Agreement between Self-Reports and Medical Records in the Reporting of Pregnancy Outcomes in Women with Type 1 Diabetes

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Abstract

Background: Since previous studies have shown that self-reports may not be as accurate as medical record data, this secondary analysis investigated the agreement between maternal self-report (SR) and medical record (MR) documentation of pregnancy outcomes (e.g., length of hospitalization, neonatal birth weight, complications) in women with type 1 diabetes (T1D).

Methods: An online SR follow-up study to evaluate long-term reproductive-health outcomes was conducted with women who previously participated in randomized controlled trials (RCT) to test a preconception counseling (PC) intervention (READY-Girls) as adolescents and included a matched comparison group of women with T1D who did not receive the PC intervention as teens. Data were collected from SR surveys and MR review. Agreement between SR and MR entries was assessed using kappa coefficients and intra-class correlation coefficients (ICC).

Results: Of the 101 women with T1D (51 RCT, 50 matched controls) who were recruited for long-term follow-up (98.0% Caucasian, age range 18-34 years, 82.0% had some college, T1D duration range 0-30 years), 18 (17.8%) reported ever being pregnant and for 8 (47.1%) of these women access to MR was obtained for 14 pregnancies. These women had a mean±SD age of 28.57±1.91 years at entry and duration of T1D of 19.75±5.45 years. Perfect agreement (kappa=1.0) was observed between SR and MR for type of delivery and neonatal birth weight >9 pounds. Excellent agreement was observed for neonatal weight (ICC=.951). Lower levels of agreement were found between SR and MR for duration of post-delivery hospitalization [maternal (ICC=.269); neonatal (ICC=.657)] and maternal and neonatal complications (kappa=.639 and kappa=.025, respectively).

Conclusion: There was excellent agreement for several pregnancy outcomes in these women with T1D. Self-report of pregnancy outcomes should be verified by medical record data, especially time-related variables which could be more difficult to recall, suggesting that method of reporting for these types of variables may not be interchangeable.

Keywords: Type 1 diabetes, medical records, self-report, pregnancy outcomes

Introduction

Women with diabetes have increased risks associated with pregnancy [1,2]. Accurate data regarding pregnancy outcomes is imperative in this population. Therefore, selection of an ac-
curate data collection method for these outcomes became a primary concern of our study, and an aim of evaluation. Our program of research is based on the following premise. To reduce the risks of complications with reproductive health in women with diabetes, the American Diabetes Association (ADA) recommends preconception counseling (PC) to start at puberty as part of standard care and it includes the following: 1) counseling about diabetes and pregnancy, specifically the malformation risk associated with unplanned pregnancies and poor metabolic control, and 2) effective use of contraception until metabolic control is attained/maintained (A1c <7%) and conception occurs [3,4]. Preconception counseling and care has reduced the risk of congenital malformations [5,6] and has been associated with improved perinatal and maternal outcomes [6,7].

Our program of research, called READY-Girls (Reproductive-health Education and Awareness of Diabetes in Youth for Girls), was tailored for teens with type 1 diabetes (T1D), and later (2005) modified for type 2 diabetes (T2D), to promote effective family planning decisions and empower them to seek preconception care when planning a pregnancy. READY-Girls is a validated developmentally appropriate evidence-based educational PC intervention for adolescent girls starting at puberty [8-10]. Three randomized controlled trials (RCT) at multiple diabetes clinic sites using the READY-Girls intervention have been conducted (1999, 2002, and 2005) on teens 13-20 years of age and results have been previously reported [11-13].

Materials and methods

This study was a secondary analysis conducted to evaluate the agreement between self-reported (SR) data and medical record (MR) data for maternal and neonatal complications. The parent study was a long-term follow-up study employing a prospective, repeated measures design which included previous participants at one site in southwestern Pennsylvania from a possible pool of 115 participants across the three READY-Girls RCTs who were at least 18 years of age and had T1D. This site is a large pediatric tertiary care research intensive institute which maintains a pre-established registry for pediatric patients with T1D born since 1955. Other participants recruited from this registry served as the comparison group. These women with T1D were matched to the READY-Girls RCT participants according to age (within 2 years), race (white, nonwhite), and duration of T1D (within 2 years). After Institutional Review Board (IRB) approval, both the previous READY-Girl participants and the potential comparison women were recruited through phone calls, mailed letters and face-to-face contact by the project nurse in the diabetes clinic. Previous consent forms from the RCTs contained statements allowing former participants to be re-contacted. The project nurse explained the study in detail, consented the participants and provided web access instructions for the follow-up surveys. Participants who identified a pregnancy in the baseline or subsequent follow-up surveys (at 6, 12 and 18 months) were contacted by letter stating the importance and rationale to a follow-up medical record review of their pregnancy. At that time, participants were asked to sign and return consent for the ascertainment of their medical records and those of their baby. However, not all participants allowed access to their medical records. Upon receipt of the consent, a research associate sent a copy of the consent and a letter to the identified hospital for the medical records for the reported pregnancy. After review of the medical record, the first research associate entered the data onto a TELEform and the second research associate visually verified the data entry after an independent review of the medical record.

Pregnancy outcomes (maternal and neonatal outcomes) were measured by self-report and medical records for each pregnancy documented. Interval or ratio level outcomes included maternal and neonatal length of hospital stay and neonatal weight (actual weight in pounds). Nominal scaled outcomes included any maternal and fetal complications, type of delivery, pregnancy outcome (i.e., live birth, stillborn, miscarriage/elective abortion), and neonatal weight > 9 pounds. Demographic and clinical data were also obtained (age, race, education, income, health insurance, living arrangements, marital status, religion, age at T1D diagnosis, duration of T1D) for sample description.

Based on a variable's level of measurement and observed data distribution, appropriate summary statistics were used to describe the total sample and subsamples based on pregnancy status and whether medical records could be obtained. Group comparative statistics (two-sample t-test or Mann-Whitney U-test for continuous type variables and contingency table analysis with chi-square test of independence with exact
estimation of p-values for categorical variables) were used to compare the subsamples based on self-reported pregnancy status and whether maternal medical records were able to be obtained for women who were ever pregnant. Agreement between mother’s self-report and the report in the medical record for selected pregnancy outcomes and maternal and neonatal complications was assessed using either unweighted or weighted kappa coefficients for categorical outcomes and intra-class correlation coefficients (ICC) for continuous type outcomes. Empirical work suggest poor agreement for kappa coefficients less than .40, fair to good agreement for kappa coefficients ranging from .40 to .75, and excellent agreement for kappa coefficients greater than .75, with kappa coefficients equal to 1 indicating perfect agreement.

Results
Of the 101 participants with T1D in the follow-up study, 18 (17.8%) self-reported ever being pregnant, with a total of 35 pregnancies being reported at the baseline assessment or during the 18-month follow-up period of the parent study. Women who reported ever being pregnant were demographically and clinically similar to women who reported never being pregnant (p≥.05), except with regards to marital status, living arrangements, age at entry into the follow-up study, and duration of T1D (p<.05). As anticipated, women who reported ever being pregnant were more likely to be married (p<.001), currently having a husband or boyfriend (p=.011), living with their husband (p<.001) and/or biological children (p<.001), and were older (p<.001) and had T1D longer (p<.001) than women who reported never being pregnant.

As reported in Table 1, the women (n=8) providing both SR and MR data were similar to those women providing only SR data (n=10), except for marital status where women providing both sources of data were more likely to be married (p=.036). Examining the self-reported information regarding the 35 pregnancies, most (n=24, 77.4%) were self-reported as deliveries greater than 20 weeks. The distribution of possible pregnancy outcomes was similar between those women providing MR data and those who did not (p=.481).

Focusing on women with both data sources, we examined the agreement in the reporting of selected pregnancy outcomes (see Table 2). Of the 14 pregnancies for which MR data were obtained, most were (n=13, 92.9%) deliveries greater than 20 weeks. The remaining pregnancy was self-reported.

Table 1. Comparisons between participants who have ever been pregnant with and without medical record (MR) data on selected characteristics provided through self-report (SR) (n=18).

| Characteristic                        | Total (n=18) | SR only (n=10) | SR and MR (n=8) | Test Statistic | p-value |
|---------------------------------------|-------------|---------------|----------------|---------------|---------|
|                                      | Mean±SD     | Mean±SD       | Mean±SD        | t-test        |         |
| Race (Caucasian)                      |             |               |                |               |         |
| (n=18)                                |             |               |                |               |         |
|                                      | 18 (100)    | 10 (100)      | 8 (100)        | --            | --      |
| Age at First Pregnancy (years) (n=17) | 25.00±3.89  | 25.00±3.81    | 25.00±4.24    | t=0.000       | p=1.000 |
| Age at Entry (years)                  | 28.39±4.07  | 28.10±5.32    | 28.57±1.91    | t=-0.358      | p=.726  |
| Age at T1D Diagnosis (years)          | 8.72±4.90   | 8.60±4.95     | 8.87±5.17     | t=-0.115      | p=.910  |
| Duration of T1D (years)               | 19.67±6.17  | 19.60±7.00    | 19.75±5.45    | t=-0.050      | p=.961  |
| Marital Status                        |             |               |                |               |         |
| Never Married                         | 5 (27.8)    | 5 (50.0)      | 0 (0)          | X²=6.188      | p=.036  |
| Married                               | 12 (66.7)   | 5 (50.0)      | 7 (87.5)       |               |         |
| Divorced/Separated/Widowed            | 1 (5.6)     | 0 (0)         | 1 (12.5)       |               |         |
| Highest Level of Education            |             |               |                | X²=1.800      | p=.671  |
| High School                           | 3 (16.7)    | 1 (10.0)      | 2 (25.0)       |               |         |
| Some College/Trade School             | 6 (33.3)    | 4 (40.0)      | 2 (25.0)       |               |         |
| College                               | 3 (16.7)    | 1 (10.0)      | 2 (25.0)       |               |         |
| Graduate school                       | 6 (33.3)    | 4 (40.0)      | 2 (25.0)       |               |         |
| Income (n=17)                         |             |               |                | X²=5.627      | p=.071  |
| <$40,000                              | 6 (35.3)    | 5 (55.6)      | 1 (12.5)       |               |         |
| $40,000-$70,000                       | 3 (17.6)    | 0 (0)         | 3 (37.5)       |               |         |
| ≥$70,000                              | 8 (47.1)    | 4 (44.4)      | 4 (50.0)       |               |         |
| Health Insurance (n=16)               |             |               |                | X²=0.356      | p=.511  |
| Private                               | 12 (75.0)   | 7 (70.0)      | 5 (83.3)       |               |         |
| Medicaid/Medical Assistance/Medicare  | 4 (25.0)    | 3 (30.0)      | 1 (16.7)       |               |         |

T1D=Type 1 diabetes
as an elective abortion greater than 20 weeks; however, the pregnancy was listed as a stillbirth with a major brain/spinal anomaly in the medical record (kappa=.482). Perfect agreement (kappa=1.000) between SR and the MR report was observed for the type of delivery and whether the baby’s weight was greater than 9 pounds. Although mothers on average tended to underreport their baby’s weight relative to that reported in the medical record, the agreement in reporting the neonatal actual weight was excellent (ICC=.951). In contrast, mothers on average reported longer lengths of hospital stay following delivery for both their baby as well as themselves compared to the medical record, with good agreement for the baby’s length of stay (ICC=.657) but poor agreement for their own post-delivery hospital stay (ICC=.296). Regarding maternal complications, the same frequency distributions were observed for both SR and MR with good agreement between mother’s SR and MR (kappa=.639). Although the same frequency distributions were observed for neonatal complications for SR and MR, very poor agreement was observed (kappa=.025).

Discussion

Accuracy of reporting outcomes surrounding pre-pregnancy, pregnancy and delivery is necessary in determining the cost-effectiveness and efficacy of preconception counseling programs in women with diabetes. While maternal reporting of perinatal events is assumed to be accurate, review of medical records is the gold standard for assuring the reliability for cost data. However, medical record reviews can be costly and time-consuming. As an exploratory aim of this study, medical record reviews of maternal and neonatal outcomes of pregnancies among former READY-Girls participants and a comparison group was conducted to determine the long-term cost-benefit of READY-Girls. Since not all participants allowed access to their medical records, the purpose of this secondary analysis was to investigate agreement between self-report and medical records for maternal and neonatal outcomes in women with T1D.

Several of the pregnancy outcomes had high agreement. Similar to what we observed for neonatal weight-related outcomes as previously reported in the literature, maternal recall of neonatal birth weight was highly accurate [21]. The high to perfect level of agreement between the SR and MR data for several of the pregnancy outcomes in these participants with T1D is similar to previous studies showing excellent agreement for mode of delivery [22,23]. These findings suggest that self-report of some pregnancy outcomes, such as type of delivery and neonatal weight, can be accurate.

Pregnancy outcomes having good agreement were neonatal length of hospital stay and maternal complications. Mothers may be accurate in recalling neonatal length of hospital stay as this occurs solely post-delivery [24].

Lastly, poor agreement was noted for maternal length of hospital stay post-delivery and neonatal complications. For durations of stay in the hospital mother’s perceptions of time

| Table 2. Agreement between Self-Report and Medical Record Data for Selected Pregnancy Outcomes (n=14 pregnancies). |
|---------------------------------------------------------------|
| **Pregnancy Outcome**                  | **Self-Report Mean±SD or n (%)** | **Medical Record Mean±SD or n (%)** | **Measure of Agreement** |
| Delivery (> 20 weeks)                  | 13 (92.9)                         | 13 (92.9)                          | kappa=.482               |
| Stillborn                              | 0 (0)                             | 1 (7.1)                            |                           |
| Miscarriage/Elective Abortion          | 1 (7.1)                           | 0 (0)                              |                           |
| **Type of Delivery (n=13)**            |                                    |                                    | kappa=1.000               |
| Vaginal                               | 6 (46.2)                          | 6 (46.2)                           |                           |
| Cesarean                              | 7 (53.8)                          | 7 (53.8)                           |                           |
| **Neonatal Weight at Birth >9 pounds (n=13)** |                                    |                                    | kappa=1.000               |
| No                                     | 9 (69.2)                          | 9 (69.2)                           |                           |
| Yes                                    | 4 (30.8)                          | 4 (30.8)                           |                           |
| **Neonatal Weight at Delivery (pounds) (n=13)** | 7.67±1.70 | 7.91±1.47 | ICC=.951 |
| **Maternal Length of Stay Following Delivery (days) (n=13)** | 2.92±1.44 | 2.00±1.08 | ICC=.296 |
| **Neonatal Length of Stay (days) (n=13)** | 3.62±2.02 | 3.08±1.66 | ICC=.657 |
| **Maternal Complications (n=13)**      |                                    |                                    | kappa=.639               |
| No                                     | 4 (30.8)                          | 4 (30.8)                           |                           |
| Yes                                    | 9 (69.2)                          | 9 (69.2)                           |                           |
| **Neonatal Complications (n=13)**      |                                    |                                    | kappa=.025               |
| No                                     | 5 (38.5)                          | 5 (38.5)                           |                           |
| Yes                                    | 8 (61.5)                          | 8 (61.5)                           |                           |

ICC=intra-class coefficient correlation
may be distorted due to recall and may not be consistent with a hospital’s recording of length stay [19]. For mothers the total length of hospital stay may include periods of pre-delivery, delivery and post-delivery [22], and mothers may not be able to specifically recall time during the post-delivery period. Regarding neonatal complications, our finding of poor agreement were consistent with previous results [22]. In our study, mother’s self-report on neonatal complications was more general, and agreement might have been better had we targeted specific complications (e.g., jaundice, hypoglycemia).

This study had several limitations that should be acknowledged, including the small sample size, limited variability as to the type of pregnancy outcome (i.e., most were deliveries), and only a limited detail as to the type of maternal and neonatal complication based on mother’s self-report.

Conclusions
Self-report of pregnancy outcomes should be verified by medical record data whenever possible, especially time-related variables which could be more difficult to recall. There was excellent agreement between the self-report and medical record data. Further analysis of specific maternal/neonatal complications is needed to determine differences between the two report types.

Competing interests
The authors declare that they have no competing interests.

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Authors’ contributions

| Authors’ Contributions                  | ARF | SMS | WHH | LMcE | DB | CMP | DCP | FPB |
|----------------------------------------|-----|-----|-----|------|----|-----|-----|-----|
| Research Concept and Design            | ✓   | ✓   |     |      |    |     |     |     |
| Collection and/or assembly of data     | ✓   |     |     |      |    |     |     |     |
| Data analysis and interpretation       | ✓   | ✓   |     | ✓    |    | ✓   |     |     |
| Writing the article                    | ✓   | ✓   |     |     |    | ✓   |     |     |
| Critical revision of the article       | ✓   | ✓   | ✓   | ✓    |    | ✓   | ✓   | ✓   |
| Final approval of article              | ✓   | ✓   | ✓   | ✓    |    | ✓   | ✓   | ✓   |
| Statistical Analysis                   | ✓   | ✓   |     |      |    |     |     |     |

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