



General Assembly

January Session, 2011

Raised Bill No. 912

LCO No. 2777

* SB00912PH 041311 *

Referred to Committee on General Law

Introduced by:
(GL)

AN ACT AUTHORIZING FLAVORING AGENTS FOR PRESCRIPTION PRODUCTS.

Be it enacted by the Senate and House of Representatives in General Assembly convened:

1 Section 1. (NEW) (*Effective July 1, 2011*) (a) For purposes of this
2 section, "flavoring agent" means an additive used in food or drugs
3 when such additive: (1) Is used in accordance with good
4 manufacturing practice principles and in the minimum quantity
5 required to produce its intended effect, (2) consists of one or more
6 ingredients generally recognized as safe in food and drugs, has been
7 previously sanctioned for use in food and drugs by the state or the
8 federal government, meets United States Pharmacopeia standards or is
9 an additive permitted for direct addition to food for human
10 consumption pursuant to 21 CFR 172, (3) is inert and produces no
11 effect other than the instillation or modification of flavor, and (4) is not
12 greater than five per cent of the total weight of the product.

13 (b) A flavoring agent may be added to a prescription product by a
14 pharmacist upon the request of the prescribing practitioner, patient for
15 whom the prescription is ordered or such patient's agent.

This act shall take effect as follows and shall amend the following sections:

Section 1	<i>July 1, 2011</i>	New section
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GL *Joint Favorable*

PH *Joint Favorable*