

Transfer in care and diabetes distress in young adults with type 1 diabetes mellitus

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ABSTRACT

Introduction Diabetes distress (DD) is a serious problem in many people with diabetes and is associated with unfavorable clinical and psychosocial outcomes in children and adults. Little is known about DD in young adults (YAs) with type 1 diabetes mellitus (T1DM) who transferred to adult care. This study aimed to explore the differences between YAs with/without DD regarding transfer experiences, self-management and health-related quality of life (HRQoL).

Research design and methods Cross-sectional online questionnaire completed by YAs with T1DM after transfer. DD was measured with the short-form Problem Areas in Diabetes scale. Descriptive analyses were followed by t-tests and χ^2 tests to explore differences between the groups with/without DD. Effect sizes were calculated.

Results Of 164 respondents with mean age 22.7 (\pm 1.56) years, 60.7% was female. The total sample scored low on DD (6.52 \pm 4.67; range: 0–17), but 57 (34.8%) had a score \geq 8, indicating DD. YAs with DD felt less ready to transfer to adult care than those without DD and scored lower on alliance between pediatric and adult care and reception in adult care. They also reported poorer self-management skills and lower HRQoL in all domains of functioning.

Conclusions More than one-third YAs experienced DD after transfer; this was associated with less favorable transition, self-management and psychosocial outcomes. Transfer in care seems to be a source of DD. Systematic screening on DD and attention for YAs' worries is recommended in both pediatric and adult care.

INTRODUCTION

Emerging adulthood is a critical period with increased risk of acute and even chronic health complications, psychological problems and psychosocial issues for young adults with type 1 diabetes mellitus (T1DM).¹ Young people growing up with T1DM go through various transitions while transitioning from childhood to adulthood. Next to the developmental milestones they are expected to achieve, they are required to transfer to adult care and take up self-management of their chronic condition, which can be challenging.^{2–3} Young adults are under-represented in research on living with T1DM, although they have unique views, challenges

Significance of this study

What is already known about this subject?

► The few studies that explored diabetes distress in young adults with type 1 diabetes mellitus (T1DM) showed that diabetes distress is significantly more present in the young adult group compared with other age groups and that more diabetes distress in young adults with T1DM is associated with impaired health outcomes and less frequent blood monitoring by young adults.

What are the new findings?

► More than one-third of the young adults with T1DM experienced diabetes distress after transfer to adult care.
► Having diabetes distress was associated with less favorable transition experiences, self-management outcomes and psychosocial outcomes.
► Transfer in care may be a source of diabetes distress in young adults.

How might these results change the focus of research or clinical practice?

► Systematic screening on diabetes distress and attention for young adults' worries is recommended. Building positive relationships and adopting motivating communication styles may help clinicians to reduce diabetes distress in this population.

and needs.^{3–4} The regular developmental tasks of young adults in emerging adulthood may influence and be influenced by T1DM.² Diabetes management, for instance, is often a lower priority for young adults with T1DM during important life transitions,⁵ and glycaemic control of this group is less optimal compared with other age groups.¹

Studies repeatedly show that living with diabetes comes with a profound emotional burden.⁶ Consequently, young people with T1DM are at significant risk for psychological comorbidity: high prevalence of depressive symptoms, anxiety and eating disorders have been reported in this group.^{7–10} Apart from these clinically defined psychological diagnoses, diabetes distress is more and



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more recognized as an important indicator for mental health.^{11 12} Diabetes distress is defined as ‘the negative emotional or affective experience resulting from the challenge of living with the demands of diabetes...’.¹³ It is associated with unfavorable clinical and psychosocial outcomes and lower levels of self-management.¹² Different studies found the number of people experiencing diabetes distress to be higher than the number of people experiencing clinically established psychological disorders such as depression.^{13 14} Therefore, diabetes distress can be a more useful indicator to gain insight into well-being and needs of (young) people with diabetes, and to prevent further deterioration of mental health in this group. This is why international diabetes guidelines recommend regular screening on diabetes distress in clinical practice.¹²

Diabetes distress has been topic of research for at least 25 years now,¹³ but relatively little research has specifically addressed young adults.^{6 15} The few studies that explored this area showed that diabetes distress is significantly more present in the young adult group compared with other age groups^{5 6} and that more diabetes distress in young adults with T1DM is associated with impaired health outcomes (eg, higher hemoglobin A1c (HbA1c) levels)^{15 16} and less frequent blood monitoring by young adults.¹⁷ Garvey and colleagues found that feeling prepared for transition to adult care was associated with lower diabetes distress in young adults with T1DM.¹⁸ However, this is the only study that addressed transition and diabetes distress. Since transition is an important phase to support the development of self-management skills and to prepare young people for adult life while maintaining good quality of life, it is useful to further explore the differences between young people with and without diabetes distress regarding transfer experiences, self-management and health-related quality of life. Such insights could help to tailor self-management support to the needs of young adults. This study aimed to explore these differences in a sample of Dutch young adults (aged 18–25 years) with T1DM.

RESEARCH DESIGN AND METHODS

Participants and setting

This study used data from a larger evaluation study of transitional care for young adults with T1DM in the Netherlands.^{19 20} The study involved 12 hospitals and was conducted between April 2016 and October 2018; its study protocol has already been published,²¹ as is the evaluation study itself.¹⁹ In the current study, results from an online questionnaire on diabetes distress, health-related quality of life, self-management and transfer experiences were explored. Participants were young adults (aged 18–25 years) with a confirmed diagnosis of T1DM who had made the transfer to adult services in 2012–2014, had no cognitive impairment and were able to speak and read Dutch. In our evaluation study, we found that the care facilities could be divided into two

groups: HI-ATT (with high attention for transitional care and thus better alliance between pediatric and adult care settings) and LO-ATT (the opposite). We established that there were hardly any significant differences between the two groups related to outcomes in young adults, such as health-related quality of life and self-management. Extra analysis (not published) also showed that there were no differences in diabetes distress. That is why in this study, we did not differentiate between teams with more or less transitional care facilities.

Data collection

The participating hospitals invited potential respondents by email in October 2017. They were asked to fill out an online questionnaire on diabetes distress, health-related quality of life, self-management and transfer experiences. Most invited people had transferred to adult services in the period 2012–2014, but five hospitals also invited young adults who had transferred in 2015–2016. Reminders were sent after 2 and 4 weeks. Every third respondent received a €20 gift voucher. Those who eventually participated, consented with the use of their data and linking these to data from their electronic charts for this study.

Measures

Background characteristics

The following sociodemographic characteristics were addressed in the questionnaire: age (years); gender (male/female); highest completed educational level at time of the study (low (ie, junior vocational or secondary general low) and medium (ie, secondary general high or senior vocational) versus high (higher educational institutions)); employment status (paid or volunteer job, yes vs no) and living status (with parents vs independent). Time since transfer (first consultation in adult care) had been collected in the overarching evaluation study from participants’ medical records.

Diabetes distress

Diabetes distress was measured with the Dutch version of the validated 5-item Problem Areas in Diabetes (PAID-5) scale.^{22 23} Items are scored on a 5-point Likert scale: 0=not a problem, 1=minor problem, 2=moderate problem, 3=somewhat serious problem, 4=serious problem. The theoretical range is 0–20; higher scores indicate more stress. Cronbach’s α in the current study was 0.89. For the categorization of groups into ‘with diabetes distress’ and ‘without diabetes distress’, a score of eight points or higher on the sum score was classified as having diabetes distress.²²

Transfer experiences

Transfer experiences were measured using the validated 20-item On Your Own Feet—Transfer Experiences Scale.²⁴ Items are scored on a 5-point Likert scale: 1=strongly disagree, 2=disagree, 3=do not agree/do not disagree, 4=agree, 5=strongly agree. Higher scores indicate more positive transfer experiences. Cronbach’s α for the total

scale was 0.91 in the current study. Subscales included: reception in adult care (four items, $\alpha=0.88$, alliance between pediatric and adult care (five items, $\alpha=0.87$), preparation for the transfer (three items, $\alpha=0.70$), readiness to transfer (six items, $\alpha=0.81$) and youth involvement (two items, $\alpha=0.60$).

Self-management

Self-management skills were measured with the validated 12-item Partners in Health scale.²⁵ Items are scored on a 9-point Likert scale, where a higher score indicates better self-management skills. Cronbach's α in the current study was 0.74.

Health-related Quality of Life

Health-related Quality of Life (HRQoL) was measured using the Dutch version of the validated 23-item Pediatric Quality of Life Inventory for Young Adults.²⁶ Items are scored on a 5-point Likert scale: 0=never, 1=almost never, 2=sometimes, 3=often, 4=almost always. Each answer is reversely scored and rescaled to a 0–100 scale (0=100, 1=75, 2=50, 3=25, and 4=0). Higher scores indicate better HRQoL. Cronbach's α for the total scale was 0.94 in the current study. Subscales included: physical health (eight items, $\alpha=0.88$, emotional functioning (five items, $\alpha=0.86$), social functioning (five items, $\alpha=0.80$) and school/work functioning (five items, $\alpha=0.80$).

Data analysis

Descriptive analyses were performed to describe the study sample's background characteristics. Differences on background characteristics between non-responders and responders were calculated using t-tests and Pearson χ^2 tests. The study sample was divided into two groups on the basis the PAID-5 scale score: (1) with diabetes distress (score ≥ 8) and (2) without diabetes distress (score < 8). Differences between these two groups were explored with t-tests and Pearson χ^2 tests and effect sizes (Cohen's *d*) were calculated; descriptive analyses (frequencies) were used to explore diabetes distress within these groups (at item level). Missing data were excluded listwise.

RESULTS

Response and background characteristics

Four hundred and thirty-five young adults with T1DM were invited to participate in the study. Of these, 177 (40.7%) filled out the questionnaire and 164 (37.7%) provided information about diabetes distress. Non-responders did not differ from the responders in age (mean \pm SD in years: 23.6 \pm 1.40 vs 23.9 \pm 1.49; $p=0.158$), but they were more often men (64.0% vs 39.3%; $p<0.001$). A description of the study sample is given in table 1.

Diabetes distress

The total study sample ($n=164$) scored low on problem areas in diabetes (6.52 \pm 4.67, range: 0–17) but 57 of them (34.8%) had a score ≥ 8 , indicating diabetes distress. These young adults with diabetes distress had a mean

Table 1 Background characteristics of total study sample ($n=164$)

Characteristic	Mean (\pm SD) or Frequency (%)
Age (years) (range 19–28 years)	22.7 (\pm 1.56)
Time since transfer* (3–6 years)	4.7 (\pm 1.13)
Gender†	
Female	99 (60.7%)
Male	64 (39.3%)
Educational level†	
Low/middle	125 (76.7%)
High	38 (23.3%)
Paid or volunteer job†	
Yes	137 (84.0%)
No	26 (16.0%)
Living†	
With parents	99 (60.7%)
Independent (on own)	64 (39.3%)

* $n=112$.

† $n=163$.

distress score of 11.88 (\pm 2.99) vs 3.66 (\pm 2.27) in those without diabetes distress ($p<0.001$). 'Worrying about the future and the possibility of serious complications' was most often mentioned as a serious problem by those with diabetes distress (67%; figure 1). Those without diabetes distress also worried about this, but only 5% considered it a serious problem. 'Feeling that diabetes is taking up too much of your mental and physical energy every day' was relatively often mentioned as a serious problem as well by 58% of those with diabetes distress. Comparisons of those with and those without diabetes distress on the PAID-5 showed substantial and significant differences between both groups (figure 1).

Differences on background characteristics and outcomes

Interestingly, there were hardly any differences in background characteristics between young adults with and without diabetes distress (table 2). However, the latter were significantly younger ($p=0.038$).

As for transfer experiences, young adults with diabetes distress overall had less positive experiences ($p=0.002$). More specifically, they rated the reception in adult care ($p=0.013$) and the alliance between pediatric and adult care ($p=0.029$) lower and felt less ready to transfer to adult care ($p<0.001$). They also reported worse outcomes on self-management ($p<0.001$) and overall HRQoL ($p<0.001$) compared with young adults without diabetes distress. HRQoL of those with diabetes distress was significantly lower on all domains, that is, physical functioning, emotional functioning, social functioning and school/work functioning. Effect sizes differed from medium to large for all outcome measures. Large effect sizes were

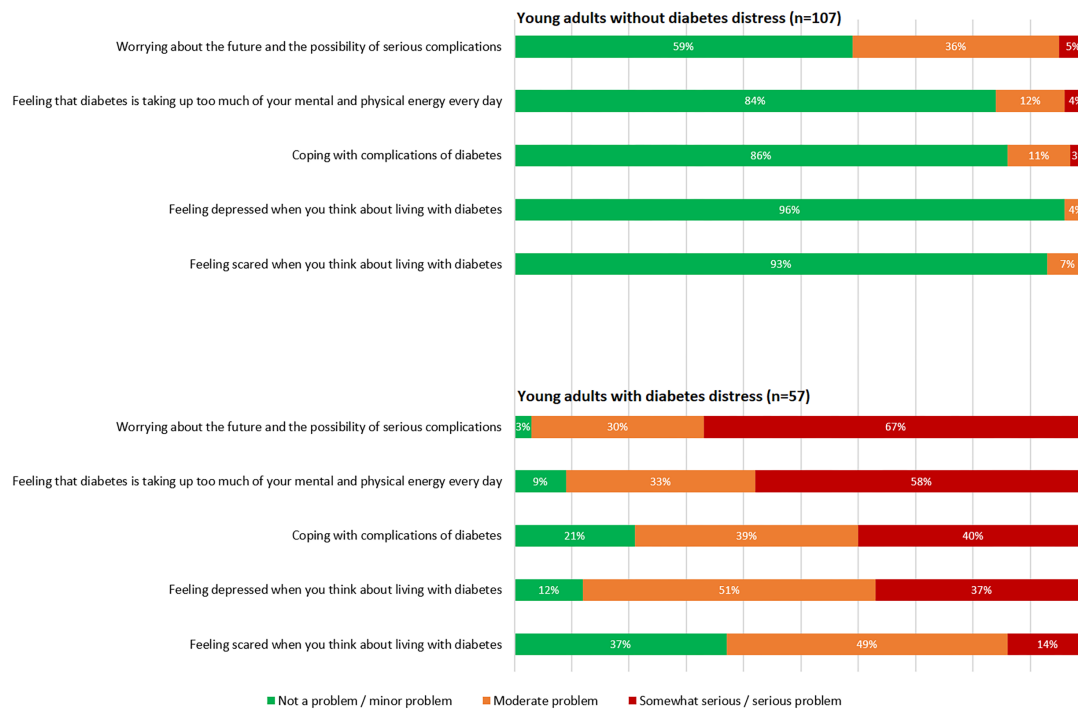


Figure 1 PAID-5 scores in young adults with and without diabetes distress. PAID, Problem Areas in Diabetes scale.

particularly found for emotional, social and school/work functioning (table 2).

DISCUSSION

General discussion

This study found substantial differences in transfer experiences, self-management and HRQoL between young adults with T1DM with and without diabetes distress. Those without diabetes distress reported better transfer experiences and better self-management skills and higher HRQoL compared with those with diabetes distress. Young adults with diabetes distress were slightly older; no other differences in background characteristics were found.

Almost 35% of the participants reported diabetes distress. The estimated prevalence of diabetes distress in other studies is 28% in adolescents and emerging adults with T1DM²⁷ and 32% in young adults with T1DM,²⁸ which is comparable to the present study. The outcome that those with diabetes distress were older than those without is congruent with the finding of Lašaitė and colleagues (2016) that emerging young adults more often experienced psychological distress compared with adolescents.²⁷ A study in over 19-year-olds with T1DM found that younger adults more often experienced diabetes distress than older adults.²⁹ Probably, the developmental challenges of young adulthood add to the distress experienced from T1DM itself, which warrants specific attention from healthcare professionals. Wentzell and colleagues indeed concluded in their integrative review that diabetes distress 'is embedded within the developmental challenges specific to living with T1DM

during this phase [i.e. emerging adulthood]'.⁶ In this light, support of parents is also an important topic. Our study revealed that less young adults who still lived with their parents had diabetes distress compared with those who lived independently. Shaw and colleagues (2021) recently emphasized the (often overlooked) role of parents in promoting young adults' well-being.³⁰

Worrying about the future and the possibility of serious complications were most often reported as (serious) problems by people with T1DM, in our study and in previous studies as well.¹³ In general, distress in people with T1DM seems to be related to emotions (worries, feelings and so on) rather than difficulties in practically managing the condition and its treatment.³¹ However, insights into specific sources of distress in young adults with T1DM is lacking. In this study, the feeling that diabetes is taking up too much mental and physical energy every day, is the second most often mentioned problem, both by respondents with and those without diabetes distress. This problem was also highlighted in our study on HRQoL in these young adults, of whom one quarter reported fatigue due to T1DM.³² Ways to counteract the feeling that diabetes management takes up too much mental and physical energy would benefit young adults with T1DM. Nevertheless, Barry-Menkhaus and colleagues recently emphasized that much work is still needed to find pragmatic and efficient ways to improve self-management of young adults with T1DM.³³ Another systematic review also highlights the need to develop effective interventions for this group.³⁴

As for the differences in outcomes between young adults with and without diabetes distress, the latter

Table 2 Differences between young adults with and without diabetes distress (mean (\pm SD) or frequency (%))

	Young adults with diabetes distress (n=57)	Young adults without diabetes distress (n=107)	Effect size	P value
<i>Background characteristics</i>				
Age	23.00 (\pm 1.67)	22.47 (\pm 1.48)	0.33	0.038
Time since transfer (3–6 years)*	4.44 (\pm 1.25)	4.82 (\pm 1.03)	0.33	0.087
Gender†				
Female	39 (68.4%)	60 (56.6%)	–	0.141
Male	18 (31.6%)	46 (43.4%)		
Educational level‡				
Low/middle	44 (78.6%)	81 (75.7%)	–	0.681
High	12 (21.4%)	26 (24.3%)		
Paid or volunteer job‡				
Yes	46 (82.1%)	91 (85.0%)	–	0.318
No	10 (17.9%)	16 (15.0%)		
Living‡				
With parents	30 (53.6%)	69 (64.5%)	–	0.188
On own	26 (46.4%)	38 (35.5%)		
<i>Transfer experiences (OYOF-TES)</i>				
Reception in adult care	3.72 (\pm 0.993)	4.04 (\pm 0.675)	0.38	0.013
Alliance between pediatric and adult care	2.92 (\pm 0.899)	3.23 (\pm 0.857)	0.35	0.029
Preparation for the transfer	2.88 (\pm 0.925)	3.02 (\pm 0.931)	0.15	0.323
Readiness to transfer	3.76 (\pm 0.710)	4.19 (\pm 0.576)	0.67	<0.001
Youth involvement	3.12 (\pm 1.02)	3.39 (\pm 0.955)	0.27	0.101
Total score	66.88 (\pm 13.94)	73.35 (\pm 11.25)	0.51	0.002
<i>Self-management</i>				
PIH	76.09 (\pm 8.81)	82.22 (\pm 7.91)	0.73	<0.001
<i>HRQOL (PedsQL-YA)</i>				
Physical functioning	79.11 (\pm 15.30)	87.21 (\pm 15.51)	0.53	0.002
Emotional functioning	52.81 (\pm 17.70)	80.28 (\pm 17.43)	1.56	<0.001
Social functioning	78.51 (\pm 16.26)	90.23 (\pm 12.69)	0.80	<0.001
School/work functioning†	61.40 (\pm 19.43)	75.99 (\pm 16.70)	0.81	<0.001
Total score	69.41 (\pm 14.35)	83.96 (\pm 13.16)	1.06	<0.001

*n=112: Young adults with diabetes distress n=41, without diabetes distress n=71.

†n=163: Young adults with diabetes distress n=57, without diabetes distress n=106.

‡n=163: Young adults with diabetes distress n=56, without diabetes distress n=107.

HRQOL, health-related quality of life; OYOF-TES, On Your Own Feet Transfer Experiences Scale; PedsQL-YA, Pediatric Quality of Life Inventory—Young Adult; PIH, Partners in Health.

reported more positive outcomes on all measures. They had better transfer experiences, more self-management skills and higher HRQoL. It is important to note that it is still not clear whether less positive transfer experiences add to diabetes distress or – vice versa – whether diabetes distress makes young adults more negative towards past experiences, including transfer. The only previous study that explored transition and diabetes distress showed that feeling prepared for the transfer to adult care was associated with lower diabetes distress in young adults.¹⁸ However, our results suggest that rather than preparation

for transfer, reception in adult care, alliance between pediatric and adult care and readiness to transfer seem to be important for young people with T1DM. This suggests that the transfer in care adds to stress experiences by young adults. More research is needed to clarify the sources of diabetes distress in young adults with T1DM.

The finding that young adults without diabetes distress score significantly better on self-management than do those with diabetes distress is not surprising. Self-management encompasses three tasks: medical management, role management and emotional management.³⁵

An association between lower self-care skills (medical management) and diabetes distress is consistently found in other age groups.¹³ Less is known about the effect of diabetes distress on social participation (ie, the degree to which someone takes up roles in society in different areas like school, work, relationships and so on), but we do know that young adults with social participation rates comparable to those of their healthy peers, generally report lower HRQoL.³⁶ Social functioning and school/work functioning were indeed significantly and substantially lower in the group with diabetes distress in this study. More research is needed to further understand the relation between diabetes distress and social participation of young adults with T1DM. Emotional management involves, among other things, stress management, positive thinking and mental well-being.³⁷ The largest negative effect of diabetes distress was indeed found for emotional functioning. More diabetes distress, therefore, could impair one's self-management skills.

The question is how to support young adults with T1DM who have diabetes distress. Apart from more insight into their sources of stress and support needs, studies researching distress in adults can provide recommendations that might be valid for young adults as well. First, as mentioned above, it is essential to regularly screen young adults for diabetes distress and have attention for and discuss any worries. Second, it is important to consider what healthcare professionals can do to counteract diabetes distress in young adults with T1DM.¹³ A recent systematic review suggested that experiencing good communication and positive experiences with healthcare professionals is found to be associated with reduced diabetes distress and better self-management outcomes.³⁸ Previous studies in young people with chronic conditions have linked positive relationships with and trust in healthcare professionals with more positive outcomes as well.^{39 40} Fisher and colleagues emphasize specific communication styles that healthcare professionals can apply to reduce diabetes distress.³⁸ These recommendations are valid for both pediatric and adult care professionals. Transitional care based on good collaboration between pediatric and adult care and harboring a warm reception in adult care could contribute to alleviate the burden for young adults.²⁰

Strengths and limitations

This study addressed a gap in current literature and provided unique insights into diabetes distress around transfer in care. Another strength is the nationwide representation of young adults with T1DM. A possible limitation is that respondents were more often women; some studies suggest there is a significant association between higher diabetes distress and female gender in emerging adults, but the relationship between diabetes distress and gender is not clear.⁶ Therefore, we cannot tell if and how the underrepresentation of males in our study may have influenced our results. Another limitation is that the cross-sectional study design did not allow

for exploration of causality; longitudinal studies are needed to identify determinants of diabetes distress in young adults with T1DM. Also, while the response rate in this study was rather low, it appears to be comparable to similar (post transition) studies in young people with T1DM.⁴¹ Finally, this study did not include any information about diabetes control and the impact on transfer, because the survey and available medical record data were collected in different time periods. Our evaluation study¹⁹ revealed that mean HbA1c scores did not change over time ($p=0.836$); they were elevated across the whole study period with no significant differences between pediatric and adult care. It seems relevant to study the association between diabetes control and diabetes distress in future, longitudinal studies.

CONCLUSION

In this study, more than one-third of the young adults with T1DM experience diabetes distress after transfer to adult care. Differences in diabetes distress are substantial and so are the consequences of having diabetes distress. Those with diabetes distress had less positive transfer experiences, suggesting that the transfer in care could be a source of diabetes distress in young adults. Specific attention is required for reception in adult care, alliance between pediatric and adult care and readiness to transfer. Young adults with diabetes distress also reported less self-management skills and lower HRQoL compared with those without diabetes distress. Structured screening of diabetes distress and attention for and addressing young adults' worries is recommended in both pediatric and adult care.

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Competing interests None declared.

Patient consent for publication Not applicable.

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Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement All data relevant to the study are included in the article.

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