Instructions

The purpose of this form is to provide readers of your manuscript with information about your other interests that could influence how they receive and understand your work. The form is designed to be completed electronically and stored electronically.

Identifying information.

The work under consideration for publication.

2.

This section asks for information about the work that you have submitted for publication. The time frame for this reporting is that of the work itself, from the initial conception and planning to the present. The requested information is about resources that you received, either directly or indirectly (via your institution), to enable you to complete the work. Checking "No" means that you did the work without receiving any financial support from any third party -- that is, the work was supported by funds from the same institution that pays your salary and that institution did not receive third-party funds with which to pay you. If you or your institution received funds from a third party to support the work, such as a government granting agency, charitable foundation or commercial sponsor, check "Yes".

3.

Relevant financial activities outside the submitted work.

This section asks about your financial relationships with entities in the bio-medical arena that could be perceived to influence, or that give the appearance of potentially influencing, what you wrote in the submitted work.

Report all sources of revenue paid (or promised to be paid) directly to you or your institution on your behalf over the 36 months prior to submission of the work. This should include all monies from sources with relevance to the submitted work, not just monies from the entity that sponsored the research.

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Intellectual Property.

This section asks about patents and copyrights, whether pending, issued, licensed and/or receiving royalties.

Relationships not covered above.

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Pending: The patent has been filed but not issued **Issued:** The patent has been issued by the agency

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Royalties: Funds are coming in to you or your institution due to your patent

Fischer

Section 1.	ldentifying Inforn	nation				
1. Given Name (Fi Nicholas	rst Name)	2. Surname (Last Name) Fischer		3. Date 04-June-2018		
4. Are you the corresponding author? Yes Volume No						
5. Manuscript Title Survival in males		ric adenocarcinoma correlat	es with mutant p53 resi	idual transcriptional activity		
6. Manuscript Ider 121364-INS-CME	ntifying Number (if you kı ED-RV-3	now it)				
	I					
Section 2.	The Work Under C	onsideration for Publica	tion			
Did you or your ins	titution at any time					
Are there any rel	evant conflicts of inter	est? ☐ Yes 🗸 No				
	out the appropriate info be removed by pressin		more than one entity p	press the "ADD" button to add a row.		
Name of Institut		Grant? Personal Non-	Financial Other? C	omments		
Section 3.	Relevant financial	activities outside the su	bmitted work.			
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Name of Entity		Grant	Financial Other? C	omments		



Section 4. Intellect	ual Property Patents & Copyrights				
If yes, please fill out the app	nether planned, pending or issued, broadly relevant to the work? Yes No ropriate information below. If you have more than one entity press the "ADD" button to add a lby pressing the "X" button.	row.			
Patent?	Pending? Issued? Licensed? Royalties? Licensee? Comments				
Section F					
Section 5. Relation	ships not covered above				
Are there other relationships or activities that readers could perceive to have influenced, or that give the appearance of potentially influencing, what you wrote in the submitted work?					
Yes, the following relationships/conditions/circumstances are present (explain below):					
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At the time of manuscript acceptance, journals will ask authors to confirm and, if necessary, update their disclosure statements. On occasion, journals may ask authors to disclose further information about reported relationships.					
Section 6. Disclosure Statement					
	res, this form will automatically generate a disclosure statement, which will appear in the box				
Mr. Fischer has nothing to	isclose.				

Evaluation and Feedback

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Prodeus

Section 1.	Identifying Inform	ation				
1. Given Name (Fi Aaron	rst Name)	2. Surname (Last Na Prodeus	me)		3. Date 04-June-2018	
4. Are you the corresponding author? Yes V No						
5. Manuscript Title Survival in males with glioma and gastric adenocarcinoma correlates with mutant p53 residual transcriptional activity						
6. Manuscript Identifying Number (if you know it) 121364-INS-CMED-RV-3						
Section 2.						
	The Work Under Co	onsideration for P	ublication			
Did you or your in:	stitution at any time					
Are there any rel	evant conflicts of intere	est? Yes ✓	No			
		ormation below. If yo		one entit	ty press the "ADD" button to add a	row.
	Excess rows can be removed by pressing the "X" button. Name of Institution (Company) Personal Non-Financial Company Company					
Name of Institut	ion/Company	Grant? Personal Fees?	Support?	Other	Comments	
Section 2						
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If yes, please fill out the appropriate information below.						
Name of Entity		Grant? Personal Fees?	Non-Financial Support?	Other?	Comments	



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If yes, please fill out the ap	whether planned, pending or issued, broadly relevant to the work? Yes No propriate information below. If you have more than one entity press the "ADD" button to add a row. ed by pressing the "X" button.			
Patent ?	Pending? Issued? Licensed? Royalties? Licensee? Comments			
Continue E				
Section 5. Relatio	onships not covered above			
	ps or activities that readers could perceive to have influenced, or that give the appearance of nat you wrote in the submitted work?			
Yes, the following relat	tionships/conditions/circumstances are present (explain below):			
✓ No other relationships/conditions/circumstances that present a potential conflict of interest				
•	acceptance, journals will ask authors to confirm and, if necessary, update their disclosure statements. ask authors to disclose further information about reported relationships.			
Section 6. Disclosi	ure Statement			
	sures, this form will automatically generate a disclosure statement, which will appear in the box			
Dr. Prodeus has nothing to	o disclose.			

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Gariépy

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Patent?	Pending? Issu	ued Royalties	? Licensee?	Comments	
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Dr. Gariépy has nothing to	disclose.				

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

	Item No	Recommendation
✓ Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the
		abstract
		(b) 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		
✓ 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	(a) Give the eligibility criteria, and the sources and methods of selection of
		participants. Describe methods of follow-up
		(b) For matched studies, give matching criteria and number of exposed and
		unexposed
✓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	(a) Describe all statistical methods, including those used to control for
		confounding
		(b) Describe any methods used to examine subgroups and interactions
		(c) Explain how missing data were addressed
		(d) If applicable, explain how loss to follow-up was addressed
		(\underline{e}) Describe any sensitivity analyses
Results		
✓ Participants	13*	(a) 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
		(b) Give reasons for non-participation at each stage
		(c) Consider use of a flow diagram
✓ Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical, social) and
		information on exposures and potential confounders
		(b) Indicate number of participants with missing data for each variable of interest
		(c) 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	(a) 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

		(c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period
✓ Other analyses	17	Report other analyses done—eg analyses of subgroups and interactions, and sensitivity analyses
Discussion		
✓Key results	18	Summarise key results with reference to study objectives
✓ Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or
		imprecision. Discuss both direction and magnitude of any potential bias
✓Interpretation	20	Give a cautious overall interpretation of results considering objectives, limitations,
		multiplicity of analyses, results from similar studies, and other relevant evidence
✓ Generalisability	21	Discuss the generalisability (external validity) of the study results
Other information		
√ Funding	22	Give 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.