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Erratum

Structural, topological, and functional characterization of transmembrane proteins TMEM213, 207, 116, 72 and 30B provides a potential link to ccRCC etiology: Am J Cancer Res. 2023; 13(5): 1863-1883

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In this article, we would like to correct the following three errors. The first error was in the Abstract. We would like to change "We confirmed the membrane-bound status of all selected TMEMs, assigned TMEM213, and 207 to early endosomes, TMEM72 to early endosomes and plasma membrane, TMEM116 and 30B to the endoplasmic reticulum" into "We confirmed the membrane-bound status of all selected TMEMs, assigned TMEM213, TMEM-116 and 30B to the endoplasmic reticulum, TMEM207 to early endosomes, TMEM72 to early endosomes and plasma membrane". The other errors were in the Discussion. We would like to change "TMEM213_ORF1, predicted to be localized in ER, was shown here to be confined to early endosomes, although the second isoform, lacking one AAG codon, was observed in both early endosomes and cytoplasm. TMEM207, 116 and 30B were localized in ER" into "TMEM213_ORF1, predicted to be localized in ER, was shown here to be confined to ER, although the second isoform, lacking one AAG codon, was observed in both ER and cytoplasm. Similarly, TMEM116 and 30B were localized in ER. TMEM207 and TMEM72 were found in early endosomes" and "In summary, TMEM213 protein localizes in early endosomes, contains a signal peptide on N-terminus, its C-terminus is directed towards cytoplasm, and it may contribute to processes like glycosylation and regulation of vasculature in ccRCC tumors" into "In summary, TMEM213 protein localizes in ER, contains a signal peptide on N-terminus, its N-terminus is directed towards cytoplasm, and it may contribute to processes like glycosylation and regulation of vasculature in ccRCC tumors". So, we would like to publish this Erratum to reflect these changes. The authors sincerely apologize for inconvenience caused by unintentional negligence.

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