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Trends and socioeconomic disparities in diabetes prevalence and quality of care among Israeli children; 2011–2018

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Abstract

Background: Despite Israel's universal health coverage, disparities in health services provision may still exist. We aimed to assess socioeconomic disparities in diabetes prevalence and quality of care among Israeli children, and to assess whether these changed over time.

Methods: We used repeated cross-sectional analyses in the setting of the National Program for Quality Indicators in Community Healthcare. The data were derived from electronic medical records from Israel's four health maintenance organizations. The study population included all Israeli children aged 2–17 years in 2011–2018 (2018: $N = 2,404,856$). Socio-economic position (SEP) was measured using Central Bureau of Statistics data further updated by a private company (Points Business Mapping Ltd), and grouped into 4 categories, ranging from 1 (lowest) to 4 (highest). We used logistic regression to assess the association of SEP with diabetes prevalence, diabetes clinic visits, hemoglobin A1C (HbA1C) testing, and poor glycemic control ($HbA1c > 9\%$), and assessed whether these changed over time.

Results: Diabetes prevalence increased with age and SEP, with a total of 3019 children with diabetes. SEP was positively associated with visiting a specialized diabetes clinic (age and sex adjusted Odds Ratio ($aOR_{SEP\ 4\ vs.\ 1}$) 2.45, 95% Confidence Interval (CI) 1.67–3.69)). Although children in higher SEPs were less likely to undergo HbA1c testing ($aOR_{SEP\ 4\ vs.\ 1}$ 0.54, 95% CI 0.40–0.72), they were also less likely to have poor glycemic control ($aOR_{SEP\ 4\ vs.\ 1}$ 0.25, 95% CI 0.18–0.34). Disparities were especially apparent among children aged 2–9 (6.5% poor glycemic control in SEP 4 vs. 38.2% in SEP 1). Poor glycemic control decreased over time, from 44.0% in 2011 to 34.1% in 2018.

Conclusions: While poor glycemic control rates among children have improved, they remain high compared to rates in adults. Additionally, substantial socioeconomic gaps remain. It is eminent to study the causes of these disparities and develop policies to improve care provided to children in the lower SEP levels, to promote health equity.

Keywords: Socioeconomic disparities, Pediatric diabetes, Diabetes prevalence, Quality of care

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Introduction

Several multi-national registries have documented increasing rates of type 1 diabetes mellitus (T1DM) in children [1, 2]. Most European countries have shown an increase in pediatric T1DM incidence rates [3]. However, some countries with high incidence rates, including Sweden, Norway and Finland, have seen a plateau in incidence rates [4–6]. In addition, large variability exists in T1DM prevalence and incidence between countries [2].

Good glycemic control is crucial in children with T1DM, as this reduces the risk of long-term complications, including end-stage renal disease, blindness, and lower-limb amputation [7]. Poor glycemic control has been reported to be associated with lower socioeconomic position (SEP) in studies from the US and Europe [7–9]. It is unclear whether this association stems from problems in access to care, or from other factors associated with low SEP.

Routine care for the management of diabetes in children and adolescents differs from the care provided to adults with diabetes, since the epidemiology, pathophysiology, developmental considerations, and response to therapy in childhood diabetes are different from diabetes in adults [10]. Pediatric diabetes care should be provided by a multidisciplinary team trained in childhood diabetes management. This care should include diabetes self-management education, psycho-social therapy and medical nutrition [10].

In Israel, all residents have universal health coverage. A previous Israeli study found that in adults lower socioeconomic position was associated with poor glycemic control [11]. An additional study found higher rates of poor glycemic control in Arab children compared to Jewish children [12]. However, it is unclear whether there is an association between SEP and diabetes care on a population-level, in a country with health care for all. We therefore aimed to assess socioeconomic disparities in diabetes prevalence and quality of care among children in Israel, and to assess whether diabetes prevalence and quality of care changed over time.

Methods

All Israeli residents are entitled to a standardized basket of medical services, which are provided by four health maintenance organizations (HMOs), under the National Health Insurance Law. All residents must be associated with an HMO of their choice, and are free to transfer between HMOs [13]. The Israeli Quality Indicators in Community Healthcare (QICH) program was created to monitor and evaluate the quality of services provided by the four HMOs. The QICH program was initiated in 2002 and was adopted as a national program in 2004 [14]. Our study used QICH data to assess the prevalence

of diabetes among Israeli children and the quality of pediatric diabetes care.

Population

The study population included all Israeli residents aged 2–17 years in 2011–2018. The data includes children with complete membership for a given year. Children who switched from one HMO to another during the calendar year, those deceased during the year, and those residing abroad for more than two years were not included. Transfers between HMOs were infrequent, with less than 2% of the population transferring annually, [15] and the other categories were negligible. Information regarding diabetes prevalence was available for children ages 2–17. In total, 2,404,856 children were included for 2018, of which 3019 were children with diabetes.

Data

The QICH program received anonymized and aggregated data for 2011–2018, originating from patient electronic medical records from the four HMOs. Our study consisted of physician visits, visits to pediatric diabetes centers, and laboratory testing. We used four indicators regarding childhood diabetes. The first assessed the prevalence of diabetes among children aged 2–17 years. Children were considered to have diabetes in a specific calendar year if they purchased at least three prescriptions for insulin medications in different months, in the previous year. According to the latest report of Israel's National Registry of Insulin-Dependent (Type 1) Diabetes in ages 0–17, 92.5% of new diabetes cases in children were type 1, and only 5% were type 2 [16]. The national registry defines new cases of diabetes, according to the date of the first insulin injection. In addition, most Israeli children with type 2 diabetes are not treated with insulin. (personal communication) Therefore, our definition of diabetes refers to type 1 diabetes. The following variables assessed the quality of care provided to children with diabetes, and were only assessed among children who were defined as having diabetes in the previous measure. The first assessed the proportion of children with diabetes who visited a pediatric diabetes clinic at least once in the past year. The two additional indicators pertain to hemoglobin A1c (HbA1c) testing. We assessed the proportion of children with diabetes that were tested for HbA1c, as well as the proportion of children who had poor glycemic control (HbA1c > 9%).

Socioeconomic position (SEP)

SEP was established using small statistical areas (SSA) from the 2008 Israeli census. SSAs include 3000–4000 people, and are created in a way that maintains homogeneity in terms of the socio-demographic distribution of the population. The SSAs were categorized into 20

groups, ranging from 1 (lowest) -20 (highest) [17]. Israel's central bureau of statistics (CBS) used information regarding housing conditions, education levels, daily functioning, employment, and household income to obtain a measure of SEP for each SSA. As the most recent available data from the CBS was from 2008, we used data updated by the POINTS Location Intelligence Company to improve the timeliness of the SEP measure [18]. The POINTS Company used more recent information on education, income, living conditions, and demographic data to update the SSAs received from the census data, and grouped them into 10 SEP categories, ranging from 1 (lowest) to 10 (highest). We furthered grouped these into 4 categories, for ease of reporting, as follows: SEP 1–3: I, 4–5: II, 6–7: III, 8–10: IV. SEP was available from 2014.

Statistical analysis

We report the age and socio-economic distribution of children and adolescents with diabetes as well as the general population of children and adolescents aged 2–17. Data regarding children in the general population included children aged 2–19, due to limitations in data availability. We report the prevalence of diabetes, diabetes clinic visits, HbA1c testing and HbA1c over 9%, stratified by age and SEP in 2018. We used univariable logistic regression to assess whether the association between prevalence of diabetes, diabetes clinic visits, HbA1c testing and HbA1c over 9%, and SEP (as a continuous variable) displayed a linear trend.

We used multivariable logistic regression to assess the association between SEP and diabetes prevalence, diabetes clinic visits, HbA1c testing and HbA1c over 9%, adjusting for age, and sex. Finally, we evaluated whether prevalence of diabetes, diabetes clinic visits, HbA1c testing and HbA1c over 9% changed over time, overall, stratified by age group and stratified by SEP, using logistic regression. Trends over time stratified by SEP were restricted to 2014–2018, as SEP was only available from 2014. We based statistical significance on a *p*-value of 0.05, and reported 95% confidence intervals. Analysis was performed using Stata V.14.0 (StataCorp, College Station, Texas, USA).

Results

Among children with diabetes, nearly 80% were 10 years of age or older, while only 2.5% were between ages 2 and 4 (Table 1). The distribution of SEP was similar among children with diabetes and in the general population. SEP was missing for 4.9% of children with diabetes, and 4.7% of children in the general population.

Diabetes prevalence increased as SEP increased, among children aged 15 and older, while there was no statistically significant increase among younger children

Table 1 Distribution of age, sex and socio-economic position among 2–17-year-old children with diabetes, and the general population, in Israel, 2018

	Children with Diabetes		General Population	
	N	%	N	%
Age				
2–4	77	2.6	504,819	21.0
5–9	538	17.8	789,602	32.8
10–14	1294	42.9	719,259	29.9
15–17*	1110	36.8	391,176	16.3
Sex				
Male	1522	50.4	1,234,005	51.3
Female	1497	49.6	1,170,851	48.7
SEP**				
1	750	24.8	640,800	26.6
2	850	28.2	651,256	27.1
3	812	26.9	666,441	27.7
4	458	15.2	333,508	13.9
Total	3019		2,404,856	

* Among the general population this included children ages 15–19

** SEP – socio economic position was missing for 149 children with diabetes (4.9%), and 112,851 children in the general population (4.7%)

(Table 2). The prevalence of diabetes clinic visits declined with age, while the prevalence of diabetes clinic visits increased as SEP increased (linear trend OR: 1.28; 95% CI 1.15–1.43). HbA1c testing was highest among children in the lowest SEP, for all age groups, and decreased as SEP increased (overall linear trend OR: 0.76; 95% CI:0.70–0.82). We found substantial SEP disparities in children with HbA1c over 9%. While only 6.5% of children aged 2–9 years in SEP 4 had poor glycemic control, 38.1% of children in SEP 1 had HbA1c over 9%. In children aged 10–14 and 15–17, the disparities were less dramatic, but the prevalence of poor glycemic control in children in SEP 1 remained over twice that in SEP 4 (53.3 vs. 19.1 and 41.8 vs. 14.4 respectively) (linear trend across all SEP OR: 0.59; 95% CI 0.54–0.65).

Controlling for age and sex, diabetes prevalence was positively associated with SEP (Table 3). Children in the highest SEP (SEP 4) had higher odds of visiting a specialized diabetes clinic compared to children in the lowest SEP (SEP 1)(aOR_{SEP 4 vs 1} 2.45, 95% CI 1.67–3.69). Children in the highest SEP had lower odds of undergoing HbA1c testing compared to those in the lowest SEP (AOR_{SEP 4 vs 1} 0.45, 95% CI 0.34–0.59). However, among those who were tested, the odds of having poor glycemic control (HbA1C > 9%) was significantly lower in the highest SEP compared to the lowest SEP (AOR_{SEP 4 vs 1} 0.20, 95% CI 0.14–0.28).

Diabetes prevalence rates remained stable over time among children aged 14 and younger, however there

Table 2 Prevalence of diabetes, diabetes clinic visits, HbA1c testing and HbA1c over 9% by SEP and age, 2018

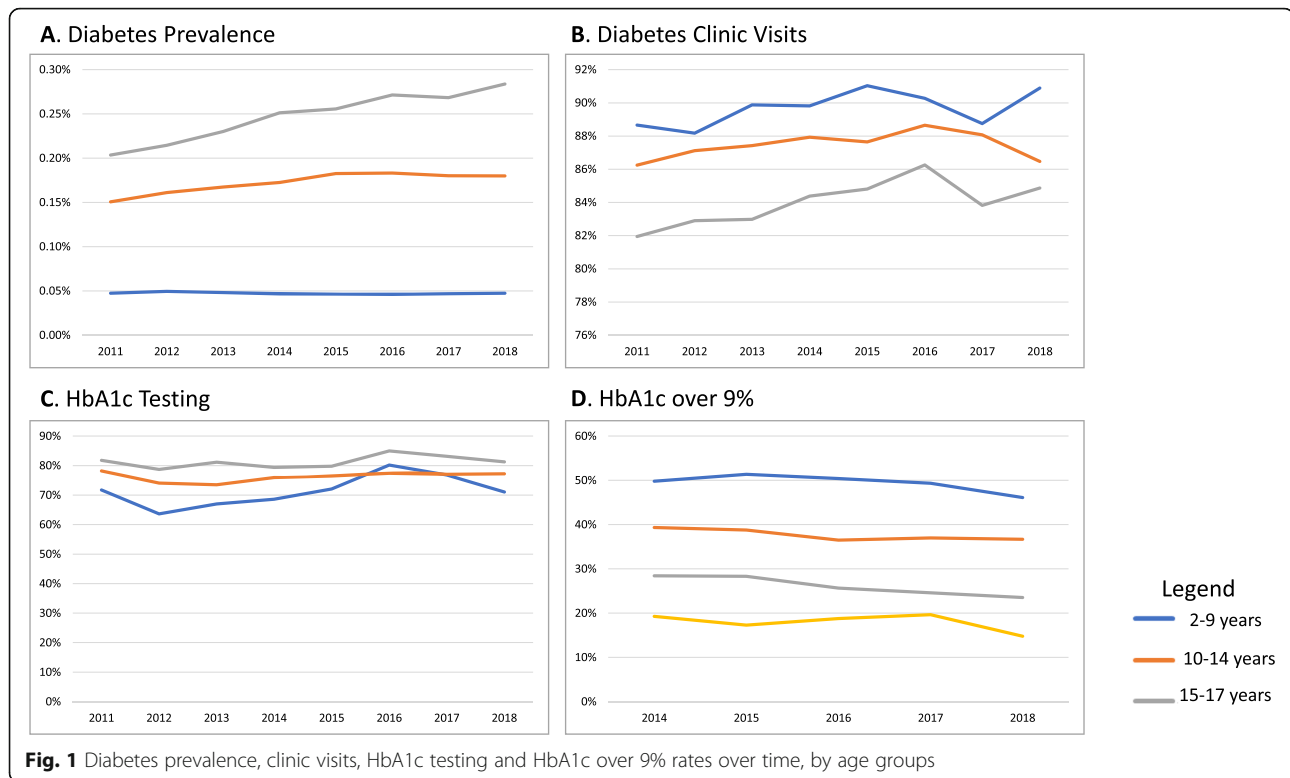
	Total	SEP 1 %	SEP 2 %	SEP 3 %	SEP 4 %	OR* (linear trend)	P-value (trend)
Diabetes Prevalence							
Age groups							
2–9	0.05	0.04	0.05	0.04	0.06	1.05	0.198
10–14	0.18	0.17	0.18	0.18	0.18	1.02	0.409
15–17	0.28	0.24	0.30	0.29	0.32	1.04	0.004
Total (n = 2,267,785)	0.13	0.12	0.13	0.12	0.14	1.04	0.045
Diabetes clinic visits							
Age groups							
2–9	90.89	91.10	89.27	91.19	92.16	1.06	0.674
10–14	86.48	83.48	87.39	87.47	90.67	1.21	0.023
15–17	84.86	78.23	84.26	85.71	94.48	1.47	< 0.001
Total (n = 3019)	86.78	83.07	86.59	87.56	92.36	1.28	< 0.001
HbA1c testing							
Age groups							
2–9	71.06	80.82	70.06	65.41	60.78	0.73	< 0.001
10–14	77.20	81.08	81.09	72.70	67.88	0.77	< 0.001
15–17	81.26	85.61	83.02	79.59	72.39	0.77	< 0.001
Total (n = 3019)	77.44	82.67	79.53	73.77	67.90	0.76	< 0.001
HbA1c over 9%							
Age groups							
2–9	24.49	38.14	26.61	11.54	6.45	0.48	< 0.001
10–14	38.94	53.33	40.99	26.05	19.08	0.57	< 0.001
15–17	33.26	41.81	36.80	26.07	14.41	0.65	< 0.001
Total (n = 2338)	34.05	46.13	36.69	23.54	14.79	0.59	< 0.001

*We used univariable logistic regression to assess whether the association between prevalence of diabetes, diabetes clinic visits, HbA1c testing and HbA1c over 9%, and SEP (as a continuous variable) displayed a linear trend. OR reflects a one-unit change in SEP. Results in bold indicate statistical significance at the 0.05 level

Table 3 Multivariable models assessing odds of diabetes prevalence, diabetes clinic visits, HbA1c testing and HbA1c > 9%, among Israeli children aged 2–17, 2018

	Diabetes Prevalence			Diabetes clinic visits			HbA1c testing			HbA1c over 9%		
	OR ¹	95% CI	p-value	OR ¹	95% CI	P-value	OR ¹	95% CI	p-value	OR ¹	95% CI	p-value
Age												
2–9	Ref			Ref			Ref			Ref		
10–14	3.79	3.44–4.18	< 0.001	0.68	0.49–0.94	0.020	1.41	1.13–1.76	0.003	2.08	1.59–2.74	< 0.001
15–19	5.98	5.40–6.62	< 0.001	0.56	0.40–0.78	< 0.001	1.85	1.46–2.35	< 0.001	1.64	1.24–2.17	< 0.001
Sex												
Male	Ref			Ref						Ref		
Female	1.05	0.97–1.13	0.222	0.88	0.71–1.10	0.270	1.09	0.91–1.30	0.357	1.08	0.90–1.30	0.432
SEP												
1	Ref			Ref			Ref			Ref		
2	1.15	1.04–1.27	0.005	1.32	1.00–1.73	0.050	0.81	0.63–1.05	0.11	0.67	0.54–0.84	< 0.001
3	1.10	1.00–1.22	0.057	1.44	1.08–1.91	0.012	0.59	0.46–0.75	< 0.001	0.35	0.27–0.45	< 0.001
4	1.21	1.08–1.36	0.001	2.45	1.67–3.69	< 0.001	0.45	0.34–0.59	< 0.001	0.20	0.14–0.28	< 0.001

¹Models adjusted for age, sex, and SEP. Results in bold indicate statistical significance at the 0.05 level



was a slight increase among children aged 15–17 ($P_{\text{trend}} < 0.001$) (Fig. 1a). There was no statistically significant change in the overall prevalence of diabetes clinic visits ($P_{\text{trend}} = 0.51$) (Fig. 1b). HbA1c testing increased slightly over time ($P_{\text{trend}} = 0.02$) (Fig. 1c). We found a substantial reduction in the prevalence of children with HbA1c over 9% over time, across all age groups ($P_{\text{trend}} < 0.001$) (Fig. 1d). Throughout the entire time period we examined, children aged 10–14 had the highest prevalence of poor glycemic control.

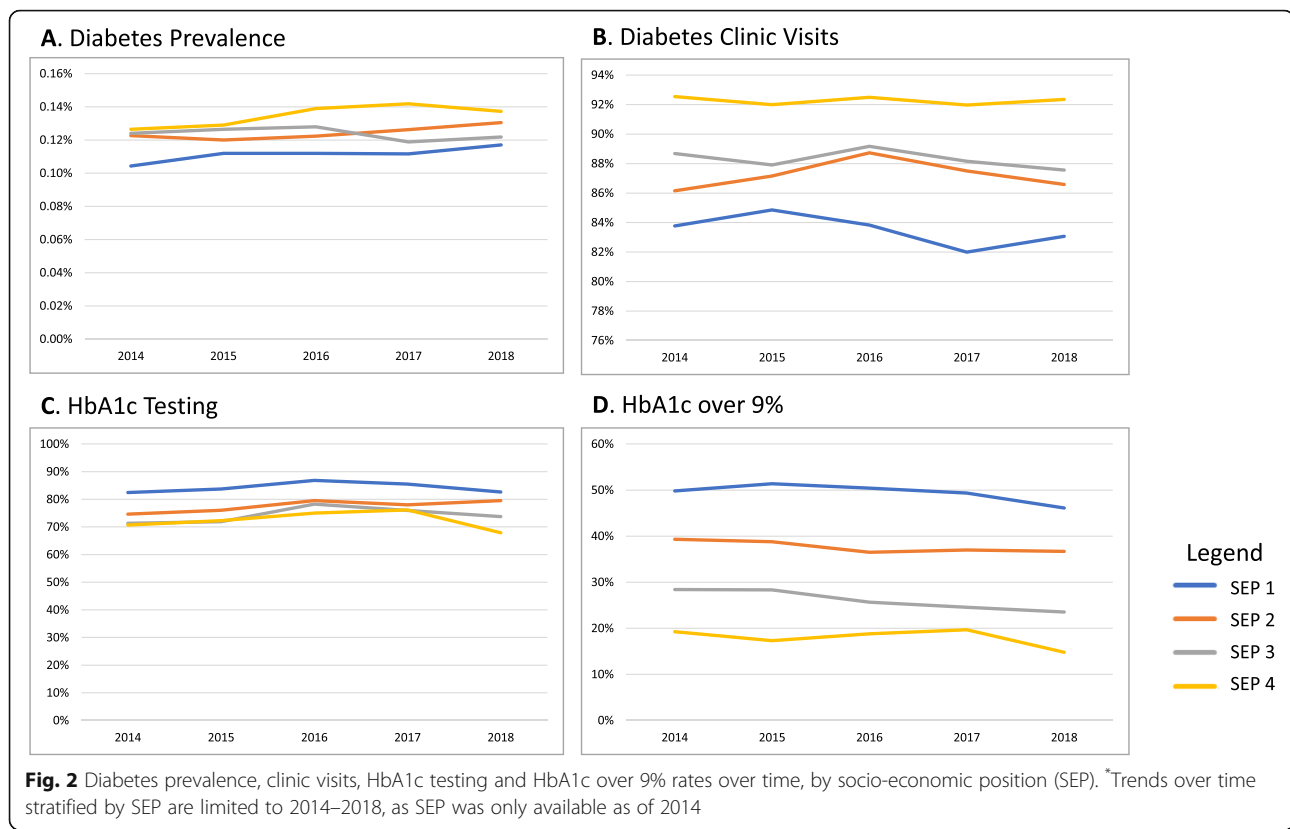
We also assessed trends over time by socio-economic position. As SEP was available as of 2014, these trends are available for 2014–2018. When we assessed trends over time by socio-economic position, we found that the changes in diabetes prevalence were not statistically significant for any of the SEP categories except for a slight increase in the lowest SEP ($P_{\text{trend}} = 0.05$) (Fig. 2a). There were no statistically significant changes in diabetes clinic visits prevalence in any SEP categories (Fig. 2b). HbA1c testing increased in all SEPs, except for SEP 4, in which it decreased. However, the change was only statistically significant in SEP 2 ($P_{\text{trend}} = 0.01$) (Fig. 2c). HbA1c decreased across all SEP categories, however the decrease was only statistically significant in children from SEP 3 ($P_{\text{trend}} = 0.02$) (Fig. 2d).

Discussion

We found substantial socioeconomic disparities in the odds of HbA1c testing as well as the odds of having poor glycemic control. While children in higher socioeconomic positions were less likely to be tested, they were considerably less likely to have poor glycemic control. Diabetes prevalence has increased slightly over time among older children, alongside an overall increase in HbA1c testing, and an overall reduction in rates of poor glycemic control. Despite the improvement in the rate of poor glycemic control, the rate remains substantially higher than the rate of uncontrolled diabetes in adults which is around 10% [18].

Our findings regarding an increased rate of poor glycemic control among children aged 10–14, are similar to those found in other studies [19, 20]. Several reasons have been suggested for the increased rate of poor glycemic control during puberty, including a reduction in insulin-stimulated glucose metabolism among pubertal children, as well as behavioral changes, such as parents becoming less involved in the diabetes management of their children during puberty [19].

Low socioeconomic status is associated with poor glycemic control in children with T1DM, even among children with health insurance [7–9, 21, 22]. Several reasons have been suggested for this association, including the burden of out-of-pockets expenses that occur even among insured individuals, co-payments and lost wages,



as well as additional barriers in the access to specialist care [22]. In addition lower SEP is associated with poorer self-management behaviors which may lead to poor glycemic control [23]. In Israel, there is universal health coverage under the National Health Insurance (NHI) law, and all residents are entitled to the same access to care [13]. However, even in countries that provide universal coverage, the use of health-services differs between people with less income and education compared to those who are wealthier and better educated [24]. Our findings indicate that while children from lower SEPs are less likely to visit a specialized pediatric diabetes clinic, they are more likely to have their HbA1c levels monitored. Therefore, it is unlikely that the poor glycemic control that we found among children in the lowest SEP is due to a barrier in the access to care. It is time-consuming and difficult to obtain good glycemic control in children with diabetes, and families with financial burdens may not be able to dedicate the time and resources necessary to do so. In addition, although all residents are entitled to care under the National Health Insurance Law, specialized pediatric diabetes clinics are not distributed uniformly throughout Israel [25]. Children in peripheral areas may face substantial additional burdens when trying to access these clinics. Programs that include additional support for families from lower SEPs, may be helpful in improving the rate

of good glycemic control [20]. Such support may include frequent contact from both medical and nursing staff, allowing rapid troubleshooting when issues arise regarding changes in insulin regimen.

The T1DM prevalence rate found in our study is slightly lower than the most recently reported prevalence rates in countries such as the Netherlands (0.17%), the UK (0.20%), and the US (0.19%) [26–28]. However, the ascertainment of T1DM varies in different studies. Increasing trends in the incidence and prevalence of T1DM have been reported [26, 27, 29]. In recent years, several countries have reported a halt in this increase [4–6]. Our study found a slow increase from 0.10% in 2011 to 0.12% in 2014, no change in T1DM prevalence through 2017, and an additional increase to 0.13% in 2018. Data from Israel's National Registry of Incident T1DM cases from 2015 indicate that there was a continuous increase in the incidence of childhood T1DM cases [16]. Longer follow up time is necessary to better understand whether our results indicate the beginning of a plateau or a continued albeit slow increase.

Our study has several strengths. First, the data in our study includes all Israeli children aged 2–17 years between 2011 and 2018, and as such our study was not prone to selection biases or sampling errors. The quality of the data provided by the QICH program is exceptionally high. Data are checked at three levels. First there is

an internal audit conducted by each HMO. Then the QICH program directorate performs a data audit, and finally there is an external process audit to ensure the validity of the procedures involved. Our study has several limitations which should be considered. First, we only have detailed area-level SEP information as of 2014, and we therefore only had limited follow up time to assess changes over time in the effect of SEP on diabetes prevalence and quality of care. Furthermore, we did not have information regarding additional factors that are known to influence glycemic control including duration of diabetes, ethnicity, and parental knowledge regarding diabetes. Finally, as we did not have individual-level data, we were unable to assess whether children who regularly visited a specialized diabetes clinic did in fact have better glycemic control compared to children who did not.

Based on data from multiple countries and continents – the adolescent period is characterized, in general, by poor metabolic control in patients with T1DM [19, 20, 22, 23]. Our data allows the identification of those who succeed in managing their disease well during this vulnerable period – in order to highlight the unique qualities typical of these patients. Gaining such insights may allow the identification of tools to promote better diabetes care for adolescents. Various factors may contribute to the increased rate of poor glycemic control among children from low SEP, including financial barriers to care, limited health education, and poor family communication skills. In order to further understand the causes of these disparities, additional research, using individual level data, is necessary to assess the association between specialized clinics visits and glycemic control, the associations between SEP and insulin pump therapy, and the association of the latter with clinical outcomes. These may be helpful in the planning and implementation of targeted interventions, that will close the gaps in T1DM care in children.

Additionally, in order to get a better understanding of the factors behind these disparities, qualitative research can be used. Focus groups with children with T1DM and their parents, physicians, and other caregivers will enable deepening the understanding regarding barriers to high quality care. Potential barriers include access barriers (such as taking time off work for appointments, cost and complexity of transportation to appointments), language and cultural barriers, as well as health literacy and family-provider communication concerns.

The policy recommendations may vary depending on the outcome of the above-mentioned research. If a lack of family communication is found to play a major role, interventions aimed at improving parent-adolescent teamwork in diabetes management tasks have been found to be beneficial in improving glycemic control while reducing family conflict [30, 31]. Some of these interventions can be integrated into routine diabetes clinic visits. In order to

improve family-provider communication, family-centered rounds may improve communication and shared decision-making between physicians and families [32]. Family-centered care is defined as interdisciplinary care in which professionals from a variety of disciplines work collaboratively to develop a unified care plan, with active participation from patients and their families [33]. If access barriers are found to be substantial, financial incentives for both parents and youth to promote adherence to intervention programs were found to substantially improve retention [31]. Small sums provided frequently were found to be most effective [34]. Finally, incentives to HMO's for improving health outcomes, may yield innovative strategies for screening and identifying T1DM patients who are at high risk for nonadherence and adopting personalized interventions which will improve their health outcomes.

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Authors' contributions

YLW and OM Conceived and designed the study; YLW MK RCM Collected the data; YLW MK RCM and OM Contributed data or analysis tools; YLW Performed the analysis; YLW Wrote the paper, with considerate input from MK RCM and OM. All authors read and approved the final manuscript.

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Availability of data and materials

The datasets analyzed during the current study are not publicly available due to legal restrictions but are available from the corresponding author on reasonable request.

Ethics approval and consent to participate

Not Applicable.

Consent for publication

Not Applicable.

Competing interests

The authors declare that they have no competing interests.

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References

1. Incidence and trends of childhood Type 1 diabetes worldwide 1990–1999. *Diabet Med.* 2006;23(8):857–66 Available from: <http://doi.wiley.com/10.1111/j.1464-5491.2006.01925.x> [cited 2019 26 Mar].
2. Patterson C, Guariguata L, Dahlquist G, Soltész G, Ogle G, Silink M. Diabetes in the young – a global view and worldwide estimates of numbers of children with type 1 diabetes. *Diabetes Res Clin Pract.* 2014;103(2):161–75.
3. Patterson CC, Harjutsalo V, Rosenbauer J, Neu A, Cinek O, Skrivarhaug T, et al. Trends and cyclical variation in the incidence of childhood type 1 diabetes in 26 European centres in the 25 year period 1989–2013: a multicentre prospective registration study. *Diabetologia.* 2019;62(3):408–17.
4. Berhan Y, Waernbaum I, Lind T, Möllsten A, Dahlquist G. Group* for the SCDS. Thirty Years of Prospective Nationwide Incidence of Childhood Type

- 1 Diabetes. *Diabetes*. 2011;60(2):577–81 Available from: <http://www.ncbi.nlm.nih.gov/pubmed/21270269> [cited 2019 26 Mar].
5. Skrivvarhaug T, Stene LC, Drivvoll AK, Strøm H, Joner G. Incidence of type 1 diabetes in Norway among children aged 0–14 years between 1989 and 2012: has the incidence stopped rising? Results from the Norwegian childhood diabetes registry. *Diabetologia*. 2014;57(1):57–62.
6. Harjutsalo V, Sund R, Knip M, Groop P-H. Incidence of Type 1 Diabetes in Finland. *JAMA*. 2013;310(4):427 Available from: <http://jama.jamanetwork.com/article.aspx?doi=10.1001/jama.2013.8399> [cited 2019 26 Mar].
7. Gallegos-Macias AR, Macias SR, Kaufman E, Skipper B, Kalishman N. Relationship between glycemic control, ethnicity and socioeconomic status in Hispanic and white non-Hispanic youths with type 1 diabetes mellitus. *Pediatr Diab*. 2003;4(1):19–23 Available from: <http://doi.wiley.com/10.1034/j.1399-5448.2003.00020.x> [cited 2019 24 Mar].
8. Tahirovic H, Toromanovic A. Glycemic control in diabetic children: role of mother's knowledge and socioeconomic status. *Eur J Pediatr*. 2010;169(8):961–4 Available from: <http://link.springer.com/10.1007/s00431-010-1156-0> [cited 2019 25 Mar].
9. Galler A, Lindau M, Erner A, Thalemann R, Raile K. Associations between media consumption habits, physical activity, socioeconomic status, and glycemic control in children, adolescents, and young adults with type 1 diabetes. *Diab Care*. 2011;34(11):2356–9 Available from: <http://www.ncbi.nlm.nih.gov/pubmed/21926289> [cited 2019 25 Mar].
10. Association AD. 13. Children and Adolescents: Standards of Medical Care in Diabetes-2020. Vol. 43, *Diabetes care*; 2020. p. S163–S182.
11. Wilf-Miron R, Peled R, Yaari E, Shem-Tov O, Weinner VA, Porath A, et al. Disparities in diabetes care: role of the patient's socio-demographic characteristics. *BMC Public Health* [Internet]. 2010;10(1):729. Available from: <http://bmcpublichealth.biomedcentral.com/articles/10.1186/1471-2458-10-729> [cited 2019 24 Feb].
12. Zuckerman-Levin N, Dabaja-Younis H, Ameer E, Cohen M, Maor Y, Shehadeh N. Effect of socioeconomic status and ethnicity on glycemic control in Arab and Jewish youth with type 1 diabetes mellitus. *Rambam Maimonides Med J*. 2018 Oct 4;9(4):e0030.
13. Gross R, Rosen B, Shirom A. Reforming the Israeli health system: findings of a 3-year evaluation. *Health Policy*. 2001;56(1):1–20.
14. Jaffe DH, Shmueli A, Ben-Yehuda A, Paltiel O, Calderon R, Cohen AD, et al. Community healthcare in Israel: quality indicators 2007–2009. *Isr J Health Policy Res*. 2012;1(1):3.
15. Clarfield AM, Manor O, Bin NG, Shvarts S, Azzam ZS, Afek A, et al. Health and health care in Israel: an introduction. *Lancet*. 2017;389:2503–13.
16. Israel Center for Disease Control. Israel National Registry of T1DM children aged 0–17 (Hebrew Report) 2015 [Internet]. Available from: https://www.health.gov.il/PublicationsFiles/diabetes_0-17_2015.pdf.pdf [cited 2020 22 Apr].
17. Central Bureau of Statistics Israel. Characterization and Classification of Geographical Units by the Socio-Economic Level of the Population 2008. Jerusalem; 2013 Jun. Report No.: 1530.
18. Calderon-Margalit R, Abu-Ahmed W, Ben-Yehuda A, Horwitz E, Krieger M, Manor O, et al. National Program for Quality Indicators in Community Healthcare in Israel 2017 Report [Internet]. 2017. Available from: <https://en.israelhealthindicators.org/publications> [cited 2020 22 Apr].
19. Gerstl E-M, Rabl W, Rosenbauer J, Gröbe H, Hofer SE, Krause U, et al. Metabolic control as reflectet by HbA1c in children, adolescents and young adults with type-1 diabetes mellitus: combined longitudinal analysis including 27,035 patients from 207 centers in Germany and Austria during the last decade. *Eur J Pediatr*. 2008;167(4):447–53 Available from: <http://link.springer.com/10.1007/s00431-007-0586-9> [cited 2019 25 Mar].
20. Scottish Study Group for the Care of the Young Diabetic SSG for the C of the Y. Factors influencing glycemic control in young people with type 1 diabetes in Scotland: a population-based study (DIABAUD2). *Diab Care*. 2001;24(2):239–44 Available from: <http://www.ncbi.nlm.nih.gov/pubmed/11213872> [cited 2019 31 Mar].
21. Hassan K, Loar R, Anderson BJ, Heptulla RA. The role of socioeconomic status, depression, quality of life, and glycemic control in type 1 diabetes mellitus. *J Pediatr*. 2006 Oct 1;149(4):526–31.
22. Pettiti DB, Klingensmith GJ, Bell RA, Andrews JS, Dabelea D, Imperatore G, et al. Glycemic Control in Youth with Diabetes: The SEARCH for Diabetes in Youth Study. *J Pediatr*. 2009;155(5):668–72 e3.
23. Guo J, Whittemore R, He G-P. The relationship between diabetes self-management and metabolic control in youth with type 1 diabetes: an integrative review. *J Adv Nurs*. 2011;67(11):2294–310 Available from: <http://doi.wiley.com/10.1111/j.1365-2648.2011.05697.x> [cited 2020 31 May].
24. Adler NE, Newman K. Socioeconomic Disparities In Health: Pathways And Policies. *Health Aff*. 2002;21(2):60–76 Available from: <http://www.healthaffairs.org/doi/10.1377/hlthaff.21.2.60> [cited 2019 25 Mar].
25. Childhood Diabetes Clinics in Israel. The Juvenile Diabetes Research Foundation in Israel. <https://jdrf.org.il/wp-content/uploads/2018/04/%D7%9E%D7%A8%D7%A4%D7%90%D7%95%D7%A%D7%99%D7%9C%D7%93%D7%99%D7%9D-.pdf>. Accessed 22 Apr 2020.
26. Fazeli Farsani S, Souverein PC, van der Vorst MMJ, Knibbe CAJ, Herings RMC, de Boer A, et al. Increasing trends in the incidence and prevalence rates of type 1 diabetes among children and adolescents in the Netherlands. *Pediatr Diab*. 2016;17(1):44–52 Available from: <http://doi.wiley.com/10.1111/pedi.12232> [cited 2019 24 Mar].
27. Dabelea D, Mayer-Davis EJ, Saydah S, Imperatore G, Linder B, Divers J, et al. Prevalence of Type 1 and Type 2 Diabetes Among Children and Adolescents From 2001 to 2009. *JAMA*. 2014;311(17):1778 Available from: <http://jama.jamanetwork.com/article.aspx?doi=10.1001/jama.2014.3201> [cited 2019 24 Mar].
28. Care processes and outcomes Audits RCPCH National Paediatric Diabetes Audit NPDA. 2020. Available from: https://www.rcpch.ac.uk/sites/default/files/2020-03/final_npda_core_report_2018-2019.pdf [cited 2020 26 May].
29. Mayer-Davis EJ, Lawrence JM, Dabelea D, Divers J, Isom S, Dolan L, Imperatore G, Linder B, Marcovina S, Pettitt DJ, Pihoker C. Incidence trends of type 1 and type 2 diabetes among youths, 2002–2012. *N Engl J Med*. 2017 Apr 13;376:1419–29. [cited 2019 25 Mar].
30. Anderson BJ, Brackett J, Ho J, LMB L. An office-based intervention to maintain parent-adolescent teamwork in diabetes management: Impact on parent involvement, family conflict, and subsequent glycemic control. *Diab Care*. 1999;22(5):713–21 Available from: <https://pubmed.ncbi.nlm.nih.gov/10332671/> [cited 2020 7 Jul].
31. Wysocki T, Harris MA, Buckloh LM, Mertlich D, Lochrie AS, Taylor A, et al. Randomized, Controlled Trial of Behavioral Family Systems Therapy for Diabetes: Maintenance and Generalization of Effects on Parent-Adolescent Communication. *Behav Ther*. 2008;39(1):33–46 Available from: <https://pubmed.ncbi.nlm.nih.gov/18328868/> [cited 2020 7 Jul].
32. Modi AC, Pai AL, Hommel KA, Hood KK, Cortina S, Hilliard ME, et al. Pediatric self-management: A framework for research, practice, and policy [Internet]. Vol. 129, *Pediatrics*. Am Acad Pediatr; 2012. p. e473–85. Available from: www.pediatrics.org/cgi/doi/10.1542/peds.2011-1635 [cited 2020 2 Jul].
33. Sisterhen LL, Blaszak RT, Woods MB, Smith CE. Defining family-centered rounds [Internet]. Vol. 19, *Teaching and Learning in Medicine*. Lawrence Erlbaum Assoc. 2007:319–22 Available from: <https://pubmed.ncbi.nlm.nih.gov/17594229/> [cited 2020 7 Jul].
34. Cutler DM, Everett W. Thinking outside the pillbox - Medication adherence as a priority for health care reform. Vol. 362, *New Engl J Med Massach Med Soc*; 2010. p. 1553–1555. Available from: <http://www.nejm.org/doi/abs/10.1056/NEJMp1002305> [cited 2020 7 Jul].

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