

## PEER-REVIEW REPORT

**Name of journal:** World Journal of Clinical Oncology

**Manuscript NO:** 54280

**Title:** Mutational analysis of Ras hotspots in urothelial carcinoma of bladder patients

**Reviewer's code:** 00722239

**Position:** Editorial Board

**Academic degree:** MD, PhD

**Professional title:** Associate Professor

**Reviewer's Country/Territory:** Japan

**Author's Country/Territory:** India

**Manuscript submission date:** 2020-01-20

**Reviewer chosen by:** Le Zhang

**Reviewer accepted review:** 2020-03-30 11:04

**Reviewer performed review:** 2020-04-05 05:29

**Review time:** 5 Days and 18 Hours

<b>Scientific quality</b>	<input type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Very good <input checked="" type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
<b>Language quality</b>	<input type="checkbox"/> Grade A: Priority publishing <input type="checkbox"/> Grade B: Minor language polishing <input checked="" type="checkbox"/> Grade C: A great deal of language polishing <input type="checkbox"/> Grade D: Rejection
<b>Conclusion</b>	<input type="checkbox"/> Accept (High priority) <input type="checkbox"/> Accept (General priority) <input type="checkbox"/> Minor revision <input checked="" type="checkbox"/> Major revision <input type="checkbox"/> Rejection
<b>Re-review</b>	<input type="checkbox"/> Yes <input type="checkbox"/> No
<b>Peer-reviewer statements</b>	Peer-Review: <input checked="" type="checkbox"/> Anonymous <input type="checkbox"/> Onymous Conflicts-of-Interest: <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No

## SPECIFIC COMMENTS TO AUTHORS



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The authors have analysed the mutational status at the hotspot regions of H-Ras, K-Ras and N-Ras genes by PCR-RFLP and DNA sequencing method using tumor tissues of 87 cases of urothelial carcinoma of bladder (UCB). Although their focus and aim of the study are interesting, the quality of submitted manuscript is insufficient for publication. This manuscript requires extensive editing including English usage. Introduction section is too long and redundant. It should be shortened and clarify the aim of study. The section of materials and methods is also too long. As their methods of investigation, both of PCR-RFLP and DNA sequencing, are well known, details of its methods could be omitted. Their results include clinicopathological summary and point mutation detection in Ras, K-Ras and N-Ras genes but results of analysis for correlation of clinicopathological factors and point mutation were not performed.