# nature research

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# **Reporting Summary**

Nature Research wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Research policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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For a	ıll statistical ana	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.				
n/a	Confirmed					
	The exact	sample size $(n)$ for each experimental group/condition, given as a discrete number and unit of measurement				
	A stateme	nt on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly				
		ical test(s) used AND whether they are one- or two-sided on tests should be described solely by name; describe more complex techniques in the Methods section.				
$\boxtimes$	A description of all covariates tested					
$\boxtimes$	A descripti	on of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons				
	igwedge A full desc AND variat	description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)				
		or null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i> ) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted ive <i>P values as exact values whenever suitable.</i>				
$\boxtimes$	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings					
$\boxtimes$	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes					
$\boxtimes$	$\square$ Estimates of effect sizes (e.g. Cohen's $d$ , Pearson's $r$ ), indicating how they were calculated					
		Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.				
Sof	tware and	d code				
Polic	y information a	about <u>availability of computer code</u>				
Da	ta collection	No software was used				
Da	ta analysis	Microsoft Excel 16.16.26: Graphpad Prism 7.05/e: 7FN2 Core: Imagel / Java 1.8.0 172: Amira v2020.2: Quantstudio 3: STAR v2.1.3: Cell				

#### Data

Policy information about availability of data

All manuscripts must include a <u>data availability statement</u>. This statement should provide the following information, where applicable:

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

- Accession codes, unique identifiers, or web links for publicly available datasets

Ranger v.3.1.0; Seurat v3.1.1 (R package); R Programming (3.5.2).

- A list of figures that have associated raw data
- A description of any restrictions on data availability

Single cell RNA sequencing GEO deposition: GSE163701. Other data that support the findings of this study are available from the corresponding author upon request. There are no restrictions on data availability.

Field-spe	cific reporting	
Please select the or	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.  Behavioural & social sciences	
	he document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
Life scier	ices study design	
	close on these points even when the disclosure is negative.	
Sample size	For animal experiments, N=5 mice (or more) / group with experiments repeated twice with similar result is what we adhered to for experiments done in main display items. These numbers are standard in mouse research in this field / with the methods used.	
Data exclusions	There were no data exclusions.	
Replication	Animal experiments were repeated at least twice with similar result	
Randomization	Allocation was random.	
Blinding	Investigators analyzing the data were blinded to data allocation performed by another investigator.	
Materials & expression or method list  Materials & expression of the control of t	ChIP-seq  cell lines  Sign of the problem of the pr	
Antibodies		
Antibodies used	Anti-E-cadherin R&D AF748; anti-N-Cadherin Abcam ab98952; anti-CXCL12 sc-28876; anti-fibronectin ab23750; anti-Ki67 RM-9106-S0, 14-5698 and Cell Signaling 9129; anti-PDGFRα ab51875; anti-PDGFRβ ab32570; anti-GFP GTX26673.	
Validation	Each of these antibodies are routinely used in the field. Each has been extensively used and validated by numerous studies by our group and others. This information is available online from the manufacturers for each.	
Eukaryotic c	ell lines	
Policy information	about <u>cell lines</u>	
Cell line source(s	LNCaP , HMVP2, RM1, MycCap	

ATCC, cell culture, tumor growth assays

All lines are negative for mycoplasma

N/A

Authentication

Mycoplasma contamination

Commonly misidentified lines (See <u>ICLAC</u> register)

## Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals C57BL/6 background mice, both sexes, various ages

Wild animals No wild animals involved.

Field-collected samples No field-collected animals involved.

Ethics oversight UTHealth Animal Care and Use Committee

Note that full information on the approval of the study protocol must also be provided in the manuscript.

### Human research participants

Policy information about studies involving human research participants

Population characteristics Males of various ages were included as published in our previous work Zhang T, et al. and Kolonin CXCL1 mediates obesity-associated adipose stromal cell trafficking and function in the tumor microenvironment. Nature Comm. 2016;7:11674-90.

Recruitment Described in our publication Zhang T, et al. and Kolonin CXCL1 mediates obesity-associated adipose stromal cell trafficking

and function in the tumor microenvironment. Nature Comm. 2016.

Ethics oversight MD Anderson Cancer Center

Note that full information on the approval of the study protocol must also be provided in the manuscript.

#### Clinical data

Policy information about <u>clinical studies</u>

All manuscripts should comply with the ICMJE guidelines for publication of clinical research and a completed CONSORT checklist must be included with all submissions.

Clinical trial registration N/A

Study protocol Not a clinical trial

Data collection Adipose tissue samples were collected during prostatectomy.

Outcomes Outcomes were not a part of the study.