

## **United States Department of Agriculture**

Marketing and Regulatory Programs Kristi Snell, PhD Chief Scientific Officer & VP of Research Karen Bohmert-Tatarev, Senior Director of Regulatory Affairs

Yield10 Bioscience

Animal Plant Health Inspection Service RSR number 23-096-02

Dear Dr. Snell and Dr. Bohmert-Tatarev:

Biotechnology Regulatory

Services

resistance

4700 River Road Riverdale MD 20737 Thank you for your letter dated April 5<sup>th</sup>, 2023, requesting a Regulatory Status Review (RSR) for *Camelina sativa* developed using genetic engineering (modified *Camelina sativa*). In your letter, you described that the *Camelina sativa* was modified to impart herbicide tolerance to glufosinate through the insertion of a *bar* gene and tolerance to sulfonylurea and imidazolinone herbicides through the insertion of an AHAS gene

RE: Regulatory Status Review of Camelina sativa developed using genetic engineering for herbicide

The Plant Protection Act of 2000 (7 U.S.C. §§ 7701 et seq.) provides USDA authority to oversee the detection, control, eradication, suppression, prevention, or retardation of the spread of plant pests to protect agriculture, environment, and the economy of the United States. USDA, through the Animal and Plant Health Inspection Service (APHIS), regulates the "Movement of Organisms Modified or Produced through Genetic Engineering" as described in 7 CFR part 340.

Consistent with 7 CFR 340.4, APHIS reviewed your modified Camelina sativa to determine whether it is subject to the regulations in 7 CFR part 340. Specifically, APHIS reviewed the modified Camelina sativa to determine whether there is a plausible pathway by which the Camelina sativa, or any sexually compatible relatives, would pose an increased plant pest risk relative to the plant pest risk posed by an appropriate Camelina sativa comparator. Based on information you provided, publicly available resources, and APHIS' familiarity with Camelina sativa and knowledge of the trait, phenotype, and mechanism of action, APHIS considered the (1) biology of nonmodified Camelina sativa and its sexually compatible relatives; (2) the trait and mechanism-of-action of the modification; and (3) the effect of the trait and mechanism-of-action on the (a) distribution, density, or development of the plant and its sexually compatible relatives, (b) production, creation, or enhancement of a plant pest or a reservoir for a plant pest, (c) harm to non-target organisms beneficial to agriculture, and (d) weedy impacts of the plant. APHIS did not identify any plausible pathway by which your modified Camelina sativa, or any of its sexually compatible relatives, would pose an increased plant pest risk relative to comparator Camelina sativa plants. APHIS has determined your Camelina sativa is unlikely to pose an increased plant pest risk relative to its comparators. Once APHIS determines that a plant product is unlikely to pose an increased plant pest risk relative to its comparator, and, thus, is not a plant pest or a plant that requires regulation because it is capable of introducing or disseminating a plant pest, APHIS has no authority to regulate it under 7 CFR part 340. Accordingly, your Camelina sativa is not subject to the regulations under 7 CFR part 340. APHIS' determination that this modified plant is not subject to the regulations extends to any progeny of the modified plant that is derived from crosses with other non-modified plants or other modified plants that are also not subject to the regulations in 7 CFR part 340.

Please be advised that APHIS' decision applies to the *Camelina sativa* developed using genetic engineering exactly as described in your letter. If at any time you become aware of any information that may affect our review of your modified *Camelina sativa*, including, for example, new information that shows the trait, phenotype, or mechanism of action is different than described in your letter, you must contact APHIS for further review of the plant at RSRrequests@usda.gov.

Please be advised that your plant product, while not regulated under 7 CFR part 340, may be subject to APHIS Plant Protection and Quarantine (PPQ) permit and/or quarantine requirements. For further information, you may contact the PPQ general number for such inquiries at 877-770-5990. Your plant product may also be subject to other regulatory authorities such as the U.S. Environmental Protection

Agency (EPA) or the Food and Drug Administration (FDA). Please contact EPA and FDA to enquire about the regulatory status of your product.

Date: November 14, 2023

Sincerely,

Bernadette Juarez

APHIS Deputy Administrator Biotechnology Regulatory Services Animal and Plant Health Inspection Service U.S. Department of Agriculture