deficiency Disorder Newsletter 1991; 7(3) (as cited by Second Look).

- Mikhailets ND, Balabolkin MI, Rakitin VA, Danilov 51. IP. Thyroid function during prolonged exposure to fluorides. *Probl Endokrinol* 1996; 42:6-9.
- Gupta SK, Khan TI, Gupta RC, Gupta AB, et al. 52. Compensatory hyperparathyroidism following high fluoride ingestion - a clinico-biochemical correlation. Indian Pediatr 2001; 38(2):139-46.
- 53. Balabolkin MI, Mikhailets ND, Lobovskaia RN, Chernousova NV. [The interrepationship of the thyroid and immune statuses of workers with longterm fluoride exposure.] Article in Russian. Ter Arkh 1995; 67(1):41-2.
- 54. Jooste PL, Weight MJ, Kriek JA, Louw AJ. Endemic goitre in the absence of iodine deficiency in schoolchildren of the Northern Cape Province of South Africa. Eur J Clin Nutr 1999; 53(1):8-12.
- Zhao W, Zhu H, Yu Z, Aoki K, et al. Long-term effects of various iodine and fluorine doses on the thyroid and fluorosis in mice. Endocr Regul 1998; 32(2):63-
- Trabelsi M, Guermazi F, Zeghal N. Effect of fluoride 56 on thyroid function and cerebellar development in mice. Fluoride 2001; 34(3):165-73.
- 57. Chinoy NJ, Patel TN. The influence of fluoride and/ or aluminum on free radical toxicity in the brain of female mice and beneficial effects of some antidotes [abstract]. Fluoride 2000; 33; S8 (cited in Spittle). 58
- Spittle, op. cit. Ekambaram P, Paul V. Calcium preventing locomotor 59. behavioral and dental toxicities of fluoride by decreasing serum fluoride level in rats. Environ Toxicol Pharmacol 2001; 9(4):141-6.
- Zhang Z, Shen X, Xu X. [Effects of selenium on the damage of learning-memory ability of mice induced by fluoride.] Article in Chinese. Wei Sheng Yan Jiu 2001: 30(3):144-6.
- Xylitol info: background: making of xylitol, metabolic features of xylitol, and safety. From Xylitol.com. 61. 62. Xylitol.org. Main page. From www.xylitol.org/main/
- asp.
- 63. Xylitol info: FAQ about xylitol. From Xylitol.com. 64.
- Ibid. 65. Ibid.
- Autio JT. Effect of xylitol chewing gum on salivary 66. Streptococcus mutans in preschool children. ASDC J Dent Child 2002; 69(1): 81-6, 13.
- 67. Hayes C. The effect of non-cariogenic sweeteners on the prevention of dental caries: a review of the evidence, J Dent Educ 2001: 65(10):1106-9.
- 68. Alanen P, Isokangas P, Gutmann K. Xylitol candies in caries prevention: results of a field study in Estonian children. Community Dent Oral Epidemiol 2000: 28(3):218-24.
- 69. Hildebrandt GH, Sparks BS. Maintaining mutans streptococci suppression with xylitol chewing gum. JAm Dent Assoc 2000; 131(7):909-16.
- 70. Hujoel PP, Makinen KK, Bennett CA, Isotupa KP, et al. The optimum time to initiate habitual xylitol gumchewing for obtaining long-term caries prevention. J Dent Res 1999; 78(3):797-803.
- 71. Soderling E, Isokangas P, Pienihakkinen K, Tenovuo J. Influence of maternal xylitol consumption on acquisition of mutans streptococci by infants. J Dent Res 2000; 79(3):882-7.
- 72. Makinen KK, Makinen PL, Pape HR Jr, Peldyak J, et al. Conclusion and review of the Michigan Xylitol Programme (1986-1995) for the prevention of dental caries. Int Dent J 1996; 46(1):22-34.
- 73. Makinen KK, Bennett CA, Hujoel PP, Isokangas PJ, et al. Xylitol chewing gums and caries rates: a 40-month cohort study. J Dent Res 1995; 74(12):1904-
- 74. Makinen KK, Makinen PL, Pape HR Jr, Allen P, et al. Stabilisation of rampant caries: polyol gums and arrest of dentine caries in two long-term cohort studies in young subjects. Int Dent J 1995; 45(1 Suppl 1):93-107
- 75. Roberts MC, Riedy CA, Coldwell SE, Nagahama S, et al. How xylitol-containing products affect cariogenic bacteria. J Am Dent Assoc 2002; 133(4): 435-41; guiz 492-3.
- 76. Soderling E, Isokangas P, Tenovuo J, Mustakallio S, Makinen KK. Long-term xylitol consumption and mutans streptococci in plaque and saliva. Caries Res 1991; 25(2):153-7.
- 77. Petersson LG, Birkhed D, Gleerup A, Johansson M, Jonsson G. Caries-preventive effect of dentrifices containing various types and concentrations of fluorides and sugar alcohols. Caries Res 1991; 25(1):74-9.
- Machiulskience V, Nyvad B, Baelum V. Caries 78. preventive effect of sugar-substituted chewing gum. Community Dent Oral Epidemiol 2001; 29(4):278-88.
- 79. Gales MA, Nguyen TM. Sorbitol compared with xylitol in prevention of dental caries. Ann Pharmacother 2000; 34(1):98-100.

- 80. Makinen KK, Hujoel PP, Bennett CA, Isokangas P, et al. A descriptive report of the effects of a 16-month xylitol chewing gum programme subsequent to a 40month sucrose gum programme. Caries Res 1998; 32(2):107-12.
- 81. Makinen KK, Chiego DJ Jr, Allen P, Bennett C, et al. Physical, chemical, and histologic changes in dentin caries lesions of primary teeth induced by regular use of polyol chewing gums. Acta Odontol Scand 1998; 56(3):148-56
- Calamari SE, Azcurra AI, Luna Maldonado ER, 82 Battellino L.J. et al. Effects of xylitol, sorbitol and fluoride mouthrinses on glucose clearance in adolescents. Acta Odontol Latinoam 1997; 10(1):25-36.
- Makinen KK, Pemberton D, Makinen PL, Chen CY, 83. et al. Polyol-combinant saliva stimulants and oral health in Veterans Affairs patients - an exploratory study. Spec Care Dentist 1996; 16(3):104-15.

Р

H

Fluoride's Toxicity

- Trahan L, Bourgeau G, Breton R. Emergence of 84. multiple xylitol-resistant (fructose PTS-)mutants from human isolates of mutans streptococci during growth on dietary sugars in the presence of xylitol. J Dent Res 1996; 75(11):1892-900.
- Scheinin A, Soderling E, Scheinin U, Glass RL, Kallio ML. Xylitol-induced changes of enamel microhardness paralleled by microradiographic observations. Acta Odontol Scand 1993; 51(4):241-6.
- Isogangas P, Makinen KK, Tiesko J, Alanen P. Long-86. term effect of xylitol chewing gum in the prevention of dental caries: a follow-up 5 years after termination of a prevention program. Caries Res 1993; 27(6):495-8.
- 87. Xylitol.org. Main page. From www.xylitol.org/ main.asp. 88. Bryson, op. cit.

E D

Т

R

M

0



For Temporary Relief of Occasional **Indigestion and Heartburn***

F



PHYSICIAN FORMULATED

Scientifically Advanced

Nutritional Supplements

GES-5 is a combination of nutrients designed to provide relief of intermittent heartburn and indigestion.* Alginic acid and sodium alginate react with saliva and gastric acid to form a foamy froth or "alginic raft" that floats on top of the gastric contents and prevents acid and food reflux from re-entering the esophagus.* Slippery elm and deglycyrrhized licorice act as demulcents which are soothing to the alimentary canal.*

For more information about our full product line or to place an order call:

1-800-792-2222 or (914) 834-1804 fax orders toll free to 1-888-800-8068

visit us at www.rxvitamins.com • email: info@rxvitamins.com * This statement has not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure or prevent any disease.

OPTIMAL NUTRITIONAL SUPPORT