SUPPLEMENT NO. 4 TO PART 744-ENTITY LIST-Continued

Country	Entity			License requirement	License review policy		Federal Register citation	
	alias: —Aviation Aviation 3 United Cottage	Aerospace, a.k.a., t International. Trebeck Street, M Kingdom W1J7 LS; b, The Street, Nacto ited Kingdom. IP10	ayfair, London, <i>and</i> Station on, Ipswich, Suf-	For all items subject to the EAR. (See § 744.11 of the EAR).	Presumption of de	enial	84 FR [INSERT FR PAGE NUMBER] 9/22/20.	
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Matthew S. Borman,

Deputy Assistant Secretary for Export Administration. [FR Doc. 2020–18515 Filed 9–21–20; 8:45 am] BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 514

[Docket No. FDA-2017-N-6381]

RIN 0910-AH51

Postmarketing Safety Reports for Approved New Animal Drugs; Electronic Submission Requirements; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is correcting a final rule that published in the **Federal Register** of July 29, 2020. That final rule requires electronic submission of certain postmarketing safety reports for approved new animal drugs and provides a procedure for requesting a temporary waiver of the electronic submission requirement. Table 2 of the final rule published with errors and this document corrects those errors. We are placing a corrected copy of the final rule in the docket.

DATES: Effective September 22, 2020.

FOR FURTHER INFORMATION CONTACT:

Linda Walter-Grimm, Center for Veterinary Medicine (HFV–240), Food and Drug Administration, 7519 Standish Pl., MPN4, Rm. 2666, Rockville, MD

TABLE 2—EXECUTIVE ORDER 13771 SUMMARY TABLE

[In 2016 Dollars over an infinite time horizon]

	Primary	Lower bound	Upper bound	Primary	Lower bound	Upper bound
	(7%)	(7%)	(7%)	(3%)	(3%)	(3%)
Present Value of Costs Present Value of Cost Savings Present Value of Net Costs Annualized Costs Annualized Cost Savings Annualized Net Costs	\$69,720 73,557 (3,837) 4,880 5,149 (269)			\$75,346 171,634 (96,287) 2,260 5,149 (2,889)		

Dated: August 14, 2020. Lauren K. Roth, Associate Commissioner for Policy. [FR Doc. 2020–18263 Filed 9–21–20; 8:45 am] BILLING CODE 4164–01–P

20855, 240–402–5762, Linda.Walter-Grimm@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 29, 2020, (85 FR 45505), FDA published the final rule "Postmarketing Safety Reports for Approved New Animal Drugs; Electronic Submission Requirements" with errors in table 2.

In FR Doc. 2020–15441, appearing on page 45509 in the **Federal Register** of July 29, 2020, the following corrections are made: