| - | | ORI | GINAL RESEARCH PAPER | Medicine |
|----------|--|--|---|--|
| Indian | | DIA | ORS AFFECTING GLYCEMIC CONTROL IN TYPE BETES PATIENTS IN RURAL POPULATION OF HYA REGION | KEY WORDS: Type 2 Diabetes Mellitus, Glycemic Control |
| D | r P K Baghel | | Professor, Department of Medicine, SSMC & SGMH, | Rewa, MP |
| | r Umesh Prata _l ngh* | р | Senior Resident, Department of Medicine, S *Corresponding Author | SSMC & SGMH, Rewa, MP |
| D | r Anuraag Gup | ota | Senior Resident, Department of Medicine, SSMC & S | GMH, Rewa, MP |
| ΔΒςΤΡΑCT | common cause of his prevention of organ magnitude of good Medical College and Material & Methoo 2016 to July 2017. If control was defined Results: Out of 500 Patients in the poor diabetes, illiteracy, o Conclusion: Prevale these and other chi | ospita and ISGM ISGM Detail as the Detail as the Detail as | mic control amongst patients with type 2 diabetes mellitus indica al admission and complications caused by diabetes. Good glycem nage and other complications arising from diabetes. The obje poor glycemic control and factors affecting glycemic control in ty 1H Rewa, Madhya Pradesh, India. is is hospital based cross sectional study conducted on 500 type 2 ed history, thorough clinical examination and anthropometric me three-month average fasting blood sugar (FBS) 80-130 mg/dl. ents 156 (31.2%) have good glycemic control and 344 (68.8%) emic control group have greater duration of diabetes, lack of adh y and dyslipidemia. (p value<0.05) of poor glycemic control is very high in this population. Further stude eristics on glycemic control among rural population of Vindhy diabetes outcomes and increase life-expectancy. | hic control is the main objective for the ctive of this study was to assess the ype 2 diabetic patients at Shyam Shah 2 diabetes mellitus patients from April easurement was done. Good glycemic b) patients have poor glycemic control. herence to treatment, family history of dies are needed to explore the effect of |

Introduction:

Diabetes mellitus (DM) refers to a group of common metabolic disorders that share the phenotype of hyperglycaemia with disturbance of carbohydrate, fat and protein metabolism resulting from defects in insulin secretion, insulin action or both. Several distinct types of DM are caused by a complex interaction of genetics and environmental factors. Depending on the aetiology of the DM, factors contributing to hyperglycaemia include reduced insulin secretion, decreased glucose utilization and increased glucose production.¹

Diabetes mellitus is accepted as a worldwide epidemic with an estimated increase in prevalence from 2.8% in 2000 to 4.4% by 2030.² The Indian Council of Medical Research India Diabetes Study (ICMR-INDIAB study) showed that India had 62.4 million people with diabetes in 2011. These numbers are projected to increase to 101.2 million by 2030.³ There is no cure for this disease and it requires continuing medical care and achieving good glycemic control to prevent acute complications and to reduce the risk of long-term complications because poor glycemic control is the most common cause of hospital admissions and complications in diabetic patients.⁴ However, large no of the diabetic patients did not maintain their blood glucose at optimum level, that's why the factors that affect the glycemic control significantly, should identify individually.

Materials and Methods:

The present cross-sectional study was carried out in patients from rural area attending MOPD and those admitted in Department of Medicine, S.S. Medical College and associated S.G.M. Hospital, Rewa (M.P.) from April 2016 to July 2017. A total of 500 Type 2 diabetes patients from rural population of Vindhya region were included in the study.

Inclusion Criteria

- 1. Type 2 diabetic patients living in rural area.
- 2. Age > 30 years.
- 3. Giving consent for study.

Exclusion Criteria

- 1. Type 1 diabetic patients.
- 2. Diabetic, but belong to urban area.
- 3. Not giving consent.

Diabetes was defined by ADA 2011 criteria- Plasma fasting blood glucose \geq 126 mg/dL or 2-hour plasma post-glucose value \geq 200

mg/dL or patients with classical symptoms of hyperglycaemia or hyperglycaemic crisis plus random blood glucose concentration \geq 200 mg/dL (\geq 11.1 mmol/L) or bA1c \geq 6.5%. Hypertension was diagnosed according to JNC-7 criteria, those with systolic blood pressure \geq 140 mmHg and diastolic blood pressure \geq 90 mmHg or who were taking anti-hypertensive medication were considered to 6 have hypertension. Dyslipidaemia was defined if patient had total cholesterol \geq 200 mg/dL in males, < 50 mg/dL in female and serum LDL \geq 100 mg/dL.

Glycemic control:

It is managing blood glucose level of diabetic patients at optimum level. Good glycemic control is defined as the three-month average fasting blood glucose 80-130 mg/dl and poor glycemic control is defined as the three-month average fasting blood glucose >130 mg/dl.^{\circ}

The following data were obtained directly from the patients: age, gender, marital status, educational stages, adherence to treatment, family history, time from onset of diabetes (considered the approximate date of diagnosis), type of treatment, addiction and history of high blood pressure. They underwent a thorough physical examination which included weight, height, waist circumference and BMI which were calculated. Data were completed by consulting medical reports of patients.

Data was at first arranged in Microsoft Excel 2016 Worksheet, developed by Microsoft, Redmond, Washington. Data is expressed as mean \pm standard deviation for continuously distributed variables and in absolute numbers and percentages for the discrete variables.

Results:

Out of 500 patients, 156 (31.2%) have good glycemic control and 344 (68.8%) patients have poor glycemic control.

Table 1: Socio-Demographic and Clinical Characteristic

| Variable | Frequency (n=500) | Good glycemic control (FBG 80- 130mg/dl) (n=156) | Poor glycemic control (FBG>130 mg/dl) (n=344) | P value | |
|----------|----------------------|---|--|---------|--|
| Sex | | | | | |
| Male | 277 | 80(28.88%) | 197(71.11%) | 0.2122 | |
| Female | 223 | 76(34.08%) | 147(65.91%) | | |
| | | | | | |

www.worldwidejournals.com

166

PARIPEX - INDIAN JOURNAL OF RESEARCH

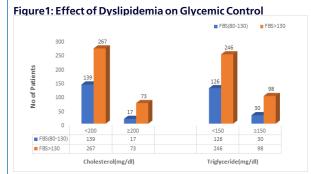
| Age grou | o (years) | | | | |
|------------|--------------|-----------------|-------------|---------|--|
| 31-40 | 30 | 13(43.33%) | 17(56.66%) | 0.3039 | |
| 41-50 | 150 | 50(33.33%) | 100(66.66%) | 1 | |
| 51-60 | 153 | 44(28.75%) | 109(71.24%) | 1 | |
| 61-70 | 90 | 26(28.88%) | 64(4.9%) | 1 | |
| >70 | 73 | 23(31.50%) | 50(68.49%) | 1 | |
| Education | al status | | | - | |
| Illiterate | 280 | 71(25.35%) | 209(74.64%) | 0.0015 | |
| Literate | 220 | 85(38.63%) | 135(61.36%) | 1 | |
| Family His | story of Dia | abetes | | | |
| Present | 187 | 48(25.66%) | 139(74.33%) | 0.0391 | |
| Absent | 313 | 108(34.50%) | 205(65.49%) | 1 | |
| Duration | of Diabete | s (years) | | | |
| <5 | 87 | 33(37.93%) | 54(62.06%) | 0.0253 | |
| 5-10 | 309 | 99(32.03%) | 210(67.96%) | 1 | |
| >10 | 104 | 24(23.07%) | 80(76.92%0 | 1 | |
| Type of Ti | reatment | | | | |
| OHA | 334 | 113(33.83%) | 221(66.16%) | <0.0001 | |
| Insulin | 82 | 32(39.02%) | 50(60.97%) | | |
| OHA+ | 15 | 6(40%) | 9(60%) | | |
| Insulin | | | | | |
| No | 69 | 5(7.2%) | 64(92.75%) | 1 | |
| Treatment | | | | | |
| | n Adheren | | | | |
| Yes | 335 | 133(39.70%) | 202(60.29%) | <0.000 | |
| No | 165 | 23(13.93%) | 142(86.66%) | 1 | |
| Addiction | (Tobacco d | chewing/smoking | | | |
| Yes | 221 | 63(28.50%) | 158(71.49%) | 0.2473 | |
| No | 279 | 93(33.33%) | 186(66.66%) | | |
| Hypertens | sion | | | | |
| Present | 83 | 20(24.09%) | 63(75.90%) | 0.1261 | |
| Absent | 417 | 136(32.61%) | 281(67.38%) | 1 | |
| Body Mas | sIndex (kg | /m2) | | | |
| <18.5 | 11 | 5(45.45%) | 69(54.54%) | 0.0202 | |
| 18.5-24.9 | 145 | 53(36.55%) | 92(63.44%) | 1 | |
| 25-29.9 | 318 | 96(30.18%) | 222(69.81%) | 1 | |
| 25 25.5 | 510 | (,-,-, | (| | |

In our study female patients have more good glycemic control than male patients (34.08% vs 28.88%). Patients age not significantly associated with glycemic control and literate patients have more good glycemic control as compared to illiterate patients (38.63% vs 25.35%).

In our study as duration of diabetes increases glycemic control decreases. Patients on OHA + Insulin have more good glycemic control as compared to OHA/Insulin alone. Patients who were on regular treatment have better glycemic control than irregular treatment (p<0.05).

Non-hypertensive patients and patients with no addiction have better glycemic control but association is not significant (p>0.05). There is a significant association (p<0.05) between lower Body Mass Index and glycemic control. As BMI increases good glycemic control decreases.

In our study there is substantially greater risk of poor glycemic control amongst patients with family history of diabetes mellitus (p<0.05).



Volume-8 | Issue-3 | March-2019 | PRINT ISSN No - 2250-1991

In our study patients with dyslipidemia have poor glycemic control as compared to patients with normal lipid profile (p value<0.05).

Table 2- Distribution of patients according to Diabetic

| Complication | Good glycemic control (FBG 80- 130mg/dl) | | P value |
|--------------------------------|--|----|---------|
| Diabetic Retinopathy (n=86) | 15 | 71 | 0.0025 |
| Diabetic Nephropathy(n=117) | 24 | 93 | 0.0044 |

In our study there is a significant association between diabetic complication and glycemic control (p value<0.05).

Discussion:

This study falls in line with the existing research that suggests that poor glycemic control among diabetic patients is largely prevalent in Indian scenario. In our study the proportion of diabetic patients having good glycemic control (31.2%) is similar to another study in which good glycemic control 30% of the patients⁹

However, there was no significant association between gender, age group, addiction, hypertension and glycemic control.

Meanwhile, in our study there was a significant association between family history of diabetes mellitus and poor glycemic control. Similar results were observed in study conducted by **Lee YH et al.**¹⁰

In our study patients with poor glycemic control was found to increase with increase in disease duration. Longer duration of diabetes is known to be associated with poor control, possibly because of progressive impairment of insulin secretion with time because of cell failure, which makes the response to diet alone or oral agents unlikely (UK Prospective Diabetes Study (UKPDS) Group,1998).¹¹

In our study higher BMI was significantly associated with poor glycemic control (p value=0.0202). Similarly, a higher proportion of patients with poor glycemic control was observed in patients whom were obese, followed by overweight.¹²⁻¹³

In Our study literate patients have good glycemic control as compared to illiterate patients (p=0.0015) which is similar to **Goudswaard et al** in which lower level of education was associated with poor glycemic control.¹⁴

In our study patients who were on regular treatment have good glycemic control as compared to patients who were on irregular treatment (p<0.0001). Similar result was observed in study conducted by **Fshea B et al.**¹⁵

In our study there is a significant association between lipid profile and glycemic control (p<0.05). Patients with hypertriglyceridemia and hypercholesterolemia have poor glycemic control as compared to patients with normal triglyceride and cholesterol level. Similar results were observed in study conducted by **Mullugeta Y** et al¹⁶.

In our study Patients on OHA + Insulin have more good glycemic control as compared to 0HA/Insulin alone (p<0.0001). This finding is inconsistent with study conducted by **Haghighatpanah M et al**¹⁷ in which patients receiving insulin + OHA or insulin as monotherapy were more likely to have poor glycemic control compared to patients who were on oral diabetes medication.

In our study patients with diabetic retinopathy and diabetic nephropathy were significantly associated with poor glycemic control as compared to patients with no complication (p<0.05).

Conclusion:

In our study the proportion of good glycemic control in rural type 2

www.worldwidejournals.com

PARIPEX - INDIAN JOURNAL OF RESEARCH

diabetic patients is very low. The present study showed that there was a significant association between glycemic control and educational status, BMI, family history, duration of diabetes, medication history, adherence to treatment, triglyceride level and cholesterol level. Based on these factors, patients at risk of poor glycemic control can be identified, and targeted interventions can be implemented for optimal outcomes.

REFERENCES:

- Classification of diabetes mellitus and other categories of glucose intolerance. In: 1. Alberti K, Zimmet P, De Fronzo R, (eds). International Textbook of Diabetes Mellitus. Chichester: John Willey and Sons Ltd, 1997:9-23. Wild S, Roglic G, Green A, et al. Global prevalence of diabetes mellitus: estimates
- 2. for the year 2000 and projections for 2030. Diabetes Care. 2004;27(5):1047-53
- Anjana RM, Pradeepa R, Deepa M, et al. Prevalence of diabetes and prediabetes (impaired fasting glucose and/or impaired glucose tolerance) in urban and rural 3 India: phase I results of the Indian council of medical research India diabetes (ICMR-INDIAB) study. Diabetologia. 2011;54(12):3022-7. IDF (2012) Diabetes atlas, 5th ed. via http://www.idf.org/diabetesatlas/5e/the-
- 4 globalburden: Accessed 2012 November 20
- American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes Care 2011;34(Suppl 1): S62-9. 5
- 6. Chobanian AV, Bakris GL, Black HR, et al. The seventh report of the joint national committee on prevention, detection, evaluation and treatment of high blood pressure: the JNC 7 report. JAMA. 2003;289(19):2560-72
- Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive summary of the third report of the national cholesterol education program (NCEP) expert panel on detection, evaluation, and treatment of high blood cholesterol in adults (adult treatment panel III). JAMA. 7 2001;285(19):2486-97.
- American Diabetes Association.6. Glycemic targets: Standards of Medical Care in 8 Diabetesd2018. Diabetes Care 2018; 41(Suppl. 1): S55-S64
- Unnikrishnan R, Anjana RM, Deepa M, Pradeepa R, Joshi SR, BhansaliA, et al; ICMR–INDIAB Collaborative Study Group. Glycemic control among individuals with 9 self-reported diabetes in India--the ICMR-INDIAB Study. Diabetes TechnolTher. 2014 Sep: 16(9): 596-603.
- Young-Hoon Lee, Min-Ho Shin, Hae-Sung Nam, Kyeong-Soo Park, Seong-Woo 10. Choi, So-Yeon Ryu, Sun-Seog Kweon Yonsei Med J. 2018 Jan 1; 59(1): 92–100. Published online 2017 Nov 29. doi: 10.3349/ymj.2018.59.1.92 Stratton I, Adler A, Neil HA. Association of glycaemia with macrovascular and
- 11. microvascular complications of type 2 diabetes (UKPDS 35): prospective observational study. BMJ. 2000; 321:405-12.
- Benoit SR, Fleming R, Tsimikas AP, Ming JI. Predictors of glycemic control among 12. patients with Type 2 diabetes: A longitudinal study. BMC Public Health. 2005 5:36-45
- Al-AkourNemeh Al A, Yousef K, Aysha MA. Glycemic control and its determinants 13. among patients with type 2 diabetes mellitus attending a teaching hospital. J Diabetes Metab. 2011; 2:4.
- 14. Goudswaard, A. N., Stolk, R. P., Zuithoff, P., & Rutten, G. E. (2004). Patient characteristics do not predict poor glycaemic control in type 2 diabetes patients treated in primary care. European Journal of Epidemiology. 2004; 19:541–545. Fseha B. Glycemic Control and its Associated Factors in Type 2Diabetic Patients in
- 15.
- Suhul Hospital, Northwest Tigray, Ethiopia. J Diabetes Metab. 2017; 8:729. Yonas Mullugeta, Rajinder Chawla, Tedla Kebede, Yesehak Worku Indian J Clin Biochem. 2012 Oct; 27(4): 363–369. Published online 2012 Jun 6. doi: 16. 10.1007/s12291-012-0225-8
- Haghighatpanah M, Nejad ASM, Haghighatpanah M, Thunga G, Mallayasamy S. Factors that Correlate with Poor Glycemic Control in Type 2 Diabetes Mellitus 17. Patients with Complications. Osong Public Health Res Perspect. 2018;9(4):167-174.