Predictors of Diabetic Ketoacidosis among Patients with Type 1 Diabetes Mellitus Seen in the Emergency Unit

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Authors’ contributions
This work was carried out in collaboration between both authors. Both authors read and approved the final manuscript.

ABSTRACT

Aims: To identify the factors that can predict the development of diabetic ketoacidosis (DKA) among patients with type 1 diabetes mellitus (T1DM) in Basrah.

Materials and Methods:
Place and Duration of Study: The study was conducted in the emergency units of Al-Faiha General Hospital and Al-Basrah General Hospital for the period from June 2013 to June 2014.
Methodology: This study was cross-sectional enrolled four hundred patients with T1DM seen in the emergency unit were selected; patients and their families were subjected to specific questionnaires prepared for this study.
Results: In this study, 400 patients with T1DM enrolled, 160 patients were having DKA (40%), 240 (60%) were not. Factors that predict DKA include age, gender, school level of both the patients and their parents, address of the patients, the early initiation of the proper insulin regimen after...
diagnosis of T1DM, presence of acute recent illness, missing insulin doses and frequency of the
dose missing, number of previous DKA attacks, using of syringes or pens as a tool of insulin
delivery, however, the most important two predictors are the source of insulin supply to patients,
presence of glucometer at homes and frequency of its uses.

**Conclusion:** This result provided evidence that multiple factors interact together to play a vital role
in the development of DKA among patients with T1DM in Basrah.

**Keywords:** Diabetes mellitus; type 1; diabetic ketoacidosis; predictors.

1. INTRODUCTION

Diabetic ketoacidosis (DKA) is a serious acute complication of diabetes mellitus. According to
American Diabetes Association, the mortality rate in patients with DKA is reported to be less than
5% [1]. However, it is reported that there has been a 4.2% decrease in the death rate per year
in the United States during 1985-2002 [2]. Mortality in patients developing DKA is predominately due to the underlying morbidities
such as sepsis or myocardial infarction [3].

The most common precipitating factor in the development of DKA is missed insulin doses [4].
Other factors that may precipitate the incidence of DKA are infections, cerebrovascular accident,
alcohol abuse, pancreatitis, myocardial infarction, trauma and drugs such as corticosteroids,
thiazides, and sympathomimetic agents.

In young patients with type 1 diabetes mellitus (T1DM), psychological problems with eating
disorders may be a contributing factor for 20% of recurrent DKA [5].

The majority of DKA cases occurs in patients with previously diagnosed diabetes [5], and only
a small proportion of patient accounts for the majority of hospitalizations [6]. Thus, it is
estimated that 50% of hospital admissions could be prevented by improved outpatient treatment
and better adherence to self-care [7].

Recent studies found higher glycosylated hemoglobin (HbA1C), female sex, older age,
family and school problems, higher insulin dose, underinsurance, and psychiatric disorders to be
associated with an increase in DKA rate [5,8,9].

The early identification and treatment of patients at risk is essential to be targeted in intervention programs aimed at preventing DKA [10].

Despite major advances in the care of diabetes, DKA remains a leading cause of hospitalization
and the leading cause of morbidity and death in children and adolescents with T1DM.

It is important to try to prevent DKA in order to reduce morbidity and mortality associated with severe metabolic decompensation [11]. This prevention can be accomplished through appropriate education, improved self-care and adherence, and consistent self-monitoring of blood glucose and ketones [12]. DKA as an initial manifestation of T1DM is less amenable to prevention [13], other than through surveillance in youth with a positive family history of diabetes, and increased public awareness of the symptoms of diabetes.

The study aimed to identify the factors that can predict the development of DKA among patients
with T1DM in Basrah.

2. MATERIALS AND METHODS

2.1 Design and Data Collection

A cross-sectional case-control study was carried out to analyze the predictors of DKA among
patients with type 1 diabetes mellitus seen in the emergency unit that involved four hundred
patients with type 1 diabetes mellitus seen in the emergency units of Al-Faiha and Al-Basrah
General Hospitals in Basrah, Southern Iraq during a one-year period (June 2013-June 2014).

The ethical committee of the Basrah College of Medicine approved the study.

2.2 Measurements

All patients are sometimes their relatives (parents, brothers, sisters) were subjected to
specific questionnaires already prepared for this study. Written informed consent was taken from
the patients or their relative if they age under 19 years.

Patients were selected according to inclusion and exclusion criteria; the inclusion criteria for
the patients were: patients who were 15 years (visiting adult emergency unit) and above and diagnosed with type 1 diabetes mellitus.

Patients who were not interested in questionnaires and those who's their first diagnosis of type 1 diabetes mellitus were as DKA during the carrying of the study in the emergency units were excluded.

T1DM was defined as insulin-dependent less than six months from diagnosis [14].

DKA was defined as a tetrad of blood glucose>200 mg/dl (11 mmol/L), ketonemia and ketonuria, venous PH<7.3 and or bicarbonate<15 mmol/L [15], in addition to the clinical features of DKA in the patients. In this study arterial, blood gas analyzer (IRMA (TRU POINT)) was used to check the PH and bicarbonate level.

2.3 Definition of Variables

Variables were defined as:

- The address of the patients: all the centered areas in Basrah are considered as urban while all the peripheral areas are considered as rural.
- In regard to the insulin starting after diagnosis of T1DM: Patients who started prescribing insulin therapy on the same day of T1DM diagnosis by the doctor are considered as immediate insulin users otherwise they are considered non-immediate insulin users.
- In regard to the regimen of insulin: in the study any patient used prandial regular human insulin three times daily with additional doses of basal insulin (including NPH insulin) at bed time is considered as basal-bolus insulin user otherwise they are considered non basal bolus insulin users.
- In the study, patients who use syringes for insulin delivery are considered as insulin syringe users otherwise they are considered non-insulin syringes users including those using insulin pens or using both syringes and insulin pens for insulin delivery.
- What is considered as "recent acute illness" in this study is that any patient who had an illness of belongs to or occurring at a time immediately before the present, and his disease is of rapid onset and or a short course.
- Regarding missed insulin dose: any doses of daily insulin missed intentionally or unintentionally were considered as missing an insulin dose.
- Previous DKA in this study, is that any patient who had DKA diagnosis documents by a doctor with hospital admission and discharging card.

2.4 Statistical Analysis

Data obtained from the questionnaires was analyzed using the Statistical Package for Social Sciences (SPSS) software version 15.0. Data are presented as a mean± standard deviation in case of continuous variables and as absolute numbers (percentage) in case of dichotomous variables with statistical analysis was carried out using Student's t test for continuous variables and Chi-square test for dichotomous variables.

The (p-value) less than 0.05 was considered significant. Univariate analysis was performed to determine each variable's association with diabetic ketoacidosis. In 400 patients, univariate analysis was performed for each of the factors in the presence or absence of diabetic ketoacidosis after the presence of diabetic ketoacidosis was used as the dependent variable. Variables found to be associated with diabetic ketoacidosis in univariate analysis were then entered into a multivariable model using logistic regression to determine the power of each variable for associated with diabetic ketoacidosis.

3. RESULTS

The baseline characteristics of all patients are given in (Table 1) including: Age, gender, duration of T1DM, school level of the patients and their parents, address, insulin started after the diagnosis, a regimen of insulin, tool for insulin delivery, recent acute illness, missed insulin dose, previous DKA, previous hospitalization, availability of glucometer at home and the frequency of its use, and finally the source of insulin supply.

Out of 400 patients included in this study 149 (37.2%) were females, and 251 (62.8%) were males. Studied all the patients had T1DM and were on insulin therapy. Their age was 15 years and above with the oldest patient was aged 45 years and with a mean age of 27.5±6.2 years.

Patients were subdivided into 2 subgroups according to their duration of T1DM, group1<3 year and group2 ≥ 3 years, whose number of patients and their percentages were 162 (40.5%), 238 (59.5%) respectively.
School level of participating patients was categorized into those who have ≤ 6 years of school achievement 209 (52.3%) and those who have >6 years school achievement 191 (47.7%). The school level of the patients' parents participating in the study categorized into: those parents who have ≤ 6 year's school achievement 257 (64.3%) and those parents who have >6 years school achievement 143 (35.7%).

Regarding the address of the patients included in this study, patients were divided into those living in rural areas 183 (45.8%) and those who live in urban areas 217 (54.2%).

According to starting on insulin therapy after diagnosis of T1DM, patients subdivided into 2 groups: group1 those who start insulin immediately after diagnosis and group2 included patients who not start insulin immediately and were found that their number and percentage 213 (53.3%), 187 (46.7%) respectively.

The regimen of the used insulin subdivided the studied patients into 2 groups: Group 1 those who used the regimen of basal-bolus insulin 186 (46.5%), group 2 those who used non basal-bolus insulin regimen (Premix insulin 70:30 twice daily or modified basal-bolus) were 214 patients (53.5%).

There were 214 (53.5%) of the studied patients were having a recent acute illness, whereas 186 (46.5%) did not have.

According to the tool used for insulin delivery, the studied patients were subdivided into: Those who used syringes only to deliver insulin 186 (46.5%), those who not used syringes only (using pens or both pens and syringes) 214 (60.3%).

According to the presence of insulin missing doses; the studied patients were subdivided into two groups: Group 1 those who had missed insulin dose 166 (41.5%), and group 2 were those who not missed insulin dose 234 (58.5%), and according to the frequency of missed insulin doses; patients were subdivided into 2 groups also: Group 1 those who had one missed insulin dose 101 (60.8%) and group 2 were those who had multiple missed insulin doses 65 (39.2%).

According to the availability of glucometer at home and frequency of its use for monitoring blood sugar, we subdivided the participated patients into 2 groups: Group 1 those who had glucometer 131 (32.8%) and group 2 were those who not had glucometer at home 265 (67.2%); and according to the frequency of glucometer using at home the studied patients subdivided into two groups: Those who use glucometer < 4 times daily to check blood sugar 52 (39.7%) and those who use glucometer ≥ 4 times daily 79 (60.3%).

According to the presence of previous DKA attack; patients were categorized into two groups: Group1 those who had previous DKA 223 (55.8%) and group 2 included those who did not have previous DKA attack 177 (44.2%); and according to the frequency of previous DKA; patients participating in this study were subdivided into: those who had <2 attacks of previous DKA 106 (47.5%) and those who had ≥2 attacks of previous DKA 117 (52.5%).

Finally, according to the source of insulin supply, patients participating in this study were subdivided into two groups: group1 those who depend on Al-Faiha Specialized Diabetes, Endocrine, and Metabolism Center (FDEMC) as a source of insulin supply 203 (50.8%) and group 2 were those who depend on the public clinic as an insulin source supply 197 (49.2%).

It was found that there was a significant relationship between the gender of the patients with T1DM and DKA development; p-value was 0.001 and the female patients mainly affected; from a total of 149 female patients with T1DM enrolled in this study; 77 developed DKA compared to 83 DKA male patients from a total of 251 male patients with T1DM involved in this study (Table 2).

Age also had a significant association with DKA development; p-value was 0.003, and young age is associated with DKA occurrence; 92 patients out a total of 193 patients aged ≤ 20 year compared to 68 patients for a total of 207 at age group> 20 year.

There is a strong relationship between duration of T1DM and DKA development; p-value=0.025.

A high effect of the educational level of both the patients and their parents on DKA developments with p-value=0.001 for both.

There was a significant effect of the patients' address on DKA developments; 105 patients out of 183 of those with T1DM living in the rural areas developed DKA, while only 55 out of 207 of those with T1DM living in urban areas developed DKA; p-value=0.001.
It was also found that there is a significant effect of the time of starting insulin therapy in patients with T1DM on DKA development, and a delay in initiation of insulin therapy after diagnosis of T1DM increase the risk of DKA development among those patients; \( p \)-value=0.001.

The initiation of a proper regimen of insulin in patients with T1DM strongly influence DKA development, with the lowest risk of DKA among those used basal-bolus regimen in which 62 patients, developed DKA from a total of 186 patients used that regimen; and the highest risk was among those who not used the regimen of basal-bolus insulin (using twice daily premix 70:30 insulin or modified basal-bolus) from a total of 214 used that regimen, 98 patients developed DKA, \( p \)-value=0.011.

The tool of insulin delivery used by patients with T1DM strongly affects the DKA development, with the highest number among those used insulin pens only (non-syringes) while the lowest number of those used syringes, \( p \)-value=0.001.

**Table 1. Study characteristics of patients**

| Variables                                | No.(%) |
|------------------------------------------|--------|
| Gender                                   |        |
| Male                                     | 251(62.8) |
| Female                                   | 149(37.2) |
| Age(Years) Mean ± SD 27.5±6.2            |        |
| \( \leq 20 \)                             | 193(48.3) |
| \( >20 \)                                | 207(51.7) |
| Duration of diabetes mellitus (Years)    |        |
| \( <3 \)                                 | 162(40.5) |
| \( \geq 3 \)                              | 238(59.5) |
| School level of patients (Years)         |        |
| \( \leq 6 \)                              | 209(52.3) |
| \( >6 \)                                 | 191(47.7) |
| School level of parents (Years)          |        |
| \( \leq 6 \)                              | 257(64.3) |
| \( >6 \)                                 | 143(35.7) |
| Address of patients                      |        |
| Rural                                    | 183(45.8) |
| Urban                                    | 217(54.2) |
| Insulin starting after diagnosis         |        |
| Immediately                              | 213(53.3) |
| Non Immediately                          | 187(46.7) |
| Regimen of insulin                       |        |
| Basal Bolus                              | 186(46.5) |
| Non Basal Bolus                          | 214(53.5) |
| Recent acute illness                     |        |
| Yes                                      | 214(53.5) |
| No                                       | 186(46.5) |
| Tool of insulin administration           |        |
| Syringes                                 | 186(46.5) |
| Non Syringes                             | 241(60.3) |
| Missed insulin dose                      |        |
| Yes                                      | 166(41.5) |
| No                                       | 234(58.5) |
| Frequency of missed insulin dose          |        |
| One Dose                                 | 101(60.8) |
| Multiple Doses                           | 65(39.2) |
| Self-monitoring of Blood Glucose (SMBG)  |        |
| Present                                  | 131(32.8) |
| Not Present                              | 265(67.2) |
| Frequency of self-monitoring of blood glucose usage (Number/Day) | |
| \( <4 \)                                | 52(39.7) |
| \( \geq 4 \)                             | 79(60.3) |
| Source of insulin                        |        |
| Al-Faiha Specialized Diabetes, Endocrine, and Metabolism Center (FDEMC) | 203(50.8) |
| Public Clinic                            | 197(49.2) |
| Previous diabetic ketoacidosis            |        |
| Yes                                      | 223(55.8) |
| No                                       | 177(44.2) |
| Number of previous diabetic ketoacidosis  |        |
| \( <2 \)                                | 106(47.5) |
| \( \geq 2 \)                             | 117(52.5) |
| DKA                                      |        |
| Yes                                      | 160(40) |
| No                                       | 240(60) |
Table 2. Univariate association of potential confounding variables with DKA (No.=400)

| Variables                                      | Diabetic ketoacidosis | Non-diabetic ketoacidosis | Total (%) | p value   | Odds ratio (95% confidence interval) |
|------------------------------------------------|-----------------------|----------------------------|-----------|-----------|--------------------------------------|
| Age years (Mean±SD)                            |                       |                            |           |           |                                      |
| ≤ 20                                          | 92                    | 101                        | 193(48.3) | 0.003     | 1.862                                |
| >20                                            | 68                    | 139                        | 207(51.7) |            | (1.242-2.791)                       |
| Sex                                            |                       |                            |           |           |                                      |
| Male N (%)                                     | 83                    | 168                        | 251(62.8) | 0.001     | 2.165                                |
| Female N (%)                                   | 77                    | 72                         | 149(37.2) |            | (1.429-3.279)                       |
| Duration of type 1 DM (Year) no (%)            |                       |                            |           |           |                                      |
| < 3                                            | 54                    | 108                        | 162(40.5) | 0.025     | 1.606                                |
| ≥ 3                                            | 106                   | 132                        | 238(59.5) |            | (1.061-2.432)                       |
| School level of patients (Years) no (%)        |                       |                            |           |           |                                      |
| ≤ 6                                            | 102                   | 107                        | 209(52.3) | 0.001     | 2.186                                |
| > 6                                            | 58                    | 133                        | 191(47.7) |            | (1.450-3.296)                       |
| School level of parents (Year) no (%)          |                       |                            |           |           |                                      |
| ≤ 6                                            | 121                   | 136                        | 257(64.3) | 0.001     | 2.373                                |
| > 6                                            | 39                    | 104                        | 143(35.7) |            | (1.525-3.692)                       |
| Address no (%)                                 |                       |                            |           |           |                                      |
| Rural                                          | 105                   | 78                         | 183(45.8) | 0.001     | 3.965                                |
| Urban                                          | 55                    | 162                        | 217(54.2) |            | (2.596-6.056)                       |
| Insulin started after diagnosis no (%)         |                       |                            |           |           |                                      |
| Immediately                                    | 46                    | 167                        | 213(53.3) | 0.001     | 0.176                                |
| Not immediately                                | 114                   | 73                         | 187(46.7) |            | (0.144-0.274)                       |
| Regimen of insulin no (%)                      |                       |                            |           |           |                                      |
| Basal bolus                                    | 62                    | 124                        | 186(46.5) | 0.011     | 1.690                                |
| Non basal bolus                                | 98                    | 116                        | 214(53.5) |            | (1.125-2.537)                       |
| Tool of insulin administration no(%)           |                       |                            |           |           |                                      |
| Syringes                                       | 52                    | 107                        | 159(39.7) | 0.017     | 1.671                                |
| Non syringes                                   | 108                   | 133                        | 241(60.3) |            | (1.101-2.536)                       |
| Recent acute illness no (%)                    |                       |                            |           |           |                                      |
| YES                                            | 109                   | 105                        | 214(53.5) | 0.001     | 2.748                                |
| No                                             | 51                    | 135                        | 186(46.5) |            | (1.807-4.178)                       |
| Missed insulin dose(s) no (%)                  |                       |                            |           |           |                                      |
| YES                                            | 81                    | 85                         | 166(41.5) | 0.001     | 1.870                                |
| NO                                             | 79                    | 155                        | 234(58.5) |            | (1.244-2.810)                       |
| Number of insulin missed doses no (%)          |                       |                            |           |           |                                      |
| Single                                         | 32                    | 69                         | 101(60.8) | 0.001     | 6.604                                |
| Multiple                                       | 49                    | 16                         | 65(39.2)  |            | (3.269-13.338)                      |
| Previous diabetic ketoacidosis no (%)          |                       |                            |           |           |                                      |
| Yes                                            | 114                   | 109                        | 223(55.8) | 0.001     | 2.978                                |
| No                                             | 46                    | 131                        | 177(44.2) |            | (1.945-4.562)                       |
| Number of previous diabetic ketoacidosis no (%)|                       |                            |           |           |                                      |
| < 2                                            | 39                    | 67                         | 106(47.5) | 0.001     | 3.068                                |
| ≥ 2                                            | 75                    | 42                         | 117(52.5) |            | (1.778-5.298)                       |
| Presence of self-monitoring blood glucose device no (%) | 23  | 108 | 131(32.8) | 0.001 | 0.205 |
| Frequency of self-monitoring blood glucose device usage(number/day) no (%) | 16 | 36 | 52(39.7) | 0.01 | 4.571 |
| Source of insulin supply no (%)                |                       |                            |           |           |                                      |
| Al-Faiha Diabetic Endocrine and Metabolic Center | 69  | 134 | 203(50.8) | 0.013 | 1.667 |
| Public Clinic                                  | 91                    | 106                        | 197(49.2) |            | (1.114-2.496)                       |
One hundred and nine (50.9%) from a total of 214 patients had acute recent illness developed DKA while only 51 (27.4%) patients developed DKA from a total of 186 patients who did not have acute recent illness, \( p\)-value = 0.001; so there is a significant correlation between DKA development among patients with T1DM and presence of recent acute illness among them.

This study also reveals a significant relationship between DKA development and missing an insulin dose, \( p\)-value = 0.001.

The importance of the frequency of the missed insulin dose also highlighted in our study and our results revealed a strong association between increased the frequency of the missed insulin doses and DKA development, \( p\)-value = 0.001.

The presence of glucometer at home and the frequency of its used for checking blood sugar has a strong effect on DKA development, with only 7 patients from a total of 79 patients who had glucometer and using it frequently developed DKA, sixteen patients developed DKA from a total of 52 patients who had a glucometer at home but used it infrequently while 137 patients from a total of 256 did not have a glucometer developed DKA, \( p\)-value = 0.01.

In this study, it was found that the source of insulin supply strongly affects the DKA development in patients with T1DM with 91 patients developed DKA from a total 197 type 1 diabetic patients took their insulin supply from public clinics while only 69 patients developed DKA from a total number of 203 took their insulin supply from FDEMC, \( p\)-value = 0.013.

With a \( p\)-value = 0.001 indicating that there is a strong relationship between presence of previous DKA attack and DKA development in patients with T1DM, from a total of 223 patients had a previous DKA attack, 114 were developed DKA at the time of the study while only 46 patients were found with DKA from a total of 136 patients with T1DM and did not have a previous DKA attack.

Finally, our study revealed that increased the number of previous DKA attacks, increase the risk of future DKA development in patients with T1DM, with a strong relationship was found in this study between number of previous DKA attacks and DKA development, from a total of 117 patients who had \( \geq \) 2 previous DKA attack, 75 patients developed DKA, while among those patients who had just < 2 previous DKA attack, which were 106 patients, only 39 patients developed DKA, \( p\)-value = 0.001.

So variables associated with DKA with statistical significance were all the studied variables. Predictors of DKA identified by multivariate analysis are shown in Table 3.

Variables that remain statistically significantly associated with diabetic ketoacidosis were: age, gender, duration of type1 diabetes mellitus, the address of the patients, insulin regimen, tool of insulin administration, presence of acute recent illness and previous diabetic ketoacidosis attacks and its number, missing of insulin and its frequency, frequency of glucometer usage at home to check blood sugar and the source of insulin supply to the patients with type1 diabetes mellitus.

4. DISCUSSION

The prevalence of T1DM is increasing worldwide [16]. DKA was universally fatal, especially before the days of insulin in the 1920s, after that, the overall mortality decreased to relatively low figures, but there is still the potential of case fatality from DKA either from acidosis or as a complication of therapy [17,18]. There was a predominance of females with DKA in our study that is similar to the universal findings [19-20]. This could be attributed to the fact that insulin purging is frequent to control weight in young women with diabetes [21,22]. Furthermore, adolescent girls with diabetes are at greater risk of developing disordered eating patterns [23]. Adolescent girls and young women are also the group at highest risk to suffer from ‘brittle’ diabetes, a form of severe unstable diabetes with increased metabolic complications, especially recurrent DKA [24].

In accordance with previous studies [17,20], teenage at diabetes onset was related to elevated incidence of DKA. On one hand, greater personal responsibility diabetes management and less parental monitoring adolescents may lead to a deterioration of metabolic control in this group. On the other hand, endocrine changes associated with puberty lead to greater insulin resistance [24]. Comparable reasons may account for the trend toward a higher incidence of DKA in patients with longer duration of diabetes.
| Variable                        | Standard error | Significance probability | Odd ratio (95% confidence interval) |
|--------------------------------|----------------|--------------------------|-------------------------------------|
| Age                            | 0.207          | 0.003                    | 1.212                               |
|                                |                |                          | (1.042-2.371)                       |
| Sex                            | 0.220          | 0.001                    | 2.121                               |
|                                |                |                          | (1.217-3.149)                       |
| Duration of type1 diabetes mellitus | 0.212          | 0.025                    | 1.106                               |
|                                |                |                          | (1.011-2.231)                       |
| School level of patients       | 0.245          | 0.001                    | 1.186                               |
|                                |                |                          | (1.430-2.286)                       |
| School level of parents        | 0.229          | 0.002                    | 2.353                               |
|                                |                |                          | (1.317-3.432)                       |
| Address                        | 0.372          | 0.001                    | 2.762                               |
|                                |                |                          | (2.566-4.053)                       |
| Regimen of insulin             | 0.207          | 0.013                    | 1.690                               |
|                                |                |                          | (1.125-2.537)                       |
| Tool of insulin administration | 0.513          | 0.012                    | 1.221                               |
|                                |                |                          | (1.101-2.122)                       |
| Recent acute illness           | 0.214          | 0.001                    | 2.748                               |
|                                |                |                          | (1.807-4.178)                       |
| Missed insulin dose            | 0.208          | 0.003                    | 1.870                               |
|                                |                |                          | (1.244-2.810)                       |
| Number of missed insulin dose(s) | 0.359          | 0.001                    | 4.612                               |
|                                |                |                          | (3.266-8.298)                       |
| Previous diabetic ketoacidosis  | 0.218          | 0.001                    | 2.978                               |
|                                |                |                          | (1.945-4.562)                       |
| Number of previous diabetic ketoacidosis | 0.279        | 0.001                    | 3.068                               |
|                                |                |                          | (1.776-5.298)                       |
| Frequency of self-monitoring blood glucose device usage | 0.497        | 0.002                    | 3.531                               |
|                                |                |                          | (1.524-8.104)                       |
| Source of insulin supply       | 0.206          | 0.012                    | 1.347                               |
|                                |                |                          | (1.104-2.146)                       |
The low educational level of the patients and their parents had significantly associated with the development of DKA in our study, and these results were also demonstrated in other universal similar studies [20,25].

As expected and supported by similar studies in neighboring countries [26-28], the patient address significantly associated with DKA occurrence and this is clearly related to the educational level of the patients and their families, their economic status and their access to the medication supply and specialized diabetic centers.

Delayed initiation of prompt insulin therapy following a diagnosis of T1DM has been associated with an increased risk of DKA development among patients with T1DM [29]; the same result was found in this study also.

It is found that the initiation of the suitable and effective insulin regime was associated with reduced DKA rate [29,30], as found in this study that the lowest DKA was among those patients using the prandial regular human insulin before each of the three-day major meals with further dose of basal insulin at bedtime (basal-bolus insulin regimen).

To our knowledge, the effect of how insulin delivered on DKA rate has not been investigated before. In our study, it has been found that using of insulin pens had been associated with an increased rate of DKA when compared with the using of the insulin syringes; this is perhaps related to the ignorance of the proper technique for the use of these pens especially among those patients with low educational levels, also in Iraq we use vial and syringes for all, except few used reusable pen, and with unavailability of insulin cartridge make patients refill them by syringes that definitely end with dosing errors because manufacturer recommendation was against the rifling of insulin cartridge.

The recent acute illness and mainly infections are important DKA precipitators in 109 patients. Most patients presented with nausea, vomiting and abdominal pain, which was interpreted as an indication to reduce or stop their insulin. This is a deep-rooted belief shared by some doctors and diabetic educators. The situation is made worse when such patients consume large amounts of sugar-rich fluid to counteract presumed hypoglycemia [31].

Insulin omission intentionally and unintentionally was found in patients with DKA in the present study. Among patients with poor compliance with insulin therapy, patients stopped insulin therapy either because they are away from insulin supply (lost, broken, etc.), not gave clear reason for stopping insulin or did not know how to handle insulin on sick day. Insulin discontinuation has long been recognized as an important precipitating cause of DKA in retrospective studies [32].

Observational studies in urban African Americans have reported that more than one-half of DKA cases in patients with diabetes were caused by noncompliance with insulin therapy [32,33]. Likewise, a retrospective study in a multiethnic population in Texas listed noncompliance with insulin injections as their most common precipitating cause of DKA [34].

Higher insulin doses were observed in the DKA group, which corresponds to the results of Rewers et al. [35]. They hypothesized that a higher reported insulin dose may represent lower endogenous insulin secretion because of longer duration of diabetes because of puberty, but it could be that in patients who miss insulin injections, a higher prescribed insulin dose may not be consistent with the actual applied insulin dose, and that what is found in this study with high number of DKA among patients with multiple missed insulin doses than those who missed a single dose.

Previous DKA and increase number of previous DKA were associated with increased DKA rate [36] and this, what is found in our study.

The goal of glucose monitoring in diabetes is to obtain useful information about the patient’s overall glucose status to normalize glucose and prevent hypoglycemia and minimize hyperglycemia through meaningful and timely interventions. As glucose control is the foundation of diabetic care, self-monitoring of blood glucose (SMBG) is the foundation of glucose monitoring. Studies have shown a direct correlation between SMBG and improve HbA1C levels and reduce DKA rate [37-39], the American Diabetes Association recommends that patient with T1DM should self-test at least three times daily. This is identical to what is found in our study of reducing the number of DKA among those who frequently used SMBG (7 patients with DKA out of a total of 72) compared with (16
Finally, to our knowledge, the effect of source of insulin supply on DKA rate has not been investigated before. In our study, we found that the source of insulin was significantly associated with the development of DKA, and patients who depend on FDEMC as a source of insulin supply were significantly had lower DKA compared with those who took insulin from the public clinic. This is may be related to the advantage of more experienced staff and multidisciplinary treatment as well as the availability of follow-up investigations and better communication with patients and the dealing with all their aspects.

Study limitations: It was a cross-sectional study; so no data available on mortality. The age of onset of T1DM was not studied, but its play a role in DKA development. The incomes of the family also play a role and was not assessed in this study. The puberty stage was not assessed in the studied patients since puberty plays an important role in the loss of control of diabetes during the transition period [40].

5. CONCLUSION

The results of this study provided evidence that multiple factors interact together to play a vital role in the development of DKA among patients with T1DM in Basrah.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

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