

# The Association of British Clinical Diabetologists (ABCD) nationwide exenatide and liraglutide audits

R.E.J. Ryder<sup>1</sup> • K.Y. Thong<sup>1</sup> • M.L. Cull<sup>1</sup> • A.P. Mills<sup>1</sup> • C. Walton<sup>2</sup> • ABCD nationwide exenatide and liraglutide audit contributors;  
<sup>1</sup>Diabetes, City Hospital, Birmingham, UK • <sup>2</sup>Diabetes, Hull Royal Infirmary, Hull, UK

Association of British Clinical Diabetologists Autumn Meeting, 10–11 November 2011, London

## Aims

- To compare use and efficacy of exenatide and liraglutide in two large scale nationwide audits of real clinical practice.

## The ABCD nationwide exenatide and liraglutide audits

	Exenatide audit	Liraglutide audit (ongoing)
Dates of data	2007–2009	2009–2010
Centres	126	64
Contributors	315	210
Patients	6717	3010

- Collected anonymised data of patients treated with exenatide or liraglutide in the UK
  - Patient demographics
  - Diabetes medications
  - HbA<sub>1c</sub>, weight
  - Lipids
  - Blood pressure
  - Adverse events and GLP-1 discontinuation.

## Methods

- Comparisons of baseline characteristics, diabetes medication use and HbA<sub>1c</sub> and weight changes using Student's t-test, Mann-Whitney U test and Chi Square test.
- Excluded liraglutide patients previously on exenatide.
- HbA<sub>1c</sub> and weight results taken at 6 months (± 6 weeks).
- Blood pressure and lipid changes also analysed.

## Results

### Baseline characteristics

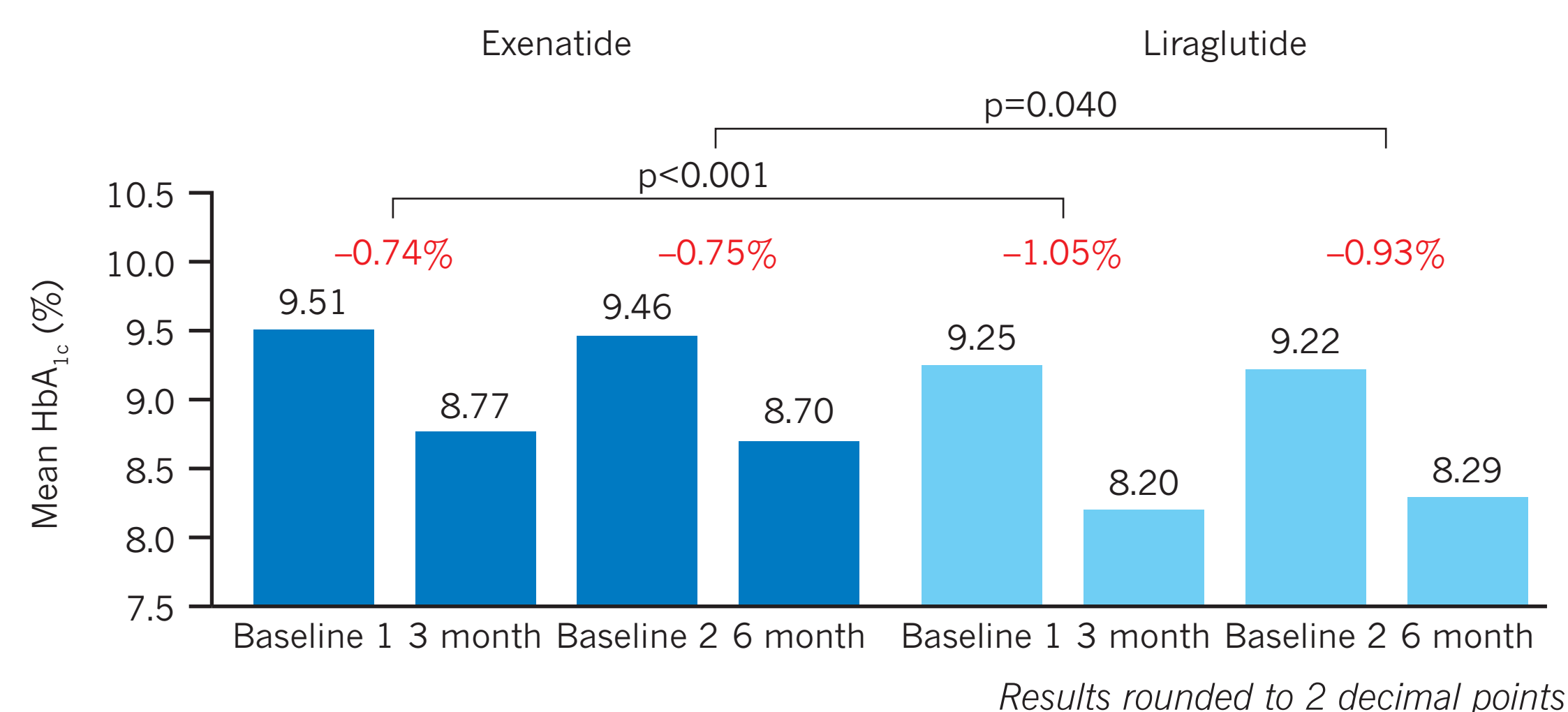
	Exenatide audit (n=6717)	Liraglutide audit (n=2303)	p value
Male (%)	54.9	54.1	0.491
Caucasian (%)	84.4	90.4	<0.001
Age (years)	54.9 (10.6)	55.4 (11.2)	0.033
Diabetes duration (years)	8 (5–13)	9 (5–13)	0.424
HbA <sub>1c</sub> (%)	9.47 (1.69)	9.32 (1.72)	0.001
Weight (kg)	113.8 (23.4)	111.1 (23.0)	<0.001
BMI (kg/m <sup>2</sup> )	39.8 (8.0)	39.1 (7.5)	<0.001
Single oral therapy (%)	12.7	12.0	0.371
Dual oral therapy (%)	28.1	28.1	0.969
≥3 oral therapy (%)	15.6	17.9	0.012
On insulin (%)	33.9	39.8	<0.001

Results with mean (SD) and median diabetes duration (inter-quartile range)

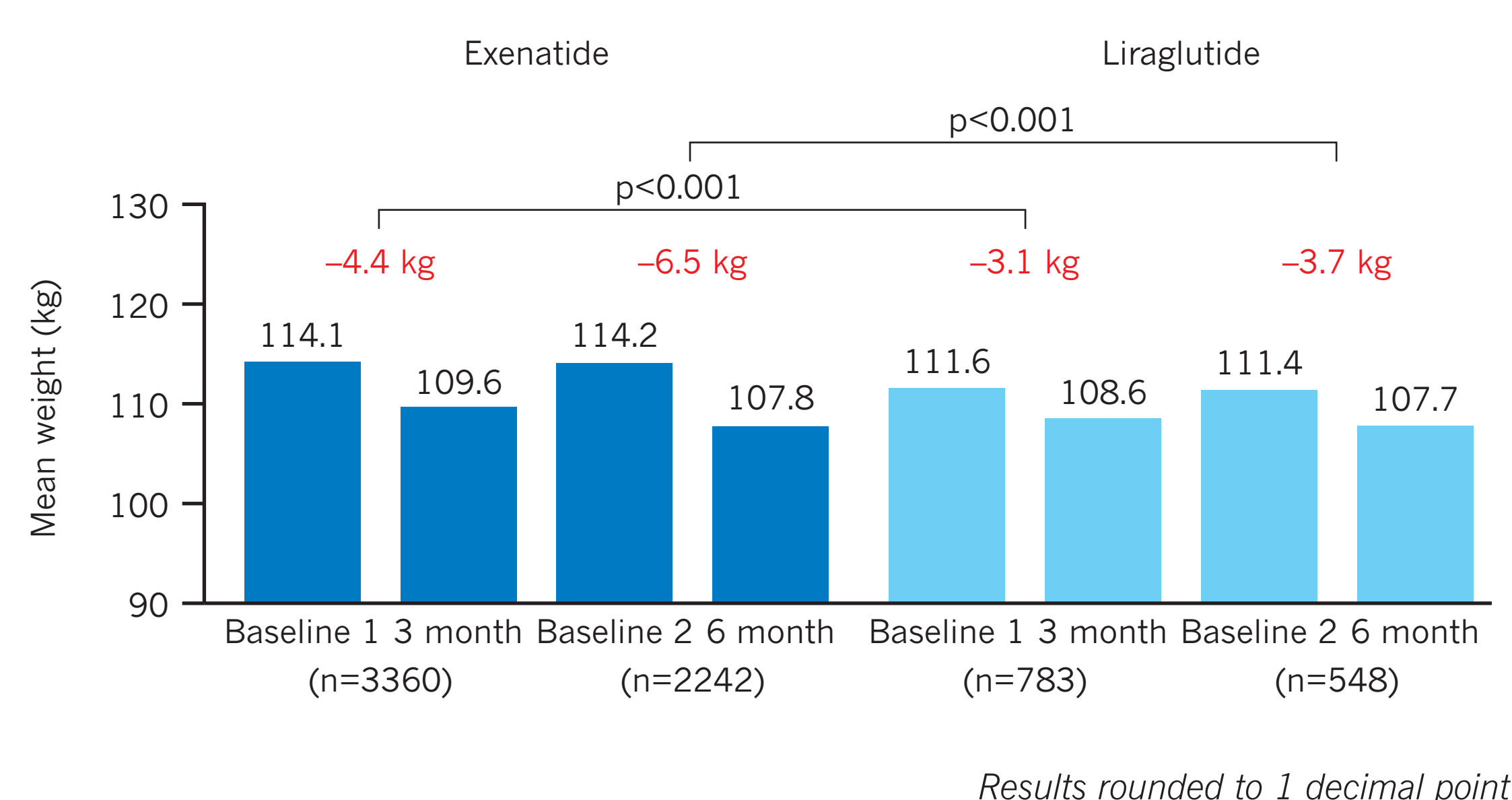
### Baseline diabetes treatment use and discontinuation

	Exenatide audit	Liraglutide audit
Metformin (%)	84.0 (0.9)	82.7 (0.7)
Sulphonylurea (%)	49.5 (6.5)	42.8 (5.3)
Thiazolidinedione (%)	27.1 (13.4)	20.5 (7.5)
Meglitinide (%)	2.0 (0.6)	1.0 (0.2)
Acarbose (%)	0.9 (0.3)	0.7 (0.3)
DPP-IV-inhibitor (%)	2.2 (1.4)	10.9 (9.3)
Exenatide (%)	–	21.9 (21.9)
Insulin (%)	33.9 (8.1)	39.6 (2.6)

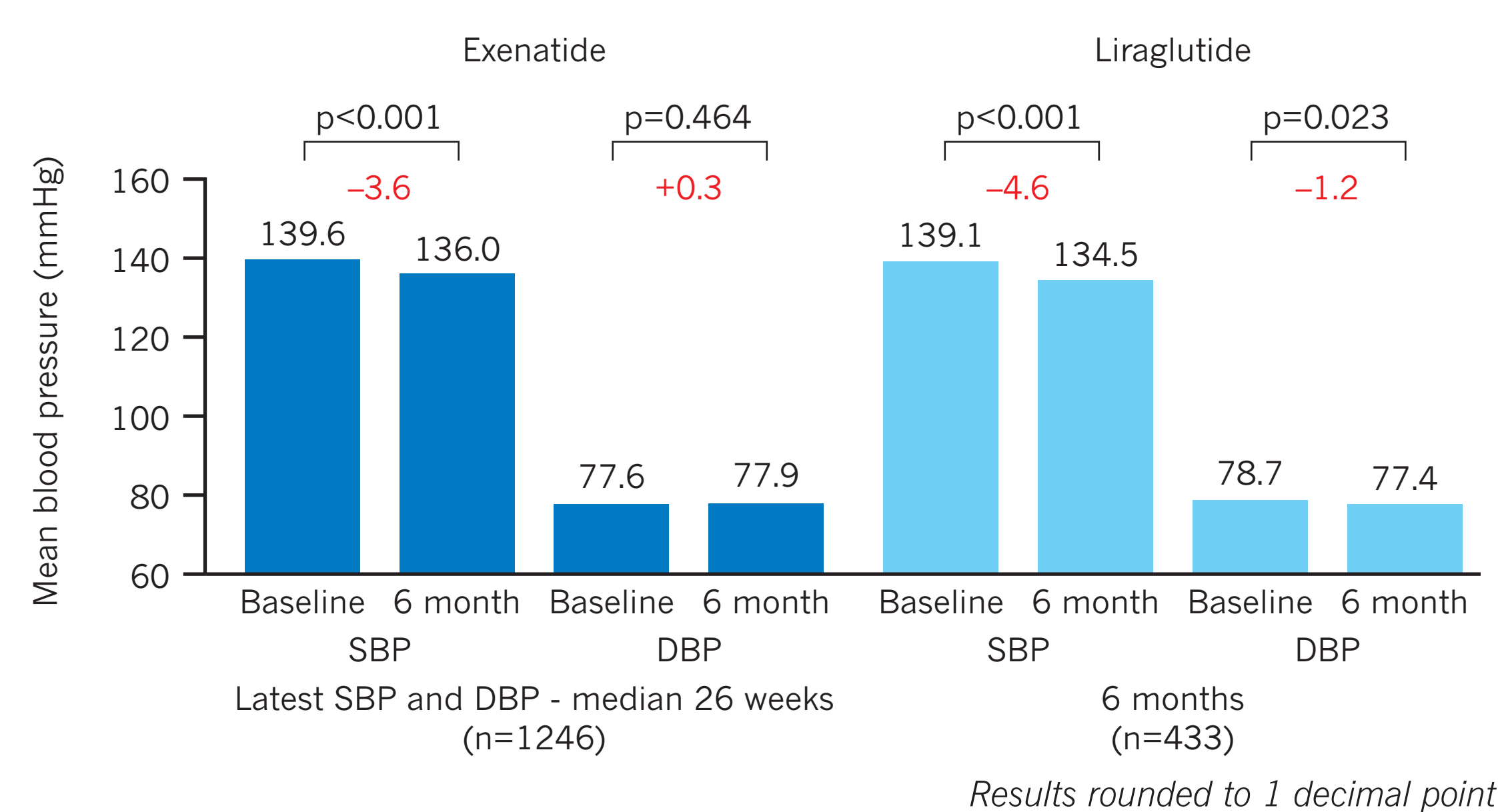
### HbA<sub>1c</sub> changes at 3 and 6 months: exenatide and liraglutide



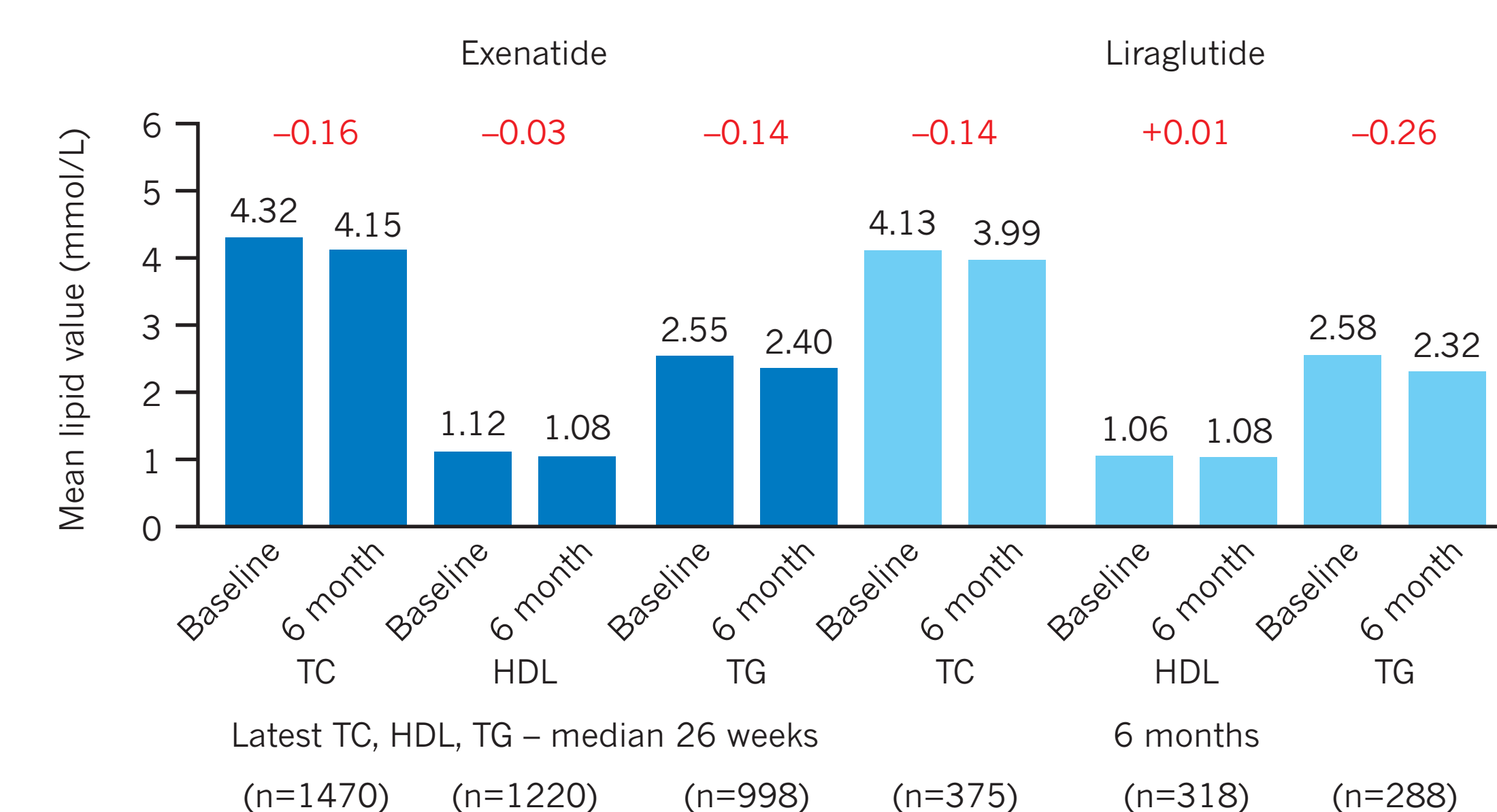
### Weight changes at 3 and 6 months: exenatide and liraglutide



### Blood pressure: exenatide and liraglutide



### Lipids: exenatide and liraglutide



Results rounded to 2 decimal points

## Conclusions

- These very large audits reveal the effectiveness of these agents in much heavier and more poorly controlled patients than those studied in clinical trials.
- Patients achieved greater HbA<sub>1c</sub> reduction but lesser weight reduction in the liraglutide audit as compared with the exenatide audit.
- However, there were lesser insulin and thiazolidinedione discontinuation but greater DPP-IV (dipeptidyl peptidase-IV) inhibitor discontinuation in the liraglutide audit. Contributors might have learnt from the previous use of exenatide to avoid over-reduction of diabetes treatment when initiating liraglutide.

## Acknowledgment

- We thank all the nationwide contributors for submitting data on patients on exenatide and liraglutide.
- The ABCD nationwide audit programme has received grants provided by Eli Lilly and Novo Nordisk. This audit was independently initiated and performed by ABCD and the authors remained independent in the analysis and the writing of this report.