

ICMJE DISCLOSURE FORM

Date: 11/14/2022

Your Name: Armin Ahmadi

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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: 11/15/2022

Your Name: Gwenaëlle Begue

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 11/15/2022

Your Name: Ana P. Valencia

Manuscript Title: The impact of nicotinamide riboside and coenzyme Q10 on physical endurance capacity and metabolic profile in patients with chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 11/16/2022

Your Name: Jennifer E. Norman

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: Click or tap to enter a date.

Your Name: 10/19/2022
Benjamin Lidgard

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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☐

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ICMJE DISCLOSURE FORM

Date: 10/31/2022

Your Name: Matthew P. Van Doren

Manuscript Title: The impact of nicotinamide riboside and coenzyme Q10 on physical endurance capacity and metabolic profile in patients with chronic kidney disease

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ICMJE DISCLOSURE FORM

Date: 11/17/2022

Your Name: David Marcinek

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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: 11/15/2022

Your Name: Sili Fan

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 3/30/2023

Your Name: Brian bennett

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 10/20/2022

Your Name: David Prince

Manuscript Title: The impact of nicotinamide riboside and coenzyme Q10 on physical endurance capacity and metabolic profile in patients with chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 3/27/2023

Your Name: Jonathan Himmelfarb

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 3/28/2023

Your Name: Jorge L. Gamboa

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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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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.

ICMJE DISCLOSURE FORM

Date: 3/27/2023

Your Name: Ian de Boer

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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: 3/29/2023

Your Name: Bryan Kestenbaum

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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ICMJE DISCLOSURE FORM

Date: 3/27/2023

Your Name: Baback Roshanravan

Manuscript Title: The impact of coenzyme Q10 and nicotinamide riboside on physical endurance capacity and metabolic profile in chronic kidney disease

Manuscript Number (if known): 167274-INS-CMED-RV-2

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		Chromadex	Donation of Nicotinamide Riboside to Univ of WA
		Tishcon	Donation of Coenzyme Q10 to Univ of WA
13	Other financial or non-financial interests	☒ None	
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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	2
Introduction			
Background and objectives	#2a	Scientific background and explanation of rationale	4 and 5
Background and objectives	#2b	Specific objectives or hypothesis	5
Methods			
Trial design	#3a	Description of trial design (such as parallel, factorial) including allocation ratio.	19
Trial design	#3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons	NA

Participants	#4a	Eligibility criteria for participants	21
Participants	#4b	Settings and locations where the data were collected	21
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 and 20
Outcomes	#6a	Completely defined prespecified primary and secondary outcome measures, including how and when they were assessed	2, 21, and 22
Sample size	#7a	How sample size was determined.	24
Sample size	#7b	When applicable, explanation of any interim analyses and stopping guidelines	NA
Randomization - Sequence generation	#8a	Method used to generate the random allocation sequence.	
19 and 20			
Randomization - Sequence generation	#8b	Type of randomization; details of any restriction (such as blocking and block size)	
19			
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	19
Randomization - Implementation	#10	Who generated the allocation sequence, who enrolled participants, and who assigned participants to interventions	NA
Blinding	#11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those assessing outcomes) and how.	19
Blinding	#11b	If relevant, description of the similarity of interventions	NA
Statistical methods	#12a	Statistical methods used to compare groups for primary and secondary outcomes	22-24
Statistical methods	#12b	Methods for additional analyses, such as subgroup	22 and

		analyses and adjusted analyses	23
Outcomes	#6b	Any changes to trial outcomes after the trial commenced, with reasons	NA
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	37
Participant flow	#13b	For each group, losses and exclusions after randomization, together with reason	37
Recruitment	#14a	Dates defining the periods of recruitment and follow-up	37
Recruitment	#14b	Why the trial ended or was stopped	NA
Baseline data	#15	A table showing baseline demographic and clinical characteristics for each group	32
Numbers analysed	#16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	32
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)	33 to 36
Outcomes and estimation	#17b	For binary outcomes, presentation of both absolute and relative effect sizes is recommended	NA
Ancillary analyses	#18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	38-40
Harms	#19	All important harms or unintended effects in each group (For specific guidance see CONSORT for harms)	45
Discussion			
Limitations	#20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	16 and 17
Interpretation	#22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	11-18

Registration	#23	Registration number and name of trial registry	3
Generalisability	#21	Generalisability (external validity, applicability) of the trial findings	11-18
Other information			
Interpretation	#22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	11-18
Registration	#23	Registration number and name of trial registry	3
Protocol	#24	Where the full trial protocol can be accessed, if available	NA
Funding	#25	Sources of funding and other support (such as supply of drugs), role of funders	3

Notes:

- 6a: 2, 21, and 22 The CONSORT checklist is distributed under the terms of the Creative Commons Attribution License CC-BY. This checklist was completed on 27. March 2023 using <https://www.goodreports.org/>, a tool made by the [EQUATOR Network](#) in collaboration with [Penelope.ai](#)

Supplemental Figure 1. Study design and sample size. Given the efficient crossover design of the study, all 25 participants received each supplement by the end of the study.

