GENERAL LICENSE NO. 6A

Transactions Related to the Exportation or Reexportation of Agricultural Commodities, Medicine, Medical Devices, Replacement Parts and Components, or Software Updates, the Coronavirus Disease 2019 (COVID-19) Pandemic, or Clinical Trials

(a) Except as provided in paragraph (c) of this general license, all transactions prohibited by the Russian Harmful Foreign Activities Sanctions Regulations, 31 CFR part 587 (RuHSR), that are ordinarily incident and necessary to: (1) the exportation or reexportation of agricultural commodities, medicine, medical devices, replacement parts and components for medical devices, or software updates for medical devices to, from, or transiting the Russian Federation; (2) the prevention, diagnosis, or treatment of COVID-19 (including research or clinical studies relating to COVID-19); or (3) ongoing clinical trials and other medical research activities that were in effect prior to March 24, 2022, are authorized.

(b) For the purposes of this general license, agricultural commodities, medicine, and medical devices are defined as follows:

(1) Agricultural commodities. For the purposes of this general license, agricultural commodities are products that fall within the term “agricultural commodity” as defined in section 102 of the Agricultural Trade Act of 1978 (7 U.S.C. 5602) and are intended for use as:

   (i) Food for humans (including raw, processed, and packaged foods; live animals; vitamins and minerals; food additives or supplements; and bottled drinking water) or animals (including animal feeds);

   (ii) Seeds for food crops;

   (iii) Fertilizers or organic fertilizers; or

   (iv) Reproductive materials (such as live animals, fertilized eggs, embryos, and semen) for the production of food animals.

(2) Medicine. For the purposes of this general license, medicine is an item that falls within the definition of the term “drug” in section 201 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321).
(3) **Medical devices.** For the purposes of this general license, a medical device is an item that falls within the definition of “device” in section 201 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321).

(c) This general license does not authorize:

(1) The opening or maintaining of a correspondent account or payable-through account for or on behalf of any entity subject to Directive 2 under E.O. 14024, *Prohibitions Related to Correspondent or Payable-Through Accounts and Processing of Transactions Involving Certain Foreign Financial Institutions*;

(2) Any debit to an account on the books of a U.S. financial institution of the Central Bank of the Russian Federation, the National Wealth Fund of the Russian Federation, or the Ministry of Finance of the Russian Federation; or

(3) Any transaction prohibited by Executive Order (E.O.) 14066 or E.O. 14068.

(d) Effective March 24, 2022, General License No. 6, dated February 24, 2022, is replaced and superseded in its entirety by this General License No. 6A.

**Note 1 to General License No. 6A.** Transactions prohibited by E.O. 14066 or E.O. 14068 include new investment in certain sectors in the Russian Federation and the importation into the United States of certain products of Russian Federation origin, such as alcoholic beverages and fish, seafood, or preparations thereof.

**Note 2 to General License No. 6A.** Nothing in this general license relieves any person from compliance with any other Federal laws or requirements of other Federal agencies.

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**Andrea M. Gacki**

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Date: 2022.03.24 15:37:32 -04'00'

Andrea M. Gacki
Director
Office of Foreign Assets Control

Dated: March 24, 2022