

#### **Instructions**

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Hou 1



Section 1.	Identifying Inform	nation		
Given Name (Fire Wanting	st Name)	2. Surname (Last Name) Hou	3. Date 16-June-2019	
4. Are you the corresponding author?		Yes ✓ No	Corresponding Author's Name Feng Bi	
5. Manuscript Title The MEK inhibitor rebound	rs enhance the efficacy	y of sorafenib against hepa	atocellular carcinoma cells through reducing p-ERK	
6. Manuscript Iden	tifying Number (if you kn	now it)		
			_	
Section 2.	The Work Under Co	onsideration for Public	cation	
any aspect of the su statistical analysis, e	ıbmitted work (including	but not limited to grants, da	a third party (government, commercial, private foundation, etc.) for ata monitoring board, study design, manuscript preparation,	
Section 3.	Relevant financial	activities outside the s	submitted work.	
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Section 4.	Intellectual Document	to Detente 9 Commit		
		ty Patents & Copyric		
Do you have any patents, whether planned, pending or issued, broadly relevant to the work? Yes Vo				

Hou 2



Section 5.		
Section 5.	Relationships not covered above	
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Dr. Hou has noth	ning to disclose.	

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Xia 1



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Zhou 1



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4. Are you the cor	responding author?	☐ Yes ✓ No	Corresponding Author's Name Feng Bi	
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Do you have any				ſ No

Zhou 2



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1 Gong



Section 1.	Identifying Inform	nation		
1. Given Name (Fir Qiyong	st Name)	2. Surname (Last Name) Gong		Date 6-June-2019
4. Are you the corresponding author?		Yes ✓ No	Corresponding Author's Name Feng Bi	
5. Manuscript Title The MEK inhibito rebound		y of sorafenib against hepa	atocellular carcinoma cells thro	ough reducing p-ERK
6. Manuscript Iden	itifying Number (if you kr	now it)		
			_	
Section 2.	The Work Under Co	onsideration for Public	cation	
any aspect of the su statistical analysis,	ubmitted work (including	but not limited to grants, da	a third party (government, comm ta monitoring board, study desig	nercial, private foundation, etc.) for in, manuscript preparation,
Section 3.	Relevant financial	activities outside the s	submitted work.	
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Section 4.	Intellectual Proper	ty Patents & Copyric	Jhts	
Do you have any	patents, whether plan	ned, pending or issued, br	oadly relevant to the work? [	Yes 🗸 No

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Section 5.		
Section 5.	Relationships not covered above	
	elationships or activities that readers could perceive to have influenced, or that give the appearance of encing, what you wrote in the submitted work?	
Yes, the follo	wing relationships/conditions/circumstances are present (explain below):	
✓ No other rela	tionships/conditions/circumstances that present a potential conflict of interest	
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	
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Dr. Gong has no	thing to disclose.	

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Nie 1



Section 1. Identifying Infor	mation	
1. Given Name (First Name) Yongzhan	2. Surname (Last Name) Nie	3. Date 16-June-2019
4. Are you the corresponding author?	Yes ✓ No	Corresponding Author's Name Feng Bi
<ol><li>Manuscript Title The MEK inhibitors enhance the effica rebound</li></ol>	cy of sorafenib against hep	atocellular carcinoma cells through reducing p-ERK
6. Manuscript Identifying Number (if you	know it)	
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Section 4. Intellectual Prope	erty Patents & Copyri	ghts
Do you have any patents, whether pla	nned, pending or issued, b	roadly relevant to the work? Yes V No

Nie 2



Section 5.	
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paten<sup>.</sup>

Tang 1



Section 1.	Identifying Inform	nation	
1. Given Name (Fi Qiulin	rst Name)	2. Surname (Last Name) Tang	3. Date 16-June-2019
4. Are you the cor	responding author?	☐ Yes ✓ No	Corresponding Author's Name Feng Bi
5. Manuscript Title The MEK inhibito rebound		y of sorafenib against hep	atocellular carcinoma cells through reducing p-ERK
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Tang 2



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Bi 1



Section 1.	Identifying Inform	nation	
1. Given Name (Figure 1)  Feng  Are you the core		2. Surname (Last Name) Bi  ✓ Yes No	3. Date 16-June-2019
5. Manuscript Title The MEK inhibito rebound	2	y of sorafenib against hepatocellular carcinoma cells	s through reducing p-ERK
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Bi 2



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