

Fair Labeling and Informed Consent Act

Proposed Legislation – 2007-2008

“Informed consumers are essential to the fair and efficient functioning of a free market economy.”
(US CODE - TITLE 15 CHAPTER 39 § 1451)

The Fair Labeling and Informed Consent Act (FLICA) would provide unambiguous information to all consumers, medical professionals, pharmacists and public health officials whenever aborted fetal or embryonic materials are used in medical products so that alternatives may be selected in advance of purchase. The legislation is consistent with existing Federal Law requiring clear, non-evasive language in consumer packaging and informed consent including, but not limited to the descriptions found in 21 CFR Part 201, Labeling; 42 USC, § 300aa-26, the Federal Food, Drug and Cosmetic Act, 502 (e), USC Title 15, Ch. 39, §1451; 41 FR 6908, Sec. 201.10 Drugs, and 502(e) of the FDAC Act which provides guidelines on labeling practices

At present, Federal Law under **42 USC, §300aa-26** requires vaccine recipients or legal guardians of children to receive Vaccine Information Statements (VIS) published by the Centers for Disease Control (CDC) prior to immunization for some, but not all vaccines. Under the National Childhood Vaccine Injury Act (NCVIA) all healthcare providers are required to provide this information for all vaccines listed in their Vaccine Injury Table (VIT) The VIS is kept updated as needed with important information for consumers.

Implementation As Easy As 1-2-3...

1. Section...

Added to 502 (e) of the Federal Food, Drug and Cosmetic Act amending the labeling requirements and one section to the Vaccine Information Statement (VIS) which is already required under Federal Law to be given to all patients prior to immunization under the National Childhood Vaccine Injury Act (NCVIA).

2. Sentences...

Added to the 42 USC §300aa Federal Regulations, which specifies and governs all guidelines for the Vaccine Information Statement

3. Words...

Three words added to the manufacturer package insert “from elective abortion” immediately following any ingredients referenced in the Description section of all package inserts that utilize aborted fetal materials, including but not limited to cells, cell lines, tissues, DNA, recombinant DNA, monoclonal antibodies, blood, proteins or components.

**IF THE FDA CAN REQUIRE FAT CONTENT ON COOKIES, WHY SHOULDN'T WE KNOW
WHAT'S IN OUR MEDICAL PRODUCTS?**

For More Information Visit Our Website At: <http://www.cogforlife.org/flica.htm>