

ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Valerie L. Darcey

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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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Valerie Darcey 5/8/23

ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Juen Guo

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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Juan Guo

ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Amber Courville

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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Amber Courville 5/8/23

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Date: 5/8/2023

Your Name: Isabelle Gallagher

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

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Isabel Zanger

ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Jason A Avery

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: W. Kyle Simmons, Ph.D.

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: John E Ingeholm

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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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13	Other financial or non-financial interests	☒ None 	
Please place an "X" next to the following statement to indicate your agreement: ☒ I certify that I have answered every question and have not altered the wording of any of the questions on this form.			

John E. Longhulen
5/10/23

ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Peter Herscovitch, M.D.

Manuscript Title: **Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial**

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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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ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Alex Martin

Manuscript Title: Restriction of dietary fat, affects brain reward regions in adults with obesity in a randomized crossover trial

Manuscript Number (if known): JCI Insight: 169759-INS-RG-RV-3

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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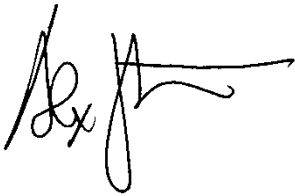
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ICMJE DISCLOSURE FORM

Date: 5/8/2023

Your Name: Kevin Hall

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Reporting checklist for randomised trial.

Based on the CONSORT guidelines.

Instructions to authors

Complete this checklist by entering the page numbers from your manuscript where readers will find each of the items listed below.

Your article may not currently address all the items on the checklist. Please modify your text to include the missing information. If you are certain that an item does not apply, please write "n/a" and provide a short explanation.

Upload your completed checklist as an extra file when you submit to a journal.

In your methods section, say that you used the CONSORT reporting guidelines, and cite them as:

Schulz KF, Altman DG, Moher D, for the CONSORT Group. CONSORT 2010 Statement: updated guidelines for reporting parallel group randomised trials

		Reporting Item	Page Number
Title and Abstract			
Title	#1a	Identification as a randomized trial in the title.	1
Abstract	#1b	Structured summary of trial design, methods, results, and conclusions	1
Introduction			
Background and objectives	#2a	Scientific background and explanation of rationale	2
Background and objectives	#2b	Specific objectives or hypothesis	2, 9
Methods			
Trial design	#3a	Description of trial design (such as parallel, factorial) including allocation ratio.	17-18
Trial design	#3b	Important changes to methods after trial commencement (such as eligibility criteria), with	N/A no method changes.

		reasons	
Participants	#4a	Eligibility criteria for participants	17
Participants	#4b	Settings and locations where the data were collected	17-18
Interventions	#5	The experimental and control interventions for each group with sufficient details to allow replication, including how and when they were actually administered	19-21
Outcomes	#6a	Completely defined prespecified primary and secondary outcome measures, including how and when they were assessed	2
Outcomes	#6b	Any changes to trial outcomes after the trial commenced, with reasons	N/A no outcomes changes.
Sample size	#7a	How sample size was determined.	25-26
Sample size	#7b	When applicable, explanation of any interim analyses and stopping guidelines	N/A no interim analyses.
Randomization - Sequence generation	#8a	Method used to generate the random allocation sequence. Page 17.	
Randomization - Sequence generation	#8b	Type of randomization; details of any restriction (such as blocking and block size) – page 17	
Randomization - Allocation concealment mechanism	#9	Mechanism used to implement the random allocation sequence (such as sequentially numbered containers), describing any steps taken to conceal the sequence until interventions were assigned	18
Randomization - Implementation	#10	Who generated the allocation sequence, who enrolled participants, and who assigned participants to interventions	18

Blinding	#11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those assessing outcomes) and how.	N/A no blinding.
Blinding	#11b	If relevant, description of the similarity of interventions	N/A
Statistical methods	#12a	Statistical methods used to compare groups for primary and secondary outcomes	25-27
Statistical methods	#12b	Methods for additional analyses, such as subgroup analyses and adjusted analyses	N/A
Results			
Participant flow diagram (strongly recommended)	#13a	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analysed for the primary outcome	Suppl. Figure 1
Participant flow	#13b	For each group, losses and exclusions after randomization, together with reason	Suppl Figure 1 & p18
Recruitment	#14a	Dates defining the periods of recruitment and follow-up	18
Recruitment	#14b	Why the trial ended or was stopped	N/A
Baseline data	#15	A table showing baseline demographic and clinical characteristics for each group	35 (Table 1)
Numbers analysed	#16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	18-19
Outcomes and estimation	#17a	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision (such as 95% confidence interval)	Tables 2 & 3
Outcomes and	#17b	For binary outcomes, presentation of both	N/A no binary

estimation		absolute and relative effect sizes is recommended	outcomes.
Ancillary analyses	#18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	Exploratory results indicated p7 & Table 3
Harms	#19	All important harms or unintended effects in each group (For specific guidance see CONSORT for harms)	19
Discussion			
Limitations	#20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	15
Generalisability	#21	Generalisability (external validity, applicability) of the trial findings	15-16
Interpretation	#22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	9
Registration	#23	Registration number and name of trial registry	17
Other information			
Interpretation	#22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	9
Registration	#23	Registration number and name of trial registry	17
Protocol	#24	Where the full trial protocol can be accessed, if available	N/A
Funding	#25	Sources of funding and other support (such as supply of drugs), role of funders	28-29

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