

The Contents of Herbal and Dietary Supplements Implicated in Liver Injury in the United States Are Frequently Mislabeled

Victor Navarro,¹ Bharathi Avula,² Ikhlas Khan,² Manisha Verma,¹ Leonard Seeff,¹ Jose Serrano,³ Andrew Stolz,⁴ Robert Fontana,⁵ and Jawad Ahmad⁶

The U.S. Drug Induced Liver Injury Network assayed the contents of herbal and dietary supplements collected from patients enrolled into its prospective study. The aim was to determine the accuracy of product labels, and to identify known hepatotoxins. Using high-performance liquid chromatography coupled with mass spectrometry to assay 272 product, 51% were found to be mislabeled; that is, to have chemical contents that did not match the label. Appearance enhancement, sexual performance, and weight loss products were most commonly mislabeled. Whether the mislabeling contributed to liver injury is under study; however, the high mislabeling rate underscores the need for more stringent regulation of supplements. (*Hepatology Communications* 2019;3:792-794).

Herbal and dietary supplements (HDS) are being used increasingly for touted “health benefits”⁽¹⁾ and are responsible for over \$36 billion in commerce in the United States each year.^(2,3) Broadly stated, HDS consist of single and multi-ingredient products that may include herbals, vitamins/minerals, and amino acids. Hepatotoxicity from HDS is being recognized more and more.⁽⁴⁾ In the US Drug Induced Liver Injury Network (DILIN) prospective study, products used as appearance and performance-enhancing (APE) supplements are the most common subclass of HDS leading to liver injury. Patients with hepatotoxicity from weight loss and other non-APE products can also sustain severe liver injury. Despite these potential risks, HDS remain widely available in

retail outlets and on the internet, with little to no required safety or efficacy testing. As a result, consumers are exposed to compounds whose biological effects are largely unknown in humans. In addition, unlabeled ingredients, whether added intentionally, accidentally, or as a byproduct of the manufacturing process, comprise yet another risk of HDS to the consumer. The HDS collected from DILIN enrollees were used to verify product labels and then estimate the rate of label inaccuracy, while searching for known hepatotoxins and pharmaceuticals among HDS implicated in causing liver injury. The purpose of this study was to determine the veracity of the labels on HDS consumed by patients enrolled in the DILIN, and to identify potential hepatotoxins in these products.

Abbreviations: APE, appearance and performance-enhancing; DILIN, US Drug-Induced Liver Injury Network; HDS, herbal and dietary supplements.

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Materials and Methods

The DILIN, a multicenter consortium funded by the National Institute for Diabetes and Digestive and Kidney Diseases, studies the etiologies, risk factors, and outcomes of patients with drug-induced liver injury (DILI). All DILIN cases undergo a causality adjudication process that determines the likelihood of liver injury due to a drug or dietary supplement and implicates the product (drug or HDS) responsible.

Between 2003 and 2015, the DILIN prospective study enrolled 341 HDS from 1,268 patients with suspected DILI; 272 of these products had labels listing their ingredients, and these underwent chemical analysis at the National Center for Natural Products Research of the University of Mississippi using ultra-high-performance liquid chromatography coupled with a quadrupole time-of-flight mass spectrometry. Of the 272 products, 96 were directly implicated as a cause for liver injury in 71 patients.

The HDS products also were analyzed for known hepatotoxins, including aflatoxins, anabolic steroids, anthraquinones, pyrrolizidine alkaloids, and pharmaceuticals. HDS were grouped according to their marketed purpose for use, such as for APE or weight loss. Mislabeling was defined as when the chemical analysis could not verify all ingredients listed on the label.

Results

Of the 272 products tested, only 132 had labels that accurately reflected the compounds identified by chemical analysis; therefore, 140 (51%) were

TABLE 1. MISLABELING IN HDS COLLECTED BY THE DILIN

Category	HDS With Labels (n)	Mislabeled HDS (n [%])
General health	53	26 (49%)
APE	46	37 (80%)
Weight loss	36	26 (72%)
Gastrointestinal symptoms	22	9 (41%)
Energy boosters	5	3 (60%)
Sexual enhancers	4	4 (100%)
Miscellaneous/Unknown	106	35 (33%)
Total	272	140 (51%)

mislabeled. These 140 products lacked at least one of the listed compounds (number of ingredients ranged from 1 to 21), and 55 products contained compounds not listed on the label, including 2 that were pharmaceuticals. The overall mislabeling rate was 51%, and rates were higher for steroidal (82%) than botanical (44%) and vitamin (49%) products. When categorized by marketed purpose of the HDS, the rates of mislabeling were highest for sexual enhancers, APE, and weight loss products (Table 1). Adulterants found in the 91 products with known hepatotoxins included anabolic steroids in 13 of 26 products, diclofenac in an agent for arthritis, and tamoxifen in a bodybuilding product. In the latter two cases, the clinical features of liver injury were consistent with the adulterants' reported biochemical and clinical liver injury pattern.

Discussion

We identified an alarmingly high rate of mislabeling in HDS collected from patients enrolled in the

ARTICLE INFORMATION:

From the ¹Department of Digestive Disease and Transplantation, Einstein Healthcare Network, Philadelphia, PA; ²National Center for Natural Products Research, the University of Mississippi, University, MS; ³Liver Diseases Branch, National Institute of Diabetes and Digestive and Kidney Diseases, Bethesda, MD; ⁴Department of Medicine, Keck School of Medicine of the University of Southern California, Los Angeles, CA; ⁵University of Michigan Medical School, Ann Arbor, MI; ⁶Department of Medicine, Icahn School of Medicine at Mount Sinai, New York, NY.

ADDRESS CORRESPONDENCE AND REPRINT REQUESTS TO:

Victor J. Navarro, M.D.
Department of Digestive Disease and Transplantation
Albert Einstein Medical Center
Suite 505 Klein

5501 Old York Road
Philadelphia, PA, 19141
E-mail: NavarroV@einstein.edu

DILIN study. Products used for APE and weight loss had the highest rates of mislabeling. In the two cases of pharmaceutical adulteration, the clinical features were highly consistent with that reported as a result of the identified adulterants.⁽⁵⁾

Our findings highlight that a large proportion of commercially available HDS are mislabeled, exposing consumers to substances unknowingly, some of which may be responsible for liver injury. At this time, we are unable to determine which of the compounds or combination of compounds are responsible for the observed liver injury. Nonetheless, the high rate of mislabeling, including identification of agents recognized to cause liver injury, may expose consumers to unrecognized risk, including interactions with prescribed medications. These preliminary findings suggest that more

stringent regulation of HDS is required to improve the accuracy and to verify product labels, to avoid potential adverse health events including liver injury.

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