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FDA Requests Comments on Genome-Edited Plants

On January 19, 2017, FDA established a [docket](#) to receive comments on the use of genome-editing techniques to produce new plant varieties that are used for human or animal food. Explaining that producers of foods from plant varieties developed using genome-editing techniques (like all food products) must ensure that the foods they offer are safe and in compliance with applicable legal requirements, FDA intends to clarify its policy for the regulation of products derived from genome-editing techniques (including new technologies that use targeted nucleases or targeted oligonucleotides to modify a plant's DNA sequence by insertion, deletion, or substitution of nucleotides at a specific site in a plant's genome).

FDA's existing policy, which provides for voluntary consultation on the safety and legality of new plant varieties derived by rDNA methods, was issued in 1992. Noting its intent to continue offering consultations for developers of new plant varieties, including those produced using genome editing, FDA is inviting comments by April 19, 2017 from developers, researchers, and other stakeholders to help inform its thinking for these products.

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