Date:9/1/21		 		 	 		 	 	
Your Name:_Xiaojing Huang_									
		 	· · ·	 	 	·	 	 	

Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adipose tissue dysfunction Manuscript number (if known): 153586-JCI-CMED-PRESUB

In the interest of transparency, we ask you to disclose all relationships/activities/interests listed below that are related to the content of your manuscript. "Related" means any relation with for-profit or not-for-profit third parties whose interests may be affected by the content of the manuscript. Disclosure represents a commitment to transparency and does not necessarily indicate a bias. If you are in doubt about whether to list a relationship/activity/interest, it is preferable that you do so.

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1	All support for the present manuscript (e.g., funding, provision of study materials, medical writing, article processing charges, etc.) No time limit for this item.	None Rockefeller University Hospital National Institute of General Medical Sciences National Center for Advancing Translational Sciences National Cancer Institute American Cancer Society American Diabetes Association private donor (Memorial Sloan Kettering Cancer Center	Pilot grant T32GM007739 UL1TR001866 P30CA008748 133831-CSDG-19-117-01-CPHPS 1-17-ACE-17
		Time frame: past	36 months
2		_x None	

	Grants or contracts from		
	any entity (if not indicated		
	in item #1 above).		
3	Royalties or licenses	_x None	
4	Consulting fees	_x None	
5	Payment or honoraria for lectures, presentations,	_x None	
	speakers bureaus,		
	manuscript writing or educational events		
6	Payment for expert testimony	_x None	
7	Support for attending meetings and/or travel	_x None	
8	Patents planned, issued or pending	_x None	
9	Participation on a Data Safety Monitoring Board or	_x None	
	Advisory Board		
10	Leadership or fiduciary role in other board, society,	_x None	
	committee or advocacy		
	group, paid or unpaid		
11	Stock or stock options	_x None	
12	Receipt of equipment, materials, drugs, medical	_x None	
	writing, gifts or other		
	services		
13	Other financial or non- financial interests	_x None	

Please place an "X" next to the following statement to indicate your agreement:
x_ I certify that I have answered every question and have not altered the wording of any of the questions on this form.

Pate:9/1/21
our Name: Olivia Maguire
Agnuscrint Title: Therapeutic andominal radiation during childhood induces chronic adinose tissue dysfunction

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Date:9/1/21
Your Name: Jeanne Walker
Manuscript Title: Therapoutic abdominal radiation during childhood induces chronic adipose tissue dysfunction

Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adipose tissue dysfunction Manuscript number (if known): 153586-JCI-CMED-PRESUB

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Date:9/1/21	
Your Name: Caroline Jiang	
Manuscript Title: Therapoutic addominal radiation during childhood induces chronic adipose tissue dysfunction	•

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Date:9/1/21	
Your Name: Thomas Carroll	

Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adipose tissue dysfunction Manuscript number (if known): 153586-JCI-CMED-PRESUB

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Date:9/1/21
Your Name:_Ji-Dung Luo
Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adipose tissue dysfunction
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Date:9/1/21	 	 	 		
Your Name: Emily Tonorezos					
- '				 	

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8	Patents planned, issued or pending	_x None	
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Date:9/1/21	
Your Name: Danielle Novetsky Friedman	
Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adipose tissue dysfunction	

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	Grants or contracts from	National Cancer Institute	U01 CA246659-A01
	any entity (if not indicated in item #1 above).	Sohn Foundation	grant
		Society of MSK Research	Grant
3	Royalties or licenses	_x None	
4	Consulting fees	_x None	
5	Payment or honoraria for	UpToDate	2014-Present
	lectures, presentations, speakers bureaus, manuscript writing or educational events	ENDO2021: Annual Meeting of the Endocrine Society	March 2021
6	Payment for expert testimony	_x None	
_	Comment for the "	A Al 1'	00/45/2040 00/40/2040/7
7	Support for attending meetings and/or travel	Agence Nationale de la Recherche (ANR), French Cohortes, Jury member	09/16/2019-09/18/2019 (Travel to Paris, France)
8	Patents planned, issued or pending	_x None	
9	Participation on a Data Safety Monitoring Board or Advisory Board	_x None	
10	Leadership or fiduciary role in other board, society, committee or advocacy group, paid or unpaid	_x None	
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	_		
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Date:9/1/21
Your Name: Paul Cohen
Manuscript Title: Therapeutic abdominal radiation during childhood induces chronic adinose tissue dysfunction

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JCI Insight Manuscript 153586-INS-CMED-RV-3

STROBE Statement—Checklist of items that should be included in reports of *cohort studies*

	Item No	Recommendation
Title and abstract	1	Indicate the study's design with a commonly used term in the title or the abstract
		Provide in the abstract an informative and balanced summary of what was done
		and what was found
Introduction		
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported
Objectives	3 🔻	State specific objectives, including any prespecified hypotheses
Methods	<u> </u>	1 3 2 21 1 21
Study design	4	Present key elements of study design early in the paper
Setting	5	Describe the setting, locations, and relevant dates, including periods of recruitment,
		exposure, follow-up, and data collection
Participants	6	
	•	participants. Describe methods of follow-up
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed n/a
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect
		modifiers. Give diagnostic criteria, if applicable
Data sources/	8*	For each variable of interest, give sources of data and details of methods of
measurement	·	assessment (measurement). Describe comparability of assessment methods if there is
		more than one group
Bias	9	Describe any efforts to address potential sources of bias
Study size	10	Explain how the study size was arrived at
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable,
		describe which groupings were chosen and why
Statistical methods	12	Describe all statistical methods, including those used to control for confounding
		(b) Describe any methods used to examine subgroups and interactions n/a
		Explain how missing data were addressed
		(d) If applicable, explain how loss to follow-up was addressed n/a
		(e) Describe any sensitivity analyses n/a
Results		
Participants	13*	Report numbers of individuals at each stage of study—eg numbers potentially
		eligible, examined for eligibility, confirmed eligible, included in the study,
		completing follow-up, and analysed
		Give reasons for non-participation at each stage
		Consider use of a flow diagram
Descriptive data	14*	Give characteristics of study participants (eg demographic, clinical, social) and
		information on exposures and potential confounders
		Indicate number of participants with missing data for each variable of interest
		Summarise follow-up time (eg, average and total amount)
Outcome data	15*	Report numbers of outcome events or summary measures over time
Main results	16	Give unadjusted estimates and, if applicable, confounder-adjusted estimates and
		their precision (eg, 95% confidence interval). Make clear which confounders were
		adjusted for and why they were included
		(b) Report category boundaries when continuous variables were categorized n/a
		(c) If relevant, consider translating estimates of relative risk into absolute risk for a
		meaningful time period n/a

Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses
Discussion		sensiti in y unuiyoes
Key results	18	Jummarise key results with reference to study objectives
Limitations	19	viscuss limitations of the study, taking into account sources of potential bias or
		imprecision. Discuss both direction and magnitude of any potential bias
Interpretation	20	Vive a cautious overall interpretation of results considering objectives, limitations,
		multiplicity of analyses, results from similar studies, and other relevant evidence
Generalisability	21	viscuss the generalisability (external validity) of the study results
Other information		
Funding	22	vive the source of funding and the role of the funders for the present study and, if
		applicable, for the original study on which the present article is based

^{*}Give information separately for exposed and unexposed groups.

Note: An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at http://www.strobe-statement.org.