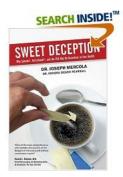


"Sweet Deception"



Author: Dr. Joseph Mercola ISBN: 0785221794 Format: Hardcover, 304 pages Publisher: Nelson Books (November 7, 2006) Date Reviewed: January 18, 2007

Among the many things you'll learn in this book, here are just a few:

Regarding Aspartame, the base ingredient in 'NutraSweet' and 'Equal' (p.37):

- ✓ It has been called '<u>The World's Best Ant Poison</u>' (p.46)
- ✓ Its meteoric rise in use in the 1980's very closely mirrors the explosion in Breast Cancer incidence (p.52)
- ✓ Formaldehyde, formed in the body when Aspartame is digested, is the chemical used to preserve cadavers (p.51)
- Phenylalanine and aspartic acid, the 2 amino acids in Aspartame, are considered "excitioxins" because of how they are incorporated into Aspartame and have been found to cause headaches, mental confusion, balance problems, and seizures (p.46)

Regarding Splenda, now America's #1 consumed artificial sweetener (pg 69),

- ✓ It was accidentally discovered while trying to create new insecticides
- ✓ It is regarded as "a chemical cousin to DDT" (p.103)
- ✓ Up to 12% of Splenda consumed (p.97) may permanently absorb into your fat cells and "continue to poison you indefinitely" (p.92).
- ✓ Digestion products of Splenda can cause liver damage and deplete antioxidants (p.100)
- \checkmark It can cross the placenta and affect the fetus during pregnancy (p.104)
- ✓ It can be VERY dangerous for diabetics because one of the sugars in Splenda (maltodextrin/dextrose) "will rapidly raise blood sugar" (p. 115).
- ✓ Splenda packets are actually 99% sugar, 1% sucralose, and costs almost 40 times more than sugar (p.113)
- ✓ Splenda is NOT good for baking because, when heated, it can break down to release a hazardous chemical called hydrogen chloride (p.119).

Regarding The FDA, the federal agency in charge of protecting you:

- ✓ Federal law prohibits the FDA from hiring experts with any potential conflicts of interest. The FDA waived this restriction more than 800 times from 1998-2000 (p.156).
- ✓ According to the Prescription Drug Fee User Act, the FDA collects "user fees" from the drug companies. These fees now make up 50% of the FDA's drug approval budget and 12% of the FDA's overall budget of \$1.3 billion (p.156).

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