

Life Sciences Reporting Summary

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Please do not complete any field with "not applicable" or n/a. Refer to the help text for what text to use if an item is not relevant to your study. [For final submission](#): please carefully check your responses for accuracy; you will not be able to make changes later.

► Experimental design

1. Sample size

Describe how sample size was determined.

A minimum of 3 independent experiments were carried out for all ex vivo and in vivo studies. For in vivo functional tests, sample sizes were 6 or 7 for therapeutic efficacy of bFGF and 10 for general long-term toxicity so as to achieve statistical difference using ANOVA or Dunnett's test with a 0.05 significance level.

2. Data exclusions

Describe any data exclusions.

Data exclusions were only allowed for animals that died before experimental termination.

3. Replication

Describe the measures taken to verify the reproducibility of the experimental findings.

All experimental findings were carried out in triplicate.

4. Randomization

Describe how samples/organisms/participants were allocated into experimental groups.

Samples and animals were chosen at random for each experimental group.

5. Blinding

Describe whether the investigators were blinded to group allocation during data collection and/or analysis.

Data collection and analysis were not blinded.

Note: all in vivo studies must report how sample size was determined and whether blinding and randomization were used.

6. Statistical parameters

For all figures and tables that use statistical methods, confirm that the following items are present in relevant figure legends (or in the Methods section if additional space is needed).

n/a Confirmed

- ☐ ☒ The exact sample size (*n*) for each experimental group/condition, given as a discrete number and unit of measurement (animals, litters, cultures, etc.)
- ☐ ☒ A description of how samples were collected, noting whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- ☐ ☒ A statement indicating how many times each experiment was replicated
- ☐ ☒ The statistical test(s) used and whether they are one- or two-sided
Only common tests should be described solely by name; describe more complex techniques in the Methods section.
- ☐ ☒ A description of any assumptions or corrections, such as an adjustment for multiple comparisons
- ☐ ☒ Test values indicating whether an effect is present
*Provide confidence intervals or give results of significance tests (e.g. *P* values) as exact values whenever appropriate and with effect sizes noted.*
- ☐ ☒ A clear description of statistics including central tendency (e.g. median, mean) and variation (e.g. standard deviation, interquartile range)
- ☐ ☒ Clearly defined error bars in all relevant figure captions (with explicit mention of central tendency and variation)

See the web collection on [statistics for biologists](#) for further resources and guidance.

► Software

Policy information about [availability of computer code](#)

7. Software

Describe the software used to analyze the data in this study.

Origin Pro 8 for ANOVA (Tukey test), GraphPad InStat 3 for Student's t-test in Fig. 7a, and Sigma plot for unpaired t-test in Fig. S9. Image J for image analysis. The mean value and standard deviation of all bar graphs were calculated and they were plotted using Graph Pad prism 7.

For manuscripts utilizing custom algorithms or software that are central to the paper but not yet described in the published literature, software must be made available to editors and reviewers upon request. We strongly encourage code deposition in a community repository (e.g. GitHub). *Nature Methods* [guidance for providing algorithms and software for publication](#) provides further information on this topic.

► Materials and reagents

Policy information about [availability of materials](#)

8. Materials availability

Indicate whether there are restrictions on availability of unique materials or if these materials are only available for distribution by a third party.

There are no unique materials with restrictions on availability.

9. Antibodies

Describe the antibodies used and how they were validated for use in the system under study (i.e. assay and species).

Two types of antibodies, anti-GFP antibody (ab6556, Abcam) and goat anti-rabbit IgG flamma 488 (RSA1241, BioActs), were used for IHC. They were used in accordance with the manufacturer's dilution recommendations. For GFP quantification assay, a GFP ELISA kit (AKR-121, Cell Biolabs) was used in accordance with the manufacturer's protocol.

10. Eukaryotic cell lines

a. State the source of each eukaryotic cell line used.

5-6bMSCs (mesenchymal stem cells) derived from human, H9C2 myoblasts derived from rat heart, and AAV293 cells established from primary embryonic human kidney transformed with human adenovirus type 5 DNA.

b. Describe the method of cell line authentication used.

The human 5-6bMSCs were obtained with Institutional Review Board (IRB) approval in Yonsei University (7001988201704BR19101). Other cell lines were authenticated by the commercial source.

c. Report whether the cell lines were tested for mycoplasma contamination.

Mycoplasma contamination was checked for the thawing portion of cell stock used for the experiments. A negative signal was observed.

d. If any of the cell lines used are listed in the database of commonly misidentified cell lines maintained by [ICLAC](#), provide a scientific rationale for their use.

No commonly misidentified cell lines were used.

► Animals and human research participants

Policy information about [studies involving animals](#); when reporting animal research, follow the [ARRIVE guidelines](#)

11. Description of research animals

Provide all relevant details on animals and/or animal-derived materials used in the study.

BALBc mice (8 weeks, male) and SD rats (8 weeks, male) were used in this study. Also, blood plasma was obtained from SD rats (9 weeks, male).

Policy information about [studies involving human research participants](#)

12. Description of human research participants

Describe the covariate-relevant population characteristics of the human research participants.

The study did not involve human research participants.