- The impact of diabetes on labour market outcomes in
- Mexico: a panel data and biomarker analysis

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Abstract

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Recent evidence for Mexico suggests important differences in health status between people with diagnosed and undiagnosed diabetes. However, there is at best scarce evidence on the economic consequences of diabetes, especially in contexts where the condition often remains undiagnosed, as is typically the case in low- and middle income countries. Using Mexican longitudinal as well as biomarker data we estimated the impact of diabetes and diabetes duration on employment probabilities, wages and working hours. We further explored how these effects differed for those aware and those unaware of their with diagnosed and undiagnosed diabetes. For the longitudinal analyses nationally representative data from 11836 men and 13745 women 15 to 64 years old were taken from three waves (2002, 2005, 2009) of the Mexican Family Life Survey. We estimated a within-between effects-fixed-effects model to account for unmeasured time-invariant confounders of diabetes while simultaneously assessing differences between subjects with and without diabetes. The within effects indicated. We found a reduction in the probability of being employed between 5.4 and 6.0 percentage points for men and women, respectively, but no effects on hours worked or wages. The between effects showed similar results. Employment probabilities fell gradually with each year since diagnosis. Using cross-sectional biomarker data, we observed that 68% of those exhibiting glycated hemoglobin (HbA1c) levels above the clinical diabetes threshold did not self-report a diagnosis, hence were undiagnosed. Nevertheless, regression analysis revealed that there was no association of diabetes with labour outcomes for undiagnosed women or men. This suggests that results based on self-reported diabetes cannot be extended to the considerable population with undiagnosed diabetes, likely because of a selection of people in worse health and with a longer diabetes duration into the diagnosed population. An earlier diagnosis and improved treatment of diabetes therefore may prevent adverse health effects and related economic hardship in Mexico.

1. Introduction

Diabetes, a disease characterized by elevated blood glucose levels due to the body's inability to use insulin properly, has in the last two decades increasingly become a problem for low- and middle-income countries (LMICs) as well as high-income countries (HICs), 35 with over two-thirds of people with diabetes living in the developing world (International Diabetes Federation 2015). In Mexico, diabetes prevalence is estimated to have grown 37 from 6.7% in 1994 to 14.4% in 2006 (Barquera, Campos-Nonato, et al. 2013) and 15.8% in 2015. Diabetes has become the number one contributor to mortality (International 39 Diabetes Federation 2015), by increasing the risk for heart disease and stroke, blindness, 40 kidney disease and nerve problems, food neurologic problems, foot ulcers and amputations 41 (Reynoso-Noverón et al. 2011). However, via effective self-management of the disease, 42 many if not all of the complications can be avoided through regular monitoring, behaviour 43 change and medication adherence, the occurrence of complications could be avoided or 44 delayed in many cases (Gregg et al. 2012; Lim et al. 2011). 45 The observed increase in diabetes incidence has been attributed to a deterioration in 46 diet and a reduction in physical activity (Barquera, Hernandez-Barrera, et al. 2008; Basu 47 et al. 2013), while genetic predisposition among Mexicans with pre-Hispanic ancestry may 48 also play a role (Williams et al. 2013). The onset of diabetes has been occurring at an 49 ever earlier age in Mexico (Bello-Chavolla et al. 2017), increasing the risk of complica-50 tions occurring during the productive lifespan. Only a minority of patients in Mexico 51 achieves adequate blood glucose control (Barquera, Campos-Nonato, et al. 2013). More-52 over, diabetes in Mexico coexist with high levels of infectious diseases, exposing is related to disease such as depression, hypertension and cardiovascular disease that burden the health system to a 'double-disease burden' increasing the pressure to identify treatment priorities and use existing resources efficiently (World Health Organization 2016). 56 Despite the catastrophic impact of diabetes on health, its economic consequences, in 57

particular in LMICs have received less attention, especially its effects on labour outcomes

(Anonymous 2007). The latter have been studied predominantly in high-income countries, 59 where substantial economic losses have been observed (Brown, Pagán, et al. 2005; Brown, 60 Perez, et al. 2011; Brown 2014; Latif 2009; Minor 2011, 2013; Minor and MacEwan 2016). 61 For LMICs less evidence is available. One study exploited a natural experiment in China 62 and found a significant reduction in income due to a recent diabetes diagnosis (Liu and 63 Zhu 2014). A study for Mexico, using cross-sectional data from 2005, found a significant (p<0.01) reduction in employment probabilities for males by 10 percentage points (p,p,0.01)and for females by 4.5 p.p. (p<0.1) (Anonymous 2007). Most existing studies rely on 66 instrumental variable (IV) estimation to address the potential endogeneity of diabetes 67 using the genetic component of diabetes based on its family history as an instrument. However, family history of diabetes may also proxy for other genetically transferred traits, including unobserved abilities, as well as intrahousehold or intergenerational dynamics 70 that impact labour outcomes directly; the validity of this IV therefore remains debatable. Panel data methods provide the opportunity to account for time-invariant unobserved individual characteristics, which may play an important role, but have not yet been used by our knowledge. Such unobservables, like for instance health endowments for instance hunger or nutrient deficiency experienced in early life, could adversely affect health as well 75 as the propensity to develop type 2 diabetes in particular later in life (Ewijk 2011; Li et al. 76 2010; Sotomayor 2013); they may also affect. Additionally, there may also be long-term 77 effects on labour outcomes—either directly through their effects on reductions in con-78 temporaneous productivity (Currie and Vogl 2013), or indirectly by limiting educational 79 attainment and human capital accumulation (Ayyagari et al. 2011). These unobservables 80 thereby present a major source of a potential bias that can be accounted for by the use of 81 panel data estimation. 82 In parallel to these identification challenges, heterogeneity in impact and measurement 83 across the population also deserves further investigation. Recent evidence from Mexico

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points to a strong positive relationship of diabetes duration with mortality due to diabetes

related with higher glycated hemoglobin (HbA1c) levels and undiagnosed diabetes had the 87 lowest diabetes related mortality risks. The latter points to potential selection issues when 88 using self-reported diabetes data to investigate economic outcomes as those. Those who 89 self-report, and hence tend to be diagnosed, are in worse health than those undiagnosed, 90 potentially leading. This can lead to an overestimation of the economic effects of diabetes, 91 in particular in populations with a large undiagnosed population, such as in many LMICs 92 (Beagley et al. 2014). So far, however, little evidence exists on the economic impact 93 according to diabetes severity and, duration or for those with undiagnosed diabetes. The objective of this study was to provide new evidence on the impact of diabetes on 95 labour outcomes, adding to previous work by paying close attention to the challenges of unobserved heterogeneity, to the chronic nature of diabetes and to the population unaware 97 of their condition (i.e. the 'undiagnosed ') undiagnosed diabetes. We used three waves of Mexican panel data the Mexican Family Life Survey (MxFLS), covering the period 2002— 2012. Applying a within-between model, which models individual fixed effects and between 100 effects separately, fixed effects model we accounted for time-invariant heterogeneity when 101 assessing the impact of self-reported diabetes and time since diagnosis on labour outcomes. 102 We also used rich and novel To assess the role of undiagnosed diabetes we used biomarker 103 data from the most recent wave of data, we assessed the role of undiagnosed diabetes last 104 wave of the MxFLS. 105

related complications (Herrington et al. 2018). A longer disease duration was found to be

106 2. Data

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This paper uses data from the Mexican Family Life Survey (MxFLS), a nationally representative longitudinal household survey, containing three waves conducted in 2002, 2005—2006 and 2009—2012. Data was collected It is the only longitudinal household survey in Mexico that provides data on a wide range of social, demographic, economic and health

characteristics (Rubalcava and Teruel 2013). Because the survey followed participants 111 moving within Mexico as well as to the US, around 90\% of the original sample have been 112 reinterviewed in the third wave. Our samples were restricted to the working age popula-113 tion (15–64) and excluded pregnant womenand those in school. Pregnant women have an 114 increased diabetes risk and may not be able to work. Since their inclusion may have biased 115 the estimates we dropped all observations of women reporting to be pregnant at the time 116 of the survey (N=764). We also dropped those reporting to be in school. The first part 117 of the analysis used all three waves and the panel structure of the data. The second part 118 used a biomarker subsample of the third wave (2009–2012). Because the biomarker sample 119 included everybody above the age of 44 but only a random subsample of those aged 44 120 or below (Crimmins et al. 2015), its age structure was older and hence its self-reported 121 diabetes prevalence higher. The analysis therefore compares with self-reported data for 122 this specific subsample only. 123

Our outcome variables of interest were employment status, weekly working hours, hourly wage, and occupation. Employment status was defined as having carried out an 125 activity that helped with the household expenses the last week and working for at least 126 four hours per week. We explicitly included informal employment and employment without 127 monetary remuneration, for instance in family businesses. Hourly wage was constructed 128 as reported monthly income from the first and second job, divided by average number 129 of weeks per month and weekly working hours. Labour income was obtained from the 130 response to questions on wages, income from piecework, tips, income from extra hours, 131 meals, housing, transport, medical benefits and other earnings, or from the response to 132 a question on aggregate labour income for the entire month. We adjusted calculated 133 wages for inflation in the year of interview and considered the log of real wages. Due 134 to a considerable number of missing or zero income reports, the sample used for the 135 wage estimation was smaller than the sample for working hours. Working hours were 136 combined from both the first and a potential second job. Descriptive statistics for the 137

entire panel sample show that 86% of men reported some form of employment compared to 37% of women (see Table 1). Interestingly, men did not report considerably higher hourly wages than women but worked more hours per week. Men also more often worked in agricultural jobs while women were more likely to be self-employed or in non-agricultural wage employment. The educational attainment of women was lower than that for men on average.

Table 1. Descriptive statistics for the panel sample (2002,2005,2009).

| | Males | | | | Females | |
|---|-------------|----------|------------|-------------|----------|------------|
| | No diabetes | Diabetes | p (t-test) | No diabetes | Diabetes | p (t-test) |
| Dependent variables | | | | | | |
| Employed | 0.87 | 0.80 | 0.00 | 0.37 | 0.26 | 0.00 |
| Hourly wage (in Mexican Peso) | 42.29 | 46.79 | 0.83 | 40.67 | 36.33 | 0.61 |
| Weekly working hours | 46.83 | 46.51 | 0.60 | 39.06 | 37.51 | 0.09 |
| Non-agricultural worker or employee | 0.51 | 0.41 | 0.00 | 0.24 | 0.13 | 0.00 |
| Agricultural worker | 0.19 | 0.13 | 0.00 | 0.02 | 0.01 | 0.00 |
| Self-employed | 0.16 | 0.26 | 0.00 | 0.09 | 0.11 | 0.04 |
| $Diabetes\ variables$ | | | | | | |
| Diabetes duration (years) | | 7.40 | | | 7.79 | |
| $Control\ variables$ | | | | | | |
| Age | 35.31 | 50.68 | 0.00 | 35.37 | 50.45 | 0.00 |
| Any medical insurance | 0.47 | 0.59 | 0.00 | 0.50 | 0.62 | 0.00 |
| City of 2,500-15,000 | 0.11 | 0.09 | 0.03 | 0.11 | 0.13 | 0.00 |
| City of 15,000-100,000 | 0.10 | 0.14 | 0.00 | 0.10 | 0.10 | 0.40 |
| City of >100,000 | 0.34 | 0.39 | 0.00 | 0.35 | 0.34 | 0.47 |
| Married | 0.53 | 0.77 | 0.00 | 0.53 | 0.66 | 0.00 |
| Number of children (age<6) in household | 1.49 | 1.14 | 0.00 | 1.60 | 1.13 | 0.00 |
| Indigenous group | 0.19 | 0.15 | 0.00 | 0.19 | 0.19 | 0.86 |
| Education | | | | | | |
| Secondary | 0.31 | 0.22 | 0.00 | 0.31 | 0.16 | 0.00 |
| High school | 0.16 | 0.07 | 0.00 | 0.14 | 0.03 | 0.00 |
| Higher education | 0.11 | 0.12 | 0.39 | 0.10 | 0.03 | 0.00 |
| Wealth index | 0.00 | 0.04 | 0.27 | -0.01 | 0.01 | 0.36 |
| N | 20391 | 994 | | 25664 | 1666 | |

Notes Mean values. Diabetes refers to self-reported diabetes.

The first part of the analysis focused on the relationship of labour outcomes with self-reported diabetes, which was based on the survey question: "Have you ever been diagnosed with diabetes?". Because the data did not distinguish between type 1 and type 2 diabetes, we assumed that the estimates represented the impact of type 2 diabetes, by far the most common type of diabetes in Mexico. As a robustness check, we re-estimated our main results categorizing diabetes into early-onset and late-onset cases, according to

the age at which diabetes was first reported in the survey. This was a similar approach to 150 Alegre-Díaz et al. (2016), who assumed that everybody diagnosed before age 35 and using 151 insulin had type 1 diabetes. Accordingly, we assumed that those first reporting a diabetes 152 diagnosis before the cut-off had type 1 diabetes while those above had type 2 diabetes. 153 Nonetheless, because we cannot warranty that this is 100% accurate as it may be unlikely 154 that both populations consisted exclusively of one type of diabetes, we preferred to think 155 of the groups as of early- and late onset groups (in the case of the within-coefficient, which 156 only takes into account incident cases). Because we did not have late-onset groups. This 157 separation also provides information about the exact age at diagnosis for all diabetes cases 158 in all three waves, effects for different age groups, as the late-onset group had an average 159 age of onset of 50 compared to 28 for the between coefficient may also stratify people with 160 diabetes into the late onset group, even though they actually had early onset diabetes but 161 only joined the sample after already having been diagnosed for several yearsearly-onset 162 group. In the pooled data, which combines all three waves, diabetes was self-reported 163 by 5% of men and 6% of women, respectively. This is consistent with other reports from 164 Mexico for the time, showing a prevalence of diagnosed diabetes of 7.5\% in 2006 in a sample 165 also including people over the age of 64 (Barquera, Campos-Nonato, et al. 2013). Apart 166 from self-reported diabetes, which was available in all rounds, we also used information 167 on the self-reported year of diagnosis as well as biometrically measured HbA1c levels for 168 a subsample of respondents from the third wave. 169 Information on the self-reported year of diagnosis reported in the third wave allowed us 170 to construct a measure of time since diagnosis. For those also present in previous waves, 171 172

we inferred the time since diagnosis by the difference between the year of the interview and the year of diagnosis. This allowed us to use panel data methods for the duration analysis as well, however limited to those reporting the year of diagnosis in the third wave.

The second part of the analysis assessed the role of measurement error associated with self-reported diabetes. This was done by also considering those with undiagnosed

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diabetes, i.e. the false negatives undiagnosed diabetes. The biometrically measured blood 177 glucose value that allowed us to identify those with undiagnosed diabetes, was available for 178 over 6000 respondents in the third wave. We chose the internationally recognized cut-off 179 of an HbA1c $\geq 6.5\%$ to define diabetes as recommended by the World Health Organi-180 zation (WHO) (World Health Organization 2011). As we show in Supplementary Table 181 S6, several observations of self-reported diabetes had HbA1c levels below the diabetes 182 threshold. We dropped those for our analysis as it was unclear if they had misreported 183 their diabetes status or had achieved these low levels as a result of their successful disease 184 management. Analysis including those cases did not lead to qualitatively different results 185 (results available on request). 186

3. Estimation strategy

To investigate the relationship between self-reported diabetes and three labour outcomes— 188 employment, wages and weekly working hours—we estimated a effects model, an extension 189 of the correlated random effects modelfirst proposed by Mundlak (1978), that explicitly 190 models both within and between effects . The within effects are identical to a fixed effects 191 model. The fixed effects (FE) model, accounting accounts for the potential bias introduced 192 by time-invariant unobservables, providing an estimate of the effect for cases that received a 193 diagnosis throughout the survey (705 incident cases compared to 970 non-changing diabetes 194 cases in the used sample). Modeling the between effect allowed us to also use information 195 from those that already had diabetes at baseline. 196

$$Y_{it} = \beta_0 + \beta_1 (D_{it} - \overline{D}_i) + \beta_2 (X_{it} - \overline{X}_i) + \beta_3 \overline{D}_i + \beta_4 \overline{X}_i + (u_i + e_{it}), \tag{1}$$

The within-effect is arrived at by fixed effects model uses only the within-person variation for identification, i.e. the difference between the diabetes indicator D_{it} and its cluster mean \overline{D}_i , so that β_1 represented the within-person variation of diabetes over time.

The same applies to the other time-varying covariates X_{it} . Cluster means of diabetes and 200 of all other time-varying covariates were also included to capture the between effect. The 201 error terms u_i and e_{it} capture the errors for the within and between variation, respectively. 202 Y_{it} was a binary variable taking a value of 1 if respondent i reported being in employment 203 at time t and 0 otherwise. Making use of the user written Stata command xthybrid, we 204 estimated effects applying a multilevel mixed-effects generalized linear model. For ease of 205 interpretation we chose to estimate a linear probability model for the effects of diabetes 206 on employment. 207

For the outcomes To estimate the effect on working hours and wages, our empirical models were estimated conditional on being in employment. Y_{it} represented the log hourly wage or the weekly working hours over the last year, for respondent i at time t.

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The regressions controlled for Because the fixed effects model accounts for any time-invariant 211 confounding, we only included time-variant confounders. We controlled for changes in the level of urbanization, education, state the educational level, the state of residence, marital 213 status, the number of children below the age of 6 in the household, a quadratic age term 214 , and calendar year dummies as well as household wealth. We also control for household 215 wealth approximated by a household asset index. We composed an indicator using prin-216 cipal component of household assets and housing following Filmer et al. (2001) (Filmer 217 and Pritchett 2001). The assets indicators asset index reflected owning a vehicle, a second 218 house, a washing machine, dryer, stove, refrigerator or furniture, any electric appliances, 219 any domestic appliances, a bicycle, farm animals, and accounted for the physical condi-220 tion of the house, proxied by the type of floor material and water access. In our main 221 regression models we did not account for body mass index (BMI). While part of the effect 222 of diabetes may be due to potential adverse effects of obesity, including BMI as a con-223 trol variable in the model would have led to biased estimates if the diagnosis of diabetes 224 also had an effect on BMI, which was likely has been shown to be the case. In general, 225 control variables should not also be potential outcome variables, hence we similarly in 226

other studies (De Fine Olivarius et al. 2015; Seuring, Suhrcke, et al. 2018; Slade 2012).
Similarly, we did not control for other chronic diseases that may have been caused by any
diseases that are likely consequences of diabetes, such as hypertension or cardiovascular
disease, as this would prevent any causal interpretation of the relationship of diabetes
with our labour market outcomes (Angrist and Pischke 2009). Stata 15 was used for all
analyses (StataCorp 2017).

233 3.1. Labour outcomes and time since diagnosis

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The chronic nature and irreversibility of diabetes provide good reason to explore the long term effects post diagnosis. To do this we estimated the following model:

$$Y_{it} = \beta_0 + \beta_1 (Dyears_{it} - \overline{Dyears}_i) + \beta_2 (X_{it} - \overline{X}_i) + \beta_3 \overline{Dyears}_i + \beta_4 \overline{X}_i + (u_i + e_{it}),$$

where $\beta_1 Dyears_n$ was continuous replaced the binary diabetes indicator of Eq 1 with a continuous variable indicating years since the diagnosis was first reported. While simultaneous inclusion of year dummies and time since diagnosis (which varies by one unit in each time period) would typically not allow separate identification of the coefficient of time since diagnosis in Eq 3.1. and Eq ??, identification here relied on the presence of people without diabetes in the sample, for which diabetes duration did not increase.

We also considered a spline function that allowed for non-linear effects over time.

$$\underline{Y_{it} = \beta_0 + \beta_1(Dsplines_{it} - \overline{Dsplines}_i) + \beta_2(X_{it} - \overline{X}_i) + \beta_3\overline{Dsplines}_i + \beta_4\overline{X}_i + (u_i + e_{it}),}$$

, with $g(Dyears_{it}) = \sum_{n=1}^{N} \delta_n \cdot max\{Dyears_{it} - \eta_{n-1}\}I_{in} \text{ and } I_{in} = 1[\eta_{n-1} \leq Dyears_{it} < \eta_n],$ with η_n being the place of the n-th node for $n = 1, 2, \dots, N$. The coefficient δ_n captures
the effect of diabetes for the n-th interval. The effects are linear if $\delta_1 = \delta_2 = \dots = \delta_n$.

Based on visual inspection (Fig. 1 on page 17) we chose three nodes located at 3, 7 and 246 12 years after diagnosis. The first three years should capture any immediate effects of the 247 diagnosis, the years four to seven any effects during time of adaptation to the disease and 248 the later terms the long term effects. We also estimated a non-linear model using dummy 249 variables for duration groups rather than splines, applying the same duration cut-offs. 250 Because the year of diagnosis was only reported in the third wave, time since diagnosis 251 was not available for those who were not interviewed in the third round. A reported 252 diagnosis in the year of the interview was counted as 'one year since diagnosis'. 253

254 3.2. Labour outcomes and biometrically measured diabetes

The biomarker analysis consisted of three steps. We first re-estimated Eq 2 to assess the relationship between self-reported diabetes with labour outcomes, but this time for the cross-sectional biomarker sample only, using the following specification:

$$Y_i = \beta_0 + \beta_1 Dsr_i + \beta_2 X_i + c_i + v_i \tag{2}$$

where v_i were community fixed effects which reflected local unobserved characteristics, such as access to healthcare, poverty and unemployment in the community. Communities (or localidades in Spanish) are the smallest administrative units (nested within municipalities) recognized by the Mexican Institute for Statistics and Geography (INEGI). We did not use household fixed effects since the average number of observations per household was close to one.

In a second step we estimated the relations between biomarker diabetes and labour outcomes, using the following equation:

$$Y_i = \beta_0 + \beta_1 Dbio_i^d + \beta_2 X_i + v_i + u_i, \tag{3}$$

where $\underline{Dbio^d}$ $\underline{Dbio_i}$ was equal to 1 if HbA1c $\geq 6.5\% \geq 6.5\%$.

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To estimate the effect of undiagnosed diabetes, we added self-reported diabetes and interacted it with the biomarker back into the equation (Eq 4).

$$Y_i = \beta_0 + \beta_1 Dsr_i + \beta_2 Dbio_i + \beta_3 Dsr_i * Dbio_i + \beta_4 X^d_i + v_i + u_i.$$

$$\tag{4}$$

Note that the interaction term This changes the interpretation of β_1 and β_2 , with

 β_1 now representing the effect on those aware of their condition but with levels below 270 the diabetes threshold; while β_2 which now reflects the effect on those with undiagnosed 271 diabetes, i.e. the respondents not self-reporting diabetes but with HbA1c levels equal to or 272 above the threshold. The interaction term β_3 shows the effect for those with self-reported 273 diabetes and levels above the threshold. 274 We further investigated the effect of the severity of diabetes on labour outcomes, re-275 placing $Dbio^d$ with $Dbio^c$, a variable that was 0 for HbA1c < 6.5% and took the actual value of HbA1c minus 6.4 for those with an $HbA1c \geq 6.5\%$ (Eq 5). This allowed us to 277 investigate the effect of a one percentage point increase in HbA1c levels for people with undiagnosed diabetes (β_2) as well as for those with self-reported diabetes above the diabetes 279 threshold (β_3) . 280

$$Y_i = \beta_0 + \beta_1 Dsr_i + \beta_2 Dbio_i^c + \beta_3 Dsr_i * HbA1c_i + \beta_4 X_i + v_i + u_i.$$
 (5)

281 4. Results

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4.1. Labour outcomes and self-reported diabetes

The results of estimating Eq 1 in Table 2 indicated significant and substantial reductions in the probability of employment for men and women with self-reported diabetes. The overall similarity of between estimates suggests that the within effect is generalizable to the entire self-reporting diabetes population, i.e. not only for representative of those that

developed diabetes after joining the survey. Additionally, it provides suggestive evidence that time-invariant unmeasured confounders may play a limited role. Employment proba-bilities were reduced by over 5-5.4 p.p. for both gendersmen and 6 p.p. for women, trans-lating into relative reductions of 14% for women and of 6% for men and 14% for women. There was no significant relationship between diabetes on one hand and working hours and wageson the other, though the between estimates suggested that men with diabetes earned generally more than their counterparts without diabetes (Table 2) with working hours or wages. Overall these results thus suggested effects at the extensive margin (employment), but not at the intensive margin (labour supply and productivity).

Dividing the diabetes population into early and late onset groups, men, and potentially also women, saw their employment probabilities negatively affected by a diabetes onset later in life (Supplementary Table S2). For women, a particularly strong effect was also found for early diabetes onset. The between estimator suggested that men and women with diabetes were less likely to be employed at older ages, but not at a younger ageIn particular, women with an early diabetes onset experienced an adverse effect. For working hours, we only found an adverse effect using the within effects estimator for early onset, which may have been spurious due to low incidence rates of diabetes. For wageseffects are less precise but may indicate increased working hours for men with an early diabetes onset, while women reduced their work hours. Finally, we found a positive effect of diabetes incidence on women. However, the within estimates for early onset cases again may be spurious. The between estimates show that especially older menwith diabetes received higher wages than those without diabeteshigher wages for women with an early diabetes onset, but no effects for men.

To assess whether diabetes affected the selection into different types of work, we investigated the role of diabetes for the probability of being in non-agricultural wage employment, agricultural employment or self-employment. The within effects estimator showed We found a reduction in the probability to work in agriculture for women, while we

Table 2. Labour outcomes and self-reported diabetes

| | Emp | loyment | Weekly work hour | 'S | Log hourly wages | | |
|---------------------------|--------------------|---------------------------------------|---|---|-----------------------------|-----------------------------|--|
| | Males | Females | Males | Females | Males | Females | |
| Diabetes (within) | -0.054** (0.025) | -0.060 -0.059** (0.024) | -0.582 - <u>0.506</u> (1.501) (0.848 1.499) | -1.990 <u>-1.998</u> (1.306 2.511) | 0.063-0.055 (0.0460.068) | 0.074-0.081 (0.0640.158) | |
| Within—Between (p value)N | 21388 | 27339 | 17616 | 9112 | 13828 | 7068 | |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, *** p < 0.05, **** p < 0.01.

found no statistically significant effects for men. The between effects suggested that men with diabetes were less likely to be employed in agriculture, but more likely to be self-employed. Women with diabetes were less likely to be employed in agricultural and non-agricultural jobs but not for men (Table 3). Disaggregating the diabetes groups further according to their age showed that most statistically significant relationships were driven by the older onset older-onset group (Supplementary Table S3). Interestingly, for For male self-employment, incidence of diabetes increased the probabilities to be self-employed in the vounger group, while it reduced the probabilities to be self-employed in the older onset group. However, especially the results for early onset diabetes in women should be interpreted earefully due to limited number of diabetes cases, older-onset group.

Table 3. Selection into types of work and self-reported diabetes.

| | Males | | | Females | | |
|------------------|------------|---------|---------------|------------|----------|---------------|
| | Non-agric. | Agric. | Self-employed | Non-agric. | Agric. | Self-employed |
| Diabetes(within) | -0.007 | -0.008 | -0.042 | -0.001 | -0.022** | -0.030^* |
| | (0.029) | (0.022) | (0.026) | (0.018) | (0.009) | (0.018) |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Because obesity is one of the main risk factors for developing diabetes and may also affect labour market outcomes, its exclusion from above models may lead to biased estimates. We therefore reestimated all regressions in this section including a binary control for obesity (BMI \geq 30) (Supplementary Table S7 and Table S8). This led to

a reduction in sample size due to the larger number of missing cases for BMI. Obesity 329 itself did not appear to be an independent predictor of any labour market outcome. The 330 estimates of the effect of diabetes remained similar for most outcomes. Only for male 331 employment probabilities, the diabetes coefficient was no longer statistically significant at 332 the 10 percent significance level. Estimating the original model without accounting for 333 obesity but using the sample with only non-missing BMI cases showed similar changes in 334 effects, suggesting that these changes were rather the result of a smaller sample size than 335 of the inclusion of obesity. 336

4.2. Labour outcomes and time since diagnosis

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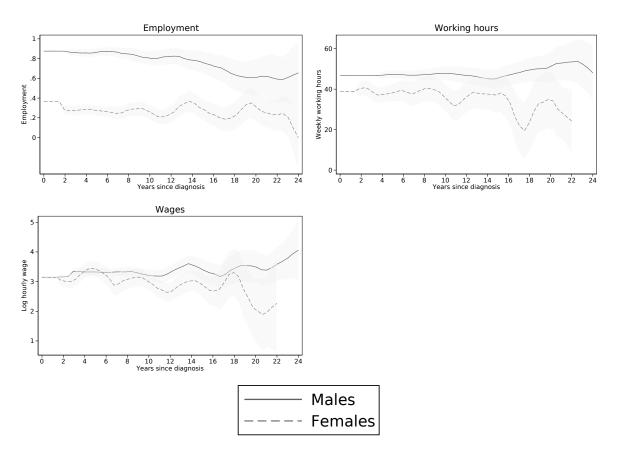
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Fig. 1 shows that the probability of employment for men steadily declined as time progressed, using a non-parametric kernel-weighted local polynomial regression. For women, a first drop-off occurred right after diagnosis; though no consistent pattern emerged thereafter. The dynamics for working hours and wages were less clear, with a possibly long term negative trend for women but not for men.

Table 4 panel A shows the results of estimating Eq 3.1., which indicated that male and female employment probabilities fell every year, with a larger effect shown by the within coefficient. For females, the within coefficient also suggested a reduction in wages

Wages were reduced for women as diabetes progressed while the between coefficient showed no association for women and a relatively small positive association with male wagesusing the linear specification. Using diabetes onset groups, there was no evidence of an effect of diabetes duration for early onset groups (see Supplementary Table S4). However, again the within results for early onset should be interpreted with caution due to the Unfortunately the very limited number of diabetes incidence cases in this group, which also cases in the early-onset group prohibited the estimation of any within effects of early diabetes onset duration for on wages and working hours. The results also indicated that the effects found in Table 4 were driven mainly by those with a diabetes onset after

Figure 1: Employment, wages, working hours and years since self reported diabetes: Kernel-weighted local polynomial regression



Notes The shaded areas indicate the 95% confidence intervals.

355 age 35.

The non-linear results for the spline function and dummy variable approach are presented in panels B and C, respectively. They suggest that the main adverse effects appeared after a prolonged time of living with diabetes; i.e. after more than seven years since diagnosis. The same was true for female wages. The lack of a statistically significant effect for the earlier years of diabetes duration may have been due to a reduction in statistical efficiency , reduced by due to the sole reliance on within-variation and the separation into duration groups and into within and between variation. Re-estimating the specifications with a random effects model that combined both types of variation into one

Table 4. Relationship between self-reported years since diagnosis and employment probabilities using continuous duration and duration splines.

| | Emplo | yment | Weekly w | ork hours | Log hou | rly wages |
|--------------------------------|-----------|--------------|--------------|-----------|---------|---------------|
| | Males | Females | Males | Females | Males | Females |
| Panel A: linear effect | | | | | | |
| Years since diagnosis (within) | -0.016*** | -0.009^* | 0.185 | 0.115 | -0.016 | -0.067** |
| | (0.006) | (0.005) | (0.334) | (0.652) | (0.018) | (0.029) |
| Panel B: splines | | | | | | |
| Years since SR diagnosis | | | | | | |
| 0-3 | -0.013 | -0.018 | 0.708 | 2.953 | -0.005 | 0.047 |
| | (0.014) | (0.016) | (0.857) | (2.700) | (0.054) | (0.124) |
| 4-7 | -0.011 | -0.002 | 0.215 | -2.517 | -0.032 | -0.131 |
| | (0.014) | (0.014) | (0.761) | (1.752) | (0.046) | (0.101) |
| 8-12 | 0.003 | -0.003 | -1.153 | 1.144 | -0.009 | -0.053 |
| | (0.021) | (0.014) | (1.252) | (1.635) | (0.065) | (0.061) |
| 13+ | -0.039*** | -0.015 | 0.720 | 0.184 | -0.007 | -0.096*** |
| | (0.014) | (0.010) | (0.943) | (1.414) | (0.057) | (0.037) |
| Panel C: dummies | , , | , , | , , | , , | , , | , , |
| 0-3 | 0.005 | -0.007 | 0.352 | 17.309* | 0.223 | -0.447 |
| | (0.052) | (0.059) | (3.123) | (9.975) | (0.186) | (0.549) |
| 4-7 | -0.031 | -0.049 | 2.860 | 10.878 | 0.047 | -0.568 |
| | (0.042) | (0.050) | (2.664) | (9.504) | (0.127) | (0.544) |
| 8-12 | -0.066 | -0.026 | -0.709 | 13.733 | -0.133 | -0.873^* |
| | (0.063) | (0.059) | (4.181) | (9.695) | (0.207) | (0.521) |
| 13+ | -0.134 | $-0.062^{'}$ | $-3.379^{'}$ | 13.309 | 0.164 | -0.882^{**} |
| | (0.098) | (0.068) | (4.715) | (9.239) | (0.284) | (0.446) |
| N | 16298 | 22427 | 10771 | 5746 | 13583 | 7391 |

Notes Panel A presents the results of the linear specifications. Panel B presents the results of the non-linear specifications. Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, *** p < 0.05, *** p < 0.01.

estimate showed that, at least for the models with dummies, there was a more or less an immediate reduction in employment probabilities that became stronger the longer a person had diabetes (see Supplementary Table S5). Note that we did not estimate models splitting diabetes in early and late onset groups, as this implied strong reductions in statistical power.

To test the robustness of the results to controlling for obesity, we estimated above models including a dummy variable for obesity (Supplementary Table S9). Overall, results remained very similar and obesity itself was not found to be affecting any labour market outcome.

4.3. Cross-sectional biomarker analysis

As reported in Supplementary Table S6, 18% of the observations in the biomarker sample 374 were false negatives, i.e. undiagnosed. Further, 2% were false positives, though the latter 375 may have included eases that received a diabetes diagnosis and managed to reduce their 376 to non-diabetes levels via medication and/reported diabetes but had an HbA1c below the 377 threshold. Because we did not know if the latter were true diabetes cases that had been 378 able to return to non-diabetic levels as a result of their treatment regime or lifestyle changes 379 (Flores-Hernández et al. 2015). Overall, or if they represented misreports, we dropped 380 these observations from the sample. Overall, 80% of the self-reports were consistent with 381 the biomarker data also had diabetes according to their HbA1c levels. Comparing the 382 health status and diabetes risk factors of the diagnosed and undiagnosed diabetes popu-383 lations suggested that those with self-reported diabetes were older and in worse health, 384 both objectively and subjectively compared to those undiagnosed. This suggests a selec-385 tion into the diagnosed group based on the severity and potentially duration of diabetes. 386 If the adverse effects of diabetes are due to its health impact, we would suspect worse 387 labour market outcomes for the diagnosed compared to the undiagnosed population.

Table 6 presents the results from estimating Eq. 2-5. Panel A confirms the earlier 389 longitudinal results using self-reported diabetes for the cross-sectional biomarker sample. 390 The results in panel B indicate that the relationship with employment became weaker 391 when using diabetes defined by the biomarker instead of self-reported diabetes, in par-392 ticular for men. Results in Panel C were obtained from estimating Eq. 4 and indicate 393 an absence of a (statistically significant) negative relationship between undiagnosed dia-394 betes (expressed in the 'Biomarker diabetes but not self-reported' coefficient) and labour 395 outcomes. The coefficients for the interaction term were mostly negative, though only 396 statistically significant in the case of female working hours. 397

To explore whether the adverse effects increased with higher HbA1c levels, we estimated

Eq. 5. The results in panel D again only suggest a negative association of a one only

Table 5. Descriptive comparison of diagnosed and undiagnosed population with diabetes.

| | | Males | | | Females | |
|---|-----------|-------------|----------|-----------|-------------|----------|
| | Diagnosed | Undiagnosed | P value | Diagnosed | Undiagnosed | P value |
| | diabetes | diabetes | (t-test) | diabetes | diabetes | (t-test) |
| Employed | 0.807 | 0.875 | 0.012 | 0.241 | 0.331 | 0.002 |
| Hourly wage | 35.931 | 30.670 | 0.120 | 36.092 | 32.638 | 0.550 |
| Usual weekly working hours | 44.341 | 46.682 | 0.104 | 34.708 | 39.681 | 0.046 |
| Age | 53.162 | 44.720 | 0.000 | 53.167 | 44.449 | 0.000 |
| Any medical insurance | 0.677 | 0.599 | 0.033 | 0.733 | 0.643 | 0.002 |
| City of 2,500-15,000 | 0.096 | 0.107 | 0.643 | 0.112 | 0.112 | 0.994 |
| City of 15,000-100,000 | 0.135 | 0.096 | 0.098 | 0.087 | 0.090 | 0.884 |
| City of >100,000 | 0.331 | 0.297 | 0.325 | 0.294 | 0.333 | 0.185 |
| Married | 0.719 | 0.643 | 0.031 | 0.633 | 0.569 | 0.035 |
| Number of children (age<6) in household | 0.950 | 1.122 | 0.073 | 0.960 | 1.257 | 0.001 |
| Indigenous group | 0.158 | 0.211 | 0.079 | 0.195 | 0.207 | 0.626 |
| Primary | 0.479 | 0.434 | 0.238 | 0.626 | 0.465 | 0.000 |
| Secondary | 0.212 | 0.231 | 0.554 | 0.132 | 0.233 | 0.000 |
| High school | 0.062 | 0.131 | 0.003 | 0.037 | 0.117 | 0.000 |
| Higher education | 0.139 | 0.113 | 0.288 | 0.030 | 0.073 | 0.003 |
| Wealth index | -0.174 | 0.117 | 0.000 | 0.012 | 0.103 | 0.157 |
| Subjective health | | | | | | |
| very good | 0.023 | 0.094 | 0.000 | 0.012 | 0.054 | 0.001 |
| good | 0.212 | 0.434 | 0.000 | 0.180 | 0.367 | 0.000 |
| fair | 0.619 | 0.442 | 0.000 | 0.643 | 0.528 | 0.000 |
| bad | 0.135 | 0.026 | 0.000 | 0.155 | 0.048 | 0.000 |
| very bad | 0.012 | 0.004 | 0.187 | 0.010 | 0.004 | 0.246 |
| Glycated hemoglobin (HbA1c) | 9.037 | 8.533 | 0.004 | 8.979 | 8.680 | 0.049 |
| Hypertension (self-reported) | 0.262 | 0.074 | 0.000 | 0.397 | 0.150 | 0.000 |
| Blood pressure | | | | | | |
| Systolic | 136.688 | 130.506 | 0.000 | 136.070 | 122.835 | 0.000 |
| Diastolic | 84.677 | 82.063 | 0.003 | 84.495 | 79.689 | 0.000 |
| Heart disease (self-reported) | 0.035 | 0.007 | 0.004 | 0.050 | 0.024 | 0.021 |
| BMI | 28.868 | 28.311 | 0.135 | 30.640 | 29.778 | 0.032 |
| Obese (BMI ≥ 30) | 0.338 | 0.311 | 0.440 | 0.469 | 0.431 | 0.225 |

were

Notes Mean values. * p < 0.10, ** p < 0.05, *** p < 0.01.

show a borderline statistically significant adverse association of 0.9 percentage points per
percentage point increase in HbA1c with female working hours for those with self-reported
diabetes. For undiagnosed diabetes, we again found no effects for female employment
probabilities.

Table 6. Biomarker results.

| | Employ | ment | Weekly wor | k hours | Log hourly | / wages |
|--|-------------------------|-------------------------|-------------------------|-------------------------|-------------------------|------------------------|
| | Males | Females | Males | Females | Males | Females |
| Panel A: Diabetes (self-reported) | | | | | | |
| Self-reported diabetes | 055057** | 050057** | -308543 | -323-2.154 | 026057 | 005 |
| | (-026.025) | (.024 .026) | (1.3211.427) | (2.1902.433) | (-066.070) | (.105 .121 |
| Panel B: Diabetes (biomarker) | | | | | | |
| Biomarker diabetes (HbA1c \geq 6.5) | 012- .013 | 032 .034* | 060018 | 1.067-1.382 | 008005 | .041 .045 |
| | (.016) | (.018) | (.844.849) | (1.4701.480) | (.045) | (.071) |
| Panel C: Self-reported and undiagnosed diabetes | | | | | | |
| Self-reported diabetes but tested negative (β_1) | 049061** | 027042 | 1.547 - 715 | 7.417 - 3.954 | .250 - .067 | 025- .034 |
| | (.059 .028) | (.051 .031) | (3.4461.574) | (4.6692.823) | (.205 .085) | (.232 .137 |
| Biomarker diabetes but not self-reported-Undiagnosed diabetes (HbA1c \geq 6.5) (β_2) | .005 006 | 020 | .174 .224 | 2.304-2.394 | .015 .014 | 049 053 |
| | (.018) | (.020) | (.960 .962) | (1.6391.647) | (.050) | (.078) |
| Panel D: HbA1c levels | (/ | (/ | (| (| () | () |
| Self-reported diabetes | 067080* | 035066 | 1.471084 | 4.974-4.463 | . 157 061 | 017 -,011 |
| | (.072 .046) | (.041 .046) | (2.6192.409) | (3.6544.592) | (.119 .107) | (.186 ,227 |
| HbA1c if ≥ 6.5 | .001 005 | 003009* | 005 150 | .235 318 | .002 .004 | 005 |
| | (.002 .005) | (.002 .006) | (.104 .253) | (.193 .463) | (.005 .014) | (.008 .019 |
| Self-reported diabetes \times HbA1c if ≥ 6.5 | .001 003 | .000010 | 207 064 | 852 .375 | 023002 | .005 .000 |
| | (.007 .012) | (.005.012) | (.294 .668) | (124 1.043) | (.014 .030) | (.022 .052 |
| N | 2749 | 3537 | 2276 | 1121 | 1787 | 866 |

Notes Results are based on community level fixed effects. Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and on lummy variable for each calendar year to account for the multiple years of data collection for the third wave. The wage and working hour models additionally control for type of work (agricultural and self employed with non-agricultural wage employment as the base - p < 0.10, **p < 0.05, ***p < 0.01.

5. Discussion

Diabetes is now one of the most common chronic diseases in low- and middle income countries, as well as high-income countries, with potential severe impacts on the health and economic well-being of those affected. Yet rigorous evidence on the economic consequences for LMICs remains scarce.

To address key methodological challenges, this paper used rich longitudinal panel data from Mexico that also contained diabetes biomarkers. The biomarker data showed alarming levels of clinically tested diabetes (27% prevalence) and indicate that a large proportion of the Mexican population (18%) is unaware of their condition and undiagnosed diabetes.

Overall, the paper found The paper provides evidence for adverse effects of self-reported diabetes on the probability of being employed, confirming earlier findings for Mexico

that used employment, working hours and wages. While earlier evidence for Mexico already exists for employment (Seuring, Goryakin, et al. 2015), this was the first time effects on working hours and wages were investigated. Furthermore, we added to the study of Seuring, Goryakin, et al. (2015) by using longitudinal instead of cross-sectional information. But, these new results also suggest a comparatively larger data to identify a causal relationship. We provided first evidence of the long term impact of diabetes on female employment probabilities. The evidence also points towards the main effects being

and explored the extend and effects of undiagnosed diabetes. We confirmed the findings 422 of Seuring, Goryakin, et al. (2015) insofar that we found an adverse effect of diabetes on 423 male employment. We further show more conclusive evidence that also women experience 424 a reduction in their employment probabilities due to diabetes. We also found that the 425 effects were mainly driven by those with a diabetes onset at a relatively later state, con-426 sisting most likely of people with of older people with most likely type 2 diabetes. This 427 was also found by Seuring, Goryakin, et al. (2015), albeit they had not considered the 428 onset of diabetes but only stratified their sample into an older and younger age group. 429 Analyses of the long term impact indicated that the employment probability fell gradu-430 ally in the years following diagnosis. The results for the using a non-linear models model 431 were less clear, potentially due to reductions in statistical power, but suggest that ad-432 verse effects became stronger with time since diagnosis. The linear effect contrasts with 433 estimates for the USA, where such an effect seemed to be absent, was absent; however 434 allowing for non-linearity revealed falling employment probabilities after 11 to 15 years for females and after 2-5 years for males (Minor 2013). For working hours and wages, our 436 results were more ambiguous with adverse effects being observed for time since diagnosis 437 on female wages only. In summary, most of the adverse effects of diabetes were found 438 at the extensive margin, mainly affecting the probability of employment, rather than the 439 intensive margin, i.e. working hours or wages. 440 Overall, a relationship of diabetes with working hours or wages is mostly absent, except 441

Overall, a relationship of diabetes with working hours or wages is mostly absent, except for the between estimator in the case of wages for men. Contrasting the between and within results suggests that this significant result may have arisen from individual selection. Although any explanation at this point is speculative, it may be that higher paid men, who also tend to be more educated and more educated individuals, were able to remain employed without experiencing wage reductions, for instance due to their particular set of skills. They may also have had access to better health care leading to better diabetes related health outcomes. Low paid workers, on the other hand, may lack access to quality

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diabetes care, making it more likely that they develop severe complications earlier (Flores-Hernández et al. 2015). They are also more likely to be in informal employment and low skilled jobs, with less job security and are thus more prone to being laid off to be replaced with healthier workers.

Because self-reported diabetes may not adequately represent the entire diabetes popu-453 lation if the share of undiagnosed diabetes is relatively large and those undiagnosed differ 454 from those diagnosed, estimates based on self-reported diabetes are likely to be biased. 455 Our results using the cross-sectional biomarker data suggest indeed that those with undi-456 agnosed diabetes were significantly healthier and younger. Consequently, diabetes based 457 on biomarkers was found to be less related to reduced employment, compared to self-458 reported diabetes, in particular for men. Further analysis showed that this was due to the 459 absence of any association between undiagnosed diabetes and employment. These results 460 are similar to those found for the USA, where no statistically significant relationship was 461 observed between undiagnosed diabetes on employment, while a significant effect of diagnosed diabetes was observed (Minor and MacEwan 2016). Our results further indicate 463 that the difference in the employment effects of diagnosed and undiagnosed diabetes was 464 not mediated by current HbA1c levels, as even undiagnosed cases with relatively high 465 HbA1c levels did not show any adverse associations with labour market outcomes. This 466 is similar to findings for Mexican-Americans in the USA, where employment outcomes 467 were unrelated to higher HbA1c levels (Brown, Perez, et al. 2011). This may stem from 468 HbA1c levels primarily being informative for the last three months, and not being the 469 only indicator for the severity of diabetes. Overall, it seems that general health differences 470 related to a longer diabetes duration and selection into the diagnosed population, for in-471 stance based on emerging diabetes related health problems, could be driving the adverse 472 economic effects among those with a self-reported diagnosis. 473

Our study had several limitations. While the within-coefficient our model accounts for any time-invariant confounding, the estimates may have been affected by unobserved time-

variant confounders. Reverse causality, where employment status affects the propensity to 476 develop or be diagnosed with diabetes, may also play a role. Existing studies that looked 477 at this particular direction of causality, however, have not found strong evidence for an 478 effect of employment status on diabetes (Bergemann et al. 2011; Schaller and Stevens 479 2015), though they were carried out in high-income countries. We do not control for the 480 effects of obesity, hypertension, self-reported health or other diseases in our models due to 481 the high probability that they are affected by diabetes themselves, which would prevent 482 a causal interpretation of the estimates. Nonetheless, robustness checks including obesity 483 indicate that our main findings remain unchanged. For the duration analysis an additional 484 limitation, imposed by the data, was that the year of diagnosis was only reported in the 485 third wave. While this still allowed us to construct an estimate of the time since diagnosis 486 for the previous waves, it restricted the analysis to those that were present in the last 487 wave, thereby excluding those that dropped out of the sample prior to the third wave. 488 Finally, we used a WHO recommended HbA1c cut-off to diagnose diabetes, due to the lack of a Mexico specific cut-off. There is some evidence that HbA1c may be affected by 490 ethnicity (Sacks 2011). Hence, if Mexican ethnicity would lead to different HbA1c levels, 491 the use of our cut-off could have led to misclassifications based on the used biomarkers. 492 Despite these limitations, our findings bear important implications. First, the impact 493 of self-reported diabetes on labor outcomes in Mexico seems mostly limited to its effect 494 on employment probabilities, though there is some indication that it could also reduce 495 wages over time for women. Second, its effect on employment was much stronger for 496 females, though the underlying reasons for this remain unclear. Potential explanations 497 are that lower working hours or wages for women make a dropout less costly. Other 498 evidence suggests that women with diabetes are in worse metabolic health compared to 499 men when they cross the diabetes threshold (Peters et al. 2015), making it more likely 500 for them to drop out. Third, caution is needed when estimates based on self-reported 501 diabetes are interpreted in terms of the entire population, i.e. extending to those with 502

undiagnosed diabetes. Ideally, studies would include a biomarker analysis, acknowledge
the differences between diagnosed and undiagnosed subpopulations between diagnosed and undiagnosed subpopulations, and
carry out a separate analysis whenever feasible. If this is not possible, study conclusions
about the effect of self-reported diabetes should be limited to this specific part of the
population, in particular in environments where the share of undiagnosed diabetes is high,
as is the case for most low- and middle income countries.

While we find no effect for undiagnosed diabetes in our cross section analysis, further 509 inquiry over time is needed. The large proportion of previously undiagnosed cases found 510 in this paper indicates that diagnosis—at least in Mexico—happens late or not at all. This 511 may reduce the possibilities to prevent complications via treatment and self-management, 512 increasing the risk of severe complications appearing premature. Earlier diagnosis and en-513 suing effective treatment may lighten the health and economic burden. Further analysis is 514 also needed to explain why the adverse economic effects are so large for women. Ultimately, 515 prevention of diabetes is of high importance. Taxation of sugar sweetened beverages may be one promising way forward, though their long-term effectiveness remains unclear. 517 Given the established links between early life heath and later life incidence Therefore more 518 research is needed to investigate the economic impact of diabetes over time. Longitudinal 519 biomarker information could be used to observe the true duration and severity of dia-520 betes as well as other chronic diseases, investments in maternal and child health may 521 be particularly attractive to address non-communicable and communicable diseases at the 522 same time that passes till a diagnosis through a doctor. This would allow for a 523 better understanding of when adverse economic effects start to appear. Further, future 524 research should investigate how the time of diagnosis and the treatment of diabetes affect 525 the occurrence of adverse labour market effects of diabetes. The results of such research 526 could inform costing studies to include more detailed information on the indirect costs 527 of diabetes; or cost-effectiveness analyses that aim to include a measure of the potential 528 benefit of the intervention to employers or society at large. 529

530 Supplementary material

Strategies to deal with inconsistent self-reporting over time

Reporting error can pose a considerable challenge in the use of self-reported data. Fortunately, the MxFLS data provide several possibilities to assess the amount of misreporting and apply corrections before estimating the labour market effects of diabetes. In what follows we describe how we have dealt with inconsistencies in self-reported diabetes over time.

Throughout the surveys, self-reported diabetes was measured by the question 'Have you ever been diagnosed by diabetes'. One of the key advantages of panel data is the repeated measurement which results in more than one data point, allowing to uncover inconsistencies for cases with multiple observations. Very little is known about inconsistencies in self-reported diabetes over time. Zajacova et al. (2010) assess the consistency of a self-reported cancer diagnosis over time in the USA. The study found that 30% of those who had reported a cancer diagnosis at an earlier point failed to report the diagnosis at a later point in time. A more recent diagnosis was found to be reported with greater consistency possibly due to increasing recall problems as time since diagnosis advanced.

When assessing the MxFLS, we also found inconsistencies in the diabetes self-reports across the three waves, with between 10–20% of those reporting diabetes in one wave not doing so in one of the subsequent waves. To improve the validity of diabetes self-reports, we were interested in reducing the amount of reporting inconsistencies.

For diabetes, the main concern with mismeasurement is related to false negatives. False
positives a lack of a diagnosis. Wrong self-reports indicating a diagnosis of diabetes are
deemed less of a problem since incentives to report diabetes when one does not have it
seem to be very limited—although we cannot exclude this. A study from China finds that
the vast majority (98%) of those who self-report diabetes are tested positive for diabetes,
while only a minority of those who are tested positive for diabetes (40%) actually self-

report the disease (Yuan et al. 2015). Our data showed a similar pattern, with a low proportion (3%) of the respondents being tested negative while self-reporting diabetes, while the majority of those who are tested positive (68%) do not self-report diabetes.

We used the above information to infer the "true" diabetes status for those with incon-559 sistent reports. For respondents present in all three waves, we corrected inconsistencies as 560 reported in Supplementary Table S1. We assumed that if diabetes was reported only once 561 in the first two waves (either in 2002 or 2005) and then not reported again in the ensu-562 ing waves, this diabetes report was likely to be false (see lines 3 and 4 in Supplementary 563 Table S1) and that the person never had received a diagnosis. If a diabetes diagnosis was 564 however reported in two of the three waves (in 2002 and 2009 but not 2005, or in 2002 565 and 2005 but not in 2009) we assumed that the respondent had diabetes in all three waves 566 (see lines 1 and 2 in Supplementary Table S1). For cases where we only had information 567 from two waves, we assumed that if a diabetes diagnosis had been reported in a prior wave 568 they also had diabetes in the ensuing wave, even if it was not reported in the latter (see lines 5 and 6 in Supplementary Table S1), given that most diabetes self-reports tend to be correct.

Table S1. Inconsistencies in diabetes self-report in MxFLS.

| | Inconsistency | Assumption | Number of observations replaced |
|---|--|--------------------------------|---------------------------------|
| 1 | Diabetes self-report only in 2002, but not in 2005 and 2009 | Has no diabetes in 2002 either | 66 |
| 2 | Diabetes self-report only in 2005, but not in 2002 and 2009 | Has no diabetes in 2005 either | 52 |
| 3 | Diabetes self-report in 2002, 2005 but not in 2009 | Has diabetes in 2009 as well | 19 |
| 4 | Diabetes self-report in 2002, 2009 but not in 2005 | Has diabetes in 2005 as well | 63 |
| 5 | Diabetes self-report in 2002, but not in 2005. Not in survey in 2009 | Has diabetes in 2005 as well | 44 |
| 6 | Diabetes self-report in 2005, but not in 2009. Not in survey in 2002 | Has diabetes in 2009 as well | 23 |

We then tested if those respondents we categorized as not having a diabetes diagnosis based on above rules were actually more likely to not have biometrically measured diabetes, using the biomarker data from wave 3. Of those with inconsistencies in their diabetes self-reports, 95 were present in the biomarker sample (46 with two self-reports (from lines 3 and 4 in Table S1) and 49 with one self-report of diabetes (from lines 1 and 2 in Supplementary Table S1)). Supplementary Figure S1 illustrates the difference between both groups and

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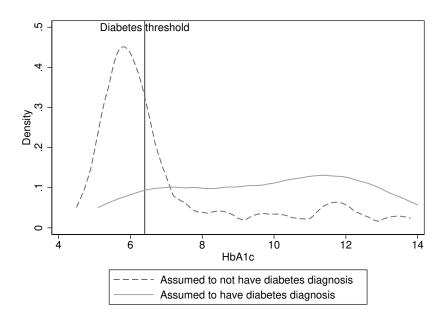
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suggests that indeed those with two self-reports of diabetes are much more likely to have
HbA1c values above the diabetes threshold. A t-test comparing the mean HbA1c for the
two groups indicates that those with two self-reports also have significantly (p<0.001)
higher HbA1c levels than those with only one self-report of diabetes (9.7% vs. 7.1%).
Further, of those with one self-report, only 30% have an HbA1c≥ 6.5% compared to 87%
of those with two self-reports. Based on these results it appears that we did minimize
misclassification of people into diabetes or no-diabetes.

Figure S1: Kernel density of HbA1c values for those with one inconsistent and two inconsistent reports.



Early versus late onset of diabetes

Table S2. Labour outcomes and self-reported diabetes by diabetes onset.

| | Emplo | yment | Weekly w | Weekly work hours | | ly wages |
|-----------------------|----------------|--------------|----------|-------------------|--------------|-------------|
| | Males | Females | Males | Females | Males | Females |
| Early onset (within) | 0.133 | -0.206** | 14.712* | -18.636* | -0.523 | 0.388*** |
| , , | (0.176) | (0.086) | (8.370) | (9.668) | (0.340) | (0.057) |
| Late onset (within) | -0.059^{**} | -0.048^{*} | -1.008 | $-1.322^{'}$ | $0.079^{'}$ | $0.067^{'}$ |
| , , | (0.025) | (0.025) | (1.513) | (2.553) | (0.067) | (0.163) |
| Early onset (between) | 0.016 | -0.052 | 1.483 | $-2.640^{'}$ | $-0.102^{'}$ | 0.286 |
| , , | (0.037) | (0.047) | (2.581) | (4.034) | (0.099) | (0.232) |
| Late onset (between) | -0.087^{***} | -0.067*** | -1.077 | -0.869 | 0.121** | -0.111* |
| , | (0.017) | (0.016) | (0.895) | (1.375) | (0.050) | (0.066) |
| N | 21388 | 27339 | 13828 | 7068 | 17616 | 9112 |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Table S3. Selection into types of work and self-reported diabetes by diabetes onset.

| | Non- | -agric. | Agrica | ulture | Self-en | nployed |
|-----------------------|------------|-----------|-----------|-----------|----------|---------|
| | Males | Females | Males | Females | Males | Females |
| Early onset (within) | 0.030 | -0.105 | -0.225 | -0.068 | 0.328** | -0.027 |
| | (0.216) | (0.074) | (0.139) | (0.047) | (0.161) | (0.048) |
| Late onset (within) | -0.008 | 0.007 | -0.002 | -0.019** | -0.053** | -0.030 |
| | (0.029) | (0.018) | (0.022) | (0.009) | (0.026) | (0.019) |
| Early onset (between) | 0.011 | -0.050 | -0.078*** | -0.006 | 0.087 | 0.005 |
| | (0.055) | (0.043) | (0.030) | (0.007) | (0.056) | (0.029) |
| Late onset (between) | -0.037^* | -0.048*** | -0.091*** | -0.009*** | 0.039* | -0.014 |
| , | (0.022) | (0.012) | (0.015) | (0.003) | (0.020) | (0.011) |
| N | 20719 | 26575 | 20719 | 26575 | 20719 | 26575 |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Table S4. Relationship between self-reported years since diagnosis and employment probabilities using continuous duration by diabetes onset.

| | Employment | | Monthly w | ork hours | Log hour | ly wages |
|-----------------------|--------------------------|--------------------------|------------------|-----------------|--------------------|---------------------|
| | Males | Females | Males | Females | Males | Females |
| Early onset (within) | -0.009 (0.017) | 0.002 (0.013) | | | | |
| Late onset (within) | -0.012** (0.005) | -0.007^* (0.004) | 0.086 (0.275) | 0.308 (0.504) | -0.006 (0.013) | -0.059*** (0.022) |
| Early onset (between) | 0.011*** (0.004) | -0.016^{***} (0.005) | -0.016 (0.012) | 0.029 (0.051) | 0.187 (0.333) | -0.401 (0.896) |
| Late onset (between) | -0.009^{***} (0.002) | -0.005^{***} (0.002) | -0.060 (0.117) | -0.144 (0.129) | 0.011** (0.006) | -0.010 (0.009) |
| N | 16308 | 22450 | 13592 | 7394 | 10778 | 5748 |

Notes The within estimator for the effects of early onset diabetes on wages and working hours could not be estimates due to no within-variation for diabetes. Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Random effects model

Table S5. Relationship between self-reported years since diagnosis and employment probabilities using continuous duration and duration splines (random effects).

| | Emplo | yment | Weekly we | ork hours | Log hourl | ly wages |
|--------------------------|------------|-----------|-----------|--------------|-------------|------------|
| | Males | Females | Males | Females | Males | Females |
| Panel A: linear effect | | | | | | |
| Years since diagnosis | -0.007*** | -0.004*** | 0.039 | -0.130 | 0.010** | -0.009 |
| | (0.002) | (0.001) | (0.102) | (0.127) | (0.005) | (0.008) |
| $Panel\ B:\ splines$ | | | | | | |
| Years since SR diagnosis | | | | | | |
| 0–3 | -0.008 | -0.015** | -0.035 | 0.507 | 0.038** | 0.034 |
| | (0.006) | (0.006) | (0.346) | (0.614) | (0.017) | (0.029) |
| 4-7 | 0.001 | 0.004 | 0.242 | $-0.570^{'}$ | -0.032 | -0.048 |
| | (0.011) | (0.011) | (0.665) | (1.062) | (0.032) | (0.052) |
| 8–12 | -0.008 | 0.002 | -0.116 | -0.080 | -0.003 | -0.074 |
| | (0.015) | (0.011) | (0.855) | (1.098) | (0.041) | (0.050) |
| 13+ | -0.012 | -0.004 | 0.035 | -0.339 | 0.029 | 0.011 |
| | (0.008) | (0.003) | (0.410) | (0.241) | (0.018) | (0.017) |
| $Panel\ C:\ dummies$ | | | | | | |
| 0–3 | -0.036* | -0.041** | -0.821 | 1.091 | 0.134** | 0.021 |
| | (0.021) | (0.021) | (1.154) | (1.826) | (0.054) | (0.083) |
| 4-7 | -0.014 | -0.056** | 0.877 | 1.200 | 0.093 | -0.003 |
| | (0.022) | (0.023) | (1.375) | (2.530) | (0.059) | (0.118) |
| 8–12 | -0.069^* | -0.043 | 0.427 | 0.302 | -0.070 | -0.148 |
| | (0.037) | (0.030) | (2.288) | (2.995) | (0.101) | (0.117) |
| 13+ | -0.121*** | -0.043 | -0.568 | -2.104 | 0.242^{*} | -0.279^* |
| | (0.045) | (0.031) | (2.280) | (3.088) | (0.126) | (0.153) |
| N | 16308 | 22450 | 13592 | 7394 | 10778 | 5748 |

Notes Panel A presents the results of the linear specifications. Panel B presents the results of the non-linear specifications. Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Table S6. Number of observations with diabetes (HbA1c $\geq 6.5\%$) and self-reported diabetes.

| | HbA1c < 6.5% | $HbA1c \ge 6.5\%$ | Total |
|-------------------------------|--------------|-------------------|-------|
| No self-reported diabetes (N) | 4544 | 1181 | 5725 |
| Row % | 79% | 21% | 100% |
| Cell % | 71% | 18% | - |
| Self-reported diabetes (N) | 129 | 554 | 683 |
| Row % | 19% | 81% | 100% |
| Cell % | 2% | 9% | - |
| Total (N) | 4673 | 1735 | 6408 |

Robustness checks

Table S7. Labour outcomes and self-reported diabetes controlling for obesity

| | Employment | | Weekly w | vork hours | Log hourly wages | |
|---|---|---------------------|--|--|------------------------------|--------------------------------|
| | Males | Females | Males | Females | Males | Females |
| $\underline{\text{Obese (BMI} \ge 30)}$ | 0.007 (0.012) | -0.005 (0.013) | $\begin{array}{c} -0.127 \\ (0.773) \end{array}$ | -1.144 (1.188) | 0.018 (0.038) | 0.082 (0.061) |
| Diabetes | $\underbrace{\begin{array}{c} -0.046 \\ (0.028) \end{array}}$ | -0.064** (0.027) | -0.689 (1.772) | $\begin{array}{c} -0.169 \\ (2.904) \end{array}$ | $\underbrace{0.036}_{0.078}$ | $\underbrace{0.033}_{(0.183)}$ |
| N_{\sim} | 17992 | 24145 | 14866 | 7929 | 11711 | 6166 |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10. ** p < 0.05. *** p < 0.01.

Table S8. Selection into types of work and self-reported diabetes controlling for obesity.

| | Males | | | Females | | | |
|------------------------|------------------------------|------------------------------|---------------------|---|---------------------|---------------------------------|--|
| | Non-agric. | Agric. | Self-employed | Non-agric. | Agric. | Self-employed | |
| Obese (BMI ≥ 30) | $0.005 \ (0.017)$ | -0.032** (0.013) | 0.036*** (0.014) | -0.021* (0.011) | 0.003 (0.004) | <u>0.010</u> (0.009) | |
| Diabetes | $\underbrace{0.010}_{0.033}$ | $\underbrace{0.002}_{0.023}$ | -0.060** (0.028) | $\underbrace{\begin{array}{c} -0.011 \\ (0.020) \end{array}}$ | -0.020** (0.010) | $\underbrace{-0.025}_{(0.021)}$ | |
| N | 17414 | 17414 | 17414 | 23458 | 23458 | 23458 | |

Notes Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, ** p < 0.05, *** p < 0.01.

Table S9. Relationship between self-reported years since diagnosis and employment probabilities using continuous duration and duration splines and controlling for obesity.

| | Employment | | Weekly work hours | | Log hourly wages | |
|--------------------------|------------|---------|-------------------|------------------|------------------|----------|
| | Males | Females | Males | Females | Males | Females |
| Panel A: linear effect | | | | | | |
| Obese (BMI ≥ 30) | 0.003 | -0.010 | 0.059 | -0.412 | 0.026 | 0.035 |
| | (0.012) | (0.014) | (0.831) | $(1.247)_{\sim}$ | (0.040) | (0.064) |
| Years since diagnosis | -0.019*** | -0.008 | 0.259 | -0.008 | -0.016 | -0.073** |
| | (0.006) | (0.006) | (0.375) | $(0.721)_{\sim}$ | (0.019) | (0.034) |
| Panel B: splines | | | | | | |
| Years since SR diagnosis | | | | | | |
| Obese (BMI ≥ 30) | 0.003 | -0.009 | 0.073 | -0.371 | 0.027 | 0.036 |
| | (0.013) | (0.014) | (0.832) | $(1.247)_{\sim}$ | (0.040) | (0.064) |
| 0-3 | -0.014 | -0.022 | 0.806 | 3.762 | -0.070 | 0.015 |
| | (0.015) | (0.017) | (1.051) | $(3.169)_{\sim}$ | (0.057) | (0.139) |
| 4-7~ | -0.003 | 0.009 | -0.293 | -3.921** | 0.035 | -0.121 |
| | (0.018) | (0.015) | (0.914) | (1.811) | (0.044) | (0.108) |
| 8-12 | -0.023 | 0.001 | -0.098 | 3.082* | -0.062 | -0.085 |
| | (0.022) | (0.016) | (1.350) | $(1.736)_{\sim}$ | (0.066) | (0.074) |
| <u>13+</u> | -0.038** | -0.024* | 0.855 | -1.128 | 0.005 | -0.065 |
| | (0.017) | (0.012) | (1.029) | $(1.421)_{\sim}$ | (0.053) | (0.063) |
| Panel C: dummies | | | | | | |
| Obese (BMI ≥ 30) | 0.005 | -0.009 | 0.044 | -0.378 | 0.026 | 0.031 |
| | (0.012) | (0.014) | (0.831) | (1.245) | (0.040) | (0.064) |
| 0-3 | 0.028 | -0.032 | 1.484 | 22.434* | 0.047 | -0.658 |
| | (0.059) | (0.065) | (3.825) | (11.579) | (0.212) | (0.622) |
| <u>4–7</u> | 0.001 | -0.054 | 2.399 | 12.909 | 0.013 | -0.793 |
| | (0.044) | (0.055) | (3.181) | (11.063) | (0.154) | (0.616) |
| <u>8–12</u> | -0.064 | 0.010 | 0.296 | 15.604 | -0.293 | -1.125* |
| | (0.069) | (0.066) | (4.994) | (11.038) | (0.247) | (0.583) |
| <u>13</u> ± | -0.208** | -0.073 | -1.966 | <u>17.459</u> * | 0.168 | -1.090** |
| | (0.105) | (0.081) | (4.975) | (10.262) | (0.256) | (0.499) |
| N _~ | 13912 | 19972 | 11622 | 6487 | 9262 | 5054 |

Notes Panel A presents the results of the linear specifications. Panel B presents the results of the non-linear specifications. Robust standard errors in parentheses. All models include variables for states, urbanization, level of education, marital status, number of children < 6, wealth, health insurance status, age squared and one dummy variable for each calendar year. * p < 0.10, *** p < 0.05, *** p < 0.01.