

# Area deprivation and demographic factors associated with diabetes technology use in adults with type 1 diabetes in Germany

Marie Auzanneau<sup>1, 2\*</sup>, Alexander J. Eckert<sup>1, 2</sup>, Sebastian M. Meyhöfer<sup>2, 3</sup>, Martin Heni<sup>4, 5</sup>, Anton Gillessen<sup>6</sup>, Lars Schwettmann<sup>7</sup>, Peter M. Jehle<sup>8</sup>, Michael Hummel<sup>9</sup>, Reinhard W. Holl<sup>1, 2</sup>

<sup>1</sup>Institute of Epidemiology and Medical Biometry, Faculty of Medicine, University of Ulm, Germany, <sup>2</sup>German Center for Diabetes Research (DZD), Germany, <sup>3</sup>Institute for Endocrinology and Diabetes, University of Lübeck, Germany, <sup>4</sup>Klinik für Innere Medizin I, Universitätsklinik Ulm, Germany, <sup>5</sup>Institute for Clinical Chemistry and Pathobiochemistry, University Hospital and Faculty of Medicine, University of Tübingen, Germany, <sup>6</sup>Department of Internal Medicine, Sacred Heart Hospital, Germany, <sup>7</sup>Division of Health Economics, Department of Health Services Research, Carl von Ossietzky University of Oldenburg, Germany, <sup>8</sup>Department of Internal Medicine I, Martin-Luther-University Halle-Wittenberg, University Medicine, Academic Hospital Paul-Gerhardt-Stift, Germany, <sup>9</sup>Forschergruppe Diabetes e.V., Helmholtz Center Munich, Germany



Specialty Section: Pediatric Endocrinology

Article type: Original Research Article

Manuscript ID: 1191138

Received on: 21 Mar 2023

Revised on: 16 Jun 2023

Journal website link: www.frontiersin.org



### Conflict of interest statement

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

### Author contribution statement

M.A. and R.W.H. designed the study. M.A. and A.J.E. analyzed the study data. M.A. created the figures, and wrote the manuscript. A.J.E., S.M.M., A.G., L.S., P.M.J., M.He, M.Hu. and R.W.H. contributed to the discussion, reviewed, and approved the manuscript.

### Keywords

diabetes, Technology, CGM, pump, age, gender, socioeconomic, deprivation

### Abstract

### Word count: 349

### Introduction

Diabetes technology improves glycemic control and quality of life for many people with type 1 diabetes (T1D). However, inequalities in access to diabetes technology exist in many countries. In Germany, disparities in technology use have been described in pediatric T1D, but no data for adults are available so far. We therefore aimed to analyze whether demographic factors and area deprivation are associated with technology use in a representative population of adults with T1D.

### Materials and methods

In adults with T1D from the German prospective diabetes follow-up registry (DPV), we analyzed the use of continuous subcutaneous insulin infusion (CSII), continuous glucose monitoring (CGM), and sensor augmented pump therapy (SAP, with and without automated insulin delivery) in 2019-2021 by age group, gender, migration background, and area deprivation using multiple adjusted regression models. Area deprivation, defined as a relative lack of area-based resources, was measured by quintiles of the German index of Multiple Deprivation (GIMD 2015, from Q1, least deprived, to Q5, most deprived districts).

#### Results

Among 13,351 adults with T1D, the use of technology decreased significantly with older age: CSII use fell from 56.1% in the 18–**<25-year age group to 3.1%** in the  $\ge$ 80-year age group, CGM use from 75.3% to 28.2%, and SAP use from 45.1% to 1.5% (all p for trend <0.001). The use of technology was also significantly higher in women than in men (CSII: 39.2% vs. 27.6%; CGM: 61.9% vs. 58.0%; SAP: 28.7% vs. 19.6%, all p <0.001), and in individuals without migration background than in those with migration background (CSII: 38.8% vs. 27.6%; CGM: 71.1% vs. 61.4%; SAP: 30.5% vs. 21.3%, all p <0.001). Associations with area deprivation were not linear: the use of each technology decreased only from Q2 to Q4.

#### Discussion

Our real-world data provide evidence that higher age, male gender, and migration background are currently associated with lower use of diabetes technology in adults with T1D in Germany. Associations with area deprivation are more complex, probably due to correlations with other factors, like the higher proportion of migrants in less deprived areas or the federal structure of the German health care system.

### Contribution to the field

Dear Dr. Addala, Please find enclosed our manuscript entitled "Area deprivation and demographic factors associated with diabetes technology use in adults with type 1 diabetes in Germany ". There is now strong evidence that diabetes technology improves glycemic control and quality of life for many people with type 1 diabetes (T1D). However, inequalities in access to diabetes technology have been reported in many countries. In Germany, disparities in technology use have been described in pediatric T1D, but no data for adults are available so far. Our real-world data provide evidence that higher age, male gender, and migration background are currently associated with lower use of diabetes technology in adults with T1D in Germany. There is a critical need to improve access to diabetes technology should be improved in Germany. We confirm that this manuscript has not been published elsewhere and is not under consideration by another journal. All authors have approved the final version of the manuscript and agree with its submission to Frontiers in endocrinology. Thank you for receiving our manuscript and considering it for review. We appreciate your time and look forward to your response. Kind regards Marie Auzanneau

### Funding information

The DPV registry and this analysis are supported by the German Center for Diabetes Research (DZD, grant number: 82DZD14E03). Further financial support for the DPV registry was provided by the German Diabetes Association (DDG), the German Robert Koch Institute (RKI).

### Ethics statements

Studies involving animal subjects Generated Statement: No animal studies are presented in this manuscript.

Studies involving human subjects Generated Statement: No human studies are presented in this manuscript.

*Inclusion of identifiable human data* Generated Statement: No potentially identifiable human images or data is presented in this study.

### Data availability statement

Generated Statement: The data analyzed in this study is subject to the following licenses/restrictions: Access to the programming code can be provided by the corresponding author upon request. For reasons of data protection, data on individual level cannot be provided. However, remote data analysis is possible. Requests to access these datasets should be directed to marie.auzanneau@uni-ulm.de.



# Area deprivation and demographic factors associated with diabetes technology use in adults with type 1 diabetes in Germany

- 1 Marie Auzanneau<sup>1,2</sup>, Alexander J. Eckert<sup>1,2</sup>, Sebastian M. Meyhöfer<sup>2,3</sup>, Martin Heni<sup>4,5</sup>, Anton
- 2 Gillessen<sup>6</sup>, Lars Schwettmann<sup>7</sup>, Peter M. Jehle<sup>8</sup>, Michael Hummel<sup>9</sup>, Reinhard W. Holl<sup>1,2</sup> for the
- **3 DPV initiative**
- <sup>4</sup> <sup>1</sup>Institute of Epidemiology and Medical Biometry, ZIBMT, University of Ulm, Ulm, Germany
- <sup>5</sup> <sup>2</sup> German Center for Diabetes Research (DZD), Neuherberg, Germany
- 6 <sup>3</sup> Institute for Endocrinology and Diabetes, University of Lübeck, Lübeck, Germany
- <sup>7</sup> <sup>4</sup> Division of Endocrinology and Diabetology, Department of Internal Medicine 1, University
- 8 Hospital Ulm, Ulm, Germany.
- <sup>9</sup> <sup>5</sup>Institute for Clinical Chemistry and Pathobiochemistry, Department for Diagnostic Laboratory
- 10 Medicine, University Hospital Tübingen, Tübingen, Germany
- <sup>6</sup>Department of Internal Medicine, Sacred Heart Hospital, Muenster, Germany
- <sup>7</sup> Division of Health Economics, Department of Health Services Research, Carl von Ossietzky
- 13 University of Oldenburg, Oldenburg, Germany
- <sup>8</sup>Department of Internal Medicine I, Martin-Luther-University Halle-Wittenberg, University
- 15 Medicine, Academic Hospital Paul-Gerhardt-Stift, Lutherstadt Wittenberg, Germany
- <sup>9</sup> Forschergruppe Diabetes e.V., Helmholtz Center Munich, Munich-Neuherberg, Germany
- 17
- 18 \* Correspondence:
- 19 Marie Auzanneau, MPH
- 20 Institute of Epidemiology and Medical Biometry, ZIBMT
- 21 University of Ulm
- 22 Albert-Einstein-Allee 41
- 23 D-89081 Ulm, Germany
- 24 Tel.: +49/731/5025483
- 25 Fax.: +49/ 731/5025309
- 26 marie.auzanneau@uni-ulm.de
- 27 ORCID ID: https://orcid.org/0000-0002-5906-6579

- 28 Keywords: diabetes, technology, CGM, pump, age, gender, socioeconomic, deprivation
- 29 Word count: 2,890 words. 1 table, 1 figure.
- 30

## 31 Abstract

32

## 33 Introduction

Diabetes technology improves glycemic control and quality of life for many people with type 1
diabetes (T1D). However, inequalities in access to diabetes technology exist in many countries. In
Germany, disparities in technology use have been described in pediatric T1D, but no data for adults
are available so far. We therefore aimed to analyze whether demographic factors and area deprivation
are associated with technology use in a representative population of adults with T1D.

39

## 40 Materials and methods

41 In adults with T1D from the German prospective diabetes follow-up registry (DPV), we analyzed the

- 42 use of continuous subcutaneous insulin infusion (CSII), continuous glucose monitoring (CGM), and
- 43 sensor augmented pump therapy (SAP, with and without automated insulin delivery) in 2019-2021
- 44 by age group, gender, migration background, and area deprivation using multiple adjusted regression
- 45 models. Area deprivation, defined as a relative lack of area-based resources, was measured by
- 46 quintiles of the German index of Multiple Deprivation (GIMD 2015, from Q1, least deprived, to Q5,
- 47 most deprived districts).
- 48

# 49 **Results**

- 50 Among 13,351 adults with T1D, the use of technology decreased significantly with older age: CSII
- use fell from 56.1% in the 18–<25-year age group to 3.1% in the  $\geq$ 80-year age group, CGM use from
- 52 75.3% to 28.2%, and SAP use from 45.1% to 1.5% (all p for trend <0.001). The use of technology
- 53 was also significantly higher in women than in men (CSII: 39.2% vs. 27.6%; CGM: 61.9% vs.
- 54 58.0%; SAP: 28.7% vs. 19.6%, all p <0.001), and in individuals without migration background than
- 55 in those with migration background (CSII: 38.8% vs. 27.6%; CGM: 71.1% vs. 61.4%; SAP: 30.5%
- vs. 21.3%, all p <0.001). Associations with area deprivation were not linear: the use of each
- 57 technology decreased only from Q2 to Q4.
- 58

## 59 **Discussion**

- 60 Our real-world data provide evidence that higher age, male gender, and migration background are
- 61 currently associated with lower use of diabetes technology in adults with T1D in Germany.
- 62 Associations with area deprivation are more complex, probably due to correlations with other factors,
- 63 like the higher proportion of migrants in less deprived areas or the federal structure of the German
- 64 health care system.
- 65

# 66 **1** Introduction

67 Over the past few years, considerable advances in diabetes technology have revolutionized the 68 management of type 1 diabetes (T1D). Not only continuous glucose monitoring systems (CGM) and 69 continuous subcutaneous insulin infusion (CSII or insulin pumps), but also innovative systems 70 connecting both devices with algorithms to facilitate automated insulin delivery (AID, or "hybrid 71 closed loop", HCL) have been increasingly used by people with T1D in high-income countries over 72 the past decade (1-3). Numerous studies indicate that the use of these different devices is associated 73 with better glycemic control (2, 4-6), less severe hypoglycemia (2, 5, 6), and improved quality of life 74 (6-8) in both children and adults with T1D. However, significant inequalities in use of modern 75 diabetes technology have been reported in many countries. In pediatric populations, persistent or 76 widening racial-ethnic and/or socioeconomic disparities in the use of CSII and CGM have been 77 described in the US (3, 4, 9–11), in Canada (12), in New-Zealand (13), in the UK (4, 14), or in 78 Germany (4, 9, 15). In adults, the use of diabetes technology is still less widespread than in children 79 and only few studies were performed. Nevertheless, ethnic disparities in the use of CSII, CGM and 80 also AID, have been described in the US (16, 17), as well as ethnic and socioeconomic disparities in 81 CSII and CGM use in the UK (18).

82 The influence of demographic or socioeconomic factors on the use of diabetes technology in adults 83 has not been analyzed to date in Germany. However, information on the actual use of the different 84 diabetes treatment devices in the entire population, including underrepresented groups, such as 85 migrants, the elderly, or the socioeconomically disadvantaged, is important. Studies focusing on 86 disadvantaged populations point out that the use of CSII and CGM helps to reduce adverse events 87 and to improve HbA1c levels in these groups and that diabetes technology has therefore the potential to reduce disparities in diabetes outcomes (19-21). Nevertheless, if those who could benefit most 88 89 from technologies have less access to it, and if these disparities increase as diabetes technologies 90 advance, disparities in diabetes outcomes are expected to worsen (22, 23). To properly assess this 91 issue, it is necessary to know accurately the current utilization rates of commercially available 92 diabetes treatment devices in different population subgroups. Therefore, we aimed to analyze recent 93 technology use in Germany in a representative population of adults with T1D by age, gender, 94 migration background, and area deprivation (as defined in the following section).

# 96 2 Materials and methods

## 97 Data source and study population

98 In this cross-sectional study, we used data from the multicenter, diabetes prospective follow-up 99 registry (DPV). As of September 2022, the DPV registry comprised demographic and clinical data of 100 about 705,000 patients with any type of diabetes, documented by 507 pediatric and adult health care 101 facilities, of which 456 are located in Germany. All participating centers transmit twice a year the 102 locally collected data in pseudonymized form to Ulm University, Germany. After plausibility checks 103 and corrections, the Ulm University aggregates the data into an anonymized database for 104 benchmarking and medical research. Data collection and analysis were both approved by the ethics 105 committee of the Medical Faculty of Ulm University (Number 314/21) and by local review boards of 106 the participating centers. In the present study, we included data documented between 2019 and 2021 107 of individuals diagnosed with T1D since at least three months, aged  $\geq 18$  years, with residence in 108 Germany. T1D was identified by a clinical diagnosis at the age of at least 6 months and the

- 109 documentation of insulin use.
- 110

## 111 Demographic variables and area deprivation

Age was divided into the following groups: 18-<25 year, 25-<40 years, 40-<60 years, 60-<80 years

and  $\geq 80$  years. Migration background was defined as place of birth outside Germany for the patient

114 or at least for one of his parents. Area deprivation was assessed using the German Index of Multiple

115 Deprivation of the year 2015 (GIMD 2015). The concept of area deprivation can be defined as a lack

- 116 of area-based resources, compared to the society in which one lives (24, 25). As described in
- 117 previous publications (24, 26), the GIMD encompassed aggregated data at district level in seven
- deprivation domains differently weighted: income (25%), occupation (25%), education (15%),

119 municipal/district revenue (15%), social capital (10%), environment (5%), and security (5%).

120 Districts were categorized into area deprivation quintiles from Q1 (lowest deprivation quintile) to Q5

121 (highest deprivation quintile). We used individual postal code of patient's residences to assign them

122 to districts and consequently to GIMD quintiles.

123

## 124 Use of diabetes technology

125 We investigated any use of insulin pump / continuous subcutaneous insulin infusion (CSII), sensor /

126 continuous glucose monitoring (CGM), and sensor augmented pump therapy (SAP) in the

observation period. SAP use was defined as simultaneous use of insulin pump and sensor, connectedor not with algorithms for automated insulin delivery (AID).

129

## 130 Statistical Analysis

131 Data documented between 2019 and 2021 were aggregated per individual as maximum (technology 132 use documented once or not during this period) or median (other variables). Using multiple logistic 133 regressions, we analyzed the proportion of individuals using CSII, CGM, and SAP by gender, age 134 group, migration background, and area deprivation. All models were adjusted for diabetes duration 135 group (0-<5 years, 5-<10 years, 10-<20 years, and  $\geq$ 20 years), and when possible for gender and age 136 groups (see above). Multiple regressions models including all factors together (gender, age group, 137 migration background, and area deprivation) were additionally performed as sensitivity analysis. In 138 addition, interactions between migration background and area deprivation were analyzed. Associations of technology use (CSII, CGM, and SAP) with HbA1c were analyzed using multiple 139 140 linear regressions in each gender, age, migration, and deprivation subgroup (stratification). All 141 models were adjusted for diabetes duration group, and when possible for gender and age groups. 142 Results of regression analyses are presented as coefficients and as adjusted proportions (least square 143 means) with 95%-confidence intervals (95%-CI). Descriptive data are given as median with lower 144 and upper quartiles for continuous variables and as percentage for binary variables. A p-value < 0.01145 in two-sided tests was considered statistically significant. Statistical analyses were conducted using 146 SAS version 9.4 (build TS1M7, SAS Institute Inc, Cary, NC).

147

# 148 **3 Results**

The study population comprised 13,351 adults with T1D, with median age of 30.9 years [lower–
upper quartile: 19.0–55.8 years] and median diabetes duration of 13.4 years [7.2–23.8 years] (Table
1). Overall, 36.4% used a CSII, 59.0% at least once a CGM (37.8% at least 90 days per year), and
27.1% both devices (22.6% SAP without AID and 4.5% SAP with AID).

# 154 **Technology use by age group**

155 The use of every technology decreased continuously and significantly with older age (p for

trend >0.001, Figure 1, Table 2). The biggest relative difference in use between two successive age

- 157 groups was for all devices between the two youngest and between the two oldest age groups (18-<25
- 158 vs. 25-<40-year-olds and 60-<80 vs.  $\geq$  80-year-olds, Figure 1). Between the two youngest age
- 159 groups, CSII use decreased from 56.1% [95%-CI: 54.5–57.7] to 32.1% [29.9–34.3], CGM use from
- 160 75.3% [74.1–76.5] to 52.8% [50.5–55.0], and SAP use from 45.1% [43.4–46.7] to 22.3% [20.5–
- 161 24.2], all differences p <0.001. Between the two oldest age groups, CSII use decreased from 12.7%
- 162 [11.4–14.1] to 3.1% [2.1–4.6], CGM use from 41.6% [39.3–43.9] to 28.2% [24.3–32.5], and SAP use
- 163 from 9.3% [8.2–10.5] to 1.5% [0.1–2.7], all differences p <0.001.
- 164

## 165 Technology use by gender

- 166 All devices were more frequently used by women than by men (all differences p <0.001, Figure 1,
- 167 Table 2). The largest difference between genders was for CSII: 39.2% [37.9–40.6] in women vs.
- 168 27.6% [26.5–28.7] in men. CGM was used by 61.9% [60.6–63.2] of the women compared to 58.0%
- 169 [56.8–59.2] of the men, and SAP by 28.7% [27.5–30.0] of the women compared to 19.6% [18.7–
- 170 20.7] of the men.
- 171

# 172 Technology use by migration background

173 Information on migration background was only documented in 5,290 of 13,351 (39.6%) individuals

- 174 (Table 1). In patients with this information, the use of every technology was significantly higher in
- 175 individuals without migration background than in those with migration background (all differences
- 176 p <0.001, Figure 1, Table 2): CSII was used by 38.8% [36.8–40.9] vs. 27.6% [25.1–30.3], CGM by
- 177 71.1% [69.2–72.9] vs. 61.4% [58.5–64.4], SAP by 30.5% [28.6–32.4] vs. 21.3% [19.1–23.6]. In
- individuals with unknown migration status, CSII was used by 30.6% [29.4–32.0], CGM by 53.9%
- 179 [52.5–55.3], and SAP by 20.7% [19.6–21.9].
- 180

## 181 **Technology use by area deprivation**

182 Associations between area deprivation and technology use were not linear (Figure 1, Table 2). The

- 183 use of every technology decreased with higher deprivation from Q2 to Q4. CGM use was also higher
- 184 in the two least deprived quintiles Q1-Q2 than in the three most deprived quintiles (Q3-Q5): 62.3%-
- 185 67.9% vs. 55.4%–57.5%.
- 186

## 187 Technology use by interaction between migration background and area deprivation

- 188 For each type of technology, results from multiple regression models including all factors together
- 189 (gender, age group, migration background, and area deprivation) were very similar to the results
- 190 presented above and all factors remained significant (p < 0.01).
- 191 Interactions between migration background and area deprivation were not significant (CSII:
- 192 p=0.794; CGM: p=0.201; CSII: p=0.782). The use of each technology was constantly higher in
- 193 patients without migration background than in those with migration background regardless of
- 194 deprivation quintile. In patients without migration background, the use of CSII varied in a nonlinear
- 195 manner across deprivation quintiles between 41.7% (Q1) and 53.1% (Q3), the use of CGM between
- 196 76.6% (Q2) and 79.2% (Q3), and the use of SAP between 35.6% (Q1) and 44.3% (Q3). In patients
- 197 with migration background, the use of insulin pump varied between 32.0% (Q1) and 41.2% (Q3), the
- use of CGM between 67.4% (Q5) and 75.0% (Q1), and the use of SAP between 26.4% (Q1) and
- 199 33.6% (Q3).
- 200

## 201 HbA1c by technology use

Adults using CSII, CGM or SAP had lower HbA1c in each gender, age, migration, and deprivation category than adults no using these devices (Table 3). All comparisons were significant, excepted in adults aged 80 or over (due to their small number, n=468), and in persons without migration background or in persons living in districts Q2 for the use of CSII or SAP (Table 3).

206

## 207 **4 Discussion**

Our analysis based on more than 13,000 adults with T1D in Germany provides real world evidence that younger age, female gender, and absence of migration background are significant facilitators for use of diabetes technology in this population. Associations with area deprivation were less clear.

- 211 Previous real-world analyses from Germany reported a higher use of diabetes technology with
- 212 younger age in pediatrics, as well as an overall lower use in adults compared to children (1, 27).
- 213 However, the impact of age on the use of diabetes technology within the adult population has not
- been investigated to date. German and international guidelines recommend the use of diabetes
- 215 technology (CSII, CGM, and also AID) for most adults, even older ones, if they desire it and if this
- use is compatible with preserving their autonomy (28, 29). Yet, our data indicate that the real-world

217 use of CSII, CGM and SAP significantly decreases with older age. Data from France also confirmed 218 a lower use of CSII with older age in adults (30). In contrast, data from the US-T1D Exchange 219 registry indicated the lowest use of both CGM and CSII in 18-25 year-olds compared to older 220 patients (3, 31). The high cost and lack of reimbursement for these technologies in the absence of 221 health insurance may explain the lower use of these technologies by young adults in the US, since 222 young adults tend to have lower incomes than their elders. In Germany, nearly all patients benefit 223 from a health insurance. Moreover, the higher initiation rate in children and adolescents in this 224 country and the continuation of technology use after childhood may contribute to the higher use in 225 young adults. Barriers related to difficulties with technology utilization seem not to play a role for 226 age differences, except perhaps in the oldest age group, in which disabilities may limit the use of 227 these devices (23, 31). Nevertheless, the impact of age on technology use in adults needs to be further 228 investigated.

229 We found a higher use of all technologies in women compared to men, with the largest difference for 230 CSII. To date, numerous studies reported a higher use of CSII or SAP, but not of CGM alone, in 231 female adolescents and adults (1, 4, 27, 30, 31, 33–35). This finding is consistent in many reports, 232 although women often report more physical barriers to technology adoption than men (23, 31). 233 Several specific indications for technology use for women exist. Current German guidelines 234 recommend for instance the use of CSII and of CGM for women before and during pregnancy (28, 235 36). CSII is also indicated in case of unsatisfying glycemic control, which is more frequent in female 236 adolescents compared to males (34). The more frequent use of a pump in young women may 237 continue with older age even if the glycemic results improve (34). In contrast to older studies, our 238 data indicate that women used a CGM more frequently than men. The greater use of SAP and AID in 239 women compared to men in the most recent years leads automatically to a higher CGM use, since a 240 CGM is part of all SAP and AID systems.

To date, only few studies have examined demographic and socioeconomic disparities in technology access in adults with T1D (16, 17, 31, 37). An analysis from the UK indicates an association between higher deprivation and lower use of CSII and CGM in adults with T1D, as well as a significant lower use of both technologies in individuals with black ethnicity compared to those with mixed or white ethnicity (18). In our analysis, differences in technology use by migration background were stronger than those by area deprivation and the use of each technology was constantly higher in adults without migration background regardless of deprivation. These results are consistent with previous findings 248 in pediatrics in Germany (15). Contrary to what is known about the situation in England (18) or the 249 United States (20, 38, 39), there is no strong correlation between migration background and regional 250 deprivation in Germany, because less migrants live in the most deprived areas (e.g. in eastern parts of 251 Germany) than in the least deprived areas (e.g. in Bavaria and in Baden-Württemberg) (40, 41). In 252 our study population with documented migration status, the highest proportion of persons with 253 migration background lived in moderately deprived area (Q3: 27.6% vs. 21.9-24.8% in the other 254 districts). In addition, almost all adults living in Germany have a statutory or private insurance that 255 reimburses most of CSII and CGM costs in case of intensive insulin therapy. Thus, in contrast to the 256 situation in the US where individuals might be disadvantaged due to their insurance status (31, 39, 257 42), economic factors should not play an important role in limiting access to technologies for T1D in 258 Germany.

259 We found, however, that the presence of migration background was significantly associated with less 260 technology use. Individuals with migration background have less often a higher qualification degree 261 than German natives (43) and some first generation migrants may have difficulties with the language 262 of the host country. This can constitute a barrier to complete the specialized education required to use 263 diabetes treatment devices (31). Initial and ongoing education and training is essential for the use of 264 diabetes technology, but it requires a number of resources, like free time, health literacy and 265 numeracy or perceived self-efficacy (6, 44). Language barriers may also exist when it comes to 266 filling out forms for reimbursement or telephone contact when technical problems with diabetes 267 devices arise (31). Finally, the choice of a specific device must be based on individual characteristics, 268 that is a person's needs, preferences and skills levels (6). In this decision-making process, the 269 subjectivity of both the patient and the provider play a role. As a consequence, provider implicit bias, 270 observed for example when the recommendation of diabetes technology unconsciously but 271 systematically disadvantages some patients due to their ethnic or socioeconomic characteristics, is 272 always possible and may also exist in Germany (38, 42).

Our results indicate better glycemic control in all adults using CSII, CGM or SAP compared to those not using these technologies. This is an argument for continuing efforts to improve access to technologies in older adults, in males and in people with migration background. However, due to the cross-sectional design of this study, these associations must be interpreted with caution and we cannot conclude on a potential causal relationship between technology use and lower HbA1c.

## 279 Strengths and limitations

280 One strength of this study is the use of the large multicenter DPV registry, which can give a good 281 insight into the real-world use of diabetes technology in adults with T1D in Germany. Even if the 282 representativeness of the registry is lower than in pediatric diabetes, the risk of selection bias in our 283 findings is relatively low and generalizations may be valid. However, given the rapid advances in 284 diabetes technology and the continued increase in its use, these analyses must be updated regularly. 285 One limitation is that socioeconomic factors were assessed at the district-level, not at the individual 286 level. Aggregated data can weaken the effect of individual socioeconomic factors on the use of 287 diabetes technology and underestimate their influence. Nevertheless, other aspects related to living 288 conditions and diabetes care, which is largely organized at the federal level, can be better reflected 289 using an area-based deprivation index. We did not account for persons who moved from one district 290 to another and thus potentially changed their deprivation category. However, only 4.6% of the 291 population have moved within Germany in 2021 (destatis.de) and only a part of this proportion may 292 have moved to a different deprivation quintile. Moreover, some of them might have moved to a more 293 deprived district, but others to a less deprived district, so that the resulting potential bias may be 294 mainly non-differential. Finally, we used a binary variable for migration background that does not 295 reflect the tremendous heterogeneity within the population. In 2021, more than a quarter of the 296 people living in Germany had a migration background (45). These persons form a very 297 heterogeneous subpopulation in terms of country of origin, time living in Germany, reasons for 298 migration, legal status, education, language skills, or access to employment. Our results do not take 299 this diversity into account and this could be the subject of future research.

## 300 Conclusion

301 Our real-world data provide evidence that higher age, male gender, and migration background are 302 associated with lower use of modern diabetes technology in adults with T1D in Germany. 303 Associations with area deprivation are more complex, probably due to correlations with other factors 304 that exert in part opposite effects, like the higher proportion of migrants in less deprived areas, or the 305 federal structure of the German health care system. There is a critical need to improve access to 306 diabetes technology in underserved groups for reducing health disparities. This can enable them to 307 benefit from the latest technological advancements and achieve better glycemic control, which has 308 the potential to ultimately improved health outcomes.

310	5	Conflict of Interest
311	The	authors declare that the research was conducted in the absence of any commercial or financial
312	rela	tionships that could be construed as a potential conflict of interest.
313		
314	6	Author Contributions
315	M.A	A. and R.W.H. designed the study. M.A. and A.J.E. analyzed the study data. M.A. created the
316	figu	rres, and wrote the manuscript. A.J.E., S.M.M., A.G., L.S., P.M.J., M.He, M.Hu. and R.W.H.
317	con	tributed to the discussion, reviewed, and approved the manuscript.
318		
319	7	Funding
320	The	e DPV registry and this analysis are supported by the German Center for Diabetes Research (DZD,
321	grai	nt number: 82DZD14E03). Further financial support for the DPV registry was provided by the
322	Ger	man Diabetes Association (DDG), the German Robert Koch Institute (RKI).
323		
324	8	Acknowledgments
325	We	thank all centers participating in the DPV initiative, especially those contributing data to this
326	ana	lysis and their patients. A list of contributing centers is available at <u>http://www.d-p-v.eu</u>
327		
328	9	Data Availability Statement
329	Acc	cess to the programming code can be provided by the corresponding author upon request. For
330	reas	sons of data protection, data on individual level cannot be provided. However, remote data
331	ana	lysis is possible.
332		
333	Tał	oles and Figures
334	Tab	le 1. Characteristics of the study population
335	Tab	le 2. Technology use: coefficients from multiple logistic regression models
336 337	Tab	<b>le 3.</b> HbA1c: results from multiple linear regression models
338	Fig	ure 1. Use of diabetes technology by age group, gender, migration background, and area
339	dep	rivation
	1	
340		

	Median (lower-	n, percent <sup>342</sup> )
	upper quartile)	3/3
Age, years	30.9 (19.0–55.8)	
Age groups		344
18 - <25 years		5,902 (44.2)
25 - <40 years		1,915 (34.3)
40 - <60 years		2,906 (21.8)
60 - <80 years		2,160 (16.2)
$\geq 80$ years		468  (3-5)  (3-7)
Sex		
Male		7,132 ( <b>34</b> 8)
Female	- T'	6,219 (46.6)
Migration background*		349
Without Migration background		4,015 (30.1)
With Migration background		1,275 (9.5)
Not documented		8,061 ( <b>69.1</b> )
Diabetes duration, years	13.4 (7.2–23.8)	
BMI **	25.7 (24.8–22.1)	352
HbA1c, %	7.65 (6.88–8.69)	252
Use of CSII		4,860 (36.4)
Use of CGM		354
Any use		7,877 (59.0)
Use $\geq$ 90 days /year		5,047 (37.8)
Use of SAP		054
All SAP		3,618 (27.1)
Only AID		601 (4.5)
All patients		13,351 (100.0)
		358

359 Unadjusted data. \* defined as birth of the patient himself or at least one of his parents outside of Germany.

 $360 \qquad ** \ Body \ Mass \ Index \ (kg/m^2).$ 

361



304		
365		
366	Table 2. Technology use: coefficients from multiple logistic regression models	

		Use of CSII	P-value	Use of CGM	P-value	Use of SAP (without AID)	P-value	Use of AID	P-value
Intercept		- 2.38	< 0.0001	- 0.93	< 0.0001	- 3.43	< 0.0001	- 17.23	< 0.0001
Diabetes	< 5 yrs	- 2.09	< 0.0001	- 0.13	0.0129	- 1.62	< 0.0001	- 0.97	< 0.0001
duration groups	5-< 10 yrs	- 1.15		- 0.10		- 0.78		- 0.52	
	10-< 20 yrs	- 0.60		- 0.17		- 0.42		- 0.42	
	$\geq$ 20 yrs	Ref.		Ref.		Ref.		Ref.	
Age groups	18 -< 25 yrs	3.50	< 0.0001	1.58	< 0.0001	3.44	< 0.0001	14.96	< 0.0001
	25 -< 40 yrs	2.67		1.00		2.77		14.14	
	40 -< 60 yrs	2.12		0.85		2.26		13.72	
	60 -< 80 yrs	1.51		0.61		1.79		13.42	
	80 -< 100 yrs	Ref.		Ref.		Ref.		Ref.	
Gender	male	- 0.53	< 0.0001	- 0.16	< 0.0001	- 0.48	< 0.0001	- 0.27	0.0015
	female	Ref.		Ref.		Ref.		Ref.	
Migration	yes	- 0.13	< 0.0001	0.34	< 0.0001	- 0.04	< 0.0001	0.39	< 0.0001
background	no	0.37		0.76		0.35		0.79	
	n.d.	Ref.		Ref.		Ref.		Ref.	
Area	Q1	0.02	0.0028	0.25	< 0.0001	0.08	< 0.0001	- 0.46	0.0124
deprivation	Q2	0.06		0.49		0.20		- 0.07	
quintiles	Q3	- 0.11		0.00		- 0.08		- 0.02	
	Q4	- 0.15		- 0.04		- 0.14		- 0.12	

Q5	Ref.	Ref.	Ref.	Ref.	
----	------	------	------	------	--

- 368 Coefficients from logistic regression models, adjusted by diabetes duration, and when possible by age groups, gender, migration background and area 369 deprivation. n.d.: not documented

- **Table 3.** HbA1c: results from multiple linear regression models

		Use of	Use of CSII		Use of CGM		P-value	Use of SAP		P-value
		no	yes		no	yes		no	yes	
Age groups	18 -< 25 yrs	8.18	7.94	< 0.0001	8.39	7.96	< 0.0001	8.18	7.90	< 0.0001
		[8.12-8.24]	[7.88-8.01]		[8.30-8.48]	[7.91–8.01]		[8.13-8.24]	[7.83 –7.97]	
	25 -< 40 yrs	8.34	7.65	< 0.0001	8.57	7.70	<0.0001	8.28	7.56	< 0.0001
		[8.23-8.45]	[7.50–7.80]		[8.45-8.69]	[7.59–7.81]		[ 8.18 - 8.38]	[7.38–7.73]	
	40 -< 60 yrs	8.05	7.65	< 0.0001	8.24	7.63	< 0.0001	8.02	7.60	< 0.0001
		[7.98-8.13]	[7.53–7.77]		[8.15-8.33]	[7.55–7.72]		[7.95-8.09]	[7.45–7.75]	
	60 -< 80 yrs	7.73	7.42	< 0.0001	7.85	7.45	< 0.0001	7.73	7.32	< 0.0001
		[7.67–7.80]	[7.29–7.55]		[7.77–7.93]	[7.36 –7.53]		[7.66–7.79]	[7.17–7.48]	
	80 -< 100 yrs	7.98	7.41	0.0611	8.01	7.80	0.1366	7.96	7.42	0.2095
		[7.84-8.11]	[6.84–7.99]		[7.86-8.17]	[7.56-8.03]		[7.83 -8.09]	[6.59-8.25]	
gender	male	8.03	7.80	< 0.0001	8.29	7.73	< 0.0001	8.03	7.74	< 0.0001
		[7.98-8.08]	[7.73–7.88]		[8.23-8.36]	[7.68–7.78]		[7.98-8.08]	[7.65–7.82]	
	female	8.18	7.73	< 0.0001	8.29	7.81	< 0.0001	8.14	7.69	< 0.0001
		[8.12-8.24]	[7.67–7.80]		[ 8.22-8.36]	[7.75–7.86]		[8.09-8.19]	[7.62–7.77]	
Migration	yes	8.46	8.02	< 0.0001	8.71	8.11	< 0.0001	8.45	7.97	< 0.0001
background		[8.33-8.59]	[7.85-8.19]		[8.53-8.88]	7.99– 8.23]		[8.33-8.57]	[7.78-8.15]	

	No	8.02	7.92	0.0609	8.21	7.91	< 0.0001	8.04	7.88	0.0054
		[7.95-8.10]	[7.85–7.99]		[8.10-8.32]	[7.85–7.96]		[7.97-8.10]	[7.80–7.96]	
	n.d.	8.06	7.60	< 0.0001	8.22	7.64	< 0.0001	8.03	7.52	< 0.0001
		[ 8.02-8.10]	[7.52–7.67]		[8.16-8.27	[7.59–7.69]		[7.99–8.07]	[7.43–7.60]	
Area	Q1	7.82	7.56	< 0.0001	7.96	7.58	< 0.0001	7.81	7.50	< 0.0001
deprivation		[7.75-7.90]	[7.46–7.66]		[7.87-8.05]	[7.51–7.65]		[7.74–7.88]	[7.39–7.61]	
quintiles	Q2	7.92	7.77	0.0265	8.16	7.73	< 0.0001	7.93	7.72	0.0033
		[7.84-8.00]	[7.67–7.87]		[8.05-8.27]	[7.65–7.80]		[7.85-8.00]	[7.61–7.83]	
	Q3	8.11	7.71	< 0.0001	8.37	7.68	< 0.0001	8.09	7.63	< 0.0001
		[8.02-8.19]	[7.59–7.82]		[8.26-8.47]	[7.60–7.77]		[8.01-8.17]	[7.50–7.76]	
	Q4	8.38	7.97	< 0.0001	8.57	7.99	< 0.0001	8.34	7.94	< 0.0001
		[8.29-8.47]	[7.83-8.10]		[8.45-8.68]	[7.89-8.09		[8.26-8.43]	[7.79 -8.09]	
	Q5	8.28	7.84	< 0.0001	8.40	7.90	< 0.0001	8.25	7.77	< 0.0001
		[8.20-8.36]	[7.73–7.95]		[8.30-8.49]	[7.81–7.99]		[8.18-8.33]	[7.64–7.89]	

373

374 Linear regression models adjusted by diabetes duration, when possible by gender and by age groups. n.d.= not documented.



# 375 11 REFERENCES

- van den Boom L, Karges B, Auzanneau M, Rami-Merhar B, Lilienthal E, Sengbusch S von, et al.
   Temporal Trends and Contemporary Use of Insulin Pump Therapy and Glucose Monitoring
   Among Children, Adolescents, and Adults With Type 1 Diabetes Between 1995 and 2017.
   *Diabetes care* (2019) 42:2050–6. doi:10.2337/dc19-0345.
- Da Silva J, Bosi E, Jendle J, Arrieta A, Castaneda J, Grossman B, et al. Real-world performance of the MiniMed<sup>TM</sup> 670G system in Europe. *Diabetes, obesity & metabolism* (2021) 23:1942–9. doi:10.1111/dom.14424.
- Foster NC, Beck RW, Miller KM, Clements MA, Rickels MR, DiMeglio LA, et al. State of Type
   1 Diabetes Management and Outcomes from the T1D Exchange in 2016-2018. *Diabetes technology & therapeutics* (2019) **21**:66–72. doi:10.1089/dia.2018.0384.
- 386
  4. Sherr JL, Hermann JM, Campbell F, Foster NC, Hofer SE, Allgrove J, et al. Use of insulin pump
  387
  388
  388
  389
  59:87–91. doi:10.1007/s00125-015-3790-6.
- Karges B, Schwandt A, Heidtmann B, Kordonouri O, Binder E, Schierloh U, et al. Association of
  Insulin Pump Therapy vs Insulin Injection Therapy With Severe Hypoglycemia, Ketoacidosis,
  and Glycemic Control Among Children, Adolescents, and Young Adults With Type 1 Diabetes. *JAMA* (2017) **318**:1358–66. doi:10.1001/jama.2017.13994.
- ElSayed NA, Aleppo G, Aroda VR, Bannuru RR, Brown FM, Bruemmer D, et al. 7. Diabetes
   Technology: Standards of Care in Diabetes-2023. *Diabetes care* (2023) 46:S111-S127.
   doi:10.2337/dc23-S007.
- 7. Polonsky WH, Hessler D, Ruedy KJ, Beck RW. The Impact of Continuous Glucose Monitoring
  on Markers of Quality of Life in Adults With Type 1 Diabetes: Further Findings From the
  DIAMOND Randomized Clinical Trial. *Diabetes care* (2017) 40:736–41. doi:10.2337/dc170133.
- 401 8. Herringshaw E. The Price of Inequality in Type 1 Diabetes Management. *Journal of the* 402 *Endocrine Society* (2022) 6:bvac051. doi:10.1210/jendso/bvac051.
- 403 9. Addala A, Auzanneau M, Miller K, Maier W, Foster N, Kapellen T, et al. A Decade of
  404 Disparities in Diabetes Technology Use and HbA1c in Pediatric Type 1 Diabetes: A
  405 Transatlantic Comparison. *Diabetes care* (2021) 44:133–40. doi:10.2337/dc20-0257.
- 406
  407
  408
  408
  409
  409
  409
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
  400
- 410 11. Lai CW, Lipman TH, Willi SM, Hawkes CP. Racial and Ethnic Disparities in Rates of
  411 Continuous Glucose Monitor Initiation and Continued Use in Children With Type 1 Diabetes.
  412 Diabetes care (2021) 44:255–7. doi:10.2337/dc20-1663.
- 413 12. Ladd JM, Sharma A, Rahme E, Kroeker K, Dubé M, Simard M, et al. Comparison of
- 414 Socioeconomic Disparities in Pump Uptake Among Children With Type 1 Diabetes in 2
- 415 Canadian Provinces With Different Payment Models. *JAMA network open* (2022) **5**:e2210464.
- 416 doi:10.1001/jamanetworkopen.2022.10464.

- 417 13. Burnside MJ, Williman JA, Davies HM, Jefferies CA, Paul RG, Wheeler BJ, et al. Inequity in access to continuous glucose monitoring and health outcomes in paediatric diabetes, a case for national continuous glucose monitoring funding: A cross-sectional population study of children with type 1 diabetes in New Zealand. *The Lancet regional health. Western Pacific* (2023)
  421 31:100644. doi:10.1016/j.lanwpc.2022.100644.
- 422 14. Ng SM, Evans ML. Widening health inequalities related to type 1 diabetes care in children and
  423 young people in the UK: A time to act now. *Diabetic medicine a journal of the British Diabetic*424 *Association* (2021) **38**:e14620. doi:10.1111/dme.14620.
- 425 15. Auzanneau M, Rosenbauer J, Maier W, Sengbusch S von, Hamann J, Kapellen T, et al.
  426 Heterogeneity of Access to Diabetes Technology Depending on Area Deprivation and
  427 Demographics Between 2016 and 2019 in Germany. *Journal of diabetes science and technology*428 (2021) 15:1059–68. doi:10.1177/19322968211028608.
- 429 16. Kanbour S, Jones M, Abusamaan MS, Nass C, Everett E, Wolf RM, et al. Racial Disparities in
  430 Access and Use of Diabetes Technology Among Adult Patients With Type 1 Diabetes in a U.S.
  431 Academic Medical Center. *Diabetes care* (2023) 46:56–64. doi:10.2337/dc22-1055.
- 432 17. Ju Z, Piarulli A, Bielick L, Marschall S, Brouillard E, Steenkamp D. Advanced Diabetes
  433 Technology Remains Underutilized in Underserved Populations: Early Hybrid Closed-Loop
  434 System Experience at an Academic Safety Net Hospital. *Diabetes technology & therapeutics*435 (2022) 24:143–7. doi:10.1089/dia.2021.0334.
- 436 18. Fallon C, Jones E, Oliver N, Reddy M, Avari P. The impact of socio-economic deprivation on
  437 access to diabetes technology in adults with type 1 diabetes. *Diabetic medicine a journal of the*438 *British Diabetic Association* (2022) **39**:e14906. doi:10.1111/dme.14906.
- 439 19. Everett EM, Wisk LE. Relationships Between Socioeconomic Status, Insurance Coverage for
  440 Diabetes Technology and Adverse Health in Patients With Type 1 Diabetes. *Journal of diabetes*441 *science and technology* (2022) 16:825–33. doi:10.1177/19322968211050649.
- 442 20. Agarwal S, Kanapka LG, Raymond JK, Walker A, Gerard-Gonzalez A, Kruger D, et al. Racial443 Ethnic Inequity in Young Adults With Type 1 Diabetes. *The Journal of clinical endocrinology*444 *and metabolism* (2020) **105**:e2960-9. doi:10.1210/clinem/dgaa236.
- 445 21. Stanley JR, Clarke AB, Shulman R, Mahmud FH. Mediating Effects of Technology-Based
  446 Therapy on the Relationship Between Socioeconomic Status and Glycemic Management in
  447 Pediatric Type 1 Diabetes. *Diabetes technology & therapeutics* (2022).
  448 doi:10.1089/dia.2022.0388.
- 449 22. Agarwal S, Simmonds I, Myers AK. The Use of Diabetes Technology to Address Inequity in
  450 Health Outcomes: Limitations and Opportunities. *Current Diabetes Reports* (2022) 22:275–81.
  451 doi:10.1007/s11892-022-01470-3.
- 452 23. Pauley ME, Berget C, Messer LH, Forlenza GP. Barriers to Uptake of Insulin Technologies and
  453 Novel Solutions. *Medical devices (Auckland, N.Z.)* (2021) 14:339–54.
  454 doi:10.2147/MDER.S312858.
- 455 24. Maier W, Scheidt-Nave C, Holle R, Kroll LE, Lampert T, Du Y, et al. Area level deprivation is
  456 an independent determinant of prevalent type 2 diabetes and obesity at the national level in
- 457 Germany. Results from the National Telephone Health Interview Surveys 'German Health
- 458 Update' GEDA 2009 and 2010. *PloS one* (2014) **9**:e89661. doi:10.1371/journal.pone.0089661.

- 459 25. Townsend P. *Poverty in the United Kingdom: A survey of household resources and standards of living*. Berkeley: Univ. of Calif. Pr (1979). 1216 p.
- 461 26. Auzanneau M, Lanzinger S, Bohn B, Kroschwald P, Kuhnle-Krahl U, Holterhus PM, et al. Area
  462 Deprivation and Regional Disparities in Treatment and Outcome Quality of 29,284 Pediatric
  463 Patients With Type 1 Diabetes in Germany: A Cross-sectional Multicenter DPV Analysis.
  464 Diabetes care (2018) 41:2517–25. doi:10.2337/dc18-0724.
- 27. van den Boom L, Auzanneau M, Woelfle J, Sindichakis M, Herbst A, Meraner D, et al. Use of
  Continuous Glucose Monitoring in Pump Therapy Sensor Augmented Pump or Automated
  Insulin Delivery in Different Age Groups (0.5 to <26 Years) With Type 1 Diabetes From 2018 to</li>
  2021: Analysis of the German/Austrian/Swiss/Luxemburg DPV Registry. *Journal of diabetes science and technology* (2023):19322968231156601. doi:10.1177/19322968231156601.
- 470 28. Holt RI, DeVries JH, Hess-Fischl A, Hirsch IB, Kirkman MS, Klupa T, et al. The management of
  471 type 1 diabetes in adults. A consensus report by the American Diabetes Association (ADA) and
  472 the European Association for the Study of Diabetes (EASD). *Diabetologia* (2021) 64:2609–52.
  473 doi:10.1007/s00125-021-05568-3.
- 474 29. Bahrmann A, Bahrmann P, Baumann J, Bauer J, Brückel E, Dreyer M, et al. S2k-Leitlinie
  475 Diagnostik, Therapie und Verlaufskontrolle des Diabetes mellitus im Alter. *Diabetologie und*476 *Stoffwechsel* (2018) 13:423–89. doi:10.1055/a-0666-0820.
- 30. Meunier L, Aguadé A-S, Videau Y, Verboux D, Fagot-Campagna A, Gastaldi-Menager C, et al.
  Age, Male Gender, and Social Deprivation Are Associated with a Lower Rate of Insulin Pump
  Therapy Initiation in Adults with Type 1 Diabetes: A Population-Based Study. *Diabetes technology & therapeutics* (2021) 23:8–19. doi:10.1089/dia.2020.0112.
- 481 31. Tanenbaum ML, Hanes SJ, Miller KM, Naranjo D, Bensen R, Hood KK. Diabetes Device Use in
  482 Adults With Type 1 Diabetes: Barriers to Uptake and Potential Intervention Targets. *Diabetes*483 *care* (2017) **40**:181–7. doi:10.2337/dc16-1536.
- 484 32. Tanenbaum ML, Commissariat PV. Barriers and Facilitators to Diabetes Device Adoption for
  485 People with Type 1 Diabetes. *Current Diabetes Reports* (2022) 22:291–9. doi:10.1007/s11892486 022-01469-w.
- 33. Shah VN, Wu M, Polsky S, Snell-Bergeon JK, Sherr JL, Cengiz E, et al. Gender differences in
  diabetes self-care in adults with type 1 diabetes: Findings from the T1D Exchange clinic registry. *Journal of diabetes and its complications* (2018) **32**:961–5. doi:10.1016/j.jdiacomp.2018.08.009.
- 490 34. Boettcher C, Tittel SR, Meissner T, Gohlke B, Stachow R, Dost A, et al. Sex differences over
  491 time for glycemic control, pump use and insulin dose in patients aged 10-40 years with type 1
  492 diabetes: a diabetes registry study. *BMJ open diabetes research & care* (2021) 9.
  493 doi:10.1136/bmjdrc-2021-002494.
- 494 35. Bak JC, Serné EH, Valk HW de, Valk NK, Kramer MH, Nieuwdorp M, et al. Gender gaps in
  495 type 1 diabetes care. *Acta Diabetologica* (2023) 60:425–34. doi:10.1007/s00592-022-02023-6.
- 496 36. Holder M, Kapellen T, Ziegler R, Bürger-Büsing J, Danne T, Dost A, et al. Diagnosis, Therapy
  497 and Follow-Up of Diabetes Mellitus in Children and Adolescents. *Experimental and clinical*498 *endocrinology & diabetes official journal, German Society of Endocrinology [and] German*499 *Diabetes Association* (2022) 130:S49-S79. doi:10.1055/a-1624-3388.

- 37. Messer LH, Addala A, Weinzimer SA. Real-World Diabetes Technology: Overcoming Barriers
  and Disparities. *Diabetes technology & therapeutics* (2023) 25:S176-S190.
  doi:10.1089/dia.2023.2511.
- 38. Odugbesan O, Addala A, Nelson G, Hopkins R, Cossen K, Schmitt J, et al. Implicit RacialEthnic and Insurance-Mediated Bias to Recommending Diabetes Technology: Insights from T1D
  Exchange Multicenter Pediatric and Adult Diabetes Provider Cohort. *Diabetes technology & therapeutics* (2022) 24:619–27. doi:10.1089/dia.2022.0042.
- 39. Majidi S, Ebekozien O, Noor N, Lyons SK, McDonough R, Gandhi K, et al. Inequities in Health
  Outcomes in Children and Adults With Type 1 Diabetes: Data From the T1D Exchange Quality
  Improvement Collaborative. *Clinical Diabetes* (2021) **39**:278–83. doi:10.2337/cd21-0028.
- 510 40. Statistisches Bundesamt. Number of foreigners in German federal states in 2021. Statista. (2022)
  511 [cited 2023 Feb 28]. Available from: https://www.statista.com/statistics/891288/foreigner512 numbers-by-state-germany.
- 513 41. Destatis. *Migration.Integration.Regionen. Gemeinsames Datenangebot von Destatis, BA und*514 BAMF [cited 2023 Feb 28]. Available from:
- 515 https://service.destatis.de/DE/karten/migration\_integration\_regionen.html.
- 42. Addala A, Hanes S, Naranjo D, Maahs DM, Hood KK. Provider Implicit Bias Impacts Pediatric
  Type 1 Diabetes Technology Recommendations in the United States: Findings from The
  Gatekeeper Study. *Journal of diabetes science and technology* (2021) 15:1027–33.
  doi:10.1177/19322968211006476.
- 520 43. Statistisches Bundesamt. "Distribution of the population with and without a migrant background
  521 in Germany as of 2019, by higher qualification degree." (2020) [cited 2023 Feb 28]. Available
  522 from: https://www.statista.com/statistics/921858/national-and-migrant-population-by-higher523 education-germany/.
- 44. Lindholm Olinder A, DeAbreu M, Greene S, Haugstvedt A, Lange K, Majaliwa ES, et al. ISPAD
   Clinical Practice Consensus Guidelines 2022: Diabetes education in children and adolescents.
   *Pediatric diabetes* (2022) 23:1229–42. doi:10.1111/pedi.13418.
- 527 45. Federal Statistical Office. *Well over one in four people in Germany had a migrant background in*528 2021 (2022) [cited 2023 Mar 8]. Available from:
- 529 https://www.destatis.de/EN/Press/2022/04/PE22\_162\_125.html.
- 530

### Figure 1.JPEG

Figure 1. Use of diabetes technology by age group, gender, migration background, and area deprivation



Area deprivation quintiles