

# Assessing prevalence of alcohol consumption in early pregnancy: Self-report compared to blood biomarker analysis.

Helen Howlett,<sup>1</sup> Shonag Mackenzie,<sup>1</sup> William K. Gray,<sup>1</sup> Judith Rankin,<sup>2</sup> Leanne Nixon<sup>1</sup>, Anthony Richardson,<sup>1</sup> Eugen-Matthias Strehle,<sup>1</sup> and Nigel W. Brown<sup>1</sup>

1. Northumbria Healthcare NHS Foundation Trust, North Tyneside General Hospital, North Shields, UK.

2. Institute of Health & Society, Newcastle University, Newcastle upon Tyne, UK

## BACKGROUND

Providing appropriate antenatal and postnatal care for women who drink alcohol in pregnancy is only possible if those at risk can be identified<sup>1</sup>. We aimed to compare the prevalence of alcohol consumption in the first trimester of pregnancy using self-report and blood biomarker analysis.

## METHODS

Six-hundred routine blood samples taken at antenatal booking in the first trimester of pregnancy, were anonymously analysed for the presence of Carbohydrate Deficient Transferrin (CDT), a validated marker of chronic alcohol exposure (normalising 2-3 weeks from the start of abstinence) and Gamma-glutamyltransferase (GGT), a liver enzyme which can be elevated for 8 weeks after alcohol exposure. In a separate cohort, data from the same antenatal booking visit was collected from medical records documenting women's self-reported alcohol consumption.

## RESULTS

\* The percentage of women who reported alcohol intake in the first trimester was 0.8%. This compared to 74.1% of women who reported consuming alcohol before pregnancy.

\* CDT analysis revealed a prevalence rate of 1.4% and GGT a prevalence rate of 3.5% in the first trimester of pregnancy.

\* Although those with elevated CDT generally had high levels of GGT, only one person was positive for CDT and GGT.

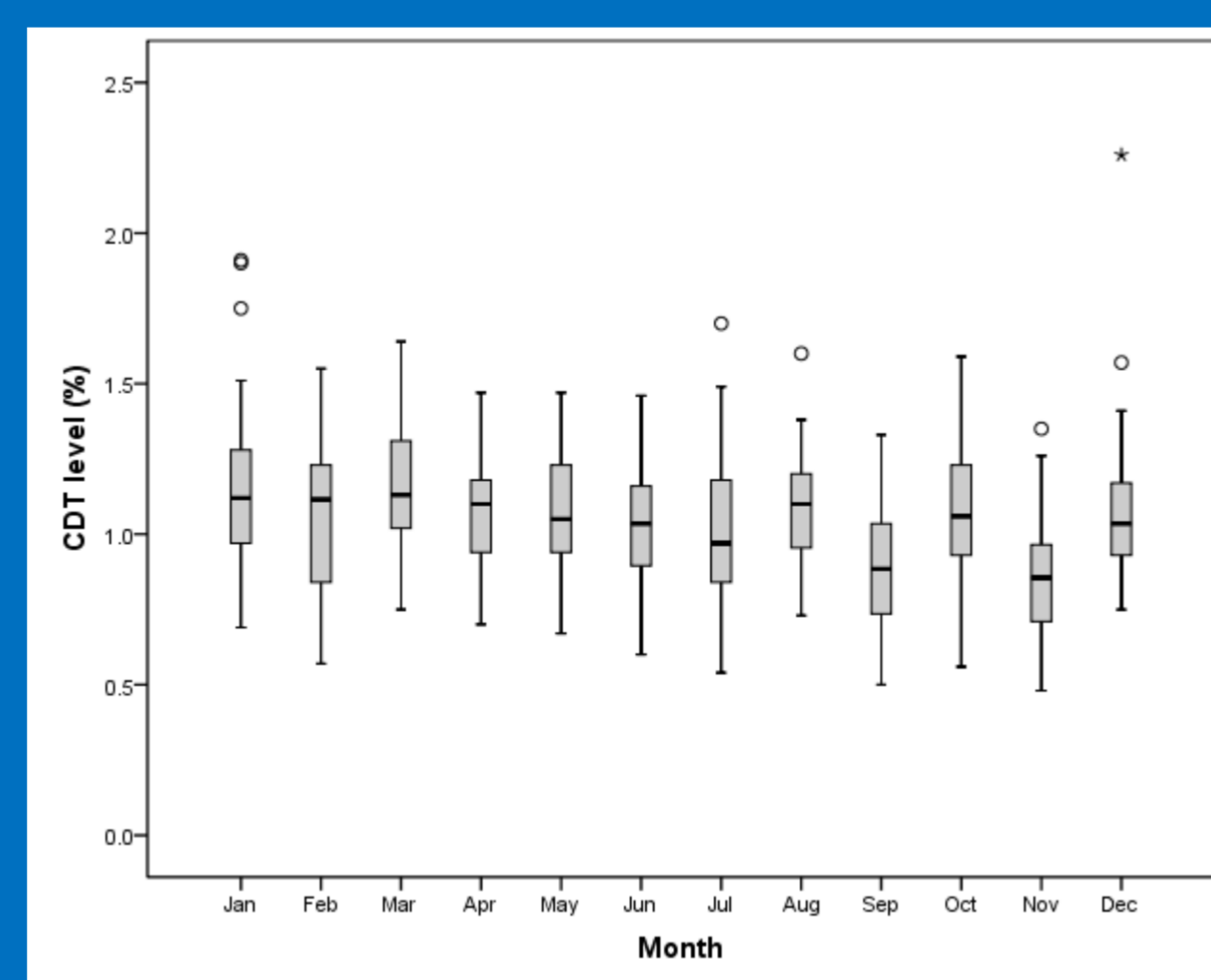


Figure 1: Box plots of CDT values per month

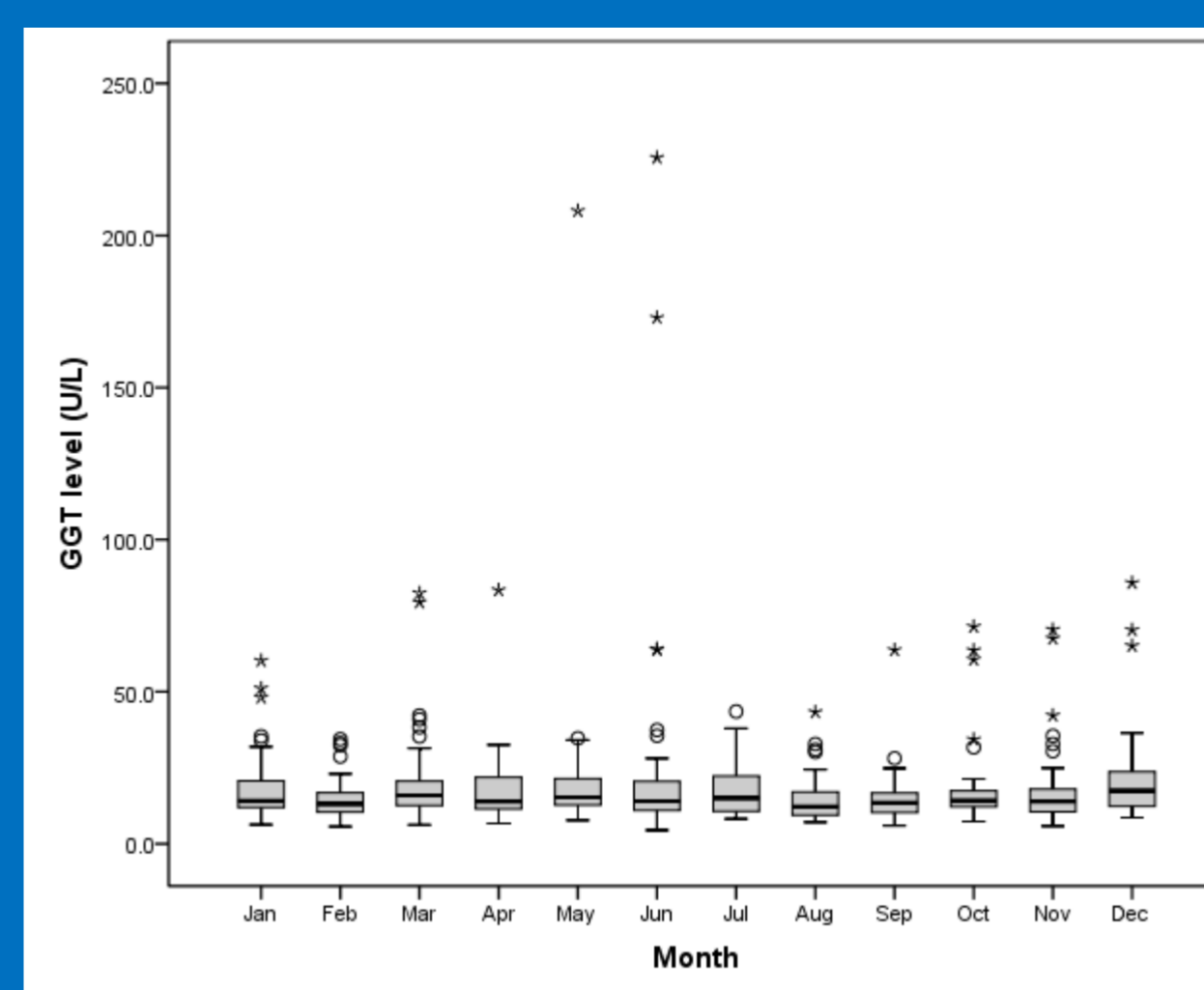


Figure 2: Box plots of GGT values per month

o = outlier, \* = extreme outlier

## Conflicts of Interest:

There were no conflicts of interest.

## CONCLUSION

Results from CDT analysis and self-report were similar, but both may under-report. Self-report may be limited by factors such as recall bias, the patient-clinician relationship, expected social norms and fear of perceived judgement<sup>2</sup>. CDT only detects sustained high level drinkers and does not identify low to moderate drinkers<sup>3</sup>. GGT appeared to lack specificity, but it may have value in supporting findings from CDT analysis. Further studies using additional blood biomarkers, or a combination of blood biomarkers and self-report, may be beneficial in detecting a more detailed drinking history in pregnancy.

## REFERENCES

1. Bakhireva LN, Savage DD. Focus on: biomarkers of fetal alcohol exposure and fetal alcohol effects. *Alcohol Res Health*. 2011;34(1):56-63.
2. Lange S, Shield K, Koren G, Rehm J, Popova S. A comparison of the prevalence of prenatal alcohol exposure obtained via maternal self-reports versus meconium testing: a systematic literature review and meta-analysis. *BMC Pregnancy Childbirth*. 2014;14:127.
3. Shipton D, Tappin D, Sherwood R, Mactier H, Aitken D, Crossley J. Monitoring population levels of alcohol consumption in pregnant women: a case for using biomarkers. *Subst Use Misuse*. 2013 Jun;48(8):569-73.

Other references available on request