ANNEX II

ANIMALS

**Form for the Submission of Prior Consultations for Animals**

1. For submissions concerning a hypothetical product, only the **General Module** of this form shall be completed.

2. Where the same genomic intervention has given rise to multiple products or lines, a single PC shall be submitted, with the common information grouped in the General Module in order to avoid repetition.

## I. DEVELOPER INFORMATION MODULE

Name and address of the entity submitting the consultation:

a. Name and address of the entity developing the product:

b. Identification of the line/product/development:

**Legal representative submitting the consultation:**

a. Name:

b. Address:

c. Telephone:

d. Email:

**Technical officer submitting the consultation:**

a. Name:

b. Address:

c. Telephone:

d. Email:

## II. GENERAL MODULE

A. Taxonomic description.

B. Select the type of product:

| | |
|---|---|
| | Real / obtained |
| | Hypothetical / under development |

C. Name assigned to the line/product/development.

D. Brief description of the objective of the intervention carried out or to be carried out.

E. Detailed description of the technique used or to be used, and all steps applied in the case presented.

F. Molecular description of the organism’s target nucleotide sequences in their state prior to application of the technique.

G. Description of the function of the sequences in their state prior to application of the technique.

H. Description of the expected/observed changes in sequence function and phenotype following application of the technique.

I. Map of any genetic construct or nucleic acid fragment used or intended to be used in the development process, detailing the genetic elements (where applicable).

## III. FINAL PRODUCT MODULE

A. Changes observed in the target sequences following application of the technique.

B. Provide experimental evidence that the product obtained contains no residual genetic constructs or nucleic acid fragments used in the genome intervention process.

## IV. PHENOTYPIC MODULE

Where the submission concerns a product, report the occurrence of any effects other than the intended phenotype.

## V. REFERENCES

A. Provide copies of any publications cited.

B. Provide a certificate or report evidencing approval of the procedures by the animal care and use committee, where animal experimentation is involved.

Argentine Republic - National Executive Branch
YEAR OF THE RECONSTRUCTION OF THE ARGENTINE NATION

**Additional Signature Page**

**Annex**

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CITY OF BUENOS AIRES
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