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Sample-size estimation

- You should state whether an appropriate sample size was computed when the study was being designed
- You should state the statistical method of sample size computation and any required assumptions
- If no explicit power analysis was used, you should describe how you decided what sample (replicate) size (number) to use

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No statistical tests were used to determine sample size. The sample number was based on previous experience with optogenetic experiments in our laboratory and the robustness of the observed effects.

Replicates

- You should report how often each experiment was performed
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- High-throughput sequence data should be uploaded before submission, with a private link for reviewers provided (these are available from both GEO and ArrayExpress)

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The 'n' number is available in the Methods and Materials Section and in the figure legends of figures 1-2-4. For the *in silico* experiments of Attract-Anchor-Drag-based (AAD) control of a spiral wave core we used 10 different simulation domains, composed of neonatal rat atrial cardiomyocytes with 17% randomly distributed cardiac fibroblasts and with natural cellular heterogeneity. Subsequent *in vitro* validation for AAD control of a spiral wave core was performed in 9 CatCh-expressing monolayers of neonatal rat atrial cardiomyocytes. AAD control of spiral wave cores of figure-of-eight type reentry was performed in 6 CatCh-expressing monolayers. AAD control of multiple spiral wave cores was performed in 3 CatCh-expressing monolayers. All monolayers included in this study showed uniform AP propagation at 1-Hz pacing and homogeneous transgene expression, as described in the Methods and Materials Section.

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Statistical reporting

- Statistical analysis methods should be described and justified
- Raw data should be presented in figures whenever informative to do so (typically when N per group is less than 10)
- For each experiment, you should identify the statistical tests used, exact values of N, definitions of center, methods of multiple test correction, and dispersion and precision measures (e.g., mean, median, SD, SEM, confidence intervals; and, for the major substantive results, a measure of effect size (e.g., Pearson's r, Cohen's d)
- Report exact p-values wherever possible alongside the summary statistics and 95% confidence intervals. These should be reported for all key questions and not only when the p-value is less than 0.05.

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We did not calculate p-values and did not perform statistical tests. The sample size is available in the Methods and Materials Section and in the figure legends 1-2-4.

(For large datasets, or papers with a very large number of statistical tests, you may upload a single table file with tests, Ns, etc., with reference to sections in the manuscript.)

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- Indicate how samples were allocated into experimental groups (in the case of clinical studies, please specify allocation to treatment method); if randomization was used, please also state if restricted randomization was applied
- Indicate if masking was used during group allocation, data collection and/or data analysis

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Not applicable.

Additional data files ("source data")

- We encourage you to upload relevant additional data files, such as numerical data that are represented as a graph in a figure, or as a summary table
- Where provided, these should be in the most useful format, and they can be uploaded as "Source data" files linked to a main figure or table
- Include model definition files including the full list of parameters used
- Include code used for data analysis (e.g., R, MatLab)
- Avoid stating that data files are "available upon request"



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Please indicate the figures or tables for which source data files have been provided: Not applicable.