

Effect of dapagliflozin on HbA1c and weight after its addition to various combinations of other diabetes medications: Association of British Clinical Diabetologist (ABCD) nationwide dapagliflozin audit

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BACKGROUND

AIMS

METHODS

The ABCD audits new pharmacotherapies for diabetes across the UK to collect real-world data on their usage, accelerate the understanding of new agents in patients in the UK and ascertain whether experience from clinical usage matches phase 3 trial data.

To evaluate the effect on HbA1c and weight after adding dapagli-

flozin to various combinations of other diabetes medications, in

The ABCD nationwide audit of dapagliflozin in real clinical use in the UK, was launched in October 2014. Anonymised data of patients treated with dapagliflozin in the UK was collected by an

ABCD members, clinicians in both primary care and secondary

care were emailed to invite them to submit clinical data on their

Those with baseline and follow-up HbA1c within a median (range)

of 6.2(4.1-9.4) months, after commencing dapagliflozin were

included. Data at baseline and first follow-up were compared

Oct 2014 - March 2016

59

156

1753

patients with type 2 diabetes in UK

online password protected questionnaire:

HbA1c, weight, BMI, Systolic BP

Patient demographics

Diabetes medications

patients treated with dapagliflozin

Data Input

Centres

Contributors

Number of patients

using student's paired t-test.

Adverse events

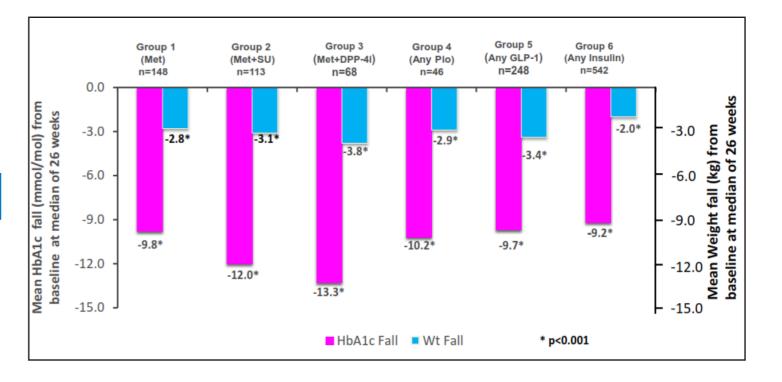
Patients were categorized into 6 groups according to their other diabetes therapies, dapagliflozin was added to : group 1 (metformin, n=148), group 2(metformin and sulphonylurea, n=113), group 3 (metformin and dipeptidyl peptidase-4 inhibitor(DPP4-i), n=68), group 4(pioglitazone with or without any other diabetes medications, n=46), group 5 (glucagon-like peptide-1 receptor agonist with or without any other diabetes medications, n=248) and group 6 (insulin with or without any other diabetes medications, n=542).

BASELINE CHARACTERISTICS

Groups	Gr 1	Gr 2	Gr 3	Gr 4	Gr 5	Gr 6
Age (years)	54.5±10.3	58.5±10.1	55.1±11.0	54.9±10.4	56.2±10.2	57.0±10.1
Sex (Males%)	48.6	57.5	45.5	65.2	53.6	52.7
T2DM Dur(years)	5.9±6.3	7.6±5.3	7.1±5.2	8.3±6.6	10.9±7.6	12.3±8.2
Wt(Kg)	101.2±26.1	99.6±24.2	94.8±18.6	111.5±27.2	108.8±23.3	104.7±20.9
BMI (Kg/m²)	35.0±7.7	34.8±7.0	34.2±7.7	37.8±7.9	38.1±7.9	36.6±6.3
HbA1c (mmol/mol)	76.7±16.3	80.8±14.8	79.6±16.2	82.0±14.7	81.5±15.2	81.8±16.2
HbA1c(%)	9.1±1.5	9.5±1.3	9.4±1.4	9.6±1.3	9.6±1.4	9.6±1.5

RESULTS

Mean(\pm SE) HbA1c fell by(-9.8(\pm 1.2)mmol/mol from 76.7(\pm 1.3) to 66.8(±1.4) mmol/mol, p<0.001) in group 1, (-12.0(±1.2) mmol/ mol from 80.8(±1.4) to 68.8(±1.3) mmol/mol, p<0.001) in group 2, (-13.3(±1.9) mmol/mol from 79.7(±1.9) to 66.3(±2.1) mmol/mol, p<0.001) in group 3, (-10.2(±2.3) mmol/mol from 82.0(±2.1) to 71.7(±2.8) mmol/mol, p<0.001) in group 4, (-9.7(±1.0) mmol/mol from 81.5(±0.9) to 71.8(±1.07) mmol/mol, p<0.001) in group 5 and (-9.2(±0.6) mmol/mol from 81.8(±0.7) to 72.6(±0.6) mmol/mol, p<0.001) in group 6.



Weight fell by $(-2.8(\pm 0.5)$ kg from $101.2(\pm 2.1)$ kg to $98.3(\pm 2.0)$ kg, p<0.001) in group 1, (-3.1(±0.4)kg from 99.6(±2.2)kg to 96.5(±2.2) kg, p<0.001 in group $2(-3.8\pm(\pm0.6)$ kg to $98.3(\pm2.2)$ kg to $90.9(\pm2.1)$ kg, p<0.001 in group 3, $(-2.9(\pm 0.6) \text{ kg from } 111.5(\pm 3.9) \text{ kg, p} < 0.001)$ in group 4, $(-3.4(\pm 1.1)$ kg from 108.8(± 1.4) kg to 105.5(± 1.4) kg, p<0.001) in group 5, (-2.0(±0.2) kg from 104.7(±0.9)kg to 102.6(±0.9) kg,p<0.001) in group 6.

The different treatment groups all had similar poor glycaemic

control (HbA1c>9%) but they varied with regard to duration of

diabetes (longest duration were insulin and GLP-1 treated patient

groups), BMI (heaviest GLP-1 and pioglitazone treated patients),

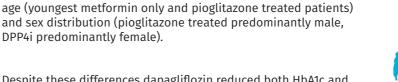
CONCLUSION

DPP4i predominantly female).

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Despite these differences dapagliflozin reduced both HbA1c and weight by clinically and statistically significant amounts in all the groups in this wide range of real world UK patients with type 2



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NHS Trust

diabetes on a variety of diabetes medications. Whilst the biggest impact was on those on DPP4i group and least for those on insulin, these differences were not statistically significant.

ACKNOWLEDGEMENT

