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Dietary Supplement Industry: Upcoming Quality Regulation Deadline

For years, dietary supplements have been scrutinized by the media for being marketed as "snake-oil" cure-alls, potentially containing components considered harmful to consumers. Under-regulation by the Food and Drug Administration (FDA) lead to concerns that these products did not fall under the same regulatory requirements as pharmaceuticals. Until now.

View the full story [here](#).

FDA to Review Medical Devices Marketed Prior to 1976

The FDA recently announced that manufacturers of 25 types of medical devices marketed prior to 1976 must submit safety and effectiveness information to the agency so that it may evaluate the risk level for each device type. Devices found by the FDA to be of high risk to consumers will be required to undergo the agency's most stringent premarket review process.

View the full story [here](#).

Recent Rumors of Contamination Urge Tougher Review of Dental Laboratories

The combination of dental appliance contamination with lead and the FDA's inspection of a number of U.S. dental laboratories has stirred domestic dental laboratories to ensure that their quality management program provides the assurance to their dental clients that they have met the requirements set forth by FDA's Title 21, Chapter I, Subchapter H - Medical Devices, Part 820 Quality System Regulation.

Read the complete article [here](#).

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