BSF2020 VIRTUAL 9 -14 NOVEMBER

Steroid Induced Hyperglycemia and Diabetes on the Haematology Ward: Are we getting it right? Re-audit following Initiation of JBDS-IP Guidelines and Teaching K Lazarus, AA Raheem, S Ross, E Nogueira, E Hui, G Zakout Northwick Park Hospital, London, UK

INTRODUCTION

- High doses of pulsed steroids are commonly integrated chemotherapeutic regimens used for treating haematologica malignancies. This can worsen glucose control in people with pre-existing diabetes (steroid induced hyperglycaemia (SIH)) o induce diabetes (steroid induced diabetes (SID)).
- The prevalence of SIH/SID in the outpatient setting has been

	Pre-Guidelines 2018 (n = 18)	Post-Guidelines 2019 (n = 18)
lean age (years)	63	71
Age range (years)	30 - 88	31 - 91
lale, female	10 male, 8 female	13 male, 5 female
laematological diagnosis	10 Lymphoma 4 Multiple Myeloma 2 ALL 1 AML 1 CLL	6 Lymphoma (5 NHL, 1 HL) 6 Myeloma 1 ALL 3 CLL 1 Myelofibrosis 1 Waldenstrom's macroglobulinemia
Average dose hydrocortisone equivalent dose/24 hours) and Average Duration	155 mg (100-625mg) 14 Days	462 mg (100-1332mg) 8.6 Days
Steroids Used (Number of Patients)	Dexamethasone (12) Prednisolone (2) Methylprednisolone (1) Prednisolone + Methypred (1) Dexamethasone + Pred (1) Dexamethasone + Hydrocortisone (1) Methylpred + Dexamethasone (1)	Dexamethasone (13) Prednisolone (5)
Percentage of patients with appropriate capillary blood glucose monitoring (as per JBDS guidelines whilst on steroids)	28%	94%

1960-2020

RESULTS

reported to be as high as 40%^{1.} The prevalence however, c SIH/SID in haematology inpatients and the effects of different steroid regimens used in various haematological malignancies on glycaemic control in inpatients are not largely known.

The Joint British Diabetes Societies for Inpatient care (JBDS-IP guidelines recommend measuring HbA1c to screen patients prio to starting steroids; however this can be misleading in patients with a paraprotein, splenomegaly, and recent blood transfusions - all frequently encountered findings in haemato-oncologi patients². Patients without pre-existing diabetes on steroid therapy should have once daily monitoring of capillary blood glucose prior lunch or evening meal, and four times daily in those with diabetes.

We evaluated the management of SIH/SID and implemented protocol for detection and management of SIH/SID on haemato-oncology ward.

Risk Factors for SID/SIH³

High steroid doses

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Prolonged steroid treatment

Obesity (BMI>30)/Polycystic ovarian syndrome

Increasing Age

Personal History of previous SIH/SID and previous history of gestational diabetes

Family History of Diabetes

METHODS

- Data from a single-centre haemato-oncology inpatient service receiving therapy as per the NICE guidelines for their respective hemato-oncologic condition was collected using a standard proforma over a 2 month period in 2018 (Group 1) and reaudited in 2019 (Group 2) after implementing 2 single-paged guidelines on SID (for patients without known diabetes) and SIH (patients with known diabetes).
- Teaching sessions were delivered and an algorithm for screening and management of SIH/SID based on JBDS-IP guidelines was introduced post Group 1 analysis.

Type 2 diabetes and steroid treatment – General Guidance⁴

- Monitoring
- Set target for Capillary Blood Glucose (CBG) e.g. 6-10mmol/L
- Consider increasing monitoring to 4 times daily Refresh diabetes education with patient If hyperglycaemia on Gliclazide – titrate to maximum of 320mg daily, with maximum 240mg in non-insulin therapies the morning Metformin – titrate to maximum of 1g BD • If on evening once daily human insulin If hyperglycaemia on insulin therapies consider switch to morning dosing • If uncontrolled hyperglycaemia or multiple daily dosing of steroid consider switch to basal analogue insulin (or alternative regimen) and involve diabetes team in hospital or community Beware of nocturnal and early morning hypoglycaemia

Pre-Guidelines 2018	Post-Guidelines 2019
28%	94%
2 018 Total = 18	2019 Total = 18
' known Diabetes 1 no Diabetes	6 known Diabetes 12 no Diabetes
 /11 – No monitoring /11 Fasting CBGs only ?Undiagnosed SID) /11 – Infrequent/delayed nonitoring ?/11 – Developed SID /7 Developed SIH 	9/12 Developed SID 5/6 Developed SIH



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- With improved CBG monitoring, 78% of patients were found to have SIH/SID in Group 2, compared to 44% in Group 1.
- 9 out of 12 patients in Group 2 without known diabetes developed SID, 5 out of 6 patients with known diabetes developed SIH.
- This could be due to increased screening and a higher average steroid dose in Group 2 compared with Group 1.
- Escalation of diabetes treatment plan including gliclazide or insulin initiation based on the protocol was also proactively and promptly started in 6 patients in Group 2 before diabetes team review, compared to none in Group 1.

CONCLUSIONS

- Awareness of SIH/SID was low among Haematology in the inpatient setting, in particular SIH.
- Our intervention led to significant improvement in screening, detection and prompt treatment of SIH/SID.
- We achieved the recommended national standard of monitoring for SIH/SIH, in 94% being appropriately screened (JBDS-IP audit standard 90%).
- Further work is needed to improve post discharge care planning. Similar practice review is currently being planned in the outpatient setting.

REFERENCES

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Poster

presented at:

British Society for

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