Presentation to primary care during the prodrome of type 1 diabetes in childhood: A case-control study using record data linkage

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Objectives: To evaluate primary care presentations during the prodrome (12 months prior to onset type-1 diabetes (T1D), with or without diabetic ketoacidosis [DKA]), to identify opportunities for earlier diagnosis.

Methods: This was a case-control study, linking 16 years of data from children (≤15 years) registered at diagnosis of T1D, and routinely collected primary care records in Wales (United Kingdom). Controls (without T1D) were matched on a 3:1 ratio. Conditional logistic regression modeling was used to compare characteristics occurring in cases (children with T1D) and controls; and cases that presented with/without DKA.

Results: A total of 1345 children with T1D (19% DKA) and 4035 controls were identified. During the 12 months prior to diagnosis, cases were 6.5 times more likely to have at least one primary care contact (P < 0.001). One to 30 days prior to diagnosis, contacts relating to blood tests, fungal conditions, respiratory tract infections (RTIs), urinary conditions, vomiting, and weight were independently associated with T1D, between 91 and 180 days prior to diagnosis. Children with a contact up to a month prior to diagnosis, relating to RTIs, antibiotic prescriptions, and vomiting, were more likely to present in DKA, as were boys (P = 0.047).

Conclusion: There are opportunities in primary care for an earlier diagnosis of T1D in childhood. These data could be used to create a predictive diagnostic tool, as a potential aid for primary care health professionals, to prevent presentation in DKA.

KEYWORDS
childhood, diabetic ketoacidosis, primary care data linkage, prodrome, type 1 diabetes mellitus

1 INTRODUCTION

Identifying children with potential new-onset type 1 diabetes (T1D) who need further investigation, out of the many children presenting with common childhood illnesses is a key challenge for general practitioners (GPs).1,2 Although worldwide the incidence of T1D in childhood is rising at a rate of 3% per year,3 it continues to be a relatively rare condition. T1D is an insidious condition, which may manifest in several ways, with subtle symptoms that can be attributed to other childhood illnesses and which may change with the developmental stage of a child. The four classic symptoms of polyuria, polydipsia, weight loss, and tiredness, may present independently, not at all, or in conjunction with other health concerns such as a viral or fungal infection, increasing the potential for misdiagnosis.

Abbreviations: ALF, anonymous linking field; CI, confidence interval; DKA, diabetic ketoacidosis; GP, general practitioner; OR, odds ratio; RTI, respiratory tract infection; SAIL, secure anonymised information linkage; T1D, type 1 diabetes; WIMD, Welsh Index of multiple deprivation.
Early diagnosis of T1D is critical to avoid children developing life-threatening diabetic ketoacidosis (DKA). Delayed diagnosis was attributed as the main cause of DKA at onset of T1D in a consensus statement from the American Diabetes Association. DKA is associated with significant morbidity, mortality and health service costs. Children presenting in DKA are likely to have poor long-term prognosis and their parents experience greater psychological distress. The rate of DKA at onset of T1D varies across the world, from 10% to 70% and has an inverse relationship with the incidence of T1D. In the United Kingdom, 25% of children diagnosed with T1D present in DKA, which is concordant with other economically developed countries. This rate has remained consistent in the United Kingdom over the past 20 years despite efforts to raise awareness of the symptoms in community and clinical settings.

A systematic review found diagnostic error as a significant risk factor for delayed diagnosis. However, most studies relied on retrospective parental or clinical recall once a child was diagnosed, with associated potential for bias. A Canadian study explored routinely collected primary care data for the 4 weeks prior to diagnosis and showed children newly diagnosed with T1D had significantly more medical encounters in the 4 weeks prior to diagnosis than matched controls, with 33% of children having had at least one medical encounter in the week preceding diagnosis. The most common reasons for the encounter were recorded as upper respiratory infections, urinary tract disorders, and gastroenteritis. A limitation of this study was that it only looked at a 4-week period prior to diagnosis, despite parents recalling that their children displayed subtle symptoms for many months prior to diagnosis. They also did not examine differences between those who presented with and without DKA.

The aim of our study was to evaluate on a country-wide basis, the extent and reasons for contact between children and primary care physicians in Wales (United Kingdom), before diagnosis of T1D, using linkage of a prospectively collected primary care dataset to a national diagnostic dataset (Brecon Group Register). By comparing data in matched controls without T1D, we explored the potential to make an earlier diagnosis, with reduced risks of DKA. To our knowledge, this is the first large-scale study to use prospectively collected primary care data to explore the pathway to diagnosis of childhood T1D, including those who present in DKA, for an extensive period prior to diagnosis.

2 | METHODS

Ethical approval was granted by London—Westminster Research Ethics Committee ref 15/LO/2054.

2.1 Study design and data sources

We conducted a matched case-control study of children aged 15 years and under using the Secure Anonymised Information Linkage (SAIL) Databank (Swansea University). The SAIL Databank contains records of over 5 million people and holds data from about 75% of Welsh GPs over the last 15 years (period of coverage varies by primary care practice as not all practices were recruited to provide data to SAIL at the same time).

GPs record details of any symptoms, diagnoses, medications, or tests, through specific electronic patient record software systems, at the time of the consultation. This routinely collected data is stored as a series of Read codes and free text. However, it is only possible to use the Read codes recorded in subsequent analyses. GPs who agree to share their data with SAIL submit full historical extracts of all coded data to SAIL annually.

Records within SAIL are anonymised and linkable with an anonymous linking field (ALF), which allows individuals to be tracked over time and across datasets, while ensuring researchers have no access to any personal identifiable data. Individuals are linked within SAIL using National Health Service number, name, gender, date of birth, and postcode. We obtained the Welsh Index of Multiple Deprivation (WIMD) quintile for individuals in the study. WIMD is an area-based measure of socioeconomic inequality. Cases and controls were required to have GP data available in SAIL for the 12 months leading up to the index date.

Cases were defined as children 15 years old and under, with T1D, resident in Wales, at the time of diagnosis, and were identified from the Brecon Group Register (Welsch Pediatric Diabetes Interest Group). The Brecon Group Register was established in 1995 and contains consented demographic data from 98% of all newly diagnosed cases of children 15 years old and under with T1D, resident in Wales, including whether they presented in DKA (pH < 7.30). There have been two major ascertainment analyses undertaken using a two-source capture-recapture model with data from pediatricians and primary care. Ascertainment of cases was 98.5% in those diagnosed from 1995 to 2005 and on a second assessment 98.1% from 1995 to 2012.

A contemporary dataset was created by linking the Brecon Group Register existing diagnostic dataset to the SAIL Databank to provide a unique dataset of children aged 0 to 15 years in Wales diagnosed with T1D in a 16-year period between January 2000 and December 2015. We aimed to identify three controls for each case. Controls were children identified from the GP data within SAIL without T1D, matched on birth date (within 1 year), gender, and primary care practice (at time of subject’s diagnosis with T1D). An index date was created for the matched controls using the date of diagnosis of the children with T1D. The resulting dataset therefore contained all primary care contact records in the 12 months prior to diagnosis for children diagnosed with T1D (including those diagnosed in DKA) and a matched sample of children without T1D.

2.2 Definition of a primary care contact

A primary care contact was defined as one primary care event per day identified by unique date and a Read code. Primary care events that did not have an interpretable Read code were included as contacts. We did not include events relating to administrative tasks, repeat prescriptions, or consultations where there was no contact with the patient.

2.3 Grouping read codes

Read codes for presenting features that might be associated with developing T1D in childhood were grouped. The research team agreed to any personal identifiable data.
which symptoms, diagnoses, examinations, tests, and prescriptions should be included within each group based on clinical experience and a review of the literature (Table 1). These variables are referred to as characteristics throughout this manuscript.

Read codes associated with medications prescribed for asthma were included as a benchmark variable. Read codes included within each grouping are listed as supplementary material (Appendix S1).

2.4 | Time periods
To evaluate the prodrome, five specific time periods prior to the index date were predetermined. These were the day of diagnosis, and 1 to 30 days, 31 to 90 days, 91 to 180 days, and 181 to 366 days prior to the day of diagnosis.

2.5 | Statistical methods
Assuming a medium effect size (odds ratio [OR] = 1.5), a moderately low prevalence rate of the presenting characteristic of 30% in the control group, matching three controls per case and using a two sided 5% alpha and 80% power, we required 322 children with T1D (using the sampsi_mcc command in STATA 15). 27

Baseline demographic data (age, gender, WIMD deprivation quintile) and number of primary care contacts in the 12 months prior to the index date were used to describe the cases and controls. A univariable conditional logistic regression model was used to compare the event of a characteristic (prodrome) associated with T1D (cases); ORs alongside 95% confidence intervals (CIs) are presented. Characteristics associated with T1D at the 10% significance level were entered into a multivariable conditional logistic regression model. Models were developed for each of the prodromal time periods. We used a cut off of less than 1% prevalence in cases to exclude characteristics from the model.

2.6 | Diabetic ketoacidosis
A secondary aim was to explore the risks associated with presenting in DKA. Given that 19% of children diagnosed with T1D were in DKA at diagnosis (260 in DKA, 1085 not in DKA) our dataset had sufficient power to observe medium effect sizes for around 20 presenting characteristics. Characteristics of presenting in DKA at diagnosis were examined and associations between the prodrome for children presenting in DKA were analyzed using the same approach as for the matched case-control study.

All analyses were performed using IBM SPSS Statistics (IBM CORP, Armonk, New York) 28 and STATA Statistical (USASTATA CORP, College Station, Texas) software, version 15. 27 Reporting adheres to STROBE for case-controls and RECORD guidelines for reporting routine data.

3 | RESULTS
Between 15 January 2000 and 8 October 2015, 3674 children were identified in the Brecon Group Register and eligible for inclusion (Figure 1). A total of 148 children were excluded due to data errors, inability to match within SAIL, and missing data. A further 2041 (55.5%) children were excluded because they did not have 12 months of GP data leading up to diagnosis. This is because the number of primary care practices submitting data to SAIL were far fewer at the turn of this century. Eligible children were comparable with those excluded in terms of age, gender, and social deprivation.

We identified 1488 cases with T1D (19% presenting in DKA) and 4464 matched controls. We excluded 143 cases (and 429 corresponding matched controls) because they had a coded diagnosis for T1D or a prescription for insulin prior to their date of diagnosis in the Brecon Group Register data. The final study population therefore comprised 5380 children (1345 children with T1D; 260 (19%) presenting in DKA) and 4035 matched controls (Figure 1). There were 57 520 primary care events relating to the 5380 children in the study population.

Demographics for the cases and controls show that that they were well matched (Table 2). The mean age of children at the time of diagnosis of T1D was 8.7 years (SD 3.7) and there were more males

### TABLE 1 Description of characteristics

| Characteristic | Includes | Excludes |
|----------------|----------|----------|
| Abdomen        | Symptoms or diagnoses (eg, colic, epigastric pain) or abdominal examination |          |
| Antibiotics    | All prescriptions for antibiotics |          |
| Blood tests    | Test requests, laboratory procedures, all test results |          |
| Constipation   | Symptoms or diagnoses (eg, constipated) or prescriptions (eg, lactulose, macrogol) | Change in bowel habit |
| Fungal         | Diagnoses (eg, candidiasis, dermatophytosis) or prescriptions (eg, micronazole, clotrimazole) |          |
| Headache       | Symptoms or diagnoses (eg, migraine, cerebral oedema) |          |
| Respiratory tract infection | Symptoms or diagnoses (eg, upper/lower respiratory tract infection, otitis media, temperature) or examinations (eg, rate of respiration) |          |
| Thirst         | Symptoms or diagnoses (eg, excessive fluid intake, polydipsia, advice about fluid intake) |          |
| Tiredness      | Symptoms or diagnoses (eg, tiredness, malaise, lethargy, fatigue) |          |
| Urinary        | Symptoms or diagnoses (eg, urinary tract infection, polyuria) or prescriptions (eg, nitrofurantoin, trimethoprim and desmopressin) or tests, results |          |
| Vomiting/nausea | Symptoms or diagnoses of nausea/vomiting | Diarrhea and vomiting combined |
| Weight         | Symptoms or diagnoses (eg, reduced appetite, weight loss, anorexia) or measures (eg, body mass index) | Weight increasing |
| Benchmark variable | Includes | Excludes |
| Asthma         | All prescriptions for asthma |          |

| Variable       | Includes | Excludes |
|----------------|----------|----------|
| Asthma         | All prescriptions for asthma |          |
than females (47%). During the 12 months prior to diagnosis, 1294 (96%) of children who developed T1D had at least one primary care contact, compared to 3261 (81%) of the matched controls (OR 6.50 [95% CI 4.84 to 8.73]; P < 0.001). Cases also had more contacts with a median of 4 (interquartile range: 2-8) compared to 3–7 in the controls.

### Table 2 - Demographics of children with type 1 diabetes and matched controls

| Demographics                        | Cases (T1 diabetes) (n = 1345) | Controls (n = 4035) |
|-------------------------------------|-------------------------------|---------------------|
| Age at diagnosis (y) mean (SD)      | 8.7 (3.7)                     | 8.7 (3.7)           |
| <2 years, N (%)                     | 23 (2.1)                      | 8.7 (3.6)           |
| 2 to <6                             | 219 (20.2)                    | 18 (6.9)            |
| 6 to <10                            | 321 (29.6)                    | 46 (17.7)           |
| 10 to 15                            | 522 (48.1)                    | 68 (26.2)           |
| Gender                              |                               |                     |
| Females                             | 1884 (46.7)                   | 628 (46.7)          |
| Males                               | 2151 (53.3)                   | 717 (53.3)          |
| Social deprivation (WIMD quintile)  |                               |                     |
| 1 (least deprived)                  | 887 (22.0)                    | 287 (21.3)          |
| 2                                   | 863 (21.4)                    | 229 (21.1)          |
| 3                                   | 823 (20.4)                    | 52 (20.0)           |
| 4                                   | 676 (16.8)                    | 181 (16.7)          |
| 5 (most deprived)                   | 786 (19.5)                    | 42 (16.2)           |
| Primary care contacts over past 12 months (incl. day of diagnosis) |                               |                     |
| None                                | 774 (19.2)                    | 51 (3.8)            |
| At least one:                        | 3261 (80.8)                   | 1294 (96.2)         |
| Median (IQR) contacts               | 3 (2-7)                       | 4 (2-8)             |

For categorical variables, data are number (%).
### Table 3

Univariable analyses of children who had at least one unique primary care contact under each characteristic by time prior to diagnosis of T1 diabetes n (%) |
| Characteristic | 1 to 30 days prior to diagnosis | 31 to 90 days prior to diagnosis | 91 to 180 days prior to diagnosis | 181 to 366 days prior to diagnosis |
|----------------|---------------------------------|---------------------------------|---------------------------------|---------------------------------|
| Abdomen        |                                |                                 |                                 |                                 |
|                 | TID cases | Control | OR (95% CI) | P value | TID cases | Control | OR (95% CI) | P value | TID cases | Control | OR (95% CI) | P value | TID cases | Control | OR (95% CI) | P value |
| Abdomen        | 15 (1.1) | 1 (<1.0) | 45.0 (5.9-340.4) | <0.001 | 13 (1.0) | 16 (<1.0) | 2.5 (12.5-53) | 0.016 | 8 (<1.0) | 21 (<1.0) | – | – | – |
| Antibiotics    | 17 (1.3) | 5 (<1.0) | 1.9 (1.5-2.4) | <0.001 | 122 (9.1) | 210 (5.2) | 1.9 (1.5-2.4) | <0.001 | 118 (8.8) | 233 (8.0) | 1.1 (0.9-1.4) | 0.339 | 151 (11.2) | 1181 (29.1) | 0.9 (0.8-1.1) | 0.138 |
| Asthma meds     | 8 (<1.0) | 10 (<1.0) | – | – | 5 (4.1) | 165 (4.1) | 1.0 (0.7-1.4) | 0.999 | 95 (7.1) | 217 (6.7) | 1.1 (0.8-1.3) | 0.661 | 126 (9.4) | 334 (8.3) | 1.1 (0.9-1.4) | 0.218 |
| Blood tests     | 248 (18.4) | 0 (0) | – | – | 120 (8.9) | 231 (10.0) | 20.7 (12.5-44.4) | <0.001 | 12 (1.0) | 28 (1.0) | – | – | 23 (1.7) | 39 (1.0) | 1.8 (1.1-3.0) | 0.029 |
| Blood (fasting tests only) | 34 (2.5) | 0 (0) | – | – | 1 (1.0) | 1 (<1.0) | – | – | 5 (<1.0) | 2 (<1.0) | – | – | 3 (<1.0) | 5 (<1.0) | – | – |
| Constipation    | 3 (<1.0) | 1 (<1.0) | – | – | 13 (1.0) | 17 (<1.0) | 2.3 (1.1-4.7) | 0.024 | 10 (<1.0) | 37 (1.0) | – | – | 7 (<1.0) | 41 (1.0) | – | – |
| Fungal          | 16 (1.2) | 2 (<1.0) | 1.9 (1.5-2.4) | <0.001 | 16 (1.0) | 9 (0.5) | 1.9 (1.5-2.4) | <0.001 | 122 (9.1) | 210 (5.2) | 1.9 (1.5-2.4) | <0.001 | 118 (8.8) | 233 (8.0) | 1.1 (0.9-1.4) | 0.339 |
| Headache        | 1 (<1.0) | 0 (0) | – | – | 8 (<1.0) | 10 (<1.0) | – | – | 8 (<1.0) | 10 (<1.0) | – | – | 6 (<1.0) | 12 (<1.0) | – | – |
| RTI             | 33 (2.5) | 7 (<1.0) | 14.1 (6.3-32.0) | <0.001 | 100 (7.4) | 168 (4.2) | 1.9 (1.5-2.5) | <0.001 | 102 (7.6) | 294 (7.3) | 1.0 (0.8-1.3) | 0.704 | 121 (9.3) | 438 (10.9) | 0.8 (0.6-1.0) | 0.046 |
| Thirst          | 144 (10.7) | 0 (<1.0) | – | – | 48 (3.6) | 0 (0) | – | – | 0 (0) | 0 (0) | – | – | 0 (0) | 3 (<1.0) | – | – |
| Tiredness       | 14 (1.0) | 0 (<1.0) | – | – | 11 (<1.0) | 0 (0) | – | – | 1 (<1.0) | 1 (<1.0) | – | – | 1 (<1.0) | 1 (<1.0) | – | – |
| Urinary         | 320 (23.8) | 0 (<1.0) | – | – | 133 (9.9) | 50 (1.2) | 9.0 (6.4-12.7) | <0.001 | 37 (2.8) | 64 (1.6) | 1.8 (1.2-2.7) | 0.007 | 41 (3.0) | 91 (2.3) | 1.4 (0.9-2.0) | 0.102 |
| Nausea/vomiting | 24 (1.8) | 0 (<1.0) | – | – | 18 (1.3) | 5 (<1.0) | 10.8 (4.0-29.0) | <0.001 | 5 (<1.0) | 13 (<1.0) | – | – | 5 (<1.0) | 10 (1.0) | – | – |
| Weight          | 118 (8.8) | 1 (<1.0) | 354.0 (59.5-235.9) | <0.001 | 72 (5.4) | 24 (<1.0) | 12.07 (7.0-20.52) | <0.001 | 22 (1.6) | 49 (1.2) | 1.4 (0.8-2.3) | 0.237 | 30 (2.2) | 71 (1.8) | 1.3 (0.8-2.1) | 0.248 |

#### 3.2 Characteristics predicting children who present in DKA at diagnosis by time period

Table 2 shows the characteristics of the individual patients by DKA at diagnosis. Table 4 shows the characteristics of the individual patients by DKA during the time period of 91 to 180 days. Table 3 shows the characteristics of the individual patients by DKA during the time period of 91 to 180 days. Table 4 shows the characteristics of the individual patients by DKA during the time period of 91 to 180 days. Table 3 shows the characteristics of the individual patients by DKA during the time period of 91 to 180 days.

#### 4.2 Predictors of children who present in DKA at diagnosis by time period

One to 30 days prior to diagnosis, 628 (46.7%) cases and 797 (63.3%) controls had at least one primary care contact and significant in 180 days. One to 30 days prior to diagnosis, 628 (46.7%) cases and 797 (63.3%) controls had at least one primary care contact and significant in 180 days. One to 30 days prior to diagnosis, 628 (46.7%) cases and 797 (63.3%) controls had at least one primary care contact and significant in 180 days. One to 30 days prior to diagnosis, 628 (46.7%) cases and 797 (63.3%) controls had at least one primary care contact and significant in 180 days.
conditions were significantly less likely to present in DKA than those who did not.

4 | DISCUSSION

This study of prospectively collected primary care data linked to a population-based pediatric diabetes diagnostic register has shown that for up to 12 months prior to diagnosis, children who went on to be diagnosed with T1D, were more likely to have contact with primary care for a range of specific reasons, than children who were not diagnosed with T1D. Furthermore, up to 30 days prior to diagnosis, children who presented in DKA, were more likely to have a primary care contact relating to RTIs, vomiting and a prescription for antibiotics. These findings together can be used to develop a prediction model for use in primary care to promote an earlier diagnosis of diabetes while reducing the risk of presentation in DKA.

4.1 | Strengths and weaknesses of the study

By analyzing data from 12 months prior to diagnosis, we have identified a longer prodrome of symptoms (up to 6 months) than is usually assumed. Our data from the Brecon Group Register is a robust, country-wide hospital recorded dataset (98% complete coverage of all known childhood-onset cases), and our prospective collection of this diagnostic dataset likely provides an accurate date of diagnosis, as it is recorded by diabetes healthcare professionals at the time of diagnosis. Another strength of our study is that it uses prospectively collected primary care data and therefore the results will be less prone to the bias and inaccuracies of recall which influence many other studies on this topic.2

One limitation of the primary care dataset is the lack of control over the quality of the data collected. In our contemporary dataset, a discrepancy in the date of diagnosis affected 10% of the sample. However, the DKA rate was unaltered by removal of these children from the analyses, so it seems unlikely that these individuals had differing characteristics which would have affected the findings. In addition, we had to exclude 55.5% of the children registered on the Brecon Cohort dataset because their GP was not submitting data to SAIL preceding diagnosis. However, all children excluded from the study were comparable with eligible children, in terms of age, gender, and social deprivation.

A final weakness of analyzing prospective, routinely collected data is that it is not possible to ascertain certain details of consultations, such as why a test was conducted, what dialogue took place at the time of the consultation, and whether a primary care physician had a suspicion of any particular condition.

4.2 | Results of the study in context

The study was conducted in a UK-context. However, early recognition of T1D in primary care, to prevent presentation in DKA at diagnosis, remains a global issue. Therefore, the results of this study are likely relevant to physicians world-wide, regardless of models of primary care.
TABLE 4
Univariable analyses of children presenting in DKA or not in DKA, with at least one contact with primary care for each characteristic n (%)

| Characteristic | Day of diagnosis | 1 to 30 days prior to diagnosis | 31 to 90 days prior to diagnosis | 91 to 180 days prior to diagnosis | 181 to 366 days prior to diagnosis |
|---------------|-----------------|--------------------------------|---------------------------------|---------------------------------|---------------------------------|
|               | Not in DKA | DKA | OR (95%CI) | P value | Not in DKA | DKA | OR (95%CI) | P value | Not in DKA | DKA | OR (95%CI) | P value | Not in DKA | DKA | OR (95%CI) | P value |
| Abdomen       | 12 (1.1) | 3 (1.2) | 1.0 (0.3-3.7) | 0.947 | 11 (1.1) | 2 (<1) | – | – | 8 (0.7) | 0 (<1) | – | – | 15 (1.4) | 0 (<1) | – | – | 33 (3) | 0 (<1) | – | – |
| Antibiotics   | 12 (1.1) | 5 (1.9) | 1.8 (0.6-5.0) | 0.296 | 85 (7.8) | 37 (14.2) | 1.9 (1.3-2.9) | 0.001 | 105 (9.7) | 13 (5.0) | 0.5 (0.3-0.9) | 0.199 | 132 (12.2) | 19 (7.3) | 0.6 (0.3-0.9) | 0.028 | 243 (22.4) | 57 (21.9) | 1.0 (0.7-1.4) | 0.869 |
| Asthma meds   | 8 (0.7) | 0 (0) | – | – | 42 (3.9) | 13 (5) | 1.3 (0.7-2.5) | 0.410 | 79 (7.3) | 16 (6.2) | 0.8 (0.5-1.1) | 0.524 | 103 (9.5) | 23 (8.8) | 0.78 (0.5-1.2) | 0.247 | 168 (15.5) | 30 (11.9) | 0.7 (0.5-1.1) | 0.108 |
| Blood tests   | 216 (19.9) | 32 (12.3) | 0.6 (0.4-0.8) | 0.005 | 108 (10.0) | 12 (4.9) | 0.4 (0.2-0.8) | 0.008 | 9 (0.8) | 3 (1.2) | 1.4 (0.4-5.2) | 0.619 | 21 (1.9) | 2 (<1) | – | – | 33 (3) | 8 (3.1) | 0.4 (0.1-1.7) | 0.209 |
| Constipation  | 2 (0.2) | 1 (1%) | – | – | 8 (0.7) | 3 (1.2) | 1.0 (0.3-3.7) | 0.947 | 11 (1.1) | 5 (1.9) | 0.6 (0.2-1.8) | 0.401 | 15 (1.4) | 6 (2.3) | 0.4 (0.2-0.8) | 0.210 | 32 (12.3) | 6 (2.3) | 0.4 (0.2-0.8) | 0.108 |
| Fungal        | 15 (1.4) | 1 (<1) | – | – | 35 (3.2) | 14 (5.4) | 1.7 (0.9-3.2) | 0.099 | 19 (1.8) | 6 (2.3) | 1.3 (0.5-3.4) | 0.552 | 18 (1.7) | 1 (<1) | – | – | 33 (3) | 6 (2.3) | 0.2 (0.1-1.7) | 0.152 |
| Headache      | 0 (0) | 1 (<1) | – | – | 6 (0.6) | 2 (<1) | – | – | 7 (0.6) | 1 (<1) | – | – | 6 (0.6) | 2 (<1) | – | – | 12 (1.1) | 0 (0) | – | – |
| ITI           | 17 (1.6) | 16 (6.2) | 4.1 (2.1-8.3) | <0.001 | 70 (6.5) | 30 (11.5) | 1.9 (1.2-3.0) | 0.006 | 83 (7.6) | 19 (7.3) | 1.0 (0.6-1.6) | 0.852 | 97 (8.9) | 24 (9.2) | 1.0 (0.7-1.7) | 0.883 | 226 (20.8) | 56 (21.9) | 1.0 (0.7-1.7) | 0.883 |
| Thirst        | 133 (12.3) | 11 (4.2) | 0.3 (0.2-0.6) | <0.001 | 40 (3.7) | 8 (3.1) | 0.8 (0.4-1.8) | 0.635 | 0 (0) | 0 (0) | – | – | 0 (0) | 0 (0) | – | – | 0 (0) | 0 (0) | – | – |
| Tiredness     | 11 (1.0) | 3 (1.2) | – | – | 8 (0.7) | 3 (1.2) | 1.6 (0.4-6.0) | 0.507 | 1 (0.1) | 0 (0) | – | – | 2 (0.2) | 0 (0) | – | – | 4 (0.4) | 1 (<1) | – | – |
| Urinary       | 278 (25.6) | 42 (16.2) | 0.6 (0.4-0.8) | 0.001 | 124 (11.4) | 9 (3.5) | 0.3 (0.1-0.6) | <0.001 | 32 (2.9) | 5 (1.9) | 0.7 (0.3-1.7) | 0.367 | 40 (3.7) | 1 (<1) | – | – | 66 (6.1) | 6 (2.3) | 0.4 (0.2-0.9) | 0.200 |
| Vomiting,nausea | 8 (0.7) | 16 (6.2) | 8.8 (3.7-20.9) | <0.001 | 9 (0.8) | 9 (3.5) | 4.3 (1.7-10.9) | 0.002 | 5 (0.5) | 0 (0) | – | – | 3 (0.3) | 2 (<1) | – | – | 10 (0.9) | 3 (1.2) | 2.8 (0.5-16.8) | 0.261 |
| Weight        | 94 (8.7) | 24 (9.2) | 1.1 (0.7-1.7) | 0.772 | 61 (5.6) | 11 (4.2) | 0.7 (0.4-1.4) | 0.372 | 21 (1.9) | 1 (<1) | – | – | 26 (2.4) | 4 (1.5) | 0.6 (0.2-1.8) | 0.404 | 52 (4.8) | 9 (3.5) | 0.6 (0.2-1.8) | 0.404 |

4.3 Implications for future research

Our results suggest that for some children there may be a prolonged period of illness prior to diagnosis. This means that we were unable to calculate the odds ratios and include these important characteristics (blood tests, antibiotics, etc.) in the models of the symptoms of T1D. This is especially true for children who presented in DKA and those that did not. Children who presented in DKA, were more likely to have had a consultation relating to RTIs, urinary tract infections and disorders, or gastrointestinal symptoms/diagnoses. This might be explained by the fact that in the Canadian study, almost 50% of the children did not see their physician within a month of diagnosis. In addition, in the Canadian study, only 50% of the children were diagnosed with a RTI, urinary tract infections and disorders, or gastrointestinal disorder in the 4 weeks prior to diagnosis of T1D. 21 In addition, in the Canadian study, almost 50% of the children did not see their physician within a month of diagnosis.

Another surprising result from our study is that boys were more likely to present in DKA. In contrast to other studies which found no gender differences. 33,34 This might be explained by the fact that in primary care, boys are less likely to seek clinical support when experiencing unusual symptoms. This means that we were unable to calculate the odds ratios and include these important characteristics (blood tests, antibiotics, etc.) in the models of the symptoms of T1D. This is especially true for children who presented in DKA and those that did not. Children who presented in DKA, were more likely to have had a consultation relating to RTIs, urinary tract infections and disorders, or gastrointestinal symptoms/diagnoses.

Overall our results suggest that for some children there may be a prolonged period prior to diagnosis. This means that we were unable to calculate the odds ratios and include these important characteristics (blood tests, antibiotics, etc.) in the models of the symptoms of T1D. This is especially true for children who presented in DKA and those that did not. Children who presented in DKA, were more likely to have had a consultation relating to RTIs, urinary tract infections and disorders, or gastrointestinal symptoms/diagnoses. This might be explained by the fact that in the Canadian study, almost 50% of the children did not see their physician within a month of diagnosis. In addition, in the Canadian study, almost 50% of the children did not see their physician within a month of diagnosis.
thirst, tiredness, urinary conditions, and vomiting/nausea) in the modeling.

5 | CONCLUSION

We conclude that there are opportunities in primary care for an earlier diagnosis of T1D in children, given their presentation with medical contacts relating to abdomen symptoms, antibiotic prescriptions, fungal conditions, RTIs, weight, urinary conditions, vomiting, and blood tests, up to 6 months prior to diagnosis. Those with primary care contacts relating to RTIs, vomiting, antibiotic prescriptions, and in particular boys, are more likely to present with DKA. This information can now be used to create a diagnostic tool for primary care physicians to help predict which children are more likely to develop T1D, thus preventing presentation in DKA.

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CONFLICTS OF INTEREST

The authors declare that no conflicts of interest associated with this manuscript.

Author contributions

All authors designed the study. D.T. conducted the matching and linking of the dataset within the SAIL databank. J.T., R.C.-J., and D.T. had full access to all the data in the study. J.T. and R.C.-J. analyzed the data. J.T. wrote the first draft of the report. All authors contributed to revision of the report and agreed to its publication. J.G. together with others established the Brecon Group Register. J.T. is the guarantor of this work, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the analysis.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of this article.

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