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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 53370

Title: Suicide attempt using potassium tablets for congenital chloride diarrhea: A case report

Reviewer's code: 02800644

Position: Editorial Board

Academic degree: MD, PhD

Professional title: Research Scientist

Reviewer's Country/Territory: United States

Author's Country/Territory: Japan

Manuscript submission date: 2019-12-17

Reviewer chosen by: Jin-Lei Wang

Reviewer accepted review: 2020-02-03 15:40

Reviewer performed review: 2020-02-10 15:59

Review time: 7 Days

SCIENTIFIC QUALITY	LANGUAGE QUALITY	CONCLUSION	PEER-REVIEWER STATEMENTS
<input type="checkbox"/> Grade A: Excellent	<input checked="" type="checkbox"/> Grade A: Priority publishing	<input type="checkbox"/> Accept	Peer-Review:
<input checked="" type="checkbox"/> Grade B: Very good	<input type="checkbox"/> Grade B: Minor language	(High priority)	<input checked="" type="checkbox"/> Anonymous
<input type="checkbox"/> Grade C: Good	polishing	<input type="checkbox"/> Accept	<input type="checkbox"/> Onymous
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of	(General priority)	Peer-reviewer's expertise on the
<input type="checkbox"/> Grade E: Do not	language polishing	<input checked="" type="checkbox"/> Minor revision	topic of the manuscript:
publish	<input type="checkbox"/> Grade D: Rejection	<input type="checkbox"/> Major revision	<input type="checkbox"/> Advanced
		<input type="checkbox"/> Rejection	<input checked="" type="checkbox"/> General
			<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input checked="" type="checkbox"/> No



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SPECIFIC COMMENTS TO AUTHORS

The authors report a rare case of CCD patient who attempted suicide with high dosage slow release KCl tablets. The clinical course of the patient is well documented, and the authors also provided a brief review for CCD in the Discussion. I believe the report is of interest to the readers of the journal. I have two short questions as follows. 1. Does the patient have mutation(s) in her SLA26A3, the responsible gene for CCD? 2. The patient showed normal kidney function. Any abnormalities in sweat secretion or lung function? Or any similarities with cystic fibrosis?

INITIAL REVIEW OF THE MANUSCRIPT

Google Search:

- ☐ The same title
- ☐ Duplicate publication
- ☐ Plagiarism
- ☐ [Y] No

BPG Search:

- ☐ The same title
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- ☐ [Y] No