

According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

OMB APPROVED
0579-0036

Interagency Report Control No. 0180-DOA-AN

Fiscal year: 2020

**UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE**

**Annual Report of Research Facility
Column E Explanation**

(TYPE OR PRINT)

This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.

1. REGISTRATION NUMBER

43-R-0014

2. Research Facility Headquarters address

*Boehringer Ingelheim Animal Health USA Inc.
3239 Satellite Blvd
Duluth, GA 30096*

3. Number of animals used in the study.

(b) (4)

4. Species (common name) of animals used in the study.

(b) (4)

5. Explain the procedure producing pain and distress.

This (b) (4) requires (b) (4) with an (b) (4) (b) (4) material in order to (b) (4) Due to the (b) (4) (b) (4)

6. Provide the scientific justification for not providing the appropriate anesthetics, analgesics, or tranquilizing drugs during procedures where the animal experienced accompanying pain or distress greater than momentary or slight.

The testing procedures for biologics (vaccines) are established by the USDA Center for Veterinary Biologics (USDA CVB) and are codified in (b) (4) (b) (4) This testing procedure is (b) (4) determine the (b) (4) are treated with Metacam (b) (4) the same time (b) (4) Metacam (b) (4) (b) (4)

7. What, if any, Federal regulations require this procedure? Cite the agency, the Code of Federal Regulations (CFR) title number, and the specific section number (e.g., APHIS, 9 CFR 113, 102):

USDA APHIS (b) (4) (b) (4)

Agency **USDA APHIS**

CFR **(b) (4)**

03 DEC 2020

According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

OMB APPROVED
0579-0036

Interagency Report Control No. 0180-DOA-AN

Fiscal year: 2020

**UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE**

**Annual Report of Research Facility
Column E Explanation**

(TYPE OR PRINT)

This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.

1. REGISTRATION NUMBER

43-R-0014

2. Research Facility Headquarters address

*Boehringer Ingelheim Animal Health USA Inc.
3239 Satellite Blvd
Duluth, GA 30096*

3. Number of animals used in the study.

(b) (4)

4. Species (common name) of animals used in the study.

(b) (4)

5. Explain the procedure producing pain and distress.

*This (b) (4) requires (b) (4) to be administered (b) (4)
(b) (4) Although (b) (4)
(b) (4) are expected this (b) (4)
(b) (4) and cause (b) (4)*

6. Provide the scientific justification for not providing the appropriate anesthetics, analgesics, or tranquilizing drugs during procedures where the animal experienced accompanying pain or distress greater than momentary or slight.

*If the (b) (4) the
(b) (4) mild. Pain relieving (b) (4)
(b) (4) and would (b) (4) relating to (b) (4)*

7. What, if any, Federal regulations require this procedure? Cite the agency, the Code of Federal Regulations (CFR) title number, and the specific section number (e.g., APHIS, 9 CFR 113, 102):

USDA APHIS (b) (4)

Agency **USDA APHIS**

CFR **(b) (4)**

03 DEC 2020

According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

OMB APPROVED
0579-0036

Interagency Report Control No. 0180-DOA-AN

Fiscal year: 2020

**UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE**

**Annual Report of Research Facility
Column E Explanation**

(TYPE OR PRINT)

This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.

1. REGISTRATION NUMBER

43-R-0014

2. Research Facility Headquarters address

*Boehringer Ingelheim Animal Health USA Inc.
3239 Satellite Blvd
Duluth, GA 30096*

3. Number of animals used in the study.

(b) (4)

4. Species (common name) of animals used in the study.

(b) (4)

5. Explain the procedure producing pain and distress.

This (b) (4) requires (b) (4) dilutions and (b) (4) may experience (b) (4) (b) (4)

6. Provide the scientific justification for not providing the appropriate anesthetics, analgesics, or tranquilizing drugs during procedures where the animal experienced accompanying pain or distress greater than momentary or slight.

(b) (4) is required to determine the (b) (4) content of the (b) (4). A study was (b) (4) to determine if (b) (4) will impact the (b) (4). The study data will be (b) (4) added to the (b) (4) (b) (4) approval is granted.

7. What, if any, Federal regulations require this procedure? Cite the agency, the Code of Federal Regulations (CFR) title number, and the specific section number (e.g., APHIS, 9 CFR 113, 102):

*USDA APHIS Per: APHIS approved Outlines of Production
(b) (4)*

Agency **USDA APHIS**

CFR

According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0579-0036. The time required to complete this information collection is estimated to average .5 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

OMB APPROVED
0579-0036

Interagency Report Control No. 0180-DOA-AN

Fiscal year: 2020

**UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE**

**Annual Report of Research Facility
Column E Explanation**

(TYPE OR PRINT)

This information is required by law (7 U.S.C. 2143 and 9 C.F.R. §2.36). Failure to report according to the regulations can result in an order to cease and desist.

1. REGISTRATION NUMBER

43-R-0014

2. Research Facility Headquarters address

*Boehringer Ingelheim Animal Health USA Inc.
3239 Satellite Blvd
Duluth, GA 30096*

3. Number of animals used in the study.

(b) (4)

4. Species (common name) of animals used in the study.

(b) (4)

5. Explain the procedure producing pain and distress.

*This (b) (4) requires (b) (4)
order to (b) (4)
(b) (4)*

6. Provide the scientific justification for not providing the appropriate anesthetics, analgesics, or tranquilizing drugs during procedures where the animal experienced accompanying pain or distress greater than momentary or slight.

The testing procedures for biologics (vaccines) are established by the USDA Center for Veterinary Biologics (USDA CVB) and are codified in 9 Code of Federal Regulations (CFR): Animals and Animal Products. This testing procedure is codified in (b) (4)

Testing is required to verify (b) (4)

According to the 9 CFR (b) (4)

(b) (4)

(b) (4)

7. What, if any, Federal regulations require this procedure? Cite the agency, the Code of Federal Regulations (CFR) title number, and the specific section number (e.g., APHIS, 9 CFR 113, 102):

USDA APHIS (b) (4)

Agency **USDA APHIS**

CFR **(b) (4)**