

Adverse Gastrointestinal Effects with Kayexalate or Kalimate: A Comprehensive Review

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Background: Patients with hyperkalemia are commonly treated with Kayexalate or Kalimate. Both drugs are associated with some fatal gastrointestinal (GI) adverse events (AEs).

Aim: To assess the clinical characteristics and outcomes of GI AEs induced by Kayexalate or Kalimate from published case reports.

Methods: We conducted a systematic review of case reports of Kayexalate or Kalimate-induced GI AEs, from PubMed, Medline, Cochrane Library, Clinical Key, and Google Scholar databases (1948 to March 31, 2020). We analyzed the clinical characteristics, GI AEs, and risk factors of enrolled patients.

Results: We identified 41 published articles describing 135 cases of GI AEs induced by Kayexalate (103 cases) or Kalimate (32 cases). The mean age of all patients was 55.5 years. Most patients were male (54.8%). As high as 55.6% preparations were administered with sorbitol whereas 44.4% preparations had no sorbitol. The average time causing GI AEs was 19.8 days. Colon was the most commonly affected site (76.3%). Drug crystals were histopathologically proven in 95.5% of the patients. Meanwhile, mortality was reported in 20.7%.

Conclusion: Kayexalate or Kalimate, without or with sorbitol combination, may be related to fatal GI damage. Uremia, hypertension, and transplantation are predisposing factors. Clinicians should be careful in prescribing Kayexalate or Kalimate to patients.

Keywords: gastrointestinal injury, hyperkalemia, Kayexalate, Kalimate

Introduction

Patients suffering from chronic kidney disease (CKD) may exhibit hyperkalemia, a commonly seen adverse event of electrolyte imbalance that can lead to fatal cardiac arrhythmias.¹ Hyperkalemia is commonly treated by a cation-exchange resin called sodium polystyrene sulfonate (Kayexalate).²⁻⁴ This drug acts as an exchange resin between sodium and potassium ions in the large intestine for accelerating potassium removal in the stool.⁵ However, its use has been reportedly related to colonic necrosis besides other fatal adverse events of gastrointestinal (GI) tract.⁶ An analog of Kayexalate, Kalimate (calcium polystyrene sulfonate), is also employed for treating hyperkalemia in clinical practice. However, there are not many incidences of Kalimate-induced GI tract injuries.⁷ Thus, our present study aimed to assess the clinical characteristics, risk factors, and outcomes of GI adverse events induced by Kayexalate or Kalimate from the published case reports in the literature.

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Materials and Methods

Study Population

We considered articles to be eligible that presented a case report or case series of GI adverse events associated with the use of Kayexalate or Kalimate. Any unfavorable or damaging effect on the GI tract in relation to the use of Kayexalate or Kalimate is considered as a GI adverse event. The age, sex, symptoms, GI injury location, and risk factors of enrolled patients were analyzed. If there is no information available for any case series from the original authors, those cases were excluded from this study.

Literature Sources

We searched databases including PubMed and Medline (n = 136), Cochrane Library (n = 35), Clinical Key (n = 203), and Google Scholar (n = 2134) (Figure 1. Flow diagram of included studies) (1948 to March 31, 2020) for the case reports, considered eligible, of GI adverse events associated with Kayexalate and Kalimate use. Language-related restrictions were not enforced in this study.

Statistical Analysis

The published reports contained 29 case reports, 11 case series, and 1 systematic review. The clinical characteristics of all enrolled patients were described using mean \pm standard deviation for continuous variables with abnormal distribution and number (percent) for categorical variables. Categorical variables were expressed as the number of cases and a percentage. These variables were compared using the *T*-test. A *P* value less than 0.05 indicates statistical significance.

Ethics Considerations

The present study was approved by the institutional review board of the Research Ethics Committee of China Medical University Hospital, in Taiwan (CMUH109-REC-01-049).

Results

A total of 2508 relevant articles were identified from our literature search. Following the application of the defined exclusion criteria, 41 articles describing 135 cases of GI adverse events induced by Kayexalate or Kalimate use

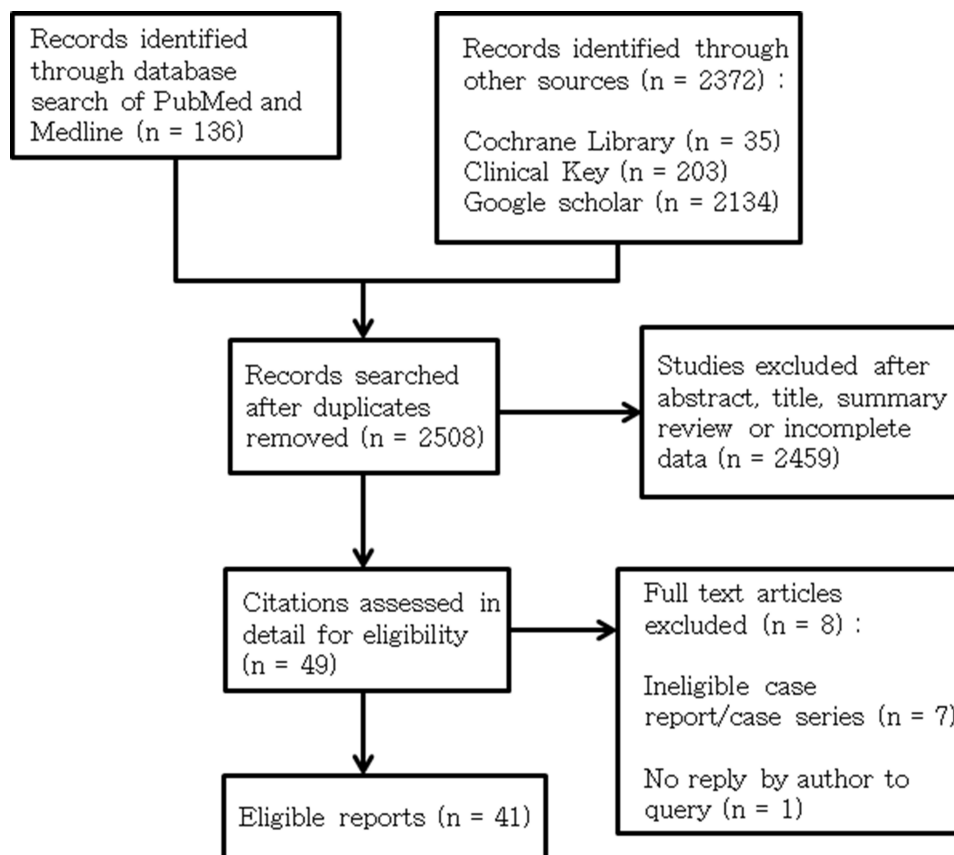


Figure 1 Flow diagram of included studies.

were enrolled in our present study (Kayexalate: 103 cases; Kalimate: 32 cases). The enrolled patients were found in 29 case reports, 11 case series, and 1 systematic review (Figure 1).⁵⁻⁵² Three cases out of the 135 reported cases in our present study were reported before 1990, 24 cases were reported during 1990 to 2000, and another 108 cases were reported after 2000.

Table 1 presents the clinical characteristics of all enrolled patients who experienced GI adverse events. The mean age of these patients was 55.5 ± 20.2 years (mean \pm standard deviation) with a range from 27 weeks of gestation to 91 years. In addition, 71.9% of them aged 55 years or more ($n = 97$), while 28.1% aged below 55 years ($n = 38$). The differences between these two age groups were found to be statistically significant ($P < 0.001$). Regarding sex distribution, 54.8% of them were male ($n = 74$), whereas 45.2% were women ($n = 61$). The differences between the male and female groups were found to be statistically significant ($P < 0.001$).

Moreover, 55.6% of all enrolled patients used Kayexalate or Kalimate with sorbitol concomitantly, whereas 44.4% used either of the drugs without sorbitol. A statistically significant difference was found between the sorbitol group and without-sorbitol group ($P < 0.001$). In the analysis of predisposing factors for hyperkalemia, 87.7% had a history of CKD ($n = 114$) and 53.1% had end-stage renal disease (ESRD) requiring dialysis ($n = 69$), 30.0% had hypertension ($n = 39$), and 18.5% had a prior solid transplant ($n = 24$). The differences were statistically significant ($P < 0.001$). The mean usage time of Kayexalate or Kalimate causing GI adverse events was 19.84 days (0.5 days to 1 year). Only 27 patients (33.3%) received Kayexalate for more than 1 month. Kayexalate or Kalimate was administered orally in 67.8% of patients, rectally in 88.5%, and via the nasogastric route in 2.5%. The difference in administration routes was statistically significant ($P < 0.001$).

The results of GI adverse events associated with Kayexalate or Kalimate use are summarized in Table 2 in our present study. The presenting GI symptoms of all enrolled patients were abdominal pain ($n = 59$, 68.6%), nausea or vomiting ($n = 6$, 7.0%), diarrhea ($n = 18$, 20.9%), and black stool or bloody stool ($n = 48$, 55.8%). A statistically significant difference was found between different symptoms ($P < 0.001$). The colon was the most commonly involved site of the GI tract ($n = 103$, 76.3%), followed by the small intestine and duodenum (25.2%), rectum (14.8%), stomach (7.4%) and esophagus (3.0%).

The differences between these various sites were significant ($P < 0.001$).

Histopathological findings associated with Kayexalate or Kalimate use were intestinal-wall ulcerations (83.7%), necrosis (60.0%), and perforation (31.9%). The differences between these histological findings were statistically significant ($P < 0.001$). In our present study, crystals of Kayexalate or Kalimate were demonstrated histopathologically in 95.5% of all patients. Meanwhile, mortality caused by GI injuries was reported in 20.7% of all patients in our present study.

Discussion

Kayexalate has been widely used for acute and chronic hyperkalemia management. It was subsequently approved by the FDA in the United States in 1958.⁷ Kalimate is a calcium-exchange resin with similar pharmacological action as that of Kayexalate, and it is also used for treating acute or chronic hyperkalemia. Kayexalate or Kalimate, which is a cation-exchange resin, can be introduced into the lower GI tract employing an enema preparation. This resin can also be introduced either orally or by nasogastric tube into the upper GI tract. When the resin is administered orally or by nasogastric tube into the stomach region, at first sodium or calcium cations are released from the resin in exchange with hydrogen ions in the stomach's acidic milieu. As the resin traverses the intestines, there is an exchange of hydrogen with potassium, followed by its excretion through the feces along with the remainder of the altered resin. Thus, the serum levels of potassium decrease over a period of hours to days.^{7,8}

Conventionally, Kayexalate was given as a water suspension. Because of the concerns of fecal impaction and constipation, it became a common practice to administer Kayexalate together with hypertonic sorbitol.² Sorbitol is a cathartic agent that has been reported to be associated with the development of intestinal injury rather than Kayexalate itself. The pathophysiological processes may be potentiated by the concomitant use of sorbitol with Kayexalate or Kalimate. Sorbitol, by elevating prostaglandin levels, causes a direct damage to the intestinal mucosa and leads to vasospasm and exacerbation of inflammation, and ultimately vascular injury.⁵ Histopathological lesions of these drugs-induced injuries vary from patchy inflammation to frank necrosis.⁸ This toxic effect of sorbitol on the GI tract has been proposed to potentially cause local ischemia and hypotension, in association with uremia and elevated renin levels in these patients, thereby adding to

Table 1 Summary of Clinicopathologic Findings of Patients with Kayexalate or Kalimate-Associated Gastrointestinal Injury in the Literature and the Present Case

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayexalate Crystals or Kalimate Crystals	Associated Conditions	Outcome		
Ziv Harel et al	58		58 ± 18	Kayexalate (41)	Sorbitol (41)	Oral route (40)	1.5	Abdominal pain/tenderness (21)	Esophagus (1)	Necrosis (30)	38	Chronic kidney disease (10)	Alive (22)		
						Per rectum (14)		Nausea/vomiting (2)	Stomach (2)					Ulceration (12)	
						Nasogastric (2)		GI bleed (9)	Small bowel (11)	Perforation (3)					Transplant (7)
								Diarrhea (3)	Cecum (4)					Coronary artery disease (8)	
									Colon (33)						CVD (1)
									Sigmoid/rectum/anus (4)					Hypertension (19)	
						Diabetes (8)									
								Chronic kidney disease (5)	Alive (11)						
						ESRD requiring dialysis (7)	Death (4)								
								Transplant (2)	Not reported (2)						
						Coronary artery disease (2)									
								CVD (4)							
						Hypertension (5)									
								Diabetes (2)							

Sung et al	I		Female (1)	63	Kayexalate	Sorbitol (1)	Oral route (1)	NA	Abdominal pain/tenderness (1) GI bleed (1)	Colon (1)	Necrosis (1) Ulceration (1)	I	ESRD requiring dialysis (1)	Alive (1)
Sumanth et al	I		Female (1)	59	Kayexalate	Sorbitol (1)	Oral route (1)	NA	GI bleed (1)	Stomach (1)	Ulceration (1)	I	ESRD requiring dialysis (1)	Alive (1)
Mazen et al	I		Male (1)	61	Kayexalate (30 g/d)	Sorbitol (1)	Oral route (1)	NA	Abdominal pain/tenderness (1) GI bleed (1)	Stomach (1) Cecum (1) Colon (1)	Ulceration (1)	I	CRF (GFR <30) (1)	Alive (1)
Gabrielle et al	I		Male (1)	58	Kayexalate	Sorbitol (1)	Per rectum (1)	14	Abdominal pain/tenderness (1) GI bleed (1)	Colon (1)	Ulceration (1) Perforation (1)	I	ESRD requiring dialysis (1) Biliary pancreatitis (1) Respiratory failure (1)	Alive (1)
Massimo et al	I		Male (1)	59	Kayexalate	Sorbitol (1)	Oral route (1)	NA	GI bleed (1)	Esophagus (1)	Ulceration (1)	I	CRF (GFR <30) (1)	Alive (1)
Ana Alves et al	I		Female (1)	83	Kayexalate	No sorbitol (1)	Oral route (1)	2	Abdominal pain/tenderness (1) GI bleed (1) Diarrhea (1)	Rectum (1)	Ulceration (1)	I	Major depressive disorder (1) Hypertension (1)	Alive (1)
L. Pirona et al	I		Male (1)	61	Kayexalate (15 g/d)	Sorbitol (1)	Oral route (1)	60	Abdominal pain/tenderness (1)	Colon (1)	Ulceration (1) Perforation (1)	I	CRF (GFR <20) (1) Hypertension (1)	Alive (1)
Nicolas et al	I		Male (1)	56	Kayexalate	No sorbitol (1)	Oral route (1)	14	Abdominal pain/tenderness (1) GI bleed (1)	Duodenum (1)	Hemorrhagic duodenitis (1) Ulceration (1)	I	ESRD and kidney transplant three weeks ago (1) Diabetes (1) Hypertension (1) Dyslipidemia (1)	Alive (1)

(Continued)

Table 1 (Continued).

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayaxalate Crystals or Kalimate Crystals	Associated Conditions	Outcome
Sheba et al	6	Male (4)	63	Kayaxalate (6)	Sorbitol (6)	Oral route (6), Per rectum (6)	7	Abdominal pain/ tenderness (1)	Jejunum (1)	Ulceration (6), Perforation (1), Stricture (1)	6	Chronic kidney disease (5)	Alive (5)
		Female (2)						GI bleed (6)	Colon (4), Rectum (4)	Polypoid lesion (1) Nodular lesion(1)		Kidney transplant and ESRD requiring dialysis (1)	Death (1)
Chou et al	1	Male (1)	30	Kayaxalate	Sorbitol (1)	Oral route (1) Per rectum (1)	3	GI bleed (1)	Colon (1)	Ulceration (1)	1	Hypertension (1) ESRD requiring dialysis (1) Ureteral urothelial carcinoma status posttransurethral resection of bladder tumor and chemotherapy	Alive (1)
Kao et al	1	Female (1)	59	Kalimate (60 g/d)	No sorbitol (1)	Oral route (1)	3	Abdominal pain/ tenderness (1) GI bleed (1)	Small bowel (1) Colon (1)	Ulceration (1) Necrosis (1) Perforation (1)	1	Hypertension (1) Diabetes (1) ESRD requiring dialysis (1) CAD (1), PAOD (1), COPD (1)	Death (1)
Mee et al	1	Male (1)	34	Kalimate	No sorbitol (1)	Oral route (1) Per rectum (1)	3	GI bleed (1)	Sigmoid/ Rectum (1)	Necrosis (1) Ulceration (1)	1	Respiratory failure (1) Hypertension (1)	Alive (1)
María et al	1	Male (1)	73	Kalimate	No sorbitol (1)	Oral route (1)	NA	GI bleed (1)	Colon (1)	Ulceration (1)	1	Dyslipidemia (1) Coronary artery disease (1) Hyperuricemia (1)	Alive (1)

Su et al	I		Female (1)	66		Kalimate (5 g/d)	No sorbitol (1)	Oral route (1)	120	Abdominal pain/tenderness (1)	Colon (1)	Necrosis (1)	I	CRF (GFR <15) (1)	Alive (1)
										GI bleed (1)	Sigmoid/rectum/anus (1)	Ulceration (1)		Hypertension (1)	
												Stenosis (1)			
Wu et al	I		Female (1)	77		Kalimate (30 g/d)	No sorbitol (1)	Oral route (1)	14	Abdominal pain/tenderness (1)	Esophagus (1)	Necrosis (1)	I	CRF (GFR <30) (1)	Alive (1)
										GI bleed (1)	Stomach (1)	Ulceration (1)		Hypertension (1)	
											Duodenum (1)				
Ana et al	I		Female (1)	83		Kayexalate	No sorbitol (1)	Oral route (1)	2	Abdominal pain (1)	Rectum (1)	Ulceration (1)	I	CRF (GFR <30) (1)	Alive (1)
										Diarrhea (1)				Hypertension (1)	
														Major depressive disorder (1)	
Sagar Patel et al	I		Male (1)	45		Kayexalate	Sorbitol (1)	Oral route (1)	4	Abdominal pain (1)	Terminal ileum	Ulceration (1)	I	Acute kidney injury (1)	Alive (1)
										Diarrhea (1)	Rectum (1)			Thymoma (1)	
										GI bleed (1)	Colon (1)				
Assanee Tongyoo et al	I		Male (1)	52		Kalimate	No sorbitol (1)	Oral route (1)	5	Abdominal pain (1)	Small bowel (1)	Obstruction (1)	0	Necrotizing fasciitis (1)	Death (1)
										Bowel ileus	Stomach (1)			Above-knee amputation (1)	
											Colon (1)			CRF (GFR <30) (1)	
Gabrielle et al	I		Male (1)	58		Kayexalate	Sorbitol (1)	Per rectum (1)	14	GI bleed (1)	Colon (1)	Ulceration (1)	I	Severe biliary pancreatitis (1)	Alive (1)
												Necrosis (1)		Acute kidney injury requiring dialysis (1)	
												Perforation (1)		Respiratory failure (1)	

(Continued)

Table 1 (Continued).

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayexalate Crystals or Kalimate Crystals	Associated Conditions	Outcome
Helena Ribeiro et al	1	Male (1)	72	Kalimate	No sorbitol (1)	Oral route (1)	3	Abdominal pain (1)	Colon (1)	Ulceration (1)	1	ESRD requiring hemodialysis (1) Ischemic heart disease (1) Arterial hypertension (1)	Alive (1)
										Necrosis (1)			
Béringère	2	Male (1)	34	Kayexalate (2)	No sorbitol (1)	Per rectum (2)	2	GI bleed (1)	Colon (2)	2	CRF (GFR <30) (2)	Alive (2)	
		Female (1)	64				3		Rectum (2)				
Alexandre	1	Male (1)	54	Kayexalate	Sorbitol (1)	Oral route (1)	2	GI bleed (1)	Stomach (1)	Ulceration (1)	1	Cardiovascular failure (1)	Alive (1)
Daphne	1	Female (1)	4	Kayexalate	Sorbitol (1)	Per rectum (1)	1	GI bleed (1)	Rectum (1)	Ulcer (1)	1	ESRD requiring hemodialysis (1) Shiga toxin-producing Escherichia coli hemolytic uremic syndrome (1)	Alive (1)
									Small bowel (1) (ileorectal anastomosis)	Polyloid lesion (1)			
Kathleen	1	Male (1)	60	Kayexalate	Sorbitol (1)	Oral route (1)	5	Abdominal pain/tenderness (1) GI bleed (1)	Colon (1)	Ulceration (1)	1	HTN (1) DM (1) CHF (1) RCC (1) CKD (1)	Alive (1)
										Necrosis (1)			
										Perforation (1)			

Chih-Chin	I	Female (1)	59	Kalimate	No sorbitol (1)	Oral route (1)	3	Abdominal pain/ tenderness (1)	Small bowel (1)	Ulceration (1)	I	ESRD (1)	Death (1)	
								GI bleed (1)	Colon (1)	Necrosis (1)				HTN (1)
										Perforation (1)				DM (1)
Philippe	I	Female (1)	73	Kalimate	No sorbitol (1)	Oral route (1)	4	Abdominal pain/ tenderness (1)	Small bowel (1)	Ulceration (1)	I	Acute colonic pseudoobstruction (1)	Death (1)	
									Colon (1)	Necrosis (1)				AKI (1)
										Perforation (1)				
EMMANUEL	I	Male (1)	53	Kayexalate	Sorbitol (1)	Oral route (1)	7	No symptoms	Esophagus (1)	Ulceration (1)	I	Lung transplantation for idiopathic pulmonary fibrosis (1)	Alive (1)	
										Gastroparesis (1)				
S Rugolotto	I	Male (1)	0.6 (27 weeks of gestation)	Kayexalate	No sorbitol (1)	Per rectum (1)	I	GI bleed (1)	Small bowel (ileum) (1)	Ulceration (1)	I	Sepsis (1)	Alive (1)	
										Necrosis (1)				ESRD (1)
Kanokrat	I	Female (1)	0.8 (38 weeks gestation)	Kalimate	No sorbitol (1)	Oral route (1)	I	Abdominal distention (1)	Small bowel (1)	Ulceration (1)	I	Pulmonary atresia (1)	Alive (1)	
								GI bleed (1)	Stomach (1)	Necrosis (1)				Acute kidney injury (1)
										Perforation (1)				
Wai Tat	I	Female (1)	79	Kalimate	No sorbitol (1)	Oral route (1)	14	Abdominal pain/ tenderness (1)	Small bowel (1)	Ulceration (1)	I	CKD (1)	Alive (1)	
								GI bleed (1)	Colon (1)	Necrosis (1)				
										Perforation (1)				

(Continued)

Table 1 (Continued).

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayexalate Crystals or Kalmate Crystals	Associated Conditions	Outcome
Mary Bui	1	Female (1)	70	Kayexalate	No sorbitol (1)	Oral route (1)	365 (one year) (a dose of 15 grams twice daily after meals)	Abdominal pain (1)	Stomach (1)	Inflammation	1	Coronary artery disease (1)	Alive (1)
								Constipation with intermittent loose bowel movements and the stool guaiac tested positive.	Small bowel (1)	A mass		ESRD requiring hemodialysis (1)	
									Colon (1) (a pseudotumor on the upper lip of the ileocecal valve)			Secondary hyperparathyroidism s/p parathyroidectomy (1)	
Abdulaziz	1	Male (1)	64	Kayexalate	No sorbitol (1)	Oral route (1)	1	Abdominal pain (1)	Small bowel (1)	Ulceration (1)	1	Tongue cancer (1)	Alive (1)
								GI bleed (1)	Colon (1)	Necrosis (1)		Septic shock (1)	
												AKI on CKD (1)	
Takeshi	1	Female (1)	77	Kalmate	Sorbitol (1)	Oral route (1)	9	Abdominal pain (1)	Colon (1)	Sigmoidovesical fistula (1)	1	Diabetic ketoacidosis	Alive (1)
								GI bleed (1)		Stenosis (1)		CAD (1)	
										Ulcer (1)			
Kohei	1	Male (1)	79	Kalmate	Sorbitol (1)	Oral route (1)	240 (8 months)	Abdominal pain (1)	Small bowel (1)	Ulcer (1)	1	DM (1)	Alive (1)
								GI bleed (1)	Colon (1)			HTN (1)	
								Diarrhea (1)				CKD (1)	
Roberto	1	Female (1)	80	Kayexalate	Sorbitol (1)	Oral route (1)	1	Abdominal pain (1)	Colon (1)	Ulceration (1)	1	CKD (1)	Alive (1)
								Diarrhea (1)		Necrosis (1)		CHF (1)	
												COPD (1)	

Smiley	10	Male (6)	64.2 (41–82, y ^o)	Kayexalate (10)	NA	NA	NA	NA	NA	NA	Small bowel (3)	Ulceration (7)	10	CKD (9)	Alive (1)	
		Female (4)										Stomach (2)				Necrosis (1)
Hussein Nassereddine	1	Male (1)	52	Kayexalate (1)	No Sorbitol (0)	NA	NA	NA	NA	NA	Esophagus (0)	Necrosis (0)	Kayexalate crystal (1)	-	Alive (1)	
											Stomach (0)	Ulceration (1)				Chronic kidney disease (1)
											Small bowel (0)	Perforation (1)				ESRD requiring dialysis (0)
											Cecum (0)					Transplant (0)
											Colon (1)					Coronary artery disease (0)
											Rectum/anus (1)					CVD (0)
																Hypertension (0)
																Diabetes (0)

(Continued)

Table 1 (Continued).

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayexalate Crystals or Kalimate Crystals	Associated Conditions	Outcome
Robert Hunt Dunlap	1	Female (1)	55	Kayexalate (1)	No Sorbitol (0)	NA	3				Kayexalate crystal (1)	-	Alive (1)
								Abdominal pain/tenderness (0)	Esophagus (0)	Necrosis (1)		Chronic kidney disease (1)	
								Nausea/vomiting (0)	Stomach (0)	Ulceration (1)		ESRD requiring dialysis (1)	
								GI bleed (1)	Small bowel (1)	Perforation (1)		Transplant (0)	
								Diarrhea (1)	Cecum (1)			Coronary artery disease (0)	
									Colon (1)			CVD (0)	
									Rectum/anus (0)			Hypertension (1)	
												Diabetes (0)	

Keigo Murakami	23	69.5		Kayaxalate (4)	Sorbitol (4)		Oral route (3), Per rectum (1)		24.8	Abdominal pain/tenderness (22)		Esophagus (0)		Necrosis (4)		Kayaxalate crystal (4)		Chronic kidney disease (0)		Alive (2)	
		Male (2)			Stomach (0)		Ulceration (4)			Nausea/vomiting (0)		Small bowel (0)		Perforation (4)		ESRD requiring dialysis (4)		Death (2)			
		Female (2)			Small bowel (0)		Cecum (0)			GI bleed (22)		Diarrhea (1)				Transplant (0)					
					Colon (3)		Rectum/anus (1)									Coronary artery disease (0)					
																CVD (0)					
																Hypertension (0)					
																Diabetes (1)					
																Chronic kidney disease (4)		Alive (15)			
																Ulceration (19)		ESRD requiring dialysis (15)		Death (4)	
																Perforation (19)		Transplant (0)			
																Small bowel (0)		Coronary artery disease (0)			
																Cecum (0)		CVD (0)			
																Colon (18)		Hypertension (0)			
																Rectum/anus (1)		Diabetes (7)			

(Continued)

Table 1 (Continued).

Authors	No. of Patients	Gender	Age, Mean (SD)	Type of Resin	Type of Suspension	Route of Administration	Time Interval (Day)	Presenting Symptoms	Location	Histopathologic Findings	Kayexalate Crystals or Kalimate Crystals	Associated Conditions	Outcome
Hussein Nassereddine	1	Male (1)	89	Kayexalate (1)	No Sorbitol (0)		NA	NA			Kayexalate crystal (1)	-	Alive (1)
						Oral route (1)			Esophagus (0)	Necrosis (0)		Chronic kidney disease (1)	
						Per rectum (1)			Stomach (0)	Ulceration (1)		ESRD requiring dialysis (0)	
						Nasogastric (0)			Small bowel (0)	Perforation (1)		Transplant (0)	
									Cecum (0)			Coronary artery disease (0)	
									Colon (1)			CVD (0)	
									Rectum/anus (0)			Hypertension (0)	
												Diabetes (1)	

Table 2 Results of Gastrointestinal Adverse Events Associated with the Use of Kayexalate or Kalimate (n = 135)

Variables	Total (Valid)	P value
Presenting symptoms, n (%)		< 0.001
Abdominal pain	59 (68.6)	< 0.001
Nausea/vomiting	6 (7.0)	0.159
GI bleeding	48 (55.8)	< 0.001
Diarrhea	18 (20.9)	0.001
Location, n (%)		< 0.001
Esophagus	4 (3.0)	0.158
Stomach	10 (7.4)	0.044
Small bowel	34 (25.2)	< 0.001
Cecum	10 (7.4)	0.156
Colon	103 (76.3)	< 0.001
Rectum	20 (14.8)	< 0.001
Histopathologic findings, n (%)		< 0.001
Necrosis	81 (60.0)	< 0.001
Ulceration	113 (83.7)	< 0.001
Perforation	43 (31.9)	0.002
Kayexalate or Kalimate crystals, n (%)	129 (95.5)	< 0.001
Associated conditions		< 0.001
Chronic kidney disease	114 (87.7)	< 0.001
ESRD requiring dialysis	69 (53.1)	< 0.001
Organ transplant	24 (18.5)	< 0.001
Coronary artery disease	23 (17.7)	0.001
Hypertension	39 (30.0)	< 0.001
Diabetes	24 (18.5)	0.013
Outcome, n (%)		< 0.001
Alive	100 (74.0)	
Death	28 (20.7)	
Not reported	7 (5.3)	

Abbreviations: GI, gastrointestinal tract; ESRD, end-stage renal disease.

the severity of the injury. The susceptibility of uremic versus non-uremic rats to injury induced by Kayexalate-sorbitol enema indicates that uremia may in fact predispose to Kayexalate-induced mucosal injury.⁹ However, an increasing number of case reports have shown that severe adverse events, such as colonic necrosis, are linked with the Kayexalate treatment without or with only a small amount of sorbitol.⁷ Our present study included numerous cases using Kayexalate or Kalimate without sorbitol who experienced similar GI adverse events.^{10,11} Moreover, Singhania N et al reported a case of colonic intestinal necrosis after co-administration of Kayexalate and activated charcoal.¹²

Furthermore, Kayexalate or Kalimate crystals were often seen to aggregate inside the damaged parts of the GI tract specimens, examined histopathologically. The presence of Kayexalate crystals in injured GI tract segments may represent a “footprint” of its use, indicating that

Kayexalate itself may be pathogenic.⁸ Previous reports have postulated several risk factors for GI adverse events, including CKD, ESRD, solid organ transplantation, immunosuppression, and postoperative status; these factors lead to GI injury associated with Kayexalate or Kalimate treatment by different mechanisms.^{5,6,8,13–16,21,22} Renal transplant recipients are at increased risk for various adverse events, including infection, graft rejection, and medication adverse effects, in the early post-transplant period.⁸ Indeed, many of these risk factors were noted in most cases reviewed in the present study. Remarkably, 87.7% of all enrolled patients in this reviewing study had a history of ESRD, acute kidney injury or CKD. As shown in Table 1, the difference between these groups of comorbidities was statistically significant.

The increased renin levels, often seen in patients with renal diseases, place them at high risk to non-occlusive mesenteric ischemia via angiotensin II-mediated vasoconstriction.¹⁶ As noticed from our findings, this risk may be elevated during the postoperative period, possibly caused by ileus-induced colonic distension (resulting in lowered blood flow to colon), simultaneous hypotension and reduced gut motility as a result of opioid use, uremia, and constipation.^{5,17,18} Patients who recently had transplantation are at a particularly elevated risk of these complications due to the administration of immunosuppressive drugs that negatively influence the normal protective and reparative capacity of GI cells.^{18,19}

The increase of inflammatory cytokines and prostaglandins may further impair local hemodynamic mechanisms, resulting in vascular injury and subsequent mucosal injury in the GI tract of individuals at high risk, such as those suffering with CKD or vascular disease, and patients’ solid organ transplant recipients. Given our present results, the pathogenesis of GI injury related to Kayexalate or Kalimate crystals is probably more complex than what we know presently and may represent a histopathologic injury in response to different heterogeneous insults.

It is difficult to define the etiology for colonic necrosis in patients with several health issues with elevated susceptibility to mucosal injury. Combined histologic, laboratory, clinical and examinations are necessary to eliminate the commonly seen causes of mucosal necrosis and also to correctly diagnose the mucosal necrosis associated with Kayexalate- or Kalimate.^{15,18,20–22,29–53} However, the observation of Kayexalate or Kalimate angulated crystals with a specific mosaic pattern is critical for diagnosing

Kayexalate- or Kalimate-related mucosal necrosis. Histologically, cholestyramine crystals (used to decrease high blood cholesterol levels or to treat itching due to biliary tract blockage) should be distinguished from Kayexalate crystals. The cholestyramine crystals are more basophilic, opaque without a mosaic pattern, and rhomboid in shape compared with the Kayexalate crystals.¹⁸ The microscopic features of Kalimate crystals and of Kayexalate crystals are the same.^{20,22}

The colon is the most common site of Kayexalate- or Kalimate-induced GI injuries. Injury caused by Kayexalate may appear in more proximal sections of the GI tract.^{15,18} As shown in Table 2, various locations of GI adverse events were statistically different, contradicting the severe adverse events in the lower GI tract from numerous case reports. Intestinal necrosis and perforation are potential complications of Kayexalate and may present in a delayed stage up to 2 months.^{28,51–53} However, Kayexalate- or Kalimate-related toxicities in the upper GI tract have been rarely described in the literature.^{20,22}

Most of these cases do have simultaneous damage to colon. This change may possibly be due to the trend for the use of oral, instead of rectal Kayexalate or Kalimate.^{5,22–28,51–53} However, regardless of the location of Kayexalate or Kalimate-associated injury, mortality remains high. The elevated mortality may be a result of aggravated damage along with the large comorbid disease burden of affected patients.⁵ In the current study, we also found safety complications associated with Kayexalate or Kalimate usage, either with or without concomitant sorbitol.^{5–53}

This study has certain limitations. First, all results were inferred from case reports and case series. Similar to other reports concerning suspected adverse drug reactions, the relationship between Kayexalate or Kalimate and the described GI adverse events presented in our review remains uncertain. Second, this study was limited by some missing information, the absence of a standardized method of reporting adverse effects, and also possible selection and publication biases. Finally, Kayexalate- or Kalimate-associated GI adverse effects that are small in severity likely never reported and thus not recorded clinically, thereby limiting the quantification of the incidence of these events and evaluating the risk factors.

Conclusions

The use of Kayexalate or Kalimate, both with and without sorbitol, may be associated with fatal GI injuries. The most common high-risk groups of Kayexalate- or

Kalimate-induced GI injuries are patients with CKD and ESRD, followed by patients with hypertension and organ transplantations. However, Kayexalate or Kalimate is given primarily to CKD or ESRD patients because those are the patients who suffer from hyperkalemia. Therefore, it would be expected that GI adverse effects of Kayexalate or Kalimate are observed in these patients. Moreover, the most common site of GI injuries induced by Kayexalate or Kalimate is the colon. Therefore, physicians should carefully assess the patient's intestinal motility and medication history, multiple medical problems, comorbidity, and surgery before prescribing Kayexalate or Kalimate for hyperkalemia management, mainly in patients with ESRD. Physicians should also consider other safer agents when appropriate. Furthermore, detecting Kayexalate or Kalimate crystals is important for pathologists to reach a correct diagnosis.

Data Sharing Statement

Data are available from the corresponding author (Dr. Jen-Wei Chou, codecol@yahoo.com.tw) for researchers who meet the criteria for access to confidential data.

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Disclosure

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