



# Renal and Glucose-Lowering Effects of Empagliflozin and Dapagliflozin in Different Chronic Kidney Disease Stages

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### Specialty section:

This article was submitted to  
Clinical Diabetes,  
a section of the journal  
Frontiers in Endocrinology

**Received:** 14 June 2019

**Accepted:** 08 November 2019

**Published:** 22 November 2019

### Citation:

Lin Y-H, Huang Y-Y, Hsieh S-H,  
Sun J-H, Chen S-T and Lin C-H  
(2019) Renal and Glucose-Lowering  
Effects of Empagliflozin and  
Dapagliflozin in Different Chronic  
Kidney Disease Stages.  
*Front. Endocrinol.* 10:820.  
doi: 10.3389/fendo.2019.00820

**Objective:** The objective of this study was to investigate the effects of sodium-glucose cotransporter 2 (SGLT2) inhibitors on renal function in different stages of chronic kidney disease (CKD).

**Design and Methods:** We conducted a retrospective cohort study using longitudinal claims data from May 2016–December 2017 from the Chang Gung Research Database. Patients who used one of the three types of SGLT2 inhibitor available at Chang Gung Memorial Hospital, namely empagliflozin 10 mg/tab (Empa10), empagliflozin 25 mg/tab (Empa25), and dapagliflozin 10 mg/tab (Dapa), were included, with the same number of matched non-users. Analysis of variance was used for continuous variables and the chi-square test was applied for categorical variables. Differences in data between two groups were analyzed using an independent *t*-test, and the basic data before and after treatment were analyzed using generalized estimating equation (GEE). The association among renal function changes was analyzed using a Cox proportional hazards model, with the results presented as unadjusted hazard ratios (HRs) with 95% confidence intervals (95% CIs).

**Results:** Among the 7,624 SGLT2 inhibitor users, 1,696 patients used Empa10, 2,654 used Empa25, and 3,274 used Dapa. Compared with non-users, dapagliflozin had the lowest risk of estimated glomerular filtration rate (eGFR) decrease over 40% from baseline within 1 year (HR 0.36, 95% CI 0.25–0.51). By using the ICD-10-CM code N179, the acute kidney injury (AKI)-related hospitalization rate was lower in Empa10 and Dapa users than in non-users (HR 0.65, 95% CI 0.49–0.86).

**Conclusion:** Lower risk of eGFR decrease over 40% and AKI-related hospitalization was found in all SGLT2 inhibitor users across the different CKD stages.

**Keywords:** sodium–glucose cotransporter 2 inhibitors, renal function, acute kidney injury, Chang Gung Research Database, glucose control



## Main Outcome Measures

The primary outcome of this study was renal function changes, including acute renal deterioration (eGFR decrease over 40% within 1 year) and AKI-related hospitalization during the follow-up period. The secondary outcome was the difference of HbA1c changes between SGLT2 inhibitor users and non-users.

## Statistical Analysis

The baseline characteristics of the three treatment groups (Empa10, Empa25, and Dapa) were compared using analysis of variance for continuous variables and the chi-square test for categorical variables. Differences in data between two groups were analyzed using an independent *t*-test, and the basic data before and after treatment were analyzed using GEE. The association among renal function changes was analyzed using a Cox proportional hazards model, with the results presented as unadjusted HRs with 95% CIs. To reduce differences in baseline characteristics among the groups, we performed propensity score matching. The baseline characteristics and Cox proportional hazards model for adjusted HRs and 95% CIs are also presented for matched pairs. All statistical analyses were performed using SAS statistical software (version 9.4; SAS Institute, Inc., Cary, NC, USA). A two-sided  $p < 0.05$  was considered statistically significant.

## RESULTS

### Study Population Characteristics

From May 1, 2016 to December 31, 2017, a total of 70,461 individuals with diabetes mellitus were registered in the CGRD. Among these patients, 7,624 patients were included as SGLT2 inhibitor users, with the same number of patients matched as non-users. Demographic characteristics such as sex and age as well as changes in biochemistry data before the first medication use and the closest data before December 2017 are summarized in **Table 1**.

### Comparison Between SGLT2 Inhibitor Users and Non-users

HbA1c decreased after at least 12 weeks in both SGLT2 inhibitor users and non-users. Regarding renal function parameters, elevated Cre and reduced eGFR were more prominent in non-users than in SGLT2 inhibitor users. The results are shown in **Table 1** and **Supplementary Table 2**. The changes in Cre and HbA1c levels and eGFR are shown in **Figure 1**, indicating significant differences in changes between the two groups ( $p < 0.001$ ).

### Comparisons Among Different SGLT2 Inhibitor User Groups

Considering SGLT2 inhibitor users according to the different drugs and dosages used, 1,696 patients used Empa10, 2,654 patients used Empa25, and 3,274 patients used Dapa. Demographic characteristics such as sex and age as well as changes in biochemistry data before the first medication use

and the closest data before December 2017 are summarized in **Table 2** and **Supplementary Table 3**.

AC and PC blood glucose and HbA1c levels decreased in all three groups, with a significant decrease in HbA1c levels in Dapa users compared with Empa25 users.

Regarding renal function parameters, Cre levels increased in all three groups, with a significant difference in change between Empa10 and Dapa users.

### Analysis of Renal Function Change in SGLT2 Inhibitor Users

Cox proportional hazard models were used to analyze changes in eGFR, and HbA1c and Cre levels among Empa and Dapa users during follow-up (**Figure 2**). A lower initial eGFR and higher Cre levels were found in Empa users than in Dapa users. Regarding changes in eGFR and Cre levels over time, significant differences between the two groups were found ( $p < 0.001$ ). We then examined changes in eGFR and Cre levels for the different SGLT2 inhibitor users. The initial Cre level was highest in Empa25 users, followed by that in Empa10 users and then in Dapa users (**Table 2**), that is, the opposite result to that for eGFR. Analysis of changes in eGFR and Cre levels over time revealed significant differences among the three groups ( $p < 0.001$ ) (**Figure 3**).

### Analysis of eGFR Decrease Over 40% and AKI-related Hospitalization in SGLT2 Inhibitor Users

Comparing eGFR decrease over 40% in SGLT2 inhibitor users and non-users, we found a lower incidence of decrease in all SGLT2 inhibitor users (HR 0.51, 95% CI 0.41–0.65) and the lowest in Dapa users (HR 0.36, 95% CI 0.25–0.51). A lower incidence of eGFR decrease over 40% in all SGLT2 inhibitor users in eGFR  $\geq 90$  mL/min/1.73 m<sup>2</sup> and 60–89 mL/min/1.73 m<sup>2</sup> subgroup (HR 0.38, 95% CI 0.26–0.55 and HR 0.64, 95% CI 0.42–0.99, respectively). But in the eGFR 60–89 mL/min/1.73 m<sup>2</sup> subgroup, only Dapa users had the decreased risk (HR 0.54, 95% CI 0.30–0.97) (**Table 3** and **Figure 4**). When followed overtime, we observed that the incidence of eGFR decrease over 40% was lower in SGLT2 inhibitor users than non-users in the 18-month follow-up (HR 0.51, 95% CI 0.41–0.65) (**Figure 5**). Similarly, the cumulative incidence initially increased in non-users compared with SGLT2 inhibitor users after the 18-month follow-up ( $p < 0.001$ ) (**Figure 6**).

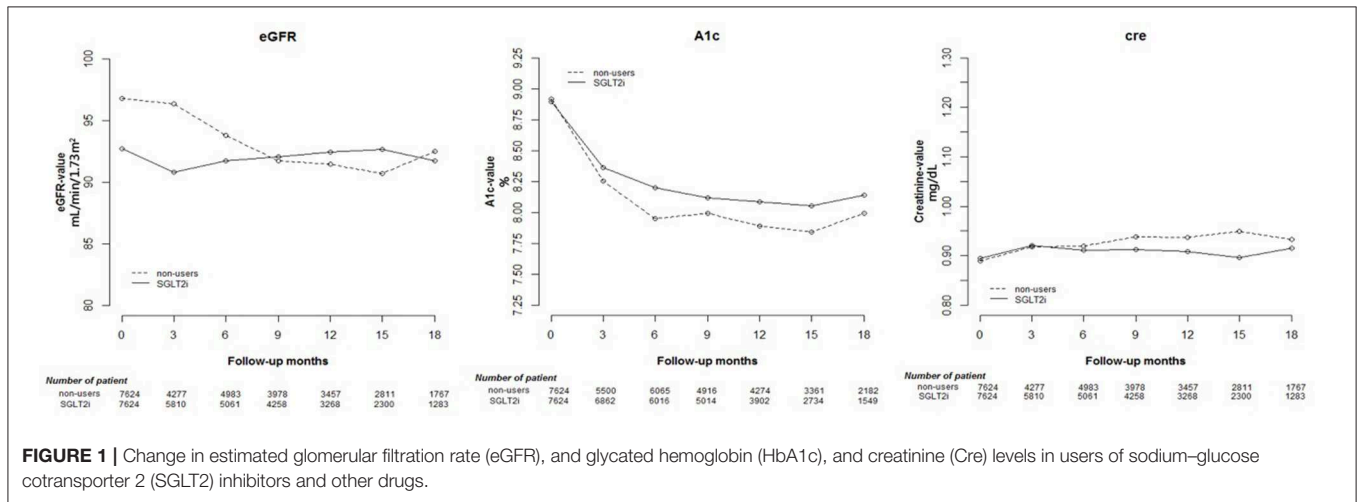
Regarding the AKI-related hospitalization rate, we observed decreased hospitalization rates in SGLT2 inhibitor users compared with non-users (HR 0.66, 95% CI 0.50–0.88) (**Figure 7**). Moreover, when comparing different renal function subgroups with eGFR of  $\geq 90$  mL/min/1.73 m<sup>2</sup> among the three SGLT2 inhibitor user groups and non-users, we found that the lower their eGFR, the higher was their AKI-related hospitalization rates (**Figure 8**). In the 18-month follow-up, there was lower AKI-related hospitalization rate observed in SGLT2 inhibitor users than non-users (HR 0.65, 95% CI 0.49–0.86) (**Figure 9**).

**TABLE 1** | Characteristics of the study population of Sodium glucose co-transporter 2 inhibitor users and non-users.

Basic statistic	SGLT2i		Matched non-users		SGLT2i vs. non-users	SGLT2i vs. non-users
	SGLT2i	Diff (1st-last)	Matched non-users	Diff (1st-last)	p-value	p-value Diff (1st-last)
No.	7,624		7,624			
Age, year	61.9 ± 11.6		61.5 ± 13.		0.061	
Sex (female), %	41.9%		42.2%		0.768	
Cre (first before drug) (mg/dL)	0.895 ± 0.354	-0.035	0.889 ± 0.382	-0.054	0.361	<0.001*
Cre (last) (mg/dL)	0.930 ± 0.432		0.943 ± 0.502		0.096	
eGFR (first before drug) (mL/min/1.73 m <sup>2</sup> )	92.692 ± 31.425	2.554	96.798 ± 38.059	4.898	<0.001*	<0.001*
eGFR (last) (mL/min/1.73 m <sup>2</sup> )	90.138 ± 32.636		91.900 ± 35.889		0.002	
HbA1c (first before drug) (%; mmol/mol)	8.890 ± 1.620 (73.839 ± 17.704)	0.706 (7.718)	8.985 ± 2.266 (74.690 ± 24.771)	0.991 (10.829)	0.015 (<0.015*)	<0.001*
HbA1c (last) (%; mmol/mol)	8.201 ± 1.407 (66.121 ± 15.376)		7.994 ± 1.729 (63.861 ± 18.905)		<0.001*	
UACR (first before drug) (mg/g)	45.445 ± 75.175	-7.778	107.529 ± 542.564	-0.868	0.336	0.391
UACR (last) (mg/g)	53.223 ± 95.205		108.397 ± 542.692		0.399	

Age, Cre, eGFR, HbA1c, and UACR presented as mean ± SD.

SGLT-2i, Sodium-glucose co-transporter 2 inhibitor; Diff, difference; Cre, creatinine; eGFR, estimated Glomerular filtration rate; HbA1c, Glycated hemoglobin; UACR, urine albumin to creatinine ratio. \*p < 0.05.



### Analysis of HbA1c Control Effect in SGLT2 Inhibitor Users

With respect to blood glucose control, a higher initial HbA1c level was found in Dapa users than in Empa users, and the trend of HbA1c change was significantly different in these two groups (Figure 2). The HbA1c change was significantly different among Dapa and Empa10 users (Figure 3). To analyze the blood glucose control effect, we set HbA1c of <7.0% as the target. We found that compared with non-users, a higher proportion of SGLT2 inhibitor users had HbA1c ≥ 7%. However, as seen in Table 1, SGLT2 inhibitor users had a lower average baseline HbA1c level

and less effective HbA1c reduction compared with non-users. In general, Dapa users had the largest decreased proportion of HbA1c ≥ 7%, whereas in the eGFR 30–59 mL/min/1.73 m<sup>2</sup> subgroup Empa had the better effect. In the eGFR15–29 mL/min/1.73 m<sup>2</sup> subgroup, the effect of decline in the proportion of HbA1c ≥ 7% worsened, and blood glucose deteriorated as well (Supplementary Table 4). Furthermore, we analyzed the effect of blood glucose control among Empa10, Empa25, and Dapa users with differing renal function, and found that Dapa users had the largest HbA1c reduction, and Emap25 had the smallest one (Supplementary Table 5).

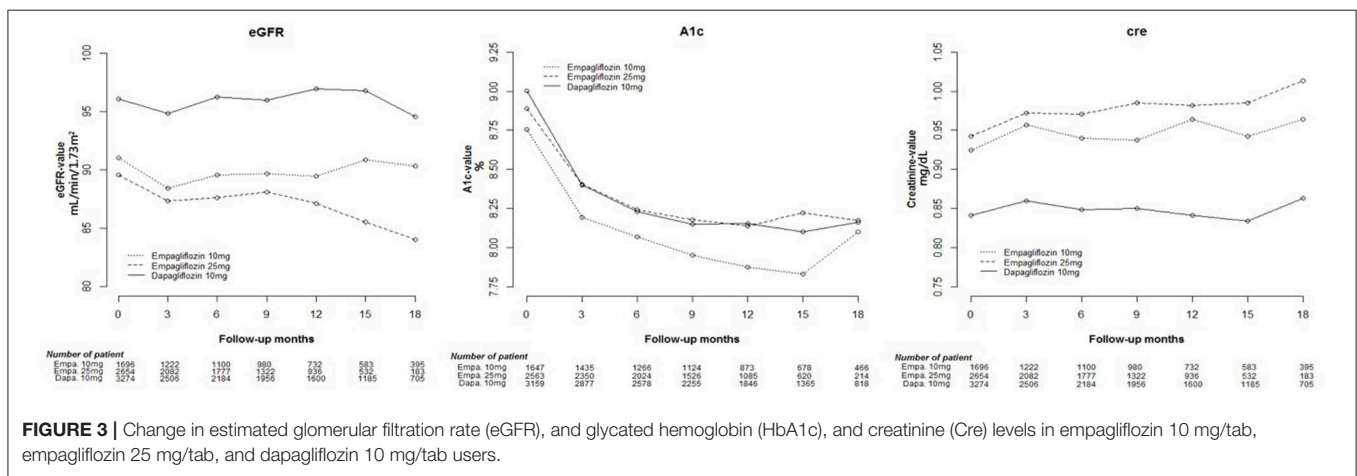
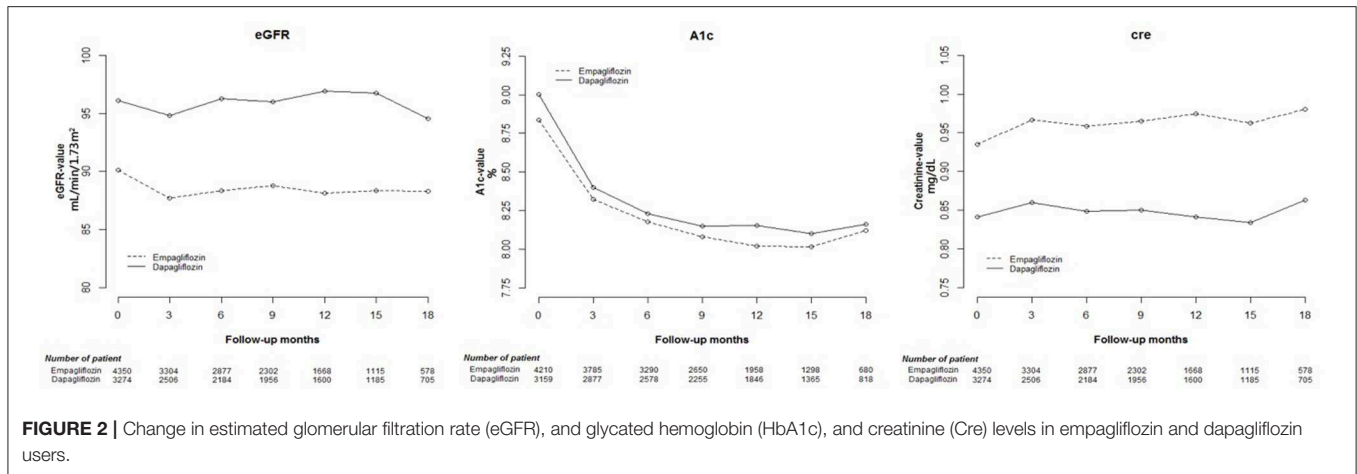
**TABLE 2 |** Characteristics of the study population of three kinds of Sodium glucose co-transporter 2 inhibitor.

Basic statistic	Empagliflozin (10 mg)	Diff (1st-last)	Empagliflozin (25 mg)	Diff (1st-last)	Dapagliflozin (10 mg)	Diff (1st-last)	Empa10 vs. Empa25 vs. Dapa p-value	Empa10 vs. Empa25 diff (varl-varf) p-value	Empa10 vs. Dapa diff (varl-varf) p-value	Empa25 vs. Dapa diff (varl-varf) p-value
N	1,696		2,654		3,274					
Age, year	63.2 ± 11.9		62.0 ± 11.7		61.2 ± 11.4		0.019 <sup>a,b,c</sup>			
Sex (female), %	42.2%		39.9%		43.5%		<0.001 <sup>b</sup>			
Cre (first before drug) (mg/dL)	0.925 ± 0.421	-0.029	0.942 ± 0.402	-0.018	0.841 ± 0.251	-0.011	<0.001 <sup>b,c</sup>	0.165	0.010*	0.163
Cre (last) (mg/dL)	0.954 ± 0.543		0.960 ± 0.492		0.852 ± 0.296		<0.001 <sup>b,c</sup>			
eGFR (first before drug) (mL/min/1.73 m <sup>2</sup> )	91.033 ± 33.474	0.628	89.551 ± 32.985	0.429	96.097 ± 28.581	-0.11	<0.001 <sup>b,c</sup>	0.689	0.145	0.217
eGFR (last) (mL/min/1.73 m <sup>2</sup> )	90.405 ± 34.715		89.122 ± 33.096		96.207 ± 32.902		<0.001 <sup>b,c</sup>			
HbA1c (first before drug) (% mmol/mol)	8.731 ± 1.621 (71.913 ± 17.721)	0.706 (7.714)	8.864 ± 1.724 (73.375 ± 18.848)	0.639 (6.991)	8.976 ± 1.553 (74.596 ± 16.980)	0.785 (8.578)	<0.001 <sup>a,b,c</sup>	0.122	0.057	<0.001*
HbA1c (last) (% mmol/mol)	8.025 ± 1.418 (64.199 ± 15.506)		8.225 ± 1.446 (66.384 ± 15.806)		8.191 ± 1.347 (66.018 ± 14.726)		<0.001 <sup>a,c</sup>			
UACR (first before drug) (mg/dL)	5.675 ± 5.086	0	47.441 ± 70.930	-12.902	25.812 ± 30.930	0.315	0.339	0.352	0.873	0.345
UACR (last) (mg/dL)	5.675 ± 5.086		60.343 ± 106.208		25.497 ± 30.873		0.358			

Age, Cre, eGFR, HbA1c, Alb(U), Creatinine(U) and UACR presented as mean ± SD.

Empa10, Empagliflozine 10 mg/tab; Empa25, Empagliflozine 25 mg/tab; Dapa10, Dapagliflozine 10 mg/tab; Diff, difference; Vari, variate initial; Varif, variate final; Cre, creatinine; eGFR, estimated Glomerular filtration rate; HbA1c, Glycated hemoglobin; AC, ante cibum (preprandial); PC, post cibum (postprandial); AST, aspartate aminotransferase; ALT, Alanine aminotransferase; Na, sodium; K, potassium; LDL-C, low-density lipoprotein cholesterol; HDL-C, High-density lipoprotein cholesterol; TG, Triglyceride; Chol, cholesterol; TBI, total bilirubin; DBI, direct bilirubin; UACR, urine albumin to creatinine ratio.

<sup>a</sup>p < 0.05 between Empa10 and Empa25, <sup>b</sup>p < 0.05 between Empa25 and Dapa10, <sup>c</sup>p < 0.05 between Empa10 and Dapa10. \*p < 0.05.



**TABLE 3 |** Incident rate of decrease in eGFR over 40% between SGLT-2 inhibitor users and non-users in different renal function group.

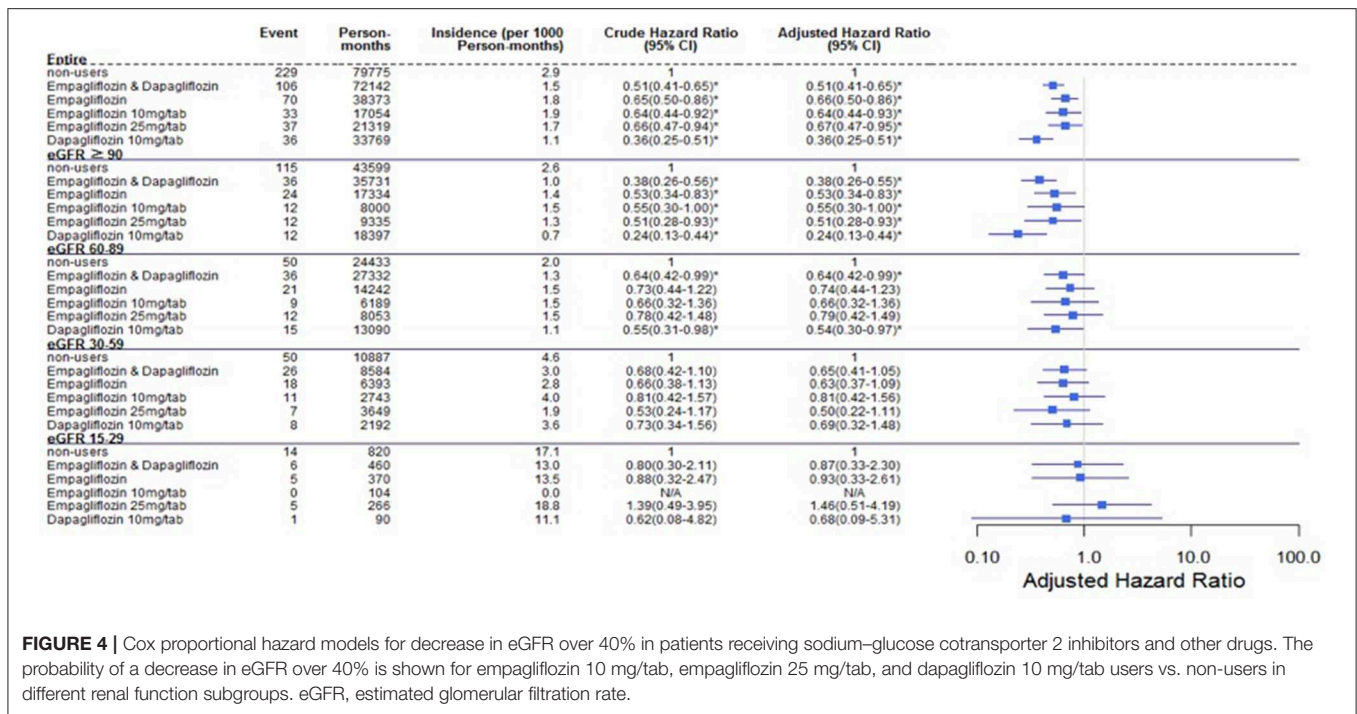
SGLT2i. Incident rate	Entire			Received Empagliflozin and Dapagliflozin			Non-users		
	Event	Person-months	Incident rate per 1,000 Person-months(95% CI)	Event	Person-months	Incident rate per 1,000 Person-months(95% CI)	Event	Person-months	Incident rate per 1,000 Person-months (95% CI)
<b>Entire cohort subgroup by eGFR</b>	335	151,916	2.21 (1.97–2.45)	106	72,142	1.47 (1.19–1.75)	229	79,775	2.87 (2.50–3.24)
≥90	151	79,330	1.90 (1.60–2.20)	36	35,731	1.01 (0.68–1.34)	115	43,599	2.64 (2.16–3.12)
60–89	86	51,765	1.66 (1.31–2.01)	36	27,332	1.32 (0.89–1.75)	50	24,433	2.05 (1.48–2.62)
30–59	76	19,471	3.90 (3.02–4.78)	26	8,584	3.03 (1.87–4.19)	50	10,887	4.59 (3.32–5.86)
15–29	20	1,280	15.62 (8.77–22.47)	6	460	13.04 (2.61–23.47)	14	820	17.07 (8.13–26.01)

## DISCUSSION

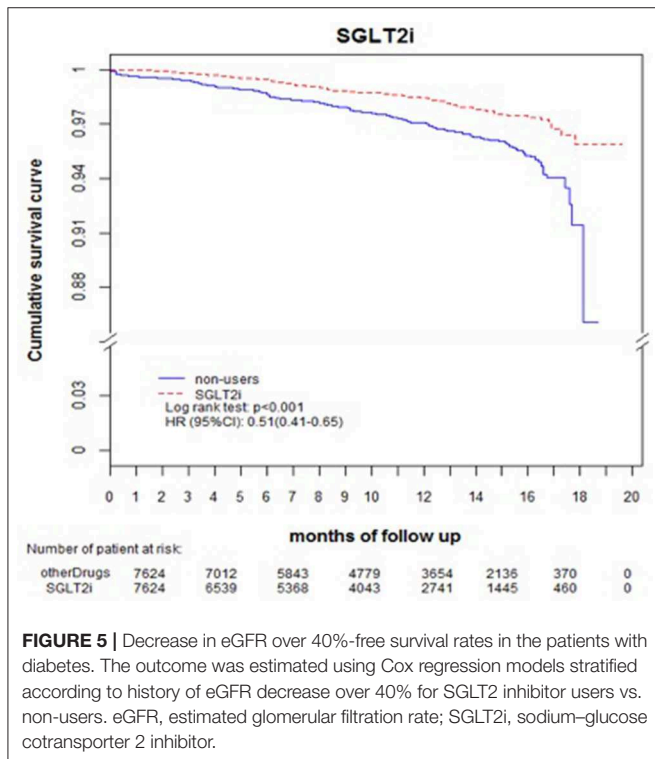
As a new class of antidiabetic drug, the SGLT2 inhibitor lowers plasma glucose levels through renal excretion (1). Because the mechanism of action of SGLT2 inhibitors is based on renal excretion of plasma glucose, whether using SGLT2 inhibitors in

patients with differing renal function would increase the rate of acute renal function deterioration or hospitalization requires further discussion. Two doses of Empa, namely 10 and 25 mg/tab, became available for use at CGMH in May and July 2016, respectively, whereas another SGLT2 inhibitor subtype, dapa10, became available at CGMH in June 2016. We stratified the Empa

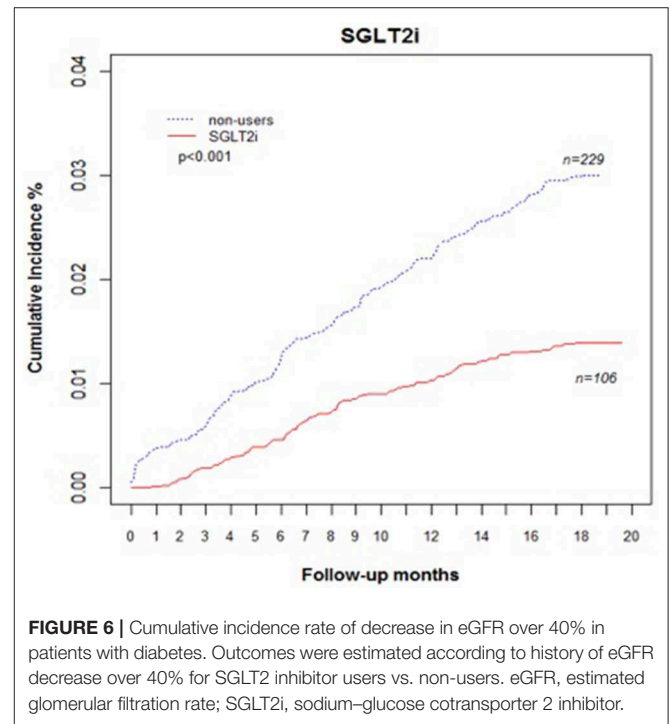




**FIGURE 4 |** Cox proportional hazard models for decrease in eGFR over 40% in patients receiving sodium–glucose cotransporter 2 inhibitors and other drugs. The probability of a decrease in eGFR over 40% is shown for empagliflozin 10 mg/tab, empagliflozin 25 mg/tab, and dapagliflozin 10 mg/tab users vs. non-users in different renal function subgroups. eGFR, estimated glomerular filtration rate.



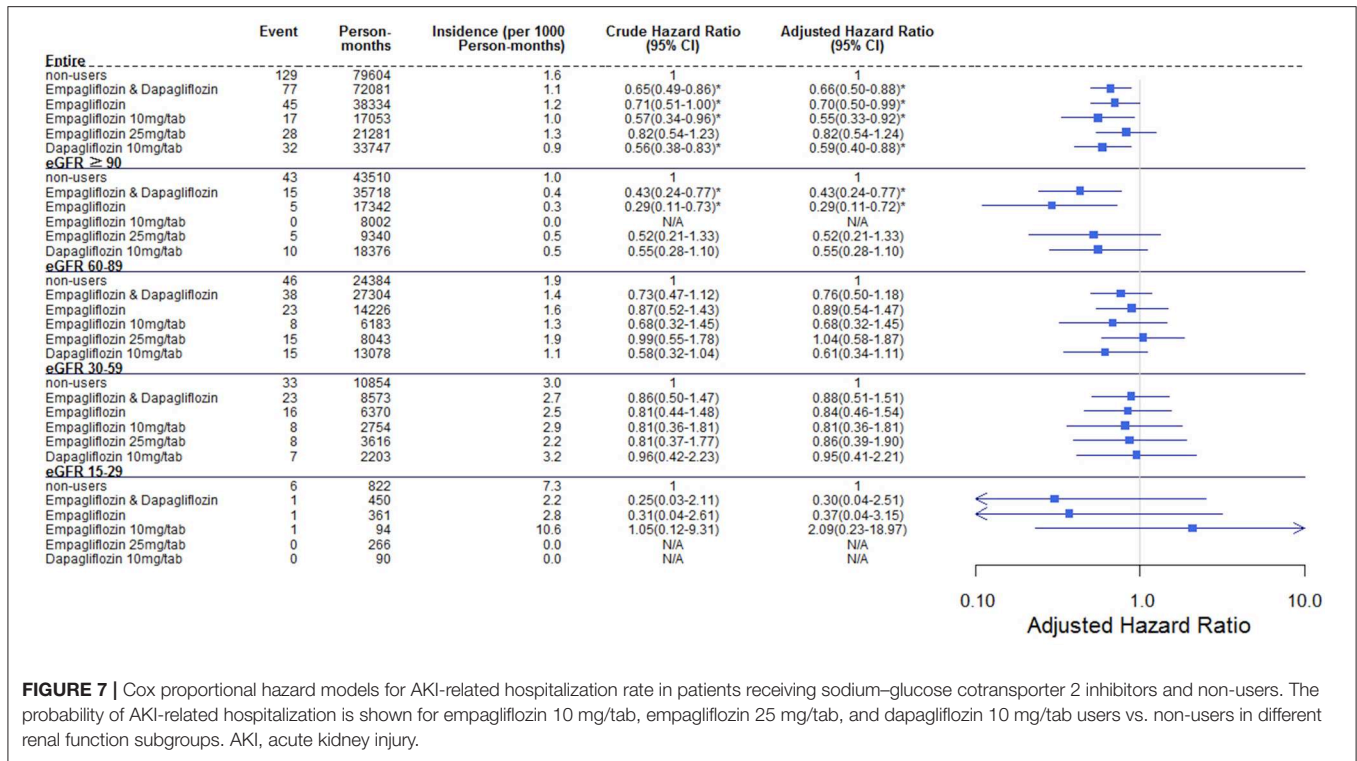
**FIGURE 5 |** Decrease in eGFR over 40%-free survival rates in the patients with diabetes. The outcome was estimated using Cox regression models stratified according to history of eGFR decrease over 40% for SGLT2 inhibitor users vs. non-users. eGFR, estimated glomerular filtration rate; SGLT2i, sodium–glucose cotransporter 2 inhibitor.



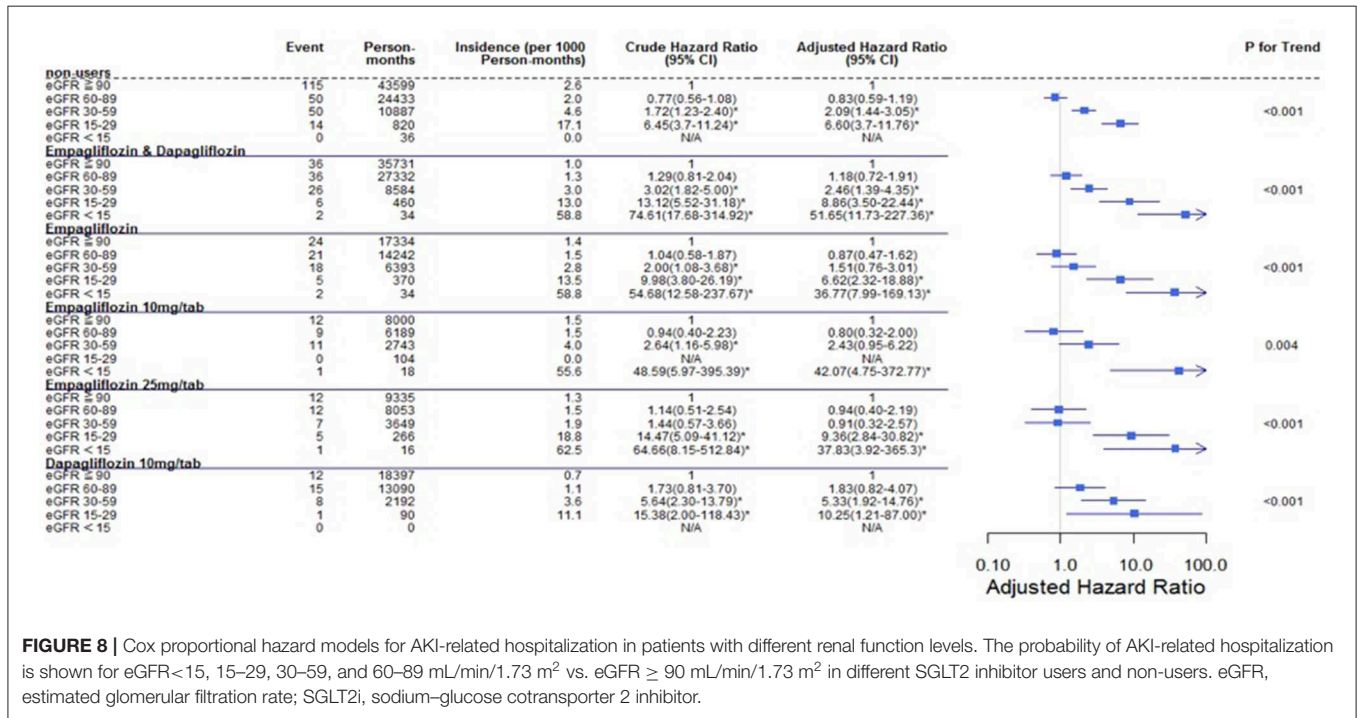
**FIGURE 6 |** Cumulative incidence rate of decrease in eGFR over 40% in patients with diabetes. Outcomes were estimated according to history of eGFR decrease over 40% for SGLT2 inhibitor users vs. non-users. eGFR, estimated glomerular filtration rate; SGLT2i, sodium–glucose cotransporter 2 inhibitor.

dose to 10 or 25 mg/day according to the design of EMPA-REG Outcome Trial (3) and the latest EMPRISE study (16). According to Levine’s review (17), the abilities of HbA1c decreasing between Empa10 and Empa25 as monotherapy are 1.44% vs. 1.43%, respectively. Besides, based on the previous reports, the glucose

lowering effect and renal protection are similar for both Empa10 or Empa25 (18). But the real world data in different CKD stage is not well-studied. The reasons to make physician choose patients who receive Empa or Dapa are dependent on the renal function (Empa could use on the patients with eGFR > 45 mL/min/1.73 m<sup>2</sup>, and Dapa could use on eGFR > 60 mL/min/1.73 m<sup>2</sup>) and



**FIGURE 7 |** Cox proportional hazard models for AKI-related hospitalization rate in patients receiving sodium–glucose cotransporter 2 inhibitors and non-users. The probability of AKI-related hospitalization is shown for empagliflozin 10 mg/tab, empagliflozin 25 mg/tab, and dapagliflozin 10 mg/tab users vs. non-users in different renal function subgroups. AKI, acute kidney injury.



**FIGURE 8 |** Cox proportional hazard models for AKI-related hospitalization in patients with different renal function levels. The probability of AKI-related hospitalization is shown for eGFR <15, 15–29, 30–59, and 60–89 mL/min/1.73 m<sup>2</sup> vs. eGFR ≥ 90 mL/min/1.73 m<sup>2</sup> in different SGLT2 inhibitor users and non-users. eGFR, estimated glomerular filtration rate; SGLT2i, sodium–glucose cotransporter 2 inhibitor.

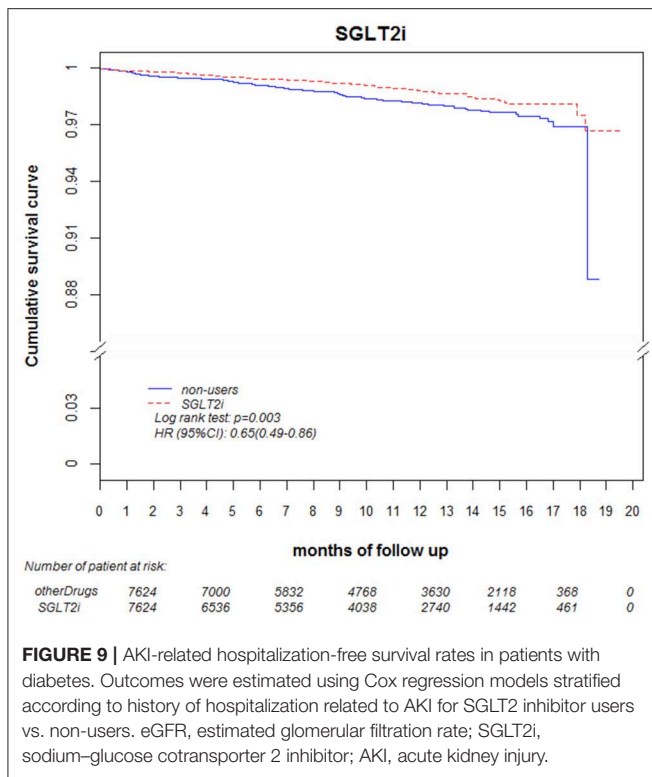
individual preference. We tried to eliminate this bias by matching initial renal function.

We therefore divided SGLT2 inhibitor users into three subgroups, namely Empa10, Empa25, and Dapa, for further comparison.

### Renal Profile

After 18-month follow-up on the renal function of SGLT2 inhibitor users and non-users, we found that Cre levels increased and eGFR reduced in both groups, with these changes being significantly lower in SGLT2 inhibitor users. Moreover, the initial





pre-treatment renal function data were statistically better in SGLT2 inhibitor users compared with non-users. In a meta-analysis, Wang et al. noted that Cre levels statistically significantly increased in patients who received SGLT2 inhibitor treatment (19), whereas, Xu et al. noted significant changes in eGFR after analyzing 47 studies evaluating the effect of SGLT2 inhibitors on renal function change (9). There are multiple potential reno-protective mechanisms of SGLT2 inhibitors including decreased hyperfiltration, reduce glomerular hypertension, lower intravascular volume, etc. (5, 11, 12). Besides, there are increased composite renal events with follow-up duration between 26 and 104 weeks, but not in the other follow-up duration noted by a meta-analysis (20). This may be the reason why there are debates surrounding the effect of SGLT2 inhibitors on renal function because the exact mechanism of SGLT2 inhibitor on kidney event is still unclear and it might be time related.

Having divided SGLT2 inhibitor users into Empa users and Dapa users, we found that Empa users had lower baseline eGFR and higher Cre levels. In the 18-month follow-up period, differences in changes in eGFR and Cre levels between the two groups were statistically significant, with renal function more obviously deteriorating after 12-months in Empa users. Sugiyama et al. noted that dapagliflozin exhibited significant renoprotective effects through renal morphology, although no large increase in eGFR or decrease in Cre levels was observed (21). Since the publication of the EMPA-REG OUTCOME study, numerous studies have supported the renoprotection of Empa (6, 22). Although only a few studies have compared empagliflozin with dapagliflozin, Huilin et al. illustrated that empagliflozin resulted

in fewer renal events than dapagliflozin (20), a result that differs from our findings. The different foothold of renal function in the two SGLT2 inhibitor user groups may explain this discrepancy.

Having divided SGLT2 inhibitor users into three subgroups, we found that Empa25 users had the highest baseline Cre and lowest eGFR, followed by Empa10 users with midrange values and then Dapa users, who had the lowest Cre levels and highest eGFR. After receiving SGLT2 inhibitors, the Cre levels increased in all three groups, with a statistically significant difference between Empa10 and Dapa users (0.029 vs. 0.011 mg/dL,  $p = 0.01$ ). Overtime, the trend of changes in Cre levels and eGFR among the three groups was statistically significant (Figure 3). Specifically, Empa25 users showed deteriorating renal function after 12-months, with a statistically significant difference relative to Dapa users.

Furthermore, there was no significant difference of UACR between SGLT2 inhibitor users and non-users. Several studies have observed that SGLT2 inhibitors have a beneficial effect of UACR reduction (23), although other results for change in UACR after SGLT2 inhibitor use are more neutral (19). Despite the beneficial effect of SGLT2 inhibitors on UACR, progression to macroalbuminuria was observed in a certain percentage of patients with type 2 diabetes (6). Moreover, in our study, baseline UACR was lower in SGLT2 inhibitor users than in non-users. Among the three subgroups of SGLT2 inhibitor users, the highest UACR was found in Empa25 users, followed by Dapa and Empa10 users. After 18-month follow up, UACR increased in Empa25 users but decreased in Dapa users. Contrast to our finding, Cherney et al. demonstrated that Empa reduced UACR and that this effect was better in patients with higher baseline UACR (8). Therefore, a different foothold may influence the effect of SGLT2 inhibitors on UACR.

### Rate of eGFR Decrease Over 40% From Baseline and AKI-related Hospitalization

A lower risk of eGFR decrease over 40% was observed in  $eGFR \geq 90$  mL/min/1.73 m<sup>2</sup> among Empa users and in  $eGFR \geq 60$  mL/min/1.73 m<sup>2</sup> among Dapa users than non-users. Perlman et al. evaluated the relationship between SGLT2 inhibitors and acute renal failure using the U.S. Food and Drug Administration (FDA) adverse event report system database and revealed that SGLT2 inhibitors were more highly related to increased acute renal failure than other drugs (13). Nadkarni et al. found no increased risk of AKI with SGLT2 inhibitor use (24). Wanner et al. analyzed AKI in differing renal function levels, with a cut-off eGFR of 60 mL/min/1.73 m<sup>2</sup> and found fewer AKI events in Empa users and patients with better renal function (6). However, Szalat et al. mentioned several possible mechanisms whereby SGLT2 inhibitors could cause acute renal failure (25).

The incidence of eGFR decrease over 40% was lower in SGLT2 inhibitor users than non-users in 18-months follow up. The benefit of reduced renal deterioration in SGLT2 inhibitor users would become apparent after a long follow-up period. Some evidence indicates that patients treated with dapagliflozin had

a higher rate of recovery from deteriorated renal function and return to the baseline level than those who were treated with other drugs or discontinued dapagliflozin (20).

SGLT2 inhibitor users did not have a higher AKI-related hospitalization rate than other drug users, nor did this rate increase over time. Among the SGLT2 inhibitor user subgroups, the trend in each group was that poorer renal function in patients led to a higher AKI-related hospitalization rate (Figure 8).

## Glucose Control

Comparing SGLT2 inhibitor users with non-users, we observed a less decrease in HbA1c levels after 18-months. This might be due to the weaker glucose lowering ability and less hypoglycemia rate of SGLT2 inhibitors compared with sulfonylurea (26, 27). Besides, in real world practice, physicians may use SGLT2 inhibitors as an add-on regimen in poorly controlled patients who usually have less response to anti-diabetic drugs.

The HbA1c level decreased in all the three SGLT2 inhibitor user subgroups (Empa10, Empa25, and Dapa), with Dapa users showing the statistically significant decrease compared with Empa25 group. Moreover, a statistically significant difference in glucose change was observed among the three groups. Johnston et al. compared several studies using Empa and Dapa and discovered that Empa25 had the best glucose-lowering effect, followed by Empa10 and then Dapa (28), a finding that differs from ours. However, our data shows that the initial HbA1c level was highest in Dapa users, followed by Empa25, with Empa10 users having the lowest HbA1c level. This finding was compatible with that of Yagi et al., who noted a better glucose-lowering effect in patients with type 2 diabetes receiving SGLT2 inhibitor treatment who had a high baseline HbA1c level (29).

This study had several limitations. First, because only Empa and dapa were currently available at CGMH and were only introduced after 2016, other SGLT2 inhibitors were not studied and follow-up time was limited. Second, because this was a non-randomized, retrospective, observational study, selection bias was possible, despite comprehensive propensity score matching and our setting the index date as the start of therapy. Third, some details of renal function could not be distinguished in the diagnostic records. However, for the majority of the participants in this large cohort study, we were still able to indicate the effects of SGLT2 inhibitors. Forth, although we tried our best to eliminate the interference of other drugs between SGLT2 inhibitors users and non-users by propensity score matching, the further evaluation of the

effects on drugs might be added on during the study period is limited.

## CONCLUSION

In real world practice, both Dapa and Empa had similar glucose-lowering effect across different CKD stages. SGLT2 inhibitor users and non-users had reduced renal function at the 18-month follow up, but SGLT2 inhibitor users exhibited lower changes compared with baseline. Moreover, SGLT2 inhibitor users had a lower incidence of eGFR decrease over 40% within 18-months without increase in the AKI-related hospitalization rate.

## DATA AVAILABILITY STATEMENT

The datasets analyzed in this manuscript are not publicly available. Requests to access the datasets should be directed to C-HL, adronlin@cgmh.org.tw.

## AUTHOR CONTRIBUTIONS

Y-HL wrote the manuscript and researched and analyzed the data. Y-YH, S-HH, J-HS, and S-TC researched the data. C-HL designed, reviewed, edited the manuscript, the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. All authors were involved in the interpretation of data, critical revision, and approval of the manuscript.

## FUNDING

This study is based in part on data from the Chang Gung Research Database provided by the Chang Gung Memorial Hospital. This work was supported by Chang Gung Memorial Hospital (grant numbers: CORPG5F0011, CMRPG3H0401, and CMRPG3H0941) and the Ministry of Science and Technology, ROC (grant numbers: NSC-MOST105-2628-B-182A-007-MY3 and NSC-MOST 105-2628-B-182-012-MY3).

## SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fendo.2019.00820/full#supplementary-material>

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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