

Increasing prevalence of gestational diabetes in women living with HIV in the UK and Ireland

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Background



<u>Current UK guidelines</u> recommend <u>selective screening 24-28 weeks</u> based on known <u>risk factors for GD</u>:

- Ethnicity
- Family history of diabetes
- Body Mass Index (>30 kg/m²)

- Risk of gestational diabetes (GD) is increasing proportionally to that of type 2 diabetes globally
- Risk factors for women living with HIV (WLWH) are poorly understood
- Conflicting evidence on the effect of antiretroviral therapy (ART) and more specifically, PI-based regimens on GD risk
- Aim of this study was to understand the characteristics of WLWH who had GD and key birth outcomes, and investigate risk factors for developing GD



Methods



- Collects population-level surveillance data from all pregnant women living with HIV in the UK and Ireland*
- Data include HIV infection history, test results, pregnancy and birth outcomes

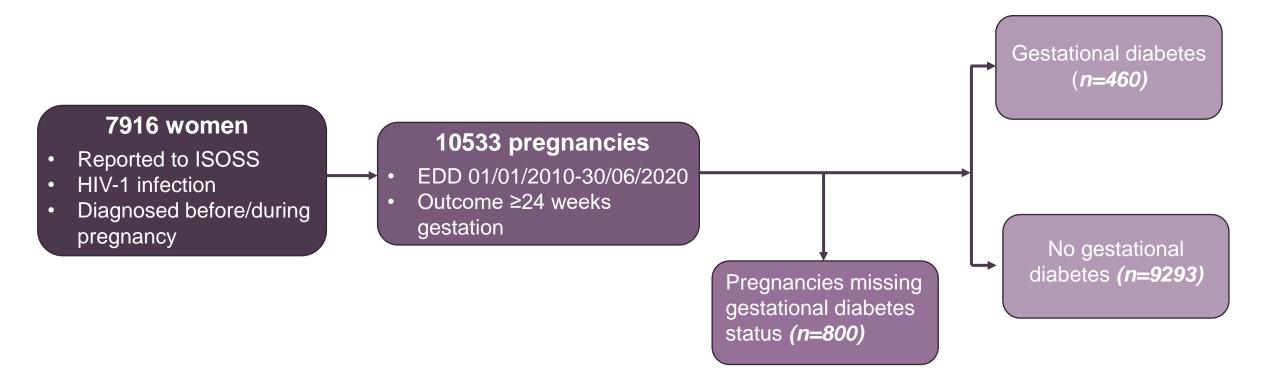
*Ireland until 2019 and England only from 2020

STATISTICAL ANALYSIS

- Multivariable logistic regression model with complete case analysis included covariates assessed for their association with GD using chi-squared tests (p<0.05)
- Regression model fitted using generalised estimating equations (GEE) to account for multiple pregnancies per woman
- Sensitivity analyses conducted to evaluate effect of universal antiretroviral therapy from 2015 on risk factors for GD

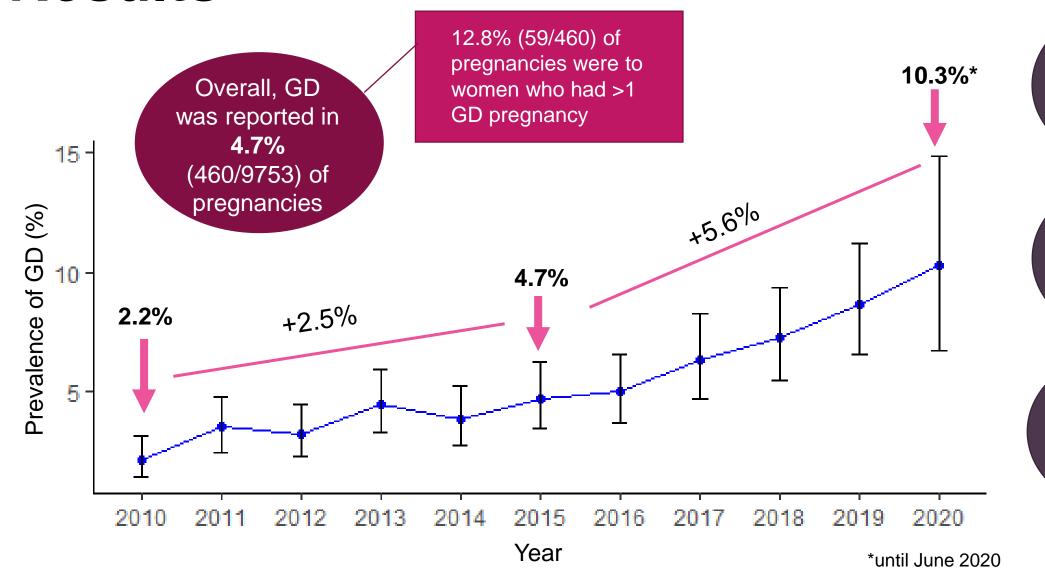


Overview of study population





Results



Median maternal age was **33 years** (q1:29 – q3:37)

71.8% of pregnancies were to Black African women

63.9% of pregnancies were to women on treatment at conception



Results – risk factors and outcomes

Risk factor	GD (<i>n</i> =460, %)	No GD (<i>n</i> =9753, %)
≥35 years	60.9	40.9
ART at conception	73.6	63.9
Asian	5.2	2.8
PI-based regimens	51.3	58.9
CD4 <350	19.0	26.8

 More women with GD were ≥35 years, on ART at conception and Asian compare to women without GD

Pregnancy Outcome	GD (<i>n</i> =460, %)	No GD (<i>n</i> =9753, %)
Stillbirth	1.30	0.60
Emergency Caesarean	33.5	23.8
Preterm delivery	15.7	10.9
Fetal Macrosomia	8.2	4.8

 Adverse pregnancy outcomes occurred more frequently for women with GD compared to women without GD



Results – statistical analysis

Risk Factor	OR	(95% CI)	GEE-aOR	(95% CI)
Year of estimated date of delivery ^a	1.15	(1.11-1.18)	1.12	(1.08-1.16)
Maternal age at delivery (years)				
<25	1		1	
25-34	1.95	(1.10-3.44)	1.76	(0.97-3.19)
≥35	4.10	(2.34-7.19)	3.28	(1.80-5.97)
Maternal ethnicity				
White	1		1	
Black African	1.62	(1.22-2.16)	1.55	(1.13-2.12)
Black Caribbean	1.65	(0.93-2.91)	1.72	(0.94-3.14)
Asian	2.86	(1.75-4.69)	2.60	(1.46-4.63)
Other	1.39	(0.75-2.58)	1.09	(0.50-2.38)
Maternal CD4 cell count (cells/μL)				
≥350	1		1	
<350	0.63	(0.49 - 0.81)	0.74	(0.57-0.96)
On treatment at conception				
No	1		1	
Yes	1.58	(1.28-1.96)	0.98	(0.77-1.25)
PI use in pregnancy				
No	1		1	
Yes	0.73	(0.61-0.89)	0.89	(0.72-1.10)

^a Year adjusted as a continuous variable in logistic regression models. Data reported by September 2020 for deliveries occurring 1 January 2010 and 30 June 2020.

- 9066 (93%) pregnancies had complete data on maternal risk factors
- Advanced maternal age (≥35 years) and Asian ethnicity remained significant risk factors in sensitivity analyses
- ART was not associated with GD risk in whole population and sensitivity analyses



Conclusion

- Overall prevalence of GD found in this study (4.7%, 95% CI: 4.3-5.2%) was slightly higher than the pooled prevalence of GD reported for Europe & America in a recent systematic review and meta-analysis (3.6% and 3.19% respectively)
- In the UK, prevalence of GD among WLWH has increased significantly in the last 10 years, even after accounting for ageing population
- Advanced maternal age was most strongly associated with increased odds of GD and this is not accounted for in current screening guidelines
- Limitations include missing data and being unable to account for weight indices such as BMI which is a known risk factor for GD in analyses



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- For any queries, please get in touch: l.bukasa@ucl.ac.uk

More information on ISOSS can be found here: www.ucl.ac.uk/isoss