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

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 of 5.4 and 6.05.9 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.

Keywords: Mexico; diabetes; biomarker; wages; fixed effects; employment, working hours

Introduction 1.

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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 as well as global problem, with over two-thirds of people with diabetes living in the 35 developing world low- and middle-income countries (LMICs) (International Diabetes Federation 2015). In Mexico, diabetes prevalence is estimated to have has grown 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 Diabetes Fed-39 eration 2015), by increasing the risk for heart disease and stroke, blindness, kidney disease 40 and nerve problems, food neurologic problems, foot ulcers and amputations (Reynoso-41 Noverón et al. 2011). However, via effective self-management of the disease, many if 42 not all of the complications can be avoided through regular monitoring, behaviour change 43 and medication adherence, the occurrence of complications could be avoided or delayed in 44 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 complications 50 occurring during the productive lifespan. Only a minority of patients in Mexico achieves 51 adequate blood glucose control (Barquera, Campos-Nonato, et al. 2013). Moreover, dia-52 betes in Mexico coexist with high levels of infectious diseases, exposing the health system to a 'double-disease burden' increasing the pressure to identify treatment priorities and use existing resources efficiently is related to diseases, including depression, hypertension 55 and cardiovascular disease that impose a heavy burden onto the health system (World 56 Health Organization 2016). Despite the catastrophic impact of diabetes on health, its economic consequences, in

particular in LMICshave received less attention, especially its effects, have received little 59 attention. This applies in particular to the evidence on the effects of diabetes on labour 60 outcomes. The latter have been studied predominantly in (Seuring, Archangelidi, et al. 61 2015). In high-income countries , where substantial economic losses have been observed 62 (Brown, Pagán, et al. 2005; Brown, Perez, et al. 2011; Brown 2014; Latif 2009; Minor 63 2011, 2013; Minor and MacEwan 2016). For less evidence is available. One A rare LMIC study exploited a natural experiment in China and found a significant reduction in income due to a recent diabetes diagnosis (Liu and Zhu 2014). A study for Mexico, using 66 cross-sectional data from 2005, found a significant (p<0.01) reduction in employment 67 probabilities for males by 10 percentage points (p.p.) and for females by 4.5 p.p. (p<0.1) 68 (Seuring, Goryakin, et al. 2015). Most existing studies rely-relied on instrumental variable (IV) estimation to address the potential endogeneity of diabetes, using the genetic 70 component of diabetes based on its family history as an instrument, to address the potential endogeneity of diabetes. However, family history of diabetes may also proxy for other genetically transferred traits, including unobserved abilities, as well as intrahousehold or intergenerational dynamics that impact labour outcomes directly; the validity of this IV therefore remains debatable. Panel data methods provide the opportunity to account for 75 time-invariant unobserved individual characteristics, which may play an important role, 76 but have but—to the best of our knowledge—have not yet been usedby our knowledge. 77 Such unobservables, like for instance health endowments for instance hunger or nutrient 78 deficiency experienced in early life, could adversely affect health as well as the propensity 79 to develop type 2 diabetes in particular later in life (Ewijk 2011; Li et al. 2010; Sotomayor 80 2013); they may also affect. Additionally, there may also be long-term effects on labour 81 outcomes—either directly through their effects on reductions in contemporaneous produc-82 tivity (Currie and Vogl 2013), or indirectly by limiting educational attainment and human 83 capital accumulation (Ayyagari et al. 2011). These unobservables thereby present a major source of a potential bias that can be accounted for by the use of panel data estimation.

In parallel to these identification challenges, heterogeneity in impact and measurement 86 across the population also deserves further investigation. Recent evidence from Mexico 87 points to a strong positive relationship of diabetes duration with mortality due to diabetes 88 related complications (Herrington et al. 2018). A longer disease duration was found to be 89 related with higher glycated hemoglobin (HbA1c) levels, and undiagnosed diabetes had the 90 lowest diabetes related mortality risks. The latter points to potential selection issues when 91 using self-reported diabetes data to investigate economic outcomes as those. Those who 92 self-report, and hence tend to be diagnosed, are in worse health than those undiagnosed. 93 potentially leading. This can lead to an overestimation of the economic effects of diabetes, in particular in populations with a large undiagnosed population, such as in many LMICs 95 (Beagley et al. 2014). So far, however, little evidence exists on the economic impact according to diabetes severity and duration or for those with undiagnosed diabetes. 97 The objective of this study was to provide new evidence on the impact of diabetes on 98 labour outcomes, adding to previous work by paying close attention to the challenges 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 effects separately, fixed effects model we accounted for time-invariant heterogeneity when

unobserved heterogeneity, to the chronic nature of diabetes and to the population unaware 100 of their condition (i.e. the 'undiagnosed ') undiagnosed diabetes. We used three waves of 101 102 103 104 assessing the impact of self-reported diabetes and time since diagnosis on labour outcomes. 105 We also used rich and novel To assess the role of undiagnosed diabetes we used biomarker 106 data from the most recent wave of data, we assessed the role of undiagnosed diabetes last 107 wave of the MxFLS. 108

109 2. Data

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This paper used data from the Mexican Family Life Survey (MxFLS), a nation-110 ally representative longitudinal household survey, containing three waves conducted in 2002, 2005–2006 and 2009–2012. The baseline sample collected information from 35,000 112 individuals from 8,400 households in 150 communities throughout Mexico. Data was collected It is the only longitudinal household survey in Mexico that provides data on a 114 wide range of social, demographic, economic and health characteristics (Rubalcava and 115 Teruel 2013). Because the survey followed participants moving within Mexico as well as 116 to the US, around 90% of the original sample have been reinterviewed in the third wave. 117 Our samples were restricted to the working age population (15–64) and excluded pregnant 118 womenand those in school. Pregnant women have an increased diabetes risk and may 119 not be able to work. Since their inclusion may have biased the estimates, we dropped all 120 observations of women reporting to be pregnant at the time of the survey (N=764). We 121 also dropped those reporting to be in school. The first part of the analysis used all three 122 wavesand, exploiting the panel structure of the data. The second part used a biomarker 123 subsample of the third wave (2009–2012). Because the biomarker sample included ev-124 erybody above the age of 44, but only a random subsample of those aged 44 or below 125 (Crimmins et al. 2015), its age structure was older and hence its self-reported diabetes 126 prevalence higher. The analysis therefore compares with self-reported data for this specific 127 subsample only. 128 Our outcome variables of interest were employment status, weekly working hours, 129 hourly wage, and occupation. Employment status was defined as having carried out an 130 activity that helped with the household expenses the last week and while working for at 131 least four hours per week. We explicitly included informal employment and employment 132

without monetary remuneration, for instance in family businesses. Hourly wage was con-

structed as reported monthly income from the first and second job, divided by average

number of weeks per month and weekly working hours. Labour income was obtained from

the response to questions on wages, income from piecework, tips, income from extra hours, 136 meals, housing, transport, medical benefits and other earnings, or from the response to a 137 question on aggregate labour income for the entire month. We adjusted calculated wages 138 for inflation in the year of interview and considered the log of real wages. Due to a consid-139 erable number of missing or zero income reports, the sample used for the wage estimation 140 was smaller than the sample for working hours. Working hours were combined from both 141 the first and a potential second job. Descriptive statistics for the entire panel sample show 142 that 86 over 80% of men with diabetes and 87% of men without diabetes reported some 143 form of employment compared to, compared to 26% of women with diabetes and 37% 144 of women without diabetes (see Table 1). Interestingly, men did not report considerably 145 higher hourly wages than women but worked more hours per week. Men also more often worked There were also little differences in working hours and wages between men and women with and without diabetes. Men worked more often 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. Similarly, those 150 without diabetes were better educated than those with diabetes. Further, the diabetes 151 sample is about 15 years older on average than the non-diabetes sample, both for men 152 and women. 153

The first part of the analysis focused on the relationship of labour outcomes with 154 self-reported diabetes, which was based on the survey question: "Have you ever been 155 diagnosed with diabetes?". Because the data did not distinguish between type 1 and type 156 2 diabetes, we assumed that the estimates represented the impact of type 2 diabetes, by 157 far the most common type of diabetes in Mexico. As a robustness check, we re-estimated 158 our main results categorizing diabetes into early-onset and late-onset cases, according to 159 the age at which diabetes was first reported in the survey. This was a similar approach to 160 Alegre-Díaz et al. (2016), who assumed that everybody diagnosed before age 35 and using 161 insulin had type 1 diabetes. Accordingly, we assumed that those first reporting a diabetes 162

diagnosis before the cut-off had type 1 diabetes while those above had type 2 diabetes. 163 Nonetheless, because we cannot warranty that this is 100% accurate (as it may be unlikely 164 that both populations consisted exclusively of one type of diabetes, we preferred to think 165 of the groups as of early- and late onset groups (in the case of the within-coefficient, which 166 only takes into account incident cases). Because we did not have late-onset groups. This 167 separation also provides information about the exact age at diagnosis for all diabetes cases 168 in all three waves, effects for different age groups, as the late-onset group had an average 169 age of onset of 50 compared to 28 for the between coefficient may also stratify people with 170 diabetes into the late onset group, even though they actually had early onset diabetes but 171 only joined the sample after already having been diagnosed for several yearsearly-onset 172 group. In the pooled data, which combines all three waves, diabetes was self-reported by 5% of men and 6% of women, respectively. This is consistent with other reports from 174 Mexico for the time, showing a prevalence of diagnosed diabetes of 7.5\% in 2006 in a sample 175 also including people over the age of 64 (Barquera, Campos-Nonato, et al. 2013). Apart from self-reported diabetes, which was available in all rounds, we also used information 177 on the self-reported year of diagnosis as well as biometrically measured HbA1c levels for 178 a subsample of respondents from the third wave. 179

Information on the self-reported year of diagnosis, reported in the third wave, allowed us to construct a measure of time since diagnosis. For those also present in previous waves, 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.

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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 diabetes, i.e. the false negativesundiagnosed diabetes. The biometrically measured blood glucose value that allowed us to identify those with undiagnosed diabetes, was available for over 6000 respondents in the third wave. We used the internationally recognized cut-off of

an HbA1c \geq 6.5% to define diabetes as recommended by the World Health Organization (WHO) (World Health Organization 2011). As we show in Supplementary Table S6, 19% of self-reported diabetes cases had HbA1c levels below the diabetes threshold. We dropped those for our analysis as it was not clear if they had misreported their diabetes status or had achieved these low levels as a result of their successful disease management. Analysis including those cases led to qualitatively similar results (results available on request).

¹⁹⁶ 3. Estimation strategy

To investigate the relationship between self-reported diabetes and three labour outcomes— 197 employment, wages and weekly working hours—we estimated a effects model, an extension 198 of the correlated random effects modelfirst proposed by Mundlak (1978), that explicitly 199 models both within and between effects. The within effects are identical to a model, 200 accounting—weekly working hours and wages—we estimated a fixed effects model. The 201 fixed effects model accounts for the potential bias introduced by time-invariant unobserv-202 ables, providing an estimate of the effect for cases that received a diagnosis throughout 203 the survey (705 incident cases compared to 970 non-changing diabetes cases in the used sample). Modeling the between effect allowed us to also use information from those that 205 already had diabetes at baseline. 206

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

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 of all other time-varying covariates were also included to capture the between effect. The error terms u_i and e_{it} capture the errors for the within and between variation, respectively.

 Y_{it} was a binary variable taking a value of 1 if respondent i reported being in employment at time t and 0 otherwise. Making use of the user written Stata command xthybrid, we estimated effects applying a multilevel mixed-effects generalized linear model. For ease of interpretation we chose to estimate a linear probability model for the effects of diabetes on employment.

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 We also included time variant confounders—as the fixed 221 effects model accounts for time invariant confounding only. We controlled for changes in 222 the level of urbanization, education, state the level of education, the state of residence, 223 marital status, the number of children below the age of 6-six in the household, a quadratic 224 age term, and calendar year dummies as well as household wealth. We also control for 225 household wealth approximated by a household asset index. We composed an indicator 226 using principal component of household assets and housing following Filmer et al. (2001) 227 (Filmer and Pritchett 2001). The assets indicators asset index reflected owning a vehicle, 228 a second house, a washing machine, dryer, stove, refrigerator or furniture, any electric ap-229 pliances, any domestic appliances, a bicycle, farm animals, and accounted for the physical 230 condition of the house, proxied by the type of floor material and water access. In our 231 main regression models we did not account for body mass index (BMI). While part of 232 the effect of diabetes may be due to potential adverse effects of obesity, including BMI 233 as a control variable in the model would have led to biased attenuated estimates if the 234 diagnosis of diabetes also had an effect on BMI, which was likely has been shown to be the 235 case. In general, control variables should not also be potential outcome variables, hence 236 we similarly-in other studies (De Fine Olivarius et al. 2015; Seuring, Suhrcke, et al. 2018; 237 Slade 2012). Similarly, we did not control for other chronic diseases that may have been 238 caused by any diseases that were likely consequences of diabetes, such as hypertension 239

or cardiovascular disease heart disease or other micro- and macro-vascular complications
(World Health Organization 2016), as this would prevent any causal interpretation of the
relationship of diabetes with our labour market outcomes (Angrist and Pischke 2009).
Nonetheless, we carried out a robustness analysis controlling for obesity. Stata 15 was
used for all analyses (StataCorp 2017).

45 3.1. Labour outcomes and time since diagnosis

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 $\beta_1 Dyears_{it}$ was continuous replaced the binary diabetes indicator of Eq 1 with
a continuous variable indicating years since the diagnosis was first reported. While
simultaneous 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. In this
case, identification 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.

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

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, ..., N. The coefficient δ_n captured
the effect of diabetes for the n-th interval. The effects are linear if $\delta_1 = \delta_2 = , ..., = \delta_n$.

Based on visual inspection (Fig. 1 on page 29) we chose three nodes located at 3, 7 and 259 12 years after diagnosis. The first three years should capture any immediate effects of the 260 diagnosis, the years four to seven any effects during time of adaptation to the disease and 261 the later terms the long term effects. We also estimated a non-linear model using dummy 262 variables for duration groups rather than splines, applying the same duration cut-offs. 263 Because the year of diagnosis was only reported in the third wave, time since diagnosis 264 was not available for those who were not interviewed in the third round. A reported 265 diagnosis in the year of the interview was counted as 'one year since diagnosis'. 266

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

$$Y_i = \beta_0 + \beta_1 D bio^{\frac{1}{2}} + \beta_2 X_i + v_i + u_i, \tag{3}$$

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

outcomes, using the following equation:

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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).

$$Y_i = \beta_0 + \beta_1 Dsr_i + \beta_2 Dbio_i + \frac{\beta_3 Dsr_i * Dbio_i + \beta_4 X_i + v_i + u_i.$$
 (4)

Note that the interaction term changes This changed the interpretation of β_1 and 282 β_2 , with β_1 now representing the effect on those aware of their condition but with levels 283 below the diabetes threshold; while β_2 reflects $\underline{Dbio_i}$, which now reflected the effect on 284 those with undiagnosed diabetes, i.e. the respondents not self-reporting diabetes but with 285 HbA1c levels equal to or above the threshold. The interaction term β_3 shows the effect 286 for those with self-reported diabetes and levels above the threshold. 287 We further investigated the effect of the severity of diabetes on labour outcomes, re-288 placing $\frac{Dbio^d}{Dbio}$ with $Dbio^c$, a variable that was 0 for HbA1c < 6.5% and took the actual value of increased by one for every percentage point increase in HbA1c for those 290 with an $HbA1c \ge 6.5\%$ (Eq 5). This allowed us to investigate the effect of a one percent-291 age point increase in HbA1c levels for people with undiagnosed diabetes (β_2) as well as 292 for those with self-reported diabetes above the diabetes threshold (β_3) . 293

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

294 4. Results

²⁹⁵ 4.1. Labour outcomes and self-reported diabetes

The results of estimating Eq 1 in Table 2 indicated significant and substantial statistically significant 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, 300 it provides suggestive evidence that time-invariant unmeasured confounders may play a 301 limited role. Employment probabilities were reduced by over 5-5.4 p.p. for both genders, 302 translating into relative reductions of 14% for women and of 6% for menmen and 5.9 303 p.p. for women. There was no significant relationship between diabetes on one hand and 304 working hours and wageson the other, though the between estimates suggested that men 305 with diabetes earned generally more than their counterparts without diabetes (Table 2). 306 Overall these results thus suggested effects at the extensive margin (employment), but not 307 at the intensive margin (labour supply and productivity) or wages. 308

Dividing the diabetes population into early and late onset groups, men, and potentially 309 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 311 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 316 were less precise but may indicated 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 diabetes higher wages for women with an early diabetes onset, but no effects for men.

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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

found no statistically significant effects for men. The between effects suggested that 327 men with diabetes were less likely to be employed in agriculture, but more likely to be 328 self-employed. Women with diabetes were less likely to be employed in agricultural and 329 non-agricultural jobs-but not for men (Table 3). Disaggregating the diabetes groups fur-330 ther according to their age showed that most statistically significant relationships were 331 driven by the older onset older-onset group (Supplementary Table S3). Interestingly, for 332 For male self-employment, incidence of diabetes increased the probabilities to be self-333 employed in the younger group, while it reduced the probabilities to be self-employed in 334 the older onset group. However, especially the results for early onset diabetes in women 335 should be interpreted carefully due to limited number of diabetes cases. older-onset group. 336

We 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 itself did not appear to be an independent predictor of any labour market outcome. The estimates of the effect of diabetes remained similar for most outcomes. Only for male employment probabilities, the diabetes coefficient was no longer statistically significant at the 10 percent significance level. Estimating the original model without accounting for obesity, but using the sample with only non-missing BMI cases, showed similar changes in effects (results available on request).

4.2. Labour outcomes and time since diagnosis

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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 the linear duration analysis, 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 wages using 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 age 35.

The non-linear results for the spline function and dummy variable approach are pre-sented 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 di-agnosis. 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 the separation into duration groups and into within and between variation. Reestimating the specifications with a random effects model that combined both types of variation into one estimate showed that, at least for the models with dum-mies, 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.

Controlling for obesity, results remained very similar and obesity itself was not found

4.3. Cross-sectional biomarker analysis

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As reported in Supplementary Table S6, 18% of the observations in the biomarker sample 382 were false negatives, i.e. undiagnosed. Further, 2\% were false positives, though the latter 383 may have included eases that received a diabetes diagnosis and managed to reduce their to non-diabetes levels via medication and/or lifestyle changes. Overall 80% of the self-reports 385 were consistent with the biomarker data. undiagnosed, which accounts for 68% of all cases 386 above the diabetes threshold. Comparing the health status and diabetes risk factors of the 387 diagnosed and undiagnosed diabetes populations suggested that those with self-reported 388 diabetes were older and in worse health, both objectively and subjectively compared to 389 those undiagnosed. This suggests a selection into the diagnosed group based on the severity 390 and potentially duration of diabetes. If the adverse effects of diabetes are due to its health 391 impact, we would suspect worse labour market outcomes for the diagnosed compared to 392 the undiagnosed population, compared to those undiagnosed. 393 Table 6 presents the results from estimating Eq. 2-5. Panel A confirms the earlier 394 longitudinal results using self-reported diabetes for the cross-sectional biomarker sample. 395 The results in panel B indicate that the relationship with employment became weaker 396 when using diabetes defined by the biomarker instead of self-reported diabetes, in par-397 ticular for men. Results in Panel C were obtained from estimating Eq. 4 and indicate 398 an absence of a (statistically significant) negative relationship between undiagnosed dia-399 betes (expressed in the 'Biomarker diabetes but not self-reported' coefficient) and labour 400 outcomes. The coefficients for the interaction term were mostly negative, though only 401 statistically significant in the case of female working hours. 402 To explore whether the adverse effects increased with higher HbA1c levels, we estimated 403 Eq. 5. The results in panel D again only suggest a negative association of a one only 404

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.

5. Discussion

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Diabetes is now one of the most common chronic diseases in low- and middle income 410 countries LMICs, as well as high-income countries high-income countries (HICs), with po-411 tential severe impacts on the health and economic well-being of those affected. Yet rigorous 412 evidence on the economic consequences for LMICs remains scarce. 413 To address key methodological challenges, this paper used rich longitudinal panel data 414 from Mexico that also contained diabetes biomarkers. The biomarker data showed alarm-415 ing levels of clinically tested diabetes (27% prevalence) and indicate that a large proportion 416 of the Mexican population (18%) is unaware of their condition has undiagnosed diabetes. 417 Overall, the paper found The paper provided evidence for adverse effects of self-418 reported diabetes on the probability of being employed, confirming earlier findings for Mexico that used employment, working hours and wages. While earlier work showed evidence for Mexico for employment (Seuring, Goryakin, et al. 2015), this paper presented, 421 by our knowledge, the first evidence on the relationship of diabetes, working hours and 422 wages. Furthermore, we added to the study of Seuring, Goryakin, et al. (2015) by using 423 longitudinal instead of cross-sectional information. But, these new results also suggest a 424 comparatively larger data to identify a causal relationship. We provided first evidence 425 of the long term impact of diabetes on female employment probabilities. The evidence 426 also points towards the main effects being in Mexico and explored the extent and effects 427 of undiagnosed diabetes. We confirmed the findings of Seuring, Goryakin, et al. (2015) 428 insofar that we found an adverse effect of diabetes on male employment. We further showed 429

more conclusive evidence that also women experience a reduction in their employment

probabilities due to diabetes. Taking into account the differences in employment between 431 men and women, the found effects translate into relative reductions of their employment 432 probabilities of 6% for men and 14% for women. We also found that the effects were mainly 433 driven by those with a diabetes onset at a relatively later state, consisting most likely of 434 people with of older people with most likely type 2 diabetes. This was also found by Seur-435 ing, Goryakin, et al. (2015) in their stratified analysis of an older and younger age group. 436 Analyses of the long term impact indicated that the employment probability fell gradually 437 in the years following diagnosis. The results for the using a non-linear models model were 438 less clear, potentially due to reductions in statistical power, but suggested that 439 adverse effects became stronger with time since diagnosis. The linear effect contrasts with 440 estimates for the USA, where such an effect seemed to be absent, was absent; however allowing for non-linearity, revealed falling employment probabilities after 11 to 15 years 442 for females and after 2-5 years for males (Minor 2013). For working hours and wages, our results were more ambiguous with adverse effects being observed for time since diagnosis on female wages only. In summary, most of the adverse effects of diabetes were found 445 at the extensive margin, mainly affecting the probability of employment, rather than the intensive margin, i.e. working hours or wages. 447 Overall, a relationship of diabetes with and working hours or wages is mostly absent, 448

Overall, a relationship of diabetes with and 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 selectionwas mostly absent. 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 have lacked access to quality diabetes care, making it more likely that they develop developed severe complications earlier (Flores-Hernández et al. 2015). They

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are also may also have been more likely to be in informal employment and low skilled jobs

, with less job security and are, and thus more prone to being laid off to be and replaced

with healthier workers.

Because We found that self-reported diabetes may not adequately represent cases were 461 not representative of the entire diabetes population if the share of undiagnosed diabetes 462 is relatively large and those undiagnosed differ from those diagnosed, estimates based 463 on self-reported diabetes are likely to be biased. Our results using the cross-sectional 464 biomarker data suggest indeed that those with undiagnosed diabetes were in Mexico. 465 A large share of people with diabetes were undiagnosed and significantly healthier and 466 younger, suggesting a selection into the diagnosed group based on the severity and duration 467 of diabetes. Consequently, diabetes based on biomarkers was found to be as defined by 468 the HbA1c threshold, was less related to reduced employment, probabilities compared to 469 self-reported diabetes, in particular for men. Further analysis showed that this was due to 470 the absence of any an association between undiagnosed diabetes and employment. These results are similar to those found for the USA, where no statistically significant relationship 472 was observed between undiagnosed diabetes on employment, while a significant effect of 473 diagnosed diabetes was observed (Minor and MacEwan 2016). Our results further indicate 474 indicated that the difference in the employment effects of diagnosed and undiagnosed 475 diabetes was not mediated by current HbA1c levels. This is similar to findings for 476 Mexican-Americans in the USA, where employment outcomes were unrelated to higher 477 HbA1c levels (Brown, Perez, et al. 2011). This may stem from A possible explanation 478 may be that HbA1c levels primarily being are primarily informative for the last three 479 months, and not being the only are not the only or best indicator for the severity of 480 diabetes. Overall, it seems that general health differences related to a longer diabetes 481 duration with its related health consequences, and selection into the diagnosed population 482 , for instance based on emerging diabetes related health problems, could be have been 483 driving the adverse economic effects among those with a self-reported diagnosis. 484

Our study had several limitations. While the within-coefficient accounts our model 485 accounted for any time-invariant confounding, the estimates may have been affected by 486 unobserved time-variant confounders. Reverse causality, where employment status affects 487 the propensity to develop or be diagnosed with diabetes, may also play have played a 488 role. Existing studies that looked at this particular direction of causality, however, have 489 not found strong evidence for an effect of employment status on diabetes (Bergemann 490 et al. 2011; Schaller and Stevens 2015), though they were carried out in high-income 491 countriesHICs. We did not control for the effects of obesity, hypertension, self-reported 492 health or other diseases in our models due to the high probability that they were affected by 493 diabetes themselves, which would have made a causal interpretation of the estimates more 494 difficult. Robustness checks including obesity indicated that our main findings remained 495 mostly unchanged and that any occurring changes were likely the result of a smaller 496 sample size rather than of the inclusion of obesity. For the duration analysis an additional 497 limitation, imposed by the data, was that the year of diagnosis was only reported in the third wave. While this still allowed us to construct an estimate of the time since diagnosis 499 for the previous waves, it restricted the analysis to those that were present in the last 500 wave, thereby excluding those that dropped out of the sample prior to the third wave. 501 Finally, we used a WHO recommended HbA1c cut-off to diagnose diabetes, due to the 502 lack of a Mexico specific cut-off. There is some evidence that HbA1c may be affected by 503 ethnicity (Sacks 2011). Hence, if Mexican ethnicity would lead to different HbA1c levels, 504 the use of our cut-off could have led to misclassification based on the used biomarkers. 505 Despite these limitations, our findings bear important implications. First, the impact 506 of self-reported diabetes on labor outcomes in Mexico seems seemed mostly limited to 507 508

of self-reported diabetes on labor outcomes in Mexico seems seemed mostly limited to its effect on employment probabilities, though there is some indication that it could also reduce wages over time for women. Second, its effect on employment was much stronger for females, though the underlying reasons for this remain unclear. Potential explanations are that lower working hours or wages for women make a dropout less costly. Other

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evidence suggests that women with diabetes are in worse metabolic health compared to 512 men when they cross the diabetes threshold (Peters et al. 2015), making it more likely 513 for them to drop out. Third, caution is needed when estimates based on self-reported 514 diabetes are interpreted in terms of the entire population, i.e. extending to those with 515 undiagnosed diabetes. Ideally, studies would include a biomarker analysis, acknowledge 516 the differences between diagnosed and undiagnosed subpopulations, undiagnosed subpopulations, and 517 carry out a separate analysis whenever feasible. If this is not possible, study conclusions 518 about the effect effects of self-reported diabetes should be limited to this specific part of 519 the population, in particular in environments. This is of particular importance in LMICs 520 where the share of undiagnosed diabetes is high, as is the case for most low- and middle 521 income countriesoften high. 522

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While we find no effect for undiagnosed diabetes in our cross section analysis, further inquiry over time is needed. The large proportion of previously undiagnosed cases indicates that diagnosis—at least in Mexico—happens found in this paper indicates that diagnosis—at least in Mexico—happens late or not at all. This may reduce the possibilities to prevent complications via treatment and self-management, increasing the risk of severe complications appearing premature. Earlier diagnosis and ensuing effective treatment may lighten the health and economic burden. Further analysis is also needed to explain why the adverse economic effects are so large for women. Ultimately, 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. Given the established links between early life heath and later life incidence. Therefore, more research is needed to investigate the economic impact of diabetes over time. Longitudinal biomarker information could be used to observe the true duration and severity of diabetes as well as other chronic diseases, investments in maternal and child health may be particularly attractive to address non-communicable and communicable diseases at the same timethe time that passes till a medical diagnosis. This would allow for a better understanding of when

adverse economic effects start to appear. Further, future research should investigate how
time of diagnosis and treatment of diabetes affect the occurrence of adverse labour market
effects of diabetes. The results of such research could allow costing studies to include more
detailed information on the indirect costs of diabetes; or inform cost-effectiveness analyses
that aim to include a measure of the potential benefit of the intervention to employers or
society at large.

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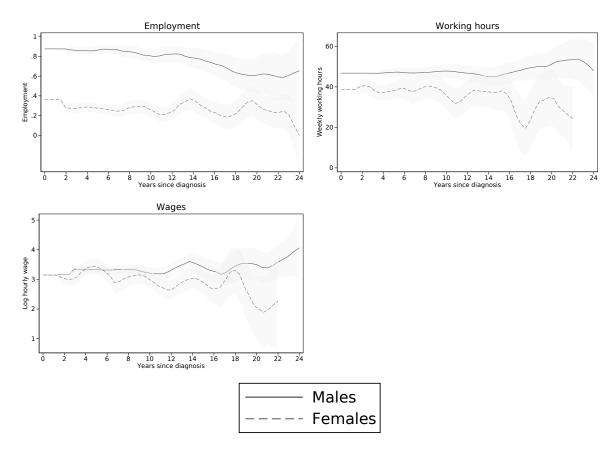
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Figures

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.

687 Tables

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.

Table 2. Labour outcomes and self-reported diabetes

	Employment		Weekly working ho	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- 1.998 (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.

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 - <u>0.006</u> (0.029)	-0.008 (0.022)	-0.042 <u>-0.043</u> (0.026)	-0.001 (0.018)	-0.022^{**} (0.009)	-0.030 -0.029 (0.018)
N	20719	20719	20719	26575	26575	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 4. Relationship between self-reported years since diagnosis and employment probabilities using continuous duration and duration splines.

	Emplo	yment	Weekly wor	king 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.

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

		Males			Females	
	Diagnosed diabetes	Undiagnosed diabetes	P value (t-test)	Diagnosed diabetes	Undiagnosed diabetes	P value (t-test)
Employed	0.8070.811	0.8750.877	0.0120.019	0.2410.233	0.3310.329	0.002
Hourly wage	35.931 35.280	30.670 30.939	0.1200.220	36.092 37.242	32.638 32.822	0.5500.495
Usual weekly working hours	44.341 44.562	46.682	0.1040.166	34.708 31.838	39.681 39.788	0.0460.004
Age	53.162 53.258	44.72045.530	0.000	53.167 53.544	44.449 45.388	0.000
Any medical insurance	0.6770.691	0.5990.589	0.0330.009	0.7330.717	0.6430.645	0.0020.025
City of 2,500-15,000	0.0960.092	0.1070.105	0.6430.593	$\frac{0.112}{0.116}$	$\frac{0.112}{0.114}$	0.9940.916
City of 15,000-100,000	0.135 0.147	0.0960.090	0.0980.021	0.087 0.079	0.0900.093	0.8840.447
City of >100,000	0.3310.332	0.2970.290	$0.325\widetilde{0.267}$	$0.294\widetilde{0.292}$	0.3330.329	$0.185\widetilde{0.250}$
Married	0.7190.751	0.6430.663	0.0310.018	0.6330.629	0.5690.588	0.0350.221
Number of children (age < 615) in household	0.9500.972	1.122 1.138	0.0730.110	0.960 0.934	1.257 1.250	0.001
Indigenous group	0.1580.171	0.2110.216	0.0790.159	0.1950.192	0.2070.209	0.6260.534
Primary	0.4790.484	0.4340.450	0.2380.406	0.6260.635	0.4650.479	0.000
Secondary	0.212	0.2310.230	0.5540.594	$\frac{0.132}{0.126}$	0.2330.230	0.000
High school	0.0620.060	0.1310.115	0.0030.022	0.0370.031	0.117 0.105	0.000
Higher education	$\frac{0.139}{0.147}$	0.1130.109	0.2880.147	0.0300.025	$\frac{0.073}{0.071}$	0.003
Wealth index	-0.174 -0.213	$\frac{0.117}{0.141}$	0.000	0.0120.033	0.1030.104	0.1570.314
Subjective health		100000		100000		100000
very good	0.0230.014	0.0940.092	0.000	$\frac{0.012}{0.013}$	0.0540.044	0.0010.010
good	0.2120.184	0.4340.431	0.000	0.1800.173	0.3670.370	0.000
fair	0.6190.664	0.4420.446	0.000	0.6430.635	0.5280.533	0.0000.002
bad	$\frac{0.135}{0.129}$	0.0260.027	0.000	$\frac{0.155}{0.170}$	$\frac{0.048}{0.047}$	0.000
very bad	0.012 0.009	0.004	0.1870.374	0.0100.009	0.004	0.2460.344
Glycated hemoglobin (HbA1c)	9.0379.635	8.5338.531	0.0040.000	8.9799.781	8.6808.699	0.0490.000
Hypertension (self-reported)	$\frac{0.262}{0.258}$	0.0740.078	0.000	0.3970.384	$\frac{0.150}{0.157}$	0.000
Blood pressure		100000		100000		
Systolic	136.688 136.475	130.506 130.981	0.0000.001	136.070 136.426	122.835 123.516	0.000
Diastolic	84.677 84.562	82.063 82.448	0.0030.025	84.495 84.912	79.689 80.019	0.000
Heart disease (self-reported)	0.0350.032	0.0070.008	0.0040.013	0.0500.041	0.0240.025	0.0210.178
BMI	28.868 28.989	28.311 28.385	0.1350.128	30.640 30.573	29.778 30.058	0.0320.234
Obese (BMI ≥ 30)	0.3380.374	0.3110.333	0.4400.301	0.4690.500	0.4310.470	0.2250.388

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

Table 6. Biomarker results.

	Employ	ment	Weekly work	ing hours	Log hourly	v 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	025034
	(.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	(/	(/	(((/	(7
Self-reported diabetes	067~.080*	035066	1.471084	4.974-4.463	.157 061	017011
	(.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 × HbA1c if ≥ 6.5	.001 003	.000 010	207064	-852.375	023002	.005 000
	(.007 .012)	(.005 .012)	(294,668)	(-4241.043)	(.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 one dummy 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.

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 are a lack of a diagnosis. Wrong self-reports indicating a diagnosis of diabetes we deemed less of a problem since incentives to report diabetes when one does not have it falsely report a diabetes diagnosis seem to be very limited—although we cannot exclude this. A study from China finds found that the vast majority (98%) of those who self-report diabetes are self-reported diabetes were tested positive for diabetes, while only a minority

of those who are were tested positive for diabetes (40%) actually self-report self-reported
the disease (Yuan et al. 2015). Our data showed a similar pattern, with a low proportion
(32%) of the respondents being tested negative while self-reporting diabetes, while the
majority of those who are were tested positive (68%) do did not self-report diabetes.

We used the above information to infer the "true" diabetes status for those with incon-718 sistent reports. For respondents present in all three waves, we corrected inconsistencies as 719 reported in Supplementary Table S1. We assumed that if diabetes was reported only once 720 in the first two waves (either in 2002 or 2005) and then not reported again in the ensu-721 ing waves, this diabetes report was likely to be false (see lines 3 and 4 in Supplementary 722 Table S1) and that the person never had received a diagnosis. If a diabetes diagnosis was 723 however reported in two of the three waves (in 2002 and 2009 but not 2005, or in 2002 and 724 2005 but not in 2009), we assumed that the respondent had diabetes in all three waves 725 (see lines 1 and 2 in Supplementary Table S1). For cases where we only had information 726 from two waves, we assumed that if a diabetes diagnosis had been reported in a prior wave 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 729 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 the 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

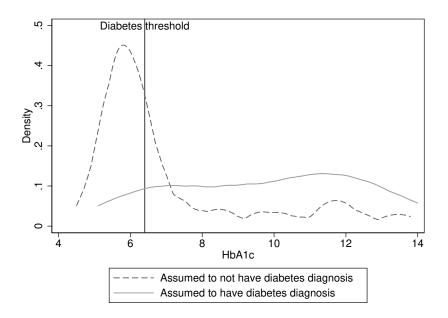
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in Supplementary Table S1)). Supplementary Figure S1 illustrates the difference between 736 both groups and suggests that indeed those with two self-reports of diabetes are were much 737 more likely to have HbA1c values above the diabetes threshold. A t-test comparing the 738 mean HbA1c for the two groups indicates indicated that those with two self-reports also 739 have had significantly (p<0.001) higher HbA1c levels than those with only one self-report 740 of diabetes (9.7% vs. 7.1%). Further, of those with one self-report, only 30% have had 741 an HbA1c $\geq 6.5\%$ 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 no 743 diabetes. 744

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.

	Employment		Weekly working hours		Log hourly wages	
_	Males	Females	Males	Females	Males	Females
Early onset(within)	0.133 (0.176)	-0.206** (0.086)	14.71214.856* (8.3708.329)	-18.636-18.250* (9.6689.515)	-0.523 -0.490 (0.340 0.337)	0.3880.375*** (0.057)
Late onset (within)	-0.059** (0.025 0.026)	-0.048 <u>0.047*</u> (0.025)	-1.008 <u>-0.936</u> (1.513 1.510)	-1.322 -1.346 (2.553)	0.079 0.071 (0.067) 0.068)	0.067-0.074 (0.1630.161)
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.

	No	on-agric.	Agric	ulture	Self-employed	
	Males	Females	Males	Females	Males	Females
Early onset (within)	0.030 (0.216)	-0.105 (0.074)	-0.225 <u>-0.226</u> (0.139)	-0.068 (0.047)	0.328** (0.161)	-0.027 (0.048)
Late onset (within)	-0.008 -0.007 (0.029)	$0.007 \\ (0.0180.019)$	-0.002 - <u>0.001</u> (0.022)	-0.019-0.018** (0.009)	-0.053-0.054** (0.026)	-0.030 -0.029 (0.019)
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		Mont	hly work hours	Log hourly wages	
	Males	Females	Males	Females	Males	Females
Early onset(within)	-0.009 <u>-0.005</u> (0.017)	$\begin{array}{c} 0.002 \ 0.007 \\ (0.013) \end{array}$				
Late onset (within)	-0.012^{**} (0.005)	-0.007-0.008** (0.004)	$\frac{0.086 - 0.088}{(0.275)}$	$\begin{array}{c} 0.308 - 0.303 \\ (0.5040.505) \end{array}$	-0.006 <u>-0.007</u> (0.013)	-0.059^{***} (0.022)
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.

746 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 wor	king hours	Log hourl	y 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

747 Robustness checks

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

	Employment		Weekly w	orking hours	Log hourly wages	
	Males	Females	Males	Females	Males	Females
Obese (BMI ≥ 30)	0.007 (0.012)	-0.005 (0.013)	(0.773)	-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)	$\underbrace{\begin{array}{c} -0.689 \\ (1.772) \end{array}}_{}$	(2.904)	$\underbrace{0.036}_{(0.078)}$	$\underbrace{0.033}_{(0.183)}$
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	-0.032**	0.036*** (0.014)	-0.021*	0.003	0.010	
Diabetes	$ \begin{array}{c} (0.017) \\ 0.010 \\ (0.033) \end{array} $	$ \begin{array}{c} (0.013) \\ 0.002 \\ (0.023) \end{array} $	$\begin{array}{c} (0.014) \\ -0.060 ** \\ (0.028) \end{array}$	$\begin{array}{c} (0.011) \\ -0.011 \\ (0.020) \end{array}$	$\begin{array}{c} (0.004) \\ -0.020** \\ \hline (0.010) \end{array}$	$ \begin{array}{c} (0.009) \\ -0.025 \\ (0.021) \end{array} $	
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 working 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)	(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)	(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)	(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)	(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)	(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)	(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	17.459*	0.168	-1.090**
	(0.105)	(0.081)	(4.975)	$(10.262)_{\sim}$	(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.