* Long In Vivo Checklist

AHA - Preclinical Animal Testing: A detailed checklist has been developed as a prerequisite for every publication involving preclinical studies. **Checklist items must be clearly presented in the manuscript, and if an item is not adhered to, an explanation should be provided.** If this information (checklist items and/or explanations) cannot be included in the main manuscript because of space limitations, please include it in an online supplement. If the manuscript is accepted, this checklist will be published as an online supplement.

This study involves animal models:

Yes

Study Design

The experimental group(s) have been clearly defined in the article, including number of animals in each experimental arm of the study.

Yes

An overall study timeline is provided.

Yes

The protocol was prospectively written

Yes

The primary and secondary endpoints are specified

N/A

For primary endpoints, a description is provided as to how the type I error multiplicity issue was addressed (e.g., correction for multiple comparisons was or was not used and why). (Note: correction for multiple comparisons is not necessary if the study was exploratory or hypothesis-generating in nature).

N/A

A description of the control group is provided including whether it matched the treated groups.

Yes

Inclusion and Exclusion criteria

Allocation concealment was used.

Inclusion and exclusion criteria for enrollment into the study were defined and are reported in the manuscript.

Yes

These criteria were set a priori (before commencing the study).

Yes

Randomization

Animals were randomly assigned to the experimental groups. If random assignment was not used, adequate explanation has been provided.

Yes

Type and methods of randomization have been described.

N/A

Methods used for allocation concealment have been reported.

N/A

Blinding

Blinding procedures with regard to masking of group/treatment assignment from the experimenter were used and are described. The rationale for nonblinding of the experimenter has been provided, if such was not performed.

Yes

Blinding procedures with regard to masking of group assignment during outcome assessment were used and are described.

Yes

If blinding was not performed, the rationale for nonblinding of the person(s) analyzing outcome has been provided.

N/A

Sample size and power calculations

Formal sample size and power calculations were conducted before commencing the study based on *a priori* determined outcome(s) and treatment effect(s), and the data are reported.

Yes

If formal sample size and power calculation was not conducted, a rationale has been provided.

Yes

Data Reporting

Baseline characteristics (species, sex, age, strain, chow, bedding, and source reported.	ee) of animals are	Yes
The number of animals in each group that were randomized, tested, and excreported. If the experimentation involves repeated measurements, the number at each time point is provided is provided for all experimental groups.		Yes
Baseline data on assessed outcome(s) for all experimental groups are repo	rted.	Yes
Details on important adverse events and death of animals during the course reported for all experimental groups.	of the experiment are	N/A
Numeric data on outcomes are provided in the text or in a tabular format in the supplementary tables, in addition to the figures.	ne main article or as	Yes
To the extent possible, data are reported as dot plots as opposed to bar grasmall sample size groups.	aphs, especially for	Yes
In the online Supplemental Material, methods are described in sufficient deta replication of the study.	il to enable full	Yes
Statistical methods		
The statistical methods used for each data set are described.		Yes
For each statistical test, the effect size with its standard error and P value is pencouraged to provide 95% confidence intervals for important comparisons.		Yes
Central tendency and dispersion of the data are examined, particularly for sr	mall data sets.	Yes
Nonparametric tests are used for data that are not normally distributed.		N/A
Two-sided P values are used.		Yes
In studies that are not exploratory or hypothesis-generating in nature, correct hypotheses testing and multiple comparisons are performed.	ions for multiple	N/A
In "negative" studies or null findings, the probability of a type II error is report	ed.	N/A
Experimental details, ethics, and funding statements		
Details on experimentation including formulation and dosage of therapeutic administration, use of anesthesia and analgesia, temperature control during opostprocedural monitoring are described.		Yes
Both male and female animals have been used. If not, the reason/justification	n is provided.	No
Statements on approval by ethics boards and ethical conduct of studies are	provided.	Yes

Yes

Date completed: 09/29/2020 16:53:27 User pid: 11987

Statements on funding and conflicts of interests are provided.