


FDA Commissioner, Dockets Management Branch (HEA-305)
Food and Drug Administration
5630 Fishers Lane, Room 1061
Rockville, MD 20852

1508 01 MAR -9 AM 07

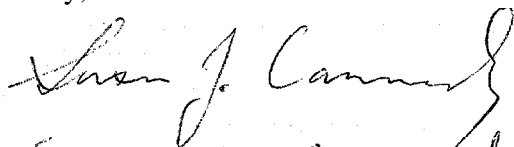
Dear FDA Commissioner:

I am writing to insist that the FDA require testing and adequate labeling of all genetically engineered and genetically modified foods.

1. The FDA **must require pre-market safety testing**. GE foods could be toxic, cause allergic reactions, have lower nutritional value and could compromise immune responses in consumers.
2. The FDA **must require mandatory pre-market environmental review**. GE crops and foods could cause irreparable damage to the environment./
3.  The FDA **must require mandatory labeling of GE foods**. Without mandatory labeling, neither consumers nor health professionals will know if an allergic or toxic reaction is the result of a GE food. Consumers will be deprived of the critical knowledge they need to hold food producers liable should any of these foods prove hazardous.
4. The FDA "Notification" policy is an insult to consumers and irresponsibly ignores strong scientific evidence of the numerous potential health and environmental risks of GE foods. These rules appear to be a decision made to convenience industry at the expense of public health, consumer information and the environment.

Thank you.

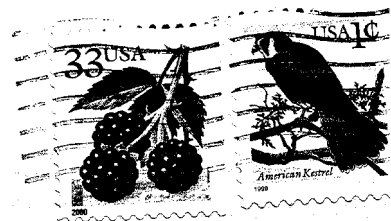
Sincerely,


Susan J. Cannedy
300 Harvey St.
Radford, VA 24141

00D_1598

C 2677

Cannedy
Harvey St.
Ford, VA 2414



FDA Commissioner
Dockets Management Branch (HFA-305)
Food and Drug Administration
5630 Fishers Lane, Room 1061
Rockville, MD 20852

