

#### **Instructions**

3.

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Section 1. Identifying Inform	ation	
1. Given Name (First Name) Bailiang	2. Surname (Last Name) Li	3. Date 19-March-2020
4. Are you the corresponding author?	Yes V No	Corresponding Author's Name Ruijiang Li
5. Manuscript Title Natural Killer Cell and Stroma Abundan Cancer	ce are Independent Progn	ostic Factors and Predict Chemotherapy Benefit in Gastric
6. Manuscript Identifying Number (if you kn	ow it)	
		-
Section 2. The Work Under Co	onsideration for Public	cation
	but not limited to grants, da	a third party (government, commercial, private foundation, etc.) for ta monitoring board, study design, manuscript preparation,
Section 3. Relevant financial	activities outside the s	ubmitted work.
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Are there any relevant conflicts of intere	est? Yes ✓ No	
Section 4. Intellectual Proper	ty Patents & Copyrig	phts
Do you have any patents, whether plant	ned, pending or issued, br	oadly relevant to the work? Yes V No



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Jiang \_\_\_\_\_\_



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1. Given Name (First Name) Yuming	2. Surname (Last Name) Jiang	3. Date 19-March-2020
4. Are you the corresponding author?	Yes 🗸 No	Corresponding Author's Name Ruijiang Li
5. Manuscript Title Natural Killer Cell and Stroma Abundan Cancer	nce are Independent Progn	ostic Factors and Predict Chemotherapy Benefit in Gastric
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Jiang 2



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Jiang 3



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Did you or your institution <b>at any time</b> recei any aspect of the submitted work (including statistical analysis, etc.)? Are there any relevant conflicts of intere	but not limited to grants, dat		
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Place a check in the appropriate boxes i of compensation) with entities as descri clicking the "Add +" box. You should rep Are there any relevant conflicts of intere	ibed in the instructions. Us port relationships that were	e one line for each entity; ac	dd as many lines as you need by
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Fisher 1



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## **TREND Statement Checklist**

Paper Item		Descriptor		
Section/ Topic	No		$\checkmark$	Pg#
Title and Abst	ract			
Title and	1	Information on how unit were allocated to interventions	V	
Abstract		Structured abstract recommended	V.	
		Information on target population or study sample	V	
Introduction				
Background	2	Scientific background and explanation of rationale	4	
_		Theories used in designing behavioral interventions	1	
Methods	•			
Participants	3	Eligibility criteria for participants, including criteria at different levels in		
		recruitment/sampling plan (e.g., cities, clinics, subjects)	8	
		Method of recruitment (e.g., referral, self-selection), including the	17	
		sampling method if a systematic sampling plan was implemented		
		Recruitment setting		
		Settings and locations where the data were collected		
Interventions	4	Details of the interventions intended for each study condition and how		
		and when they were actually administered, specifically including:		
		Content: what was given?		
		Delivery method: how was the content given?	<u> </u>	
		O Unit of delivery: how were the subjects grouped during delivery?	<b>}</b>	
		Deliverer: who delivered the intervention?	<b></b>	
		Setting: where was the intervention delivered?	<b></b>	
		<ul> <li>Exposure quantity and duration: how many sessions or episodes or events were intended to be delivered? How long were they intended to last?</li> </ul>		
		<ul> <li>Time span: how long was it intended to take to deliver the intervention to each unit?</li> </ul>	}	
		<ul> <li>Activities to increase compliance or adherence (e.g., incentives)</li> </ul>		
Objectives	5	Specific objectives and hypotheses		
Outcomes	6	Clearly defined primary and secondary outcome measures		
		Methods used to collect data and any methods used to enhance the quality of measurements		
		<ul> <li>Information on validated instruments such as psychometric and biometric properties</li> </ul>	}	
Sample Size	7	How sample size was determined and, when applicable, explanation of any interim analyses and stopping rules		
Assignment	8	Unit of assignment (the unit being assigned to study condition, e.g.,		
Method		individual, group, community)		
		Method used to assign units to study conditions, including details of any		
		restriction (e.g., blocking, stratification, minimization)		
		Inclusion of aspects employed to help minimize potential bias induced due		
		to non-randomization (e.g., matching)	١	

## **TREND Statement Checklist**

Blinding	9	Whether or not participants, those administering the interventions, and	
(masking)	,	those assessing the outcomes were blinded to study condition assignment; if so, statement regarding how the blinding was accomplished and how it was assessed.	
Unit of Analysis	10	<ul> <li>Description of the smallest unit that is being analyzed to assess intervention effects (e.g., individual, group, or community)</li> <li>If the unit of analysis differs from the unit of assignment, the analytical method used to account for this (e.g., adjusting the standard error estimates by the design effect or using multilevel analysis)</li> </ul>	
Statistical Methods	11	<ul> <li>Statistical methods used to compare study groups for primary methods outcome(s), including complex methods of correlated data</li> <li>Statistical methods used for additional analyses, such as a subgroup analyses and adjusted analysis</li> <li>Methods for imputing missing data, if used</li> <li>Statistical software or programs used</li> </ul>	
Results			
Participant flow	12	<ul> <li>Flow of participants through each stage of the study: enrollment, assignment, allocation, and intervention exposure, follow-up, analysis (a diagram is strongly recommended)</li> <li>Enrollment: the numbers of participants screened for eligibility, found to be eligible or not eligible, declined to be enrolled, and enrolled in the study</li> </ul>	
		Assignment: the numbers of participants assigned to a study condition	
		<ul> <li>Allocation and intervention exposure: the number of participants assigned to each study condition and the number of participants who received each intervention</li> </ul>	
		<ul> <li>Follow-up: the number of participants who completed the follow-up or did not complete the follow-up (i.e., lost to follow-up), by study condition</li> </ul>	
		<ul> <li>Analysis: the number of participants included in or excluded from the main analysis, by study condition</li> </ul>	
		<ul> <li>Description of protocol deviations from study as planned, along with reasons</li> </ul>	
Recruitment	13	Dates defining the periods of recruitment and follow-up	
Baseline Data	14	Baseline demographic and clinical characteristics of participants in each study condition	
		Baseline characteristics for each study condition relevant to specific disease prevention research	
		Baseline comparisons of those lost to follow-up and those retained, overall and by study condition	
		<ul> <li>Comparison between study population at baseline and target population of interest</li> </ul>	
Baseline equivalence	15	Data on study group equivalence at baseline and statistical methods used to control for baseline differences	

### **TREND Statement Checklist**

Numbers analyzed	16	<ul> <li>Number of participants (denominator) included in each analysis for each study condition, particularly when the denominators change for different outcomes; statement of the results in absolute numbers when feasible</li> <li>Indication of whether the analysis strategy was "intention to treat" or, if not, description of how non-compliers were treated in the analyses</li> </ul>		
Outcomes and estimation	17	For each primary and secondary outcome, a summary of results for each estimation study condition, and the estimated effect size and a confidence interval to indicate the precision		
		<ul> <li>Inclusion of null and negative findings</li> <li>Inclusion of results from testing pre-specified causal pathways through</li> </ul>		
Ancillary analyses	18	<ul> <li>which the intervention was intended to operate, if any</li> <li>Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory</li> </ul>		
Adverse events	19	Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals)		
DISCUSSION				
Interpretation	20	Interpretation of the results, taking into account study hypotheses, sources of potential bias, imprecision of measures, multiplicative analyses, and other limitations or weaknesses of the study	\	
		Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations		
		Discussion of the success of and barriers to implementing the intervention, fidelity of implementation		
Generalizability	21	<ul> <li>Discussion of research, programmatic, or policy implications</li> <li>Generalizability (external validity) of the trial findings, taking into account the study population, the characteristics of the intervention, length of follow-up, incentives, compliance rates, specific sites/settings involved in the study, and other contextual issues</li> </ul>		
Overall Evidence	22	General interpretation of the results in the context of current evidence and current theory		

*From:* Des Jarlais, D. C., Lyles, C., Crepaz, N., & the Trend Group (2004). Improving the reporting quality of nonrandomized evaluations of behavioral and public health interventions: The TREND statement. *American Journal of Public Health*, 94, 361-366. For more information, visit: <a href="http://www.cdc.gov/trendstatement/">http://www.cdc.gov/trendstatement/</a>