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Original research

# Type 2 Diabetes Mellitus non-surgical remission: A possible mission

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## ARTICLE INFO

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# ABSTRACT

ywords:	Background: Remission of Type 2 Diabetes Mellitus (DM) has been observed throughout the last few years, yet
abetes Mellitus	factors associated with remission through non-surgical approaches are still unclear. So, the factors associated
mission	with type 2 DM remission were investigated.
ldagliptin	<i>Methods:</i> Out 670 patients, 63 patients gained non-surgical remission of type 2 DM (defined according to ADA criteria), and 396 patients served as control matched for age, sex, and BMI. The records of patients who attended Alexandria University Students Hospital between the years 2015 and 2018 were reviewed retrospectively. The collected data at first visit and 1 year after remission were history, examination, HbA1c %, oral glucose tolerance test & fasting blood glucose and lipid profile and type of treatment. <i>Results:</i> 75% of patients were females with mean age $51.5 \pm 11.34$ years. The mean duration of treatment till remission was $26.4 \pm 33.1$ months. One year after remission HbA1c & BMI were significantly decreased (P = 0.001, 0.03; respectively). However, cox proportional hazards model & ROC curve analysis showed that age < 50 years, female sex, short duration of diabetes < $5.2$ years, initial HbA1c < $8.5\%$ , HDL > $45.6$ mg/dl, and initial intervention with lifestyle modification, 2000 mg metformin and 100 mg vildagliptin are the factors associated with remission. <i>Conclusion:</i> Our mission in type 2 DM female patients < 50 years with short duration of disease < $5.2$ years, initial HbA1c < $8.5\%$ and HDL > $45.6$ mg/dl, is to achieve diabetes remission by intensive life style modification with daily metformin and vildagliptin.

## Introduction

Diabetes Mellitus (DM) is a major public health problem which is associated with serious complications that can be life threatening. Even prediabetes has shown to be associated with various complications including increased cardiovascular risk [1].

Traditionally DM has been considered a progressive incurable condition and the best targets for management were tight glycemic & metabolic control to prevent cardiovascular and other complications [2].

Studies to date have focused largely on DM remission after bariatric surgery, specifically gastric bypass and biliopancreatic diversion as more effective than conventional medical therapy in controlling hyperglycemia in severely obese patients with type 2 diabetes [3,4]. Yet, few studies demonstrated remission of type 2 DM using nonsurgical approaches. The Look AHEAD (Action for Health for Diabetes) study is perhaps the largest randomized controlled trial of an intensive lifestyle

intervention among adults with type 2 DM. Although designed principally to examine the effect of weight loss on cardiovascular disease incidence, it also determined the association of an intensive lifestyle intervention with the frequency of partial and complete type 2 DM remission [5].

## Patients and methods

We conducted a retrospective cohort study on type 2 DM patients attending Endocrinology clinic at Alexandria University Students' Hospital (AUSH) from the year 2015 till the year 2018 to describe the variables associated with remission among patients with type 2 DM who received the standards of care according to updated guidelines.

We excluded patients based on the following criteria: 1) Type 1 DM, 2) History of bariatric surgery, 3) Long-term use (> 60 days) of steroids or other hyperglycemic drugs in the year prior to baseline, 4) Organ failure (heart, renal, or liver), 5) Patients who did not complete one

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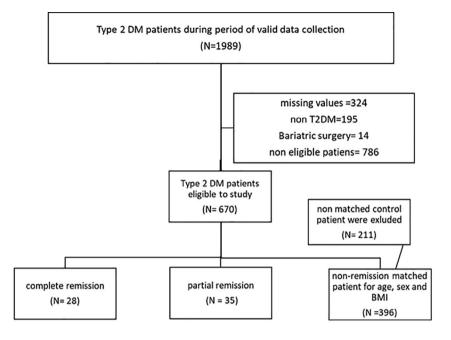


Fig. 1. Flow chart of patients enrolled in the study.

Table 1Demographic data of enrolled patients.

	Range (mean ± SD)		p-Value	
	Remission	No remission		
Female sex (%)	75 73		0.90	
Age (years)	25-60 28.5-63.5		0.19	
	(51.5 ± 11.34)	$(50.0 \pm 12.5)$		
Duration of DM (years)	0.5–9.5	4.5-21	0.02	
	$(4.20 \pm 4.13)$	$(12.20 \pm 7.63)$		
Initial body mass index	25.9-41.5	27.11-46	0.50	
(kg/m <sup>2</sup> )	$(34.22 \pm 7.31)$ $(35.8 \pm 8.69)$			
Smoker (%)	6.7	71	0.00	
Comorbidities (%)			0.01	
Hypertension	30	42		
Coronary heart disease	10	31		
Dyslipidemia	20	54		
Peripheral neuropathy	10	27		
Nephropathy	2	11		

year after remission; or 6) Patients who missed more than 2 visits during follow-up.

Definition of remission was based on the 2009 ADA consensus statement; Partial remission defined as having two or more consecutive subdiabetic HbA1c measurements, all of which were in the range of 5.7–6.4% over a period of at least 12 months and complete remission defined as having two or more consecutive normoglycemic HbA1c measurements, all of which were < 5.7% over a period of at least 12 months. Each definition of remission requires the absence of pharmacologic treatment during the defined observation period [6].

All patients around the age of 30 years were subjected to antibodies screening and C-peptide level testing to exclude type 1 DM.

We collected baseline covariates from records at the first visit including; demographic characteristics, history including history of DM (time of diagnosis and duration of treatment), and physical examination. Patients were followed up every 3 months till the end of one year after remission. The laboratory work up related to glycemic control (fasting blood sugar, oral glucose tolerance test, and HbA1c), lipid profile (LDL, HDL, VLDL, and total cholesterol) and biochemical assays (estimated glomerular filtration rate [eGFR]) were done every visit.

All patients were motivated to intensive lifestyle intervention according to ADA guidelines. The intensive intervention included caloric restriction (under supervision of a registered dietitian), moderate-intensity aerobic physical activity (goal 175 min weekly), and smoking cessation [7]. Monitoring of lifestyle adherence was assessed every visit through structured self-reporting diaries & structured personal interviews.

Descriptive statistics were used to summarize the baseline demographic and clinical data, disease characteristics and treatment options. Differences in baseline characteristics between remission and control groups were tested by independent t test (normally distributed data) or Mann– Whitney test (non-normally distributed data).

We examined unadjusted and adjusted models based on a complete case–only analysis. The adjusted model included demographic data (age at cohort entry and sex), baseline factors (BMI, eGFR, HbA<sub>1c</sub> level), comorbidities (cardiovascular disease, hypertension, dyslipidemia), and diabetes factors (oral or insulin use, drugs categories, and years since diabetes diagnosis).

We calculated the cumulative incidence of remission using the Kaplan-Meier method of estimating the cumulative probability of an event. Cox proportional hazards models were specified to identify significant factors associated with the time of remission. Receiver operating characteristic curve (ROC) was plotted to analyze a recommend cut off value of significant continuous variables.

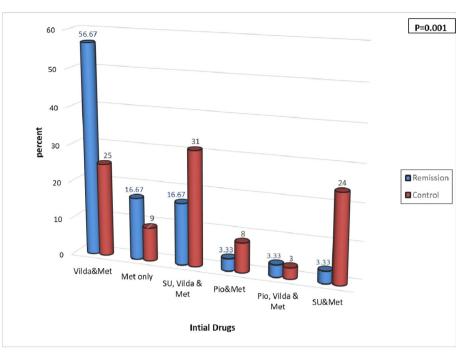
All analyses were conducted using SPSS version 22 (SPSS Inc., an IBM company; Chicago, Ill) and MedCalc<sup>®</sup> 13.3.3.0 statistical software (MedCalc Software; Mariakerke, Belgium) packages. All statistical tests were two sided; the level of significance was < 0.05.

Collection and analysis of the data in this study were ethically approved by local Institutional Review Board.

## Results

Medical records evaluated retrospectively showed that; out of 670 patients with type 2 DM, 63 patients showed DM remission; around 44% of patients developed complete remission and 56% developed partial remission (Fig. 1).

At the first visit the demographic data were included in Table 1 10%



SU: Sulphonyl urea, Vilda: Vildagliptin, Met: Metformin, Pio: Pioglitazone

#### Table 2

Comparison between the first visit and one year after remission regarding weight and metabolic profile of patients.

Mean (min-max)	First visit	Last visit	*P (2-tailed)
BWT (Kg)	94.72 ± 19.35 (57–150)	89.12 ± 16.99 (60-145)	0.003
BMI (Kg/m <sup>2</sup> )	34.22 ± 7.31 (23-55)	$30.16 \pm 6.25 (24.2-53.2)$	0.027
A1C (%)	8.87 ± 1.49 (7-12.4)	5.55 ± 0.53 (4-7)	0.0001
FPS (mg/dl)	185.77 ± 74.58 (102-415)	101.1 ± 11.24 (76–122)	0.0001
PPS (mg/dl)	263.97 ± 95.19 (144-476)	116.6 ± 25.2 (70–167)	0.0001
HDL (mg/dl)	42.93 ± 8.63 (25-66)	96.97 ± 45.19 (31-229)	0.0001
LDL (mg/dl)	$143.13 \pm 39.98 (100-246)$	48.44 ± 10.82 (31-68)	0.006
TG (mg/dl)	$146.13 \pm 94.19 (22-479)$	91.82 ± 71.23 (26-331)	0.002

Data are mean  $\pm$  standard deviation. BWT: body weight; BMI: body mass index; A1C: hemoglobin A1c, FPS: fasting plasma glucose, PPS: post prandial plasma glucose, HDL: high density lipoprotein, LDL: low density lipoprotein, TG: triglycerides.

\* Statistically significant at  $p \leq$  0.05.

suffered from ischemic heart disease, and 20% had dyslipidemia.

In the first visit, all patients who showed remission were adherent to lifestyle modification and to the prescribed oral antidiabetic drugs; the most frequently prescribed drugs in remission group were vilda-gliptin + metformin, versus sulphonyl urea + vildagliptin + metformin in control; group (56.67%, 31%, respectively) (Fig. 2). Only 26% of those patients were maintained on statin.

The mean duration of treatment till complete remission was 12.5  $\pm$  6.95 months, which was significantly lower than mean duration of treatment in partial remission (74  $\pm$  14.29 months, p < 0.001) and also in non-remission patients (84  $\pm$  24.98 months, p < 0.019).

In comparison to first visit, patients after 1 year of remission their body weight & BMI were significantly decreased; mean 94.72  $\pm$  19.35 kg Vs 89.12  $\pm$  16.99 kg (P = 0.003), and 34.22  $\pm$  7.31 kg/m<sup>2</sup> Vs 32.16  $\pm$  6.25 kg/m<sup>2</sup> (P = 0.027), respectively. Study findings were adjusted for baseline BMI. Also, there was a significant decrease in Hb A1c, fasting plasma glucose (FPG), and oral glucose tolerance test; 8.87  $\pm$  1.49% Vs 5.55  $\pm$  0.53% (P = 0.0001), 185.77  $\pm$  74.58 mg/dl Vs 101.1  $\pm$  11.24 mg/dl (P = 0.0001), and 263.97  $\pm$  95.19 mg/dl Vs 116.6  $\pm$  25.2 mg/dl (P = 0.0001), respectively (Table 1).

Moreover, lipid profile of the patients 1 year after remission has

changed dramatically.

Compared to first visit, there was a significant increase in HDL;  $42.93 \pm 8.63 \text{ mg/dl}$  Vs  $96.97 \pm 45.19 \text{ mg/dl}$  (P = 0.0001), whereas there was a significant decrease in LDL and triglycerides (TG);  $143.13 \pm 39.98 \text{ mg/dl}$  Vs  $48.44 \pm 10.82 \text{ mg/dl}$  (P = 0.006) and  $146.13 \pm 94.19 \text{ mg/dl}$  Vs  $91.82 \pm 71.23 \text{ mg/dl}$  (P = 0.002), respectively (Table 2).

As shown in Fig. 3; the median cumulative incidences of partial and complete remission were 1.4% (95% CI 1.1–1.5%) and 1.17% (95% CI 0.940–2.54%), respectively. The cumulative incidence of remission was 3.47%. Patients with DM duration less than 2.5 years showed a high incidence of remission; 4.6% (95% CI 4.1–5.3%) whereas a low incidence of remission; 0.67% (95% CI 0.4–0.8%) was found in patients with DM duration more than 9 years.

After univariate analysis of variables associated with remission (partial & complete), the cox proportional hazards models were performed which showed that in a younger age, female patients, with shorter duration of diabetes and without history of smoking or IHD, especially with lower baseline HbA1c level and higher HDL have the highest probability for diabetes remission. And they will have 7 times the chance for diabetes remission if they were initially treated with 2000 mg metformin and 100 mg vildagliptin but did not start with

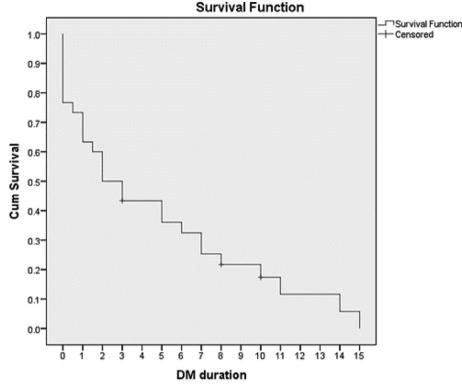


Fig. 3. Kaplan-Meier curve for cumulative incidence of type 2 DM remission.

## Table 3

The cox proportional hazards models of included type 2 diabetic patients to identify significant factors associated with remission.

	В	S.E	*Adjusted p	HR	95% CI	
					LL	UL
Age	-0.099	0.01	0.039	4.41	1.080	18.060
Female sex	0.004	0.26	0.048	1.004	0.974	1.030
nonsmoking	0.228	0.3	0.003	1.26	1.010	1.560
Diabetes duration	-0.03	0.022	0.003	6.00	0.999	8.002
History of IHD	-2.06	0.9	0.023	5.2	0.022	0.748
Initial HbA1c	-2.347	0.763	0.002	10.45	2.340	16.570
Initial HDL	0.259	0.166	0.011	0.77	0.557	1.069
2000 mg Metformin and Vildagliptin	0.185	0.12	0.012	7.204	0.951	9.523
Insulin therapy	-0.08	0.05	0.008	7.008	0.940	10.080

CI: Confidence interval; LL: Lower limit; UL: Upper limit; HR: Hazards ratio. \* Adjusted P is significant if < 0.05.

insulin therapy after adjusting for the effects of other factors, (Table 3). As shown in Fig. 4, receiver operating characteristic (ROC) curve was constructed for determining the best cut off value for prediction of type 2 DM remission detected by Youden index. The cut off and predictive values were age < 50 years (sensitivity 71.5% & specificity 61.5%), duration of DM < 5.2 years (sensitivity 89.5% & specificity 59.5%), baseline HbA1c level < 8.5% (sensitivity 58.7% & specificity 55.2%) and HDL > 45.6 mg/dl (sensitivity 62% & specificity 69%). ROC curve for the prediction of remission of DM showed that highest sensitivity was found with short duration of DM while highest specificity was found with HDL.

#### Discussion

Type 2 DM has been regarded as a progressive lifelong disease with long term dangerous complications. Yet, a growing body of evidence from clinical research has reported the remission of type 2 DM mostly after bariatric surgery. However, recently remission of type 2 DM can be achieved using non surgical approaches.

The current study showed that the median incidences of partial and complete remission among 670 patients with type 2 DM throughout 3 years were 1.4% (95% CI 1.1–1.5%) and 1.17% (95% CI 0.940–2.54%), respectively. In accordance with our finding, Karter et al, found that the incidences of remissions per 1000 person-years of partial, complete, or prolonged remission were 2.8 (2.6–2.9), 0.24 (0.20–0.28), and 0.04 (0.01–0.06), respectively [8]. Also, Uusitupa showed that remission of type 2 DM can be achieved and the most important factor affecting remission is weight loss which improves insulin sensitivity and on the long run may help to preserve  $\beta$ -cell mass [9].

The largest trial to date; the Look AHEAD multicenter randomized controlled trial, compared intensive lifestyle intervention (ILI) with diabetes support and education (DSE) among 4503 overweight & obese patients with type 2 DM across 4 years. Results showed that ILI group was more likely to experience remission (partial or complete), with prevalence of 11.5% (95% CI, 10.1%–12.8%) during the first year and 7.3% (95% CI, 6.2%–8.4%) at 4 years, compared with 2% for the DSE group at both time points (95% CIs, 1.4%–2.6% at year 1 and 1.5%–2.7% at year 4) (P < 0.001 for each). In agreement with our findings, DM remission occurred particularly in those whose diabetes is of short duration, who have lower HbA1c levels, and who do not require insulin therapy. Also, in line with our results, baseline BMI was not associated with remission in the Look AHEAD study, which found that only a change in BMI predicted remission [10].

Contrary to our results, Karter et al found that older age is associated with higher rates of remission. Their results are consistent with evidence from the Diabetes Prevention Program demonstrating that older adults over 65 years experienced the greatest risk reduction in diabetes from lifestyle modification [8].

In accordance with our results, Lean et al showed in the DiRECT trial that type 2 DM remission can occur without surgery and the most potent predictor of remission is weight loss > 15 kg and that sustained remission was linked to the extent of sustained weight loss [11].

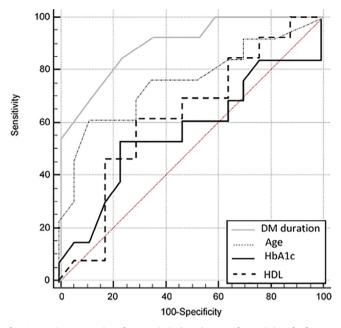


Fig. 4. Receiver operating characteristic (ROC) curve determining the best cut off value for predictors of type 2 DM remission.

The majority of patients who showed remission in our study were initially prescribed vildagliptin + metformin. After occurrence of remission, patients stopped all drugs and were maintained only on lifestyle programs. Vellanki et al, in their 4-year placebo-controlled study found that remission of type 2 DM is maintained by metformin + sitagliptin. Maintenance of remission was significantly higher in sitagliptin + metformin group compared with placebo (P = 0.015), and mean time to relapse was significantly prolonged in the metformin + sitagliptin group compared with the placebo group (P = 0.004) [12]. However, the proposed mechanisms for remission of T2DM were related to weight loss, this could not be explained by modest effect of vildagliptin and metformin on weight. We matched the cases and control for BMI and adjusted all factors against BMI in cox regression analysis. This could be explained by activation of hematopoietic stem cells in response to DPP4 inhibitors which control inflammatory and immune response pathogenesis of T2DM [13], in association with  $\beta$ -cell regeneration [14].

The limitation of the study is that diabetic patients receiving new therapeutic agents such as sodium glucose transporter 2 inhibitors (SGLT2 inhibitors) and glucagon like peptide 1 agonists (GLP1 agonists) were not discussed in our study. This is due to lack of availability during the study period as these agents have become part of standard of care only recently in Egypt after we have started the study. Moreover most of the Egyptian diabetic patients are poor and can't afford such expensive drugs.

## Conclusion

Type 2 DM remission is possible without surgical intervention. Factors influencing medical remission include; female sex, young age < 50 years, short duration since diagnosis of DM < 5.2 years, baseline HbA1c < 8.5% and HDL > 45.6 mg/dl. Therefore, our

mission in type 2 DM patients with such criteria is to achieve diabetes remission as a treatment target by intensive life style modification in association with daily metformin and vildagliptin.

#### Recommendation

A double blinded clinical trial is needed to assess factors associated with remission of T2DM using more data including new therapeutic agents for diabetes such as SGLT2 inhibitors and GLP1 agonists.

# Author disclosures

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## Authors' contributions

Authors have an equal contribution in this research.

#### **Declaration of Competing Interest**

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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