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March 24, 2003

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

Dear Sir/Madam:

As Vice President of Retail Operations & Natural Products of a health food pharmacy chain headquartered in Wadsworth, Ohio, I have been in the natural products industry for several years. Both my customers and I appreciate how the passage of the Dietary Supplement Health and Education Act of 1994 improved consumer access to dietary supplements and information about them, while increasing consumer protection against unsafe products and false and misleading claims.

It causes considerable concern that the FDA has only just begun to initiate aggressive enforcement actions under DSHEA, yet is calling for suggestions for increased legislative authority in order to better regulate the supplement industry. Should you not first give these recent efforts a chance to work, and perhaps even issue good manufacturing practices for supplements, before calling for new laws?

Simply put, I believe the agency can regulate Ephedra without dismantling DSHEA.

It is also my understanding that DSHEA increased the FDA's enforcement powers and that the FDA can seize a dietary supplement if it presents an unreasonable or significant risk of illness or injury. Furthermore, our Government can stop the sale of an entire class of dietary supplements if it poses an imminent public health hazard.

The former FDA commissioner, Dr. Jane Henney, has even stated before Congress that she believes DSHEA provides the FDA with the necessary legal authority to protect the public health. I agree with Dr. Henney in the strong support of DSHEA and do not think any additional legislative authority is necessary.

Sincerely,

Jon A. Fiume  
 Vice President  
 Retail Operations & Natural Products

JAF/tk

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