



# Individualized Glycemic Control in Type 2 Diabetic Patients in Iran: A Multi-Center Data Analysis

Abolfazl Sadeghi<sup>1</sup>, PharmD;  Yahya Bayazidi<sup>1</sup>, PhD;  Majid Davari<sup>1</sup>, PharmD, PhD; Abbas Kebriaeezadeh<sup>1</sup>, PharmD, PhD; Amin Assarian<sup>1</sup>, PharmD; Alireza Esteghamati<sup>2</sup>, MD; Sepideh Yousefi<sup>3</sup>, PharmD

<sup>1</sup>Department of Pharmacoeconomics and Pharmaceutical Administration, School of Pharmacy, Tehran University of Medical Sciences, Tehran, Iran;

<sup>2</sup>Endocrinology and Metabolism Research Center, Vali-Asr Hospital, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran;

<sup>3</sup>School of Pharmacy and Pharmaceutical Science, Islamic Azad University, Tehran, Iran

## Correspondence:

Yahya Bayazidi, PhD;  
Social Security Organization,  
Postal code: 14155-65599,  
Tehran, Iran

Tel: +98 21 66491070

Fax: +98 21 66419537

Email: ybayazidi@gmail.com

Received: 18 September 2021

Revised: 15 April 2022

Accepted: 10 June 2022

## What's Known

- Data regarding individualized glycemic targets in patients with type 2 diabetes is scant in Iran. Before calling any amendment to Iranian diabetic guidelines and reaching robust judgment, we intended to evaluate the outcomes of current treatment protocols from the perspective of individualized recommendation.

## What's New

- As a result of poor glycemic control and high incidence rate of diabetes-related complications, glycemic control outcomes in under-treatment diabetic patients were considerably out of individualized targets. About 78% (77.6-79%) of patients had not achieved the individualized glycemic targets.

## Abstract

**Background:** Clinical guidelines and expert committees have recently suggested that the hemoglobin A1C (HbA1c) should be individualized based on various criteria. Data regarding the achievement of individualized glycemic targets in type 2 diabetes mellitus (T2DM) patients is scant in Iran. We intended to provide information found on real-world outcomes from the perspective of an individualized recommendation.

**Methods:** A cross-sectional analysis was conducted in 15 diabetes centers in Iran between 2013-2017. Two steps cluster sampling selection was used to recruit 1591 patients with T2DM. Considering Ismail-Beigi's individualized strategy, the study population was categorized into five treatment intensities of HbA1c: most intensive ( $\leq 6.5\%$ ), intensive (6.5–7.0%), less intensive ( $\sim 7.0\%$ ), not intensive (7.0–8.0%), and moderated ( $\sim 8.0\%$ ). The percentage of patients who met their group individualized glycemic targets was estimated as the degree of achievement of each treatment intensity.

**Results:** The cumulative incidence rate of early microvascular, advanced microvascular, and macrovascular complications was 53%, 25%, and 34%, respectively. Besides, 78% [77.6-79%] of patients did not achieve individualized glycemic targets.

**Conclusion:** The outcome showed poor individualized glycemic control and a high incidence of diabetes complications. Considering individualized HbA1c targets for Iranian patients with T2DM is an urgent need.

Please cite this article as: Sadeghi A, Bayazidi Y, Davari M, Kebriaeezadeh A, Assarian A, Esteghamati AR, Yousefi S. Individualized Glycemic Control in Type 2 Diabetic Patients in Iran: A Multi-Center Data Analysis. Iran J Med Sci. 2023;48(3):286-291. doi: 10.30476/ijms.2022.92805.2409.

**Keywords** • Diabetes mellitus • Individualized glycemic control • Precision medicine • Diabetes complications

## Introduction

The foremost step in managing type 2 diabetes mellitus (T2DM) is targeting glycemic goals.<sup>1,2</sup> In Iran, the golden target of hemoglobin A1C (HbA1c) for managing T2DM is  $<7.0\%$ , the same as the American Diabetic Association (ADA) guideline recommendation.<sup>3,4</sup> Patients with poorly controlled blood glucose are endangered with an increased risk of atherosclerotic cardiovascular disease (CVD) and microvascular disease.<sup>5</sup> A secondary prospective cohort analysis<sup>6</sup> and a comprehensive review<sup>7</sup> have proposed the optimization treatment strategies to prevent diabetes complications. Moreover, in the last decade, clinical guidelines and expert committees have recommended that

HbA1C should be individualized based on various criteria, including duration of diabetes, diabetic complications, socio-economic characteristics, comorbid conditions, life expectancy, and history of hypoglycemia.<sup>8-10</sup> According to Ismail-Beigi and others, the glycemetic target (HbA1c) range for T2DM patients should be individualized based on age, diabetes duration, macrovascular and microvascular complications, and propensity for hypoglycemia.<sup>11</sup> Furthermore, careful consideration must be given to each patient's capacities, desires, values, living situations, support systems, cognitive status, overall prognosis, and life expectancy. Consequently, glycemetic targets should not be viewed as tight goals; they should be flexible and adaptable to changes based on patients' health and living conditions.<sup>12</sup> The result of a patient-level Monte Carlo-based Markov model also showed that individualized control was a cost-saving scenario and would generate more Quality Adjusted Life Years (QALYs) than uniform intensive control (HbA1c level <7%) for the US T2DM population from a health care sector perspective.<sup>13</sup>

Iranian diabetic patients are expected to reach 5.2 million in 2025.<sup>14</sup> A previous study reported that Iranian patients with T2DM exert a significant burden on the healthcare system.<sup>15</sup> Additionally, the Iranian diabetes population has a high prevalence of long-standing diabetes and microvascular complications.<sup>7</sup> Analyzing the real-world outcome of the traditional guideline is vital before calling for any amendment in the treatment of T2DM. To the best of our knowledge, this study is the first evaluation in Iran to compare the real-world health outcome (HbA1c results) from the perspective of a personalized strategy.

## Materials and Methods

### Study Design

Clinical data extraction was performed from paper and electronic records at 15 diabetes centers (public, semi-public [centers related to the social security insurance organization], and private centers) in five provinces in Iran including Tehran, Isfahan, Yazd, Mazandaran, and Kurdistan.

Two-stage cluster sampling was applied to recruit the required sample size: 1) province selection (clusters) and 2) diabetes-center selection in the clusters. Subjects were selected by a random sampling method based on the patient identification number. The entire population was included since the center's statistical population was small.

Subjects were eligible for inclusion in our study if they received diabetic routine care, regular glycemetic control visits, laboratory

investigations, and antidiabetic medications for at least five years. Written informed consent was taken from those aged 18 or older if they were willing/able to participate in the study.

Ethics approval was obtained from the Ethics Committee of Tehran University of Medical Sciences (IR.TUMS.PSRC.REC.1396.1991).

### Individualized Strategy

We defined individualized HbA1C targets according to the Ismail-Beigi methodology by considering multi-variables, including HbA1C, duration of diabetes, age, and history of microvascular and macrovascular complications.

Duration of diabetes is defined as the difference between the patient's age and the age of diabetes diagnosis. The short duration was determined as five to ten years; while over ten years was taken as a long duration, and advanced age was considered for subjects aged 75 or older. Macrovascular complications are regarded as the diagnosis of the following: heart failure, angina, myocardial infarction, and stroke. Advanced microvascular complications are defined as having one or more of the following: receipt of nephrology care (dialysis in the past 12 months), macroalbuminuria (the ratio of albumin to creatinine >300 with CACR test), blindness, severe neuropathy, and amputation.

Treatment intensities (HbA1C targets) were defined based on the chosen individualized recommendations as: most intensive ( $\leq 6.5\%$ ), intensive (6.5–7.0%), less intensive ( $\sim 7.0\%$ ), not intensive (7.0–8.0%), and moderated ( $\sim 8.0\%$ ). The patients were assigned to each treatment intensity, then the percentage of T2DM patients who met their treatment intensity' targets was extracted. Finally, the result of the individualized target was compared with the uniform conventional target (HbA1c<7%) as a reference to stick to the current recommendations guideline.

### Statistical Analysis

Statistical analysis was performed using the STATA software version 14 for Windows (Stata Corp., College Station, TX, US). Based on the results of normality tests, continuous variables (i.e., duration of disease, glycemetic indices) are presented either as mean $\pm$ SD, and categorical outcomes (i.e., meeting the preset glycemetic targets) are demonstrated as proportions (95% confidence interval [95% CI]) or mean $\pm$ SEM.

## Results

### Patient Characteristics

In total, 1591 patients with T2DMs were included in this study. Patient characteristics are

shown in table 1. The mean age was 60, and 58% of patients were women. The mean HbA1C in all patients for five years (2013-17) was 7.56, and 33% had a short disease duration. The cumulative incidence rates of macrovascular, microvascular, and advanced microvascular complications were 34%, 53%, and 25% over five years, respectively. The results showed an increase of 11.5% in the incidence rate of macrovascular complications and 19.43% in early microvascular complications over the period.

### Individualization of Glycemic Targets

Using Ismail-Beigi and colleagues strategy and including clinical information, the study population were stratified and simplified in the five treatment intensities (HbA1c target).<sup>11</sup> Table 2 shows the percentage of the study population from 2013 to 2017 including in each of the five treatment intensities. We calculated the number of patients in each group who had met the group target and the average of the studied population who were in individualized targets. The details of the data are presented in table 3.

**Table 1:** Clinical and demographic information of type 2 diabetes patients

Variable	Category	N (%) N=1591		
Sex	Women	925 (58.11)		
	Men	666 (41.86)		
Age (year)	20-44	111 (7)		
	45-65	986 (62)		
	66-75	382 (24)		
	>75	111 (7)		
Diabetes Duration	Short diabetes duration	522 (32.81)		
	Long diabetes duration	1069 (67.19)		
Macrovascular complications	Prevalence	Five-years	183 (11.5)	
		Incidence rate	2013	491 (30.86)
			2017	674 (42.36)
Early microvascular complications	Prevalence	Five -years	309 (19.43)	
		Incidence rate	2013	749 (47.07)
			2017	1058 (66.50)
Advanced microvascular complications	Prevalence	Five-years	180 (11.32)	
		Incidence rate	2013	313 (19.67)
			2017	493 (30.99)
HbA1c (mean±SD)	Five-years		7.56±1.22	
		2013	7.58±1.38	
		2017	7.58±1.39	

**Table 2:** Patients in each treatment intensity category

Treatment intensity (HbA <sub>1c</sub> target)	Number of patients in each category N (%)				
	2013	2014	2015	2016	2017
Most intensive (≤6.5%)	151 (9.54)	137 (8.65)	110 (6.95)	93 (5.87)	85 (5.37)
Intensive (6.5%–7.0%)	410 (25.90)	357 (22.55)	326 (20.59)	295 (18.64)	241 (15.22)
Less intensive (7.0%)	309 (19.52)	338 (21.35)	368 (23.25)	393 (24.83)	425 (26.85)
Not intensive (7.0%–8.0%)	551 (34.81)	578 (36.51)	579 (36.58)	576 (36.39)	573 (36.20)
Moderated (8.0%)	162 (10.23)	173 (10.93)	200 (12.63)	226 (14.28)	259 (16.36)

**Table 3:** The degree to which personalized HbA1c targets are met

Treatment intensity (HbA <sub>1c</sub> target)	The percentages of degree of achievement of individualized HbA1c targets*				
	2013	2014	2015	2016	2017
Most intensive (≤6.5%)	25.17	26.28	24.55	18.28	12.94
Intensive (6.5%–7.0%)	17.80	15.97	19.94	15.93	14.94
Less intensive (7.0%)	12.30	13.31	17.12	17.81	20.71
Not intensive (7.0%–8.0%)	32.12	29.93	34.89	33.85	35.43
Moderated (8.0%)	12.35	8.67	11.00	7.96	8.49
The degree of achievement	21.86	20.59	23.94	21.92	22.74
Patient percentages on conventional target (≤7%)	32	34	34	31	31

\*Percentage of patients who meet HbA1c targets in each Treatment Intensity.

### Meeting Individualized Targets

Twenty-one percent of patients (21.8 and 22.7% from 2013 to 2017) met the individualized glycemic targets. At the same time, about 32% (32 and 31% from 2013 to 2017) of the study population were within the target for uniform conventional recommendation (<7%, 53 mmol/mol) (table 3).

## Discussion

Achievement of individualized HbA1c target levels showed that only one-fifth of patients with T2DM who regularly received diabetic treatment, were adequately controlled (21.88% [22.4-21%]). However, a study in the US<sup>16</sup> showed that 29% and 30 % of T2DM patients were out of the target control based on ADA guidelines and the Ismail-Beigi strategy,<sup>11</sup> respectively.

Another study conducted by Coons and colleagues reported that the level of glycemic control would be different from targets recommended by traditional guidelines if considering the age and comorbidities of each patient. It accentuated that more studies should be done to develop a methodology to individualize the HbA1C in Canada.<sup>17</sup> These crucial factors regarding traditional guidelines and individualized recommendations bring burning questions about the goal of diabetes translational and quality improvement studies. Studies show that the prevalence of major microvascular complications of T2DM patients in Iran is high.<sup>18</sup> Besides, cardiovascular complications related to diabetes are more frequent in Iran than in other Eastern Mediterranean and Middle Eastern countries.<sup>19</sup> In the same vein, our result showed an increase of 11.5% in the incidence rate of macrovascular complications and 19.43% in early microvascular complications over the study period (table 1).

On the other hand, the Iranian diabetes guideline, similar to the ADA guideline, recommends that the HbA1C should be less than 7% for managing T2DM.<sup>5</sup> However, several studies reported that most patients with T2DM did not reach this target.<sup>20, 21</sup> It is reasonably expected that this inadequate glycemic control will increase diabetes complications, decrease the patient's quality of life, and intensify the disease burden for the patients and the healthcare system. Therefore, we suggest that diabetes guidelines be re-evaluated from various health-related facets, and individualized strategy is one of them.

The data regarding the achievement of individualized HbA1c targets in T2DM patients is scarce in Iran. We intended to provide information illustrating the pharmaceutical health outcome

from the perspective of an individualized strategy proposed by Ismail-Beigi and others.<sup>11</sup>

The central goal of recommendations to individualize glycemic targets (HbA1c) is to increase the health outcomes in the diabetes population. In fact, individualized recommendations encourage more intensive glycemic control for younger T2DM patients with fewer comorbidities to prevent macrovascular and microvascular complications in the long-term and more relaxed glycemic control for elderly patients and those with the risk of hypoglycemia or those with multiple comorbidities.<sup>4, 22, 23</sup> Moreover, as part of selecting wisely, the American Geriatric Society has warned about intensive glycemic control in older patients with diabetes by prescribing medications other than metformin.<sup>24</sup> Overtreatment in elders means using glucose-lowering medicines (except metformin) to achieve a tight HbA1c target that might surge the risk of hypoglycemia.<sup>25, 26</sup> Studies showed that older patients with T2DM might have more restricted HbA1C <7% than younger patients,<sup>27</sup> which illustrates overtreatment in some elders according to individualized recommendations.<sup>25</sup>

Comparing our results with international evidence depicts the need for emergency amendments to Iranian diabetes management and a strong recommendation for considering individualized HbA1c targets. Nevertheless, it is clear that all required conditions for implementing the individualized strategy for managing diabetes, including education, financial resources, and healthcare facilities, must be considered in advance.

Our study had several strengths and limitations. We could declare that this study reflected the outcomes of real-world ongoing clinical practices as the data were collected randomly at the patient level from diabetic clinics. Despite these merits, our study had some limitations. The range of hypoglycemic episodes in over-treated patients was not assessed, as they were not documented in patients' clinical information.

## Conclusion

The outcome showed poor individualized glycemic control and a high incidence of diabetes complications in the population with T2DM in Iran. Considering individualized HbA1c targets for Iranian patients with T2DM is an urgent need.

## Acknowledgment

We thank all the staff of the Diabetes Research Centers that helped our research team to conduct this research.

### Authors' Contribution

AS, YB, MD, AA, AE, SY: the conception and design of the work, interpretation of the data, drafting of the article. All authors approved the present version of the manuscript. All authors agreed on being accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

**Conflict of Interest:** None declared.

### References

- 1 Writing Group for the DERG, Orchard TJ, Nathan DM, Zinman B, Cleary P, Brillon D, et al. Association between 7 years of intensive treatment of type 1 diabetes and long-term mortality. *JAMA*. 2015;313:45-53. doi: 10.1001/jama.2014.16107. PubMed PMID: 25562265; PubMed Central PMCID: PMCPMC4306335.
- 2 Zabala A, Darsalia V, Holzmann MJ, Franzen S, Svensson AM, Eliasson B, et al. Risk of first stroke in people with type 2 diabetes and its relation to glycaemic control: A nationwide observational study. *Diabetes Obes Metab*. 2020;22:182-90. doi: 10.1111/dom.13885. PubMed PMID: 31576643.
- 3 American Diabetes A. Standards of medical care in diabetes--2012. *Diabetes Care*. 2012;35:S11-63. doi: 10.2337/dc12-s011. PubMed PMID: 22187469; PubMed Central PMCID: PMCPMC3632172.
- 4 American Diabetes A. Standards of medical care in diabetes--2014. *Diabetes Care*. 2014;37:S14-80. doi: 10.2337/dc14-S014. PubMed PMID: 24357209.
- 5 American Diabetes A. 6. Glycemic Targets: Standards of Medical Care in Diabetes-2018. *Diabetes Care*. 2018;41:S55-S64. doi: 10.2337/dc18-S006. PubMed PMID: 29222377.
- 6 Kaze AD, Santhanam P, Erqou S, Ahima RS, Echouffo-Tcheugui JB. Long-term variability of glycemic markers and risk of all-cause mortality in type 2 diabetes: the Look AHEAD study. *BMJ Open Diabetes Res Care*. 2020;8. doi: 10.1136/bmjdr-2020-001753. PubMed PMID: 33257421; PubMed Central PMCID: PMCPMC7705503.
- 7 Moradi Y, Baradaran HR, Djalalinia S, Chinekesh A, Khamseh ME, Dastoorpoor M, et al. Complications of type 2 diabetes in Iranian population: An updated systematic review and meta-analysis. *Diabetes Metab Syndr*. 2019;13:2300-12. doi: 10.1016/j.dsx.2019.05.018. PubMed PMID: 31235172.
- 8 Menendez Torre E, Lafita Tejedor FJ, Artola Menendez S, Millan Nunez-Cortes J, Alonso Garcia A, Puig Domingo M, et al. Recommendations for the pharmacological treatment of hyperglycemia in type 2 diabetes. *Aten Primaria*. 2011;43:202. doi: 10.1016/j.aprim.2010.07.012. PubMed PMID: 21382648; PubMed Central PMCID: PMCPMC7024946.
- 9 Handelsman Y, Mechanick JI, Blonde L, Grunberger G, Bloomgarden ZT, Bray GA, et al. American Association of Clinical Endocrinologists Medical Guidelines for Clinical Practice for developing a diabetes mellitus comprehensive care plan. *Endocr Pract*. 2011;17:1-53. doi: 10.4158/ep.17.s2.1. PubMed PMID: 21474420.
- 10 Inzucchi SE, Bergenstal RM, Buse JB, Diamant M, Ferrannini E, Nauck M, et al. Management of hyperglycemia in type 2 diabetes: a patient-centered approach: position statement of the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). *Diabetes Spectrum*. 2012;25:154-71. doi: 10.2337/diaspect.25.3.154.
- 11 Ismail-Beigi F, Moghissi E, Tiktin M, Hirsch IB, Inzucchi SE, Genuth S. Individualizing glycemic targets in type 2 diabetes mellitus: implications of recent clinical trials. *Ann Intern Med*. 2011;154:554-9. doi: 10.7326/0003-4819-154-8-201104190-00007. PubMed PMID: 21502652.
- 12 Davies MJ, D'Alessio DA, Fradkin J, Kernan WN, Mathieu C, Mingrone G, et al. Management of Hyperglycemia in Type 2 Diabetes, 2018. A Consensus Report by the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). *Diabetes Care*. 2018;41:2669-701. doi: 10.2337/dci18-0033. PubMed PMID: 30291106; PubMed Central PMCID: PMCPMC6245208.
- 13 Laiteerapong N, Cooper JM, Skandari MR, Clarke PM, Winn AN, Naylor RN, et al. Individualized Glycemic Control for U.S. Adults With Type 2 Diabetes: A Cost-Effectiveness Analysis. *Ann Intern Med*. 2018;168:170-8. doi: 10.7326/M17-0537. PubMed PMID: 29230472; PubMed Central PMCID: PMCPMC5989575.
- 14 Javanbakht M, Mashayekhi A, Baradaran HR, Haghdoost A, Afshin A. Projection of Diabetes Population Size and Associated Economic Burden through 2030 in Iran: Evidence from Micro-Simulation Markov Model and Bayesian Meta-Analysis. *PLoS One*.

- 2015;10:e0132505. doi: 10.1371/journal.pone.0132505. PubMed PMID: 26200913; PubMed Central PMCID: PMC4511591.
- 15 Peykari N, Djalalinia S, Kasaeian A, Naderimagham S, Hasannia T, Larijani B, et al. Diabetes research in Middle East countries; a scientometrics study from 1990 to 2012. *J Res Med Sci.* 2015;20:253-62. PubMed PMID: 26109972; PubMed Central PMCID: PMC4468230.
  - 16 Laiteerapong N, John PM, Nathan AG, Huang ES. Public health implications of recommendations to individualize glycemc targets in adults with diabetes. *Diabetes Care.* 2013;36:84-9. doi: 10.2337/dc11-2344. PubMed PMID: 22961575; PubMed Central PMCID: PMC3526201.
  - 17 Coons MJ, Greiver M, Aliarzadeh B, Meaney C, Moineddin R, Williamson T, et al. Is glycemc control in Canadians with diabetes individualized? A cross-sectional observational study. *BMJ Open Diabetes Res Care.* 2017;5:e000316. doi: 10.1136/bmjdr-2016-000316. PubMed PMID: 28761645; PubMed Central PMCID: PMC5530242.
  - 18 Moradi Y, Baradaran HR, Djalalinia S, Chinekesh A, Khamseh ME, Dastoorpoor M, et al. Complications of type 2 diabetes in Iranian population: An updated systematic review and meta-analysis. *Diabetes Metab Syndr.* 2019;13:2300-12. doi: 10.1016/j.dsx.2019.05.018. PubMed PMID: 31235172.
  - 19 Cho NH, Shaw JE, Karuranga S, Huang Y, da Rocha Fernandes JD, Ohlrogge AW, et al. IDF Diabetes Atlas: Global estimates of diabetes prevalence for 2017 and projections for 2045. *Diabetes Res Clin Pract.* 2018;138:271-81. doi: 10.1016/j.diabres.2018.02.023. PubMed PMID: 29496507.
  - 20 Dhanaraj E, Raval AD, Yadav R, Bhansali A, Tiwari P. Prescribing pattern of antidiabetic drugs and achievement of glycemc control in T2DM patients tertiary care hospital in North India. *International Journal of Diabetes in Developing Countries.* 2013;33:140-6. doi: 10.1007/s13410-013-0123-5.
  - 21 Lipska KJ, Yao X, Herrin J, McCoy RG, Ross JS, Steinman MA, et al. Trends in Drug Utilization, Glycemc Control, and Rates of Severe Hypoglycemia, 2006-2013. *Diabetes Care.* 2017;40:468-75. doi: 10.2337/dc16-0985. PubMed PMID: 27659408; PubMed Central PMCID: PMC5360291.
  - 22 Cheng AY, Lau DC. The Canadian Diabetes Association 2013 clinical practice guidelines-raising the bar and setting higher standards! *Can J Diabetes.* 2013;37:137-8. doi: 10.1016/j.cjcd.2013.04.005. PubMed PMID: 24070834.
  - 23 Kirkman MS, Briscoe VJ, Clark N, Florez H, Haas LB, Halter JB, et al. Diabetes in older adults. *Diabetes Care.* 2012;35:2650-64. doi: 10.2337/dc12-1801. PubMed PMID: 23100048; PubMed Central PMCID: PMC3507610.
  - 24 Caverly TJ, Fagerlin A, Zikmund-Fisher BJ, Kirsh S, Kullgren JT, Prenovost K, et al. Appropriate Prescribing for Patients With Diabetes at High Risk for Hypoglycemia: National Survey of Veterans Affairs Health Care Professionals. *JAMA Intern Med.* 2015;175:1994-6. doi: 10.1001/jamainternmed.2015.5950. PubMed PMID: 26502113.
  - 25 Lipska KJ, Ross JS, Miao Y, Shah ND, Lee SJ, Steinman MA. Potential overtreatment of diabetes mellitus in older adults with tight glycemc control. *JAMA Intern Med.* 2015;175:356-62. doi: 10.1001/jamainternmed.2014.7345. PubMed PMID: 25581565; PubMed Central PMCID: PMC4426991.
  - 26 Shi Min Ko M, Kit Lee W, Chang Ang L, Goh SY, Mong Bee Y, Ming Teh M. A Cross-Sectional study on risk factors for severe hypoglycemia among Insulin-Treated elderly type 2 diabetes Mellitus (T2DM) patients in Singapore. *Diabetes Res Clin Pract.* 2022;185:109236. doi: 10.1016/j.diabres.2022.109236. PubMed PMID: 35131380.
  - 27 Stark Casagrande S, Fradkin JE, Saydah SH, Rust KF, Cowie CC. The prevalence of meeting A1C, blood pressure, and LDL goals among people with diabetes, 1988-2010. *Diabetes Care.* 2013;36:2271-9. doi: 10.2337/dc12-2258. PubMed PMID: 23418368; PubMed Central PMCID: PMC3714503.