

# NATIONAL ENVIRONMENTAL POLICY ACT

## PRELIMINARY FINDING OF NO SIGNIFICANT IMPACT

### Regarding Deregulating a Petition (19-099-01p) Under 7 CFR part 340

from: Westhoff Vertriebsgesellschaft mbH

A1-DFR petunias

**United States Department of Agriculture  
Animal and Plant Health Inspection Service  
Biotechnology Regulatory Services**

The United States Department of Agriculture (USDA), Animal and Plant Health Inspection Service (APHIS) has developed this preliminary Finding of No Significant Impact (hereafter referred to as FONSI) to comply with the requirements of the National Environmental Policy Act (NEPA) of 1969, as amended, the Council of Environmental Quality's (CEQ) NEPA implementing regulations, and APHIS NEPA implementing procedures (7 CFR part 372). This FONSI sets forth APHIS' NEPA decision with respect to potential impacts to the human environment that could derive from a determination of nonregulated status for A1-DFR petunias.

Westhoff Vertriebsgesellschaft mbH (Westhoff), submitted a petition (19-099-01p) to the USDA APHIS, requesting that genetically engineered (GE) petunias referred to as A1-DFR petunias, and any petunia lines derived from crosses of A1-DFR petunias and conventional petunias, or nonregulated GE petunias, no longer be considered regulated articles under Title 7 of the Code of Federal Regulations part 340 (7 CFR part 340). A1-DFR petunias (23 events that contain one or more copies of the *A1* DFR gene), have been genetically engineered to express the dihydroflavonol 4-reductase (DFR) enzyme from maize (A1-DFR) allowing the plants to produce the plant pigment pelargonidin, which is a type of anthocyanin pigment, in their flower petals. A1-DFR petunias are intended to provide additional color varieties. A1-DFR petunias are currently regulated by APHIS because they were developed using genetic elements from the plant pests cauliflower mosaic virus (CaMV) and *Agrobacterium tumefaciens*, both of which are regulated articles under 7 CFR part 340.2.<sup>1</sup>

As part of evaluation of Westhoff's petition, APHIS conducted an Environmental Assessment (EA) to inform APHIS' decision regarding the regulatory status of A1-DFR petunias. The EA evaluates the potential impacts of APHIS' regulatory decision on the quality of the human environment.<sup>2</sup> The EA did not identify any significant impacts that would derive from either an approval or a denial of the petition. Therefore, the Agency has prepared this FONSI, pursuant to 40 CFR part 1508.13, which provides a summary of the EA, and the reasons why APHIS'

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<sup>1</sup> Disarmed *Agrobacterium* is commonly used in the genetic modification of plants. Disarmed means the *Agrobacterium* is non-virulent.

<sup>2</sup> Under NEPA regulations, the "human environment" includes "the natural and physical environment and the relationship of people with that environment" (40 CFR § 1508.14).

decision to issue a determination of nonregulated status for A1-DFR petunias will not have a significant impact on the human environment.

### **The Coordinated Framework and APHIS Regulatory Authority**

In 1986, the United States government issued a comprehensive regulatory policy for the regulation of products of biotechnology known as the Coordinated Framework for the Regulation of Biotechnology (Coordinated Framework) (51 FR 23302, 57 FR 22984). Since 1986, the Environmental Protection Agency (EPA), Food and Drug Administration (FDA), and USDA have regulated GE organisms consistent with the principles of this framework. These guiding principles include: (1) agencies should define those transgenic organisms subject to review to the extent permitted by their respective statutory authorities; (2) agencies should focus on the characteristics and risks of the biotechnology product, not the process by which it is created; and, (3) agencies should exercise oversight of biotechnology products only when there is evidence of “unreasonable” risk.

Within the USDA, APHIS is responsible for “protecting animal and plant health” as one of its primary strategic goals. APHIS provides leadership in ensuring the health and care of plants and animals. The agency’s strategic goals help improve agricultural productivity and competitiveness, and contribute to the national economy and the public health. The USDA asserts that all methods of agricultural production (conventional, organic, or the use of GE varieties) can provide benefits to the environment, consumers, and farm income.

APHIS regulates GE organisms to ensure that they do not pose a plant pest risk pursuant to the Plant Protection Act (PPA) of 2000, as amended (7 USC §§ 7701 et seq.) and APHIS implementing regulations at 7 CFR part 340. APHIS regulations at 7 CFR part 340 govern the importation, interstate movement, and environmental release of GE organisms that may pose a plant pest risk. A GE organism is considered a regulated article if the donor organism, recipient organism, vector, or vector agent used in engineering the organism belongs to one of the taxa listed in the regulation (7 CFR part 340.2) and is also considered a plant pest; such as *Agrobacterium tumefaciens*. A GE organism is also regulated under 7 CFR part 340 when APHIS has reason to believe that the GE organism may be a plant pest or APHIS does not have sufficient information to determine if the GE organism is unlikely to pose a plant pest risk. A GE organism is no longer subject to the PPA or to the requirements of 7 CFR part 340 when APHIS determines that a GE organism is unlikely to pose a plant pest risk.

The FDA is responsible for ensuring the safety and proper labeling of all plant-derived foods and feeds pursuant to the authority of the Federal Food, Drug, and Cosmetic Act (FFDCA), including those that are genetically engineered. The FDA policy statement concerning oversight of products derived from new plant varieties, including those genetically engineered, was published in the *Federal Register* on May 29, 1992.<sup>3</sup> Pursuant to this policy, the FDA uses a consultation process to ensure that human food and animal feed safety issues are resolved prior to commercial distribution of products of genetic engineering. To help developers of food and feed derived

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<sup>3</sup> Available at U.S. FDA: Statement of Policy - Foods Derived from New Plant Varieties; <http://www.fda.gov/Food/GuidanceRegulation/GuidanceDocumentsRegulatoryInformation/Biotechnology/ucm096095.htm>

from GE crops comply with their obligations pursuant under Federal food safety laws, the FDA encourages them to participate in a voluntary consultation process.

The EPA is responsible for regulating the sale, distribution, and use of pesticides, including pesticides that are produced by an organism through techniques of modern biotechnology pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). The EPA sets tolerance limits for residues of pesticides on and in food and animal feed, or establishes an exemption from the requirement for a tolerance, pursuant to FFDCA. In addition, the EPA regulates certain biological control organisms pursuant to the Toxic Substances Control Act (TSCA).

### **APHIS' Response to Petitions for Nonregulated Status**

APHIS regulations at 7 CFR part 340 provide that any person may submit a petition to APHIS requesting that, because the GE organism does not pose a plant pest risk, it should not be regulated by APHIS. As required by 7 CFR part 340.6, APHIS must respond to petitioners with a regulatory status decision. If APHIS determines, based on a Plant Pest Risk Assessment (PPRA) and other relevant information that the GE organism is unlikely to pose a plant pest risk, the GE organism is no longer subject to regulation under 7 CFR part 340.

### **Public Involvement**

On July 25, 2019, APHIS announced in the *Federal Register* that it was making Westhoff's petition available for public review and comment to help identify potential environmental and interrelated economic impacts that APHIS should consider in evaluation of the petition.<sup>4</sup> APHIS accepted written comments on the petition for a period of 60 days, until midnight September 23, 2019. At the end of the comment period APHIS had received a total of nine comments – seven were in support of the Westhoff petition and two were opposed to deregulation. APHIS evaluated the comments and integrated the concerns raised into the EA. All comments received on the petition are available for public review at [www.regulations.gov](http://www.regulations.gov), Docket ID: APHIS-2019-0037.

### **Environmental Assessment and Scope of Analysis**

An EA was prepared consistent with CEQ regulations (40 CFR parts 1500-1508) and USDA-APHIS NEPA implementing procedures (7 CFR part 372). APHIS developed a list of topics for consideration in the EA based on issues identified in public comments submitted on the petition and draft EA for A1-DFR petunias, other EAs and EISs evaluating petitions for nonregulated status, and the scientific literature on floriculture, plant biotechnology, and the environmental sciences. The following topics were identified as relevant to the scope of analysis (40 CFR §1508.25):

### **Commercial Production**

- Petunia Production
- Pest and Pathogen Management

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<sup>4</sup> Federal Register, Vol. 84, No. 143, July 25, 2019, p. 35849 – Westhoff Vertriebsgesellschaft mbH; Availability of Petition for Determination of Nonregulated Status of Petunias Genetically Engineered for Flower Color [Docket No. APHIS-2019-0037, [www.regulations.gov](http://www.regulations.gov)].

## **Physical Environment**

- Soils
- Water Resources
- Air Quality

## **Biological Environment**

- Soil Biota
- Animal and Plant Communities
- Gene Flow and Weediness
- Biodiversity

## **Human Health Considerations**

- Public Health and Worker Safety

## **Socioeconomic Considerations**

- Domestic Economic Environment
- International Trade

In addition to evaluation of potential direct and indirect impacts, potential cumulative impacts relative to these topics were also considered. Additionally potential impacts on threatened and endangered species, as well as adherence of the regulatory decision to executive orders, and environmental laws and regulations to which the regulatory status decision may be subject were analyzed.

## **Alternatives Evaluated in the EA**

The EA considered two alternatives in responding to Westhoff's petition, to either deny or approve the request for nonregulated status, and analyzed the potential environmental, human health, and socioeconomic impacts that may result from the two alternatives.

### *No Action: Deny the Petition and Continuation as a Regulated Article*

One of the alternatives that must be considered by APHIS is a "No Action Alternative," pursuant to CEQ regulations at 40 CFR § 1502.14. Under the No Action Alternative, APHIS would deny the petition. A1-DFR petunias and progeny derived from A1-DFR petunias would continue to be regulated articles under 7 CFR part 340. Because APHIS concluded in its PPRA that A1-DFR petunias are unlikely to pose a plant pest risk (USDA-APHIS 2020) this is not APHIS' preferred alternative. Choosing this alternative would not be an appropriate response to the petition for nonregulated status, nor satisfactorily meet the purpose and need for making a regulatory status decision pursuant to the requirements of 7 CFR part 340.

### *Preferred Alternative: Determination of Nonregulated Status for A1-DFR Petunias*

Under this alternative, A1-DFR petunias and progeny derived from this event would no longer be regulated articles under the regulations at 7 CFR part 340 because it was determined that, based on the scientific evidence before the Agency, A1-DFR petunias are unlikely to pose a plant pest risk (USDA-APHIS 2020). APHIS would no longer require authorizations for introductions of A1-DFR petunias and progeny derived from this event. This alternative best satisfies the purpose and need to respond appropriately to the petition for nonregulated status pursuant to the requirements of 7 CFR § 340.6, the Agency's statutory authority under the PPA, and the biotechnology regulatory policies described for the Coordinated Framework.

### Alternatives Considered but Dismissed from Detailed Analysis in the EA

APHIS evaluated several alternatives for consideration in the EA in light of the Agency's statutory authority under the PPA and APHIS implementing regulations at 7 CFR part 340, but dismissed these alternatives from detailed analysis in the EA. The alternatives considered are described in the EA along with the reasons for dismissal from detailed analysis.

### Environmental Consequences of APHIS' Selected Action

The EA provides analyses of the alternatives APHIS considered, to which the reader is referred for specific details. The following table briefly summarizes the potential environmental impacts of the alternatives evaluated in the EA.

<b>Summary of Potential Impacts for the Alternatives Considered</b>		
<b>Analysis</b>	<b>No Action Alternative: Continue to Regulate A1-DFR Petunias as a Plant Pest</b>	<b>Preferred Alternative: Determination of Nonregulated Status for A1-DFR Petunias</b>
Meets Purpose and Need	No	Yes
Unlikely to pose a plant pest risk	Addressed by the use of regulated field trials and past observations.	Determined by the plant pest risk assessment (USDA-APHIS 2020).
<b>Horticultural Production</b>		
Acreage and Areas of Petunia Production	Petunias are primarily grown for the retail market inside greenhouses. Michigan, Ohio, New York, and Pennsylvania are the leading producers of petunia. Petunias have consistently ranked among the five most commonly sold bedding plants. Current trends in petunia production are not anticipated to change.	A1-DFR petunias will provide an additional color variety of petunia and is expected to compete with other color varieties that are currently in production and offered for sale. A determination of nonregulated status for A1-DFR petunias is not expected to change the acreage or areas used for petunia seed and bedding plant production.
Horticultural Practices and Inputs	Horticultural practices and inputs used in petunia production would remain unchanged.	The change in color in A1-DFR petunias does not cause changes in growth habit, temperature tolerances, nutritional requirements, or other factors that would alter horticultural practices used in petunia production.
<b>Physical Environment</b>		
Soils	Growing practices and inputs used for commercial production of petunia that may impact soil resources would not change from those currently used.	The potential impacts of A1-DFR petunias production on soil quality are not expected to differ from the No Action Alternative.
Water Resources	Existing water use and water quality conditions would be expected to be unchanged.	Because A1-DFR petunias are similar to non-GE cultivated petunia, approval of the petition and subsequent commercial production of

<b>Summary of Potential Impacts for the Alternatives Considered</b>		
<b>Analysis</b>	<b>No Action Alternative: Continue to Regulate A1-DFR Petunias as a Plant Pest</b>	<b>Preferred Alternative: Determination of Nonregulated Status for A1-DFR Petunias</b>
		A1-DFR petunias would present the same potential risks to water resources as non-GE cultivated petunia varieties.
Air Quality	Current impacts to air quality associated with petunia production practices would be expected to continue unchanged.	Sources of potential impacts on air quality are the same as those under the No Action Alternative.
<b>Biological Resources</b>		
Soil Biota	Current impacts to soil biota associated with petunia production practices would be expected to continue unchanged.	A1-DFR petunias are not expected to change the practices and inputs used in petunia production that could cause new impacts to soil biota.
Animal Communities	A variety of animal and insect species feed on or use petunia. Mammals and birds may use petunias for food or feed on the insects feeding on petunias. Invertebrates can feed on petunia plants or prey upon other insects as well as using petunia for pollen and nectar sources.	A1-DFR petunias would not require any change to petunia production practices. DFR and associated pelargonidin and NPTII introduced into A1-DFR petunias present negligible risk to wildlife. Potential impacts to animal communities are not anticipated to be different compared to the No Action Alternative
Plant Communities	Because petunia cultivation typically occurs in greenhouses, the plant communities associated with petunia production are limited. Potential impacts to plant communities associated with petunia production would be expected to continue unchanged. The impacts to plant communities from petunias in commercial or residential areas is not expected to change.	Potential impacts to plant communities are not anticipated to be different compared to the No Action Alternative
Gene Flow and Weediness	Petunia does not cross with species of other genera and hybrids of closely related species are rare in nature. No plants in the genus <i>Petunia</i> are on the Federal noxious weed list nor are they listed as invasive by any state. Petunia does not spread vegetatively, and roots	A1-DFR petunias have been modified for a change in flower color only. The change in color in A1-DFR petunias does not cause changes in seed set, pollen availability, growth habit, temperature tolerances, nutritional requirements, or other factors that would alter where it can be grown or

<b>Summary of Potential Impacts for the Alternatives Considered</b>		
<b>Analysis</b>	<b>No Action Alternative: Continue to Regulate A1-DFR Petunias as a Plant Pest</b>	<b>Preferred Alternative: Determination of Nonregulated Status for A1-DFR Petunias</b>
	will not form on discarded parts of a plant under outdoor conditions (Westhoff 2019). Little evidence exists to suggest that petunia behaves as a weed.	the potential for cross pollinating compared to currently available petunia varieties.
Biodiversity	Petunia production typically occurs in greenhouses reducing any impacts on biodiversity. As an ornamental plant grown in beds, pots, and hanging baskets, petunia largely relates to biodiversity within the built environment by serving as a food source for pollinators.	A1-DFR petunias would not be expected to change growing practices, and therefore would not likely impact biodiversity any differently than conventional petunia.
<b>Human and Animal Health</b>		
Human Health	Petunias are not a food and not consumed by humans or used for animal feed. Management practices for petunia production, and the associated human health impacts, are expected to continue unchanged.	Potential impacts to human health are not anticipated to be different from those under the No Action Alternative. The EPA WPS will continue to provide the same level of protection as is currently available
<b>Socioeconomics</b>		
Domestic Economic Environment	Petunia production and use is expected to continue much as it is currently.	A determination of nonregulated status for A1-DFR petunias is not expected to adversely impact domestic petunia markets. A1-DFR petunias would provide novel colored flowers. This additional color variety is not expected to result in a significant increase in petunia demand or production in the United States.
International Trade	There would be no impacts on trade under the No Action Alternative.	A1-DFR petunias would be subject to the same international regulatory requirements as currently traded flower varieties. U.S. imports of A1-DFR petunias would no longer require authorization under 7 CFR part 340, otherwise U.S. petunia imports and exports would be unaffected by a determination of nonregulated status to A1-DFR petunias.

<b>Summary of Potential Impacts for the Alternatives Considered</b>		
<b>Analysis</b>	<b>No Action Alternative: Continue to Regulate A1-DFR Petunias as a Plant Pest</b>	<b>Preferred Alternative: Determination of Nonregulated Status for A1-DFR Petunias</b>
<b>Coordinated Framework</b>		
U.S. Regulatory Agencies	Because A1-DFR petunias do not contain a GE pesticide and there is no change to pesticide use and A1-DFR petunias are not intended for human and animal consumption a consultation with the EPA is not required and FDA’s voluntary consultation is not necessary.	Because A1-DFR petunias do not contain a GE pesticide and there is no change to pesticide use and A1-DFR petunias are not intended for human and animal consumption a consultation with the EPA is not required and FDA’s voluntary consultation is not necessary.
<b>Regulatory and Policy Compliance</b>		
ESA, CWA, CAA, SDWA, NHPA, EOs	Fully compliant	Fully compliant

**Finding of No Significant Impact**

The analysis in the EA indicates that there will not be a significant impact, individually or cumulatively, on the quality of the human environment as a result of this proposed action. I agree with this conclusion and therefore find that an EIS need not be prepared. This NEPA finding is based on the following context and intensity factors (40 CFR part 1508.27).

*Context*

The term “context” means identification of the locations and resources that could potentially be affected by the Agency’s action. The EA identified the areas in which petunia is grown and may be cultivated in the United States, and those aspects of the human environment potentially affected by the Agency’s regulatory decision. This action has the potential to affect GE and non-GE petunia production systems; environments adjacent to and associated with A1-DFR petunias production systems; and domestic and foreign horticultural markets. The areas affected by a determination of nonregulated status of A1-DFR petunias are those areas of the United States in which producers and consumers can grow A1-DFR petunias. In the United States, petunia is commercially produced in many states. Michigan, Ohio, New York, and Pennsylvania are the leading states in terms of number of producers. In 2018, there were 1,056 producers of nursery flats in the United States, 984 of hanging baskets, and 872 of pots (USDA-NASS 2019a). If all petunias produced were eventually planted in outdoor flower gardens in the United States, the planted area would be fairly small, about 419 million sq. ft. (~10,000 acres), a small area compared to the total covered area for commercial floriculture crop production of 859 million sq. ft. (USDA-NASS 2019a) and a very tiny fraction of the 319 million acres planted in principle crops in the United States (USDA-NASS 2019b).

Several GE cut flower varieties are currently produced: 19 varieties of GE carnation (*Dianthus caryophyllus*), 1 GE rose (*Rosa × hybrida*), and 1 baby’s breath (*Gypsophila* spp.) (USDA 2016).



During 2015 and 2016, bright orange-colored petunias were observed in flower boxes decorating the Helsinki railway station (Servick 2017). The cultivar at the Helsinki railways station was Bonnie Orange. Tests showed that this variety was GE (Haselmair-Gosch et al. 2018). Additionally, these tests suggested the GE petunia was the same as that developed by Meyer et al. (1987) (Meyer et al. 1987). Distributors apparently imported or bred the flowers without realizing the plants were GE varieties. On May 2, 2017, the Germany-based horticultural firm Selecta Klemm informed APHIS that it had moved a GE orange petunia into the United States (Malakoff 2017). This led to testing by USDA of numerous petunia varieties, which confirmed this particular variety and several others were GE and met APHIS' regulatory definition of a regulated article under 7 CFR part 340. On May 16, 2017, APHIS announced to the public and industry that several varieties of GE petunias had been imported into the United States and distributed interstate without proper APHIS authorization (Malakoff 2017). The USDA asked the industry supply chain to voluntarily stop sale of the unauthorized GE varieties. APHIS worked with breeders and growers represented by the American Seed Trade Association (ASTA) and AmericanHort to ensure that all the implicated GE petunia varieties were withdrawn from distribution and destroyed. The petunia industry has voluntarily removed GE petunias from commerce.

A1-DFR petunias will provide additional color varieties of petunia and is expected to compete with other color varieties that are currently in production and offered for sale in the United States. Commercial production of petunia will continue to be dictated by the domestic and import floral market demands and choices made by consumers, not only for petunias, but for other flowers that serve similar ornamental purposes as potted plants, hanging baskets, and in flower beds. A determination of nonregulated status for A1-DFR petunias is not expected to change the acreage, methods, and areas used for petunia seed and bedding plant production.

### *Intensity*

Within the context discussed above, intensity means the degree or severity of potential impacts. As recommended by CEQ (40 CFR part 1508.27), the following were considered in evaluating intensity and making this NEPA determination.

1. *Impacts that may be both beneficial and adverse.*

A determination of nonregulated status for A1-DFR petunias will have no significant environmental impact on the availability of petunia varieties. As considered and analyzed in Chapter 4 of the EA, a determination of nonregulated status for A1-DFR petunias is not expected to change the acreage, methods, and areas used for petunia seed and bedding plant production. The availability of A1-DFR petunias will not alter the areas of commercial petunia production in the United States, and there are no anticipated changes in the availability of petunia varieties on the market. A determination of nonregulated status for A1-DFR petunias will provide additional color varieties of petunia and is expected to compete with other color varieties that are currently in production and offered for sale in the United States.

2. *The degree to which the proposed action affects public health or safety.*

Petunia is not consumed by humans and is not used as animal feed, therefore, FDA's voluntary consultation is not necessary. The potential human health impacts associated

with pesticide use for the production of A1-DFR petunias would be the same as those used for conventional petunia varieties as production practices will not change. The EPA WPS will continue to provide the same level of protection as is currently available.

3. *Unique characteristics of the geographic area such as proximity to historic or cultural resources, park lands, prime farmlands, wetlands, wild and scenic rivers, or ecologically critical areas.*

The EA concluded it is unlikely that historic or cultural resources, park lands, prime farm lands, wetlands, wild and scenic areas, or ecologically critical areas would be significantly impacted by a determination of nonregulated status for A1-DFR petunias. There are no species in the *Petunia* genus that are native to the United States, although there are several introduced (naturalized) species that derived from *Petunia* plants/seed brought to the United States during the early 1900s. Introduced petunias can be found along roadsides, edges of fields, areas along railroads, cracks along urban sidewalks and roadside curbs, edges of garden beds, vacant lots, and waste ground (Hilty 2017). Hybrids of closely related *Petunia* species are rare in nature with varying degrees of fertility (Jędrzejuk et al. 2017). Therefore, invasion of park lands, wetlands, wild and scenic areas, or ecologically critical areas by A1-DFR petunias or feral hybrids is considered unlikely. APHIS conducted a PPRA and concluded that it is unlikely that A1-DFR petunias will become weedy or invasive, and that it is similarly unlikely that gene introgression from A1-DFR petunias into wild *Petunia* species will increase the weediness of any A1-DFR petunias hybrids (USDA-APHIS 2020). Consequently, a determination of nonregulated status for A1-DFR petunias is not expected to have significant impacts on historic or cultural resources, park lands, prime farmlands, wetlands, wild and scenic rivers, or ecologically critical areas.

4. *The degree to which the effects on the quality of the human environment are likely to be highly controversial.*

Approval of Westhoff's petition for nonregulated status for A1-DFR petunias is not an action considered highly controversial in nature. The EA concluded that the agronomic practices and inputs that would be used for production of A1-DFR petunias are no different than those utilized for production of current petunia varieties. Thus, the potential sources of impacts, and the nature of potential impacts on physical and biological resources that could derive from production of A1-DFR petunias are no different than that of currently cultivated petunia varieties. The change in color in A1-DFR petunias does not cause changes in growth habit, temperature tolerances, nutritional requirements, or other factors that would alter where or how it can be grown compared to non-GE petunia varieties; they present no risk to plants, animals, and other taxa. There are no novel or unique impacts on the human environment, nor any considered controversial, that would derive from approval of the petition.

5. *The degree to which the possible effects on the human environment are highly uncertain or involve unique or unknown risks.*

The potential impacts of petunia production on the human environment are well understood and thoroughly evaluated in the EA. A1-DFR petunias will provide additional color varieties of petunia and is expected to compete with other color varieties that are

currently in production and offered for sale in the United States. Commercial production of petunia will continue to be dictated by the domestic and import floral market demands and choices made by consumers, not only for petunias, but for other flowers that serve similar ornamental purposes as potted plants, hanging baskets, and in flower beds. Over a million orange petunias (presumed to be GE petunia varieties) have been sold over the last 15 years (COGEM 2017); APHIS is unaware of any reports of GE petunia populations that have formed naturalized populations, or adversely impacted naturalized populations. APHIS is unaware of any reports of GE petunia populations adversely impacting the built environment. Therefore, the impacts are not highly uncertain, and do not involve unique or unknown risks.

6. *The degree to which the action may establish a precedent for future actions with significant effects or represents a decision in principle about a future consideration.*

Approval of Westhoff's petition would not establish a precedent for future actions that would result in significant impacts on the human environment, nor would it represent a decision in principle about a future decision. Approval of the petition is based upon an independent determination of whether A1-DFR petunias are unlikely to pose a plant pest risk (USDA-APHIS 2020) pursuant to 7 CFR part 340, and an environmental analysis consistent with NEPA and CEQ implementing regulations. APHIS has reviewed and approved petitions for nonregulated status since 1992. All petitions submitted were reviewed independent of the other, and determinations of regulatory status issued in part based on plant pest risk assessments and relevant NEPA analyses specific for the GE organism subject of the petition. Each petition that APHIS receives is specific for a particular GE organism-trait combination and undergoes an independent review to determine if the regulated article may pose a plant pest risk. The requirements for petitions for nonregulated status, applicable to both APHIS and the petitioner, are described in 7 CFR part 340. These requirements have been reviewed above under the sections summarizing APHIS' regulatory authority, and APHIS' requirements to respond to petitions for nonregulated status.

7. *Whether the action is related to other actions with individually insignificant but cumulatively significant impacts.*

The EA discusses potential cumulative impacts on horticultural practices and inputs; human and animal health; physical and biological resources; as well as on socioeconomic issues. Impacts from the cultivation of A1-DFR petunias would not be considered cumulatively significant and no different from that which occurs with currently cultivated petunia varieties.

8. *The degree to which the action may adversely affect districts, sites, highways, structures, or objects listed in or eligible for listing in the National Register of Historic Places or may cause loss or destruction of significant scientific, cultural, or historic resources.*

The EA concluded that approval of the petition is not an action that would directly or indirectly alter the character or use of properties protected under the National Historic Preservation Act. It would have no impact on districts, sites, highways, structures, or objects listed in, or eligible for listing in, the National Register of Historic Places, nor cause any loss or destruction of significant scientific, cultural, or historic resources.

9. *The degree to which the action may adversely affect the endangered or threatened species or its habitat that has been determined to be critical under the Endangered Species Act of 1973.*

APHIS analyzed the potential effects of A1-DFR petunias on threatened and endangered species and critical habitat in Chapter 6 of the EA. APHIS concluded that approval of the petition for nonregulated status for A1-DFR petunias, and any subsequent commercial production of these petunia events, will have no effect on listed species or species proposed for listing, and would not affect designated habitat or habitat proposed for designation. Because of this no-effect determination, consultation under Section 7(a)(2) of the Act or the concurrences of the U.S. Fish and Wildlife Service and National Marine Fisheries Services are not required.

10. *Whether the action threatens a violation of federal, state, or local law or requirements imposed for the protection of the environment.*

The EA evaluated the federal, state, and local laws and regulations, executive orders, and policy related to Westhoff's petition. The EA concluded that approval of the petition would not lead to circumstances that resulted in non-compliance with federal, state, or local laws and regulations providing protections for environmental and human health.

### **NEPA Finding and Rationale**

I have carefully reviewed the EA prepared for this NEPA finding and the input from the public involvement process. In light of the FONSI, APHIS will implement Alternative 2 as described in the EA (Determination that A1-DFR petunias are No Longer Regulated Articles). This alternative meets APHIS' purpose and need to allow the safe development and use of GE organisms, and is consistent with the plant pest provisions of the PPA.

As stated in CEQ regulations, "the agency's preferred alternative is the alternative which the agency believes would fulfill its statutory mission and responsibilities, giving consideration to economic, environmental, technical and other factors." The Preferred Alternative has been selected for implementation taking into consideration a number of environmental, economic, and social factors. Based upon our evaluation and analysis, the Preferred Alternative is selected because (1) it allows APHIS to fulfill its statutory mission to protect the health and value of American agriculture and natural resources using a science-based regulatory framework that allows for the safe development and use of GE organisms; and (2) it allows APHIS to fulfill its regulatory obligations. As a result of the analyses conducted in the EA and summarized in this FONSI, I have concluded that granting nonregulated status to Westhoff's A1-DFR petunias will have no significant impacts on the human environment as a result of making a determination of nonregulated status.

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Bernadette Juarez  
APHIS Deputy Administrator

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Date

Biotechnology Regulatory Services  
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U.S. Department of Agriculture

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