

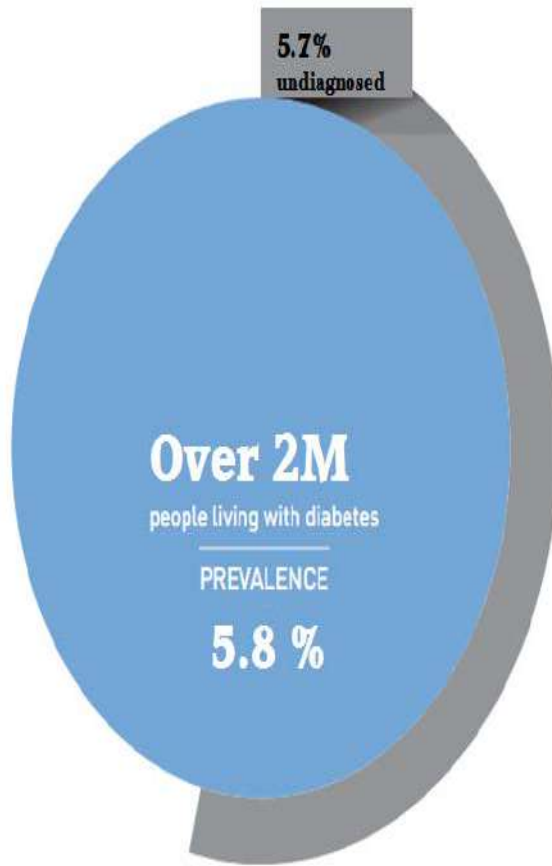
Comprehensive Management of Diabetes

PROFESSOR TINT SWE LATT

PRESIDENT

MYANMAR DIABETES ASSOCIATION

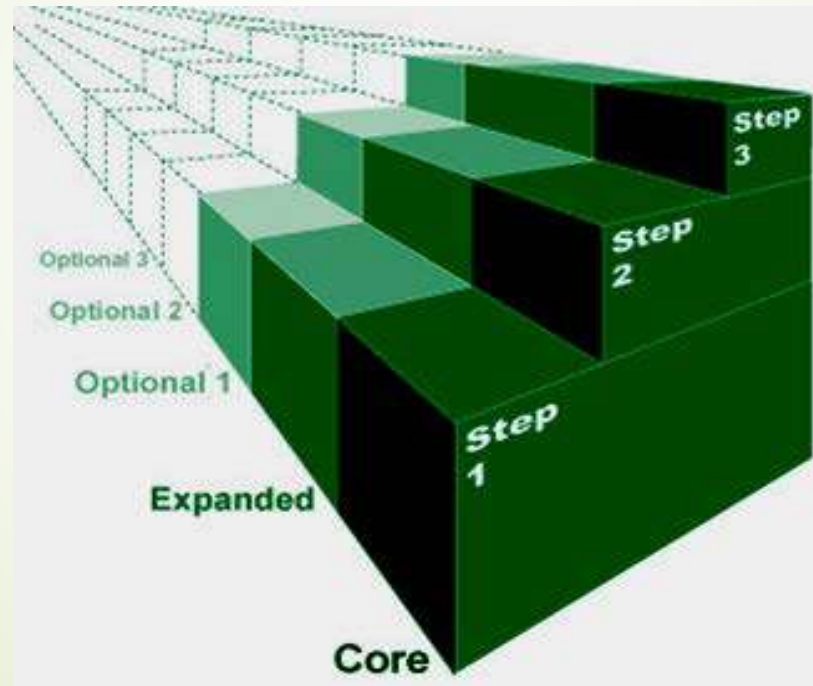
Magnitude of problem of Diabetes in MYANMAR



Diabetes in Myanmar [20-79 years]	2014
Adult Population [1000`s]	35,436
Number of People With Diabetes [1000`s]	2,051.0
Regional Prevalence [%]	5.8
Comparative Prevalence [%]	6.1
Undiagnosed Cases [1000`s]	1,292.9
Total diabetes-related deaths	60,518
Deaths under the age of 60 [%]	60.7
Total diabetes-related health expenditure [USD]	30.9

STEPS approach to Surveillance

**A framework for risk factors, diseases etc
sensitive to resource availability**





Health Behaviours

- Tobacco use
- Alcohol consumption (heavy)
- Alcohol abstainers
- Physical inactivity
- Fruit/vegetable intake




Physiological Factors

- Obesity/overweight (& mean BMI)
- Raised blood pressure (& mean systolic blood pressure)
- Raised lipid (& mean total cholesterol)
- Diabetes (& mean blood glucose)



Disease outcomes

- Heart disease
- Stroke
- Cancers
- Diabetes

- 
- Study area : 15 townships of Yangon Division (both rural and urban areas)
- Study population : age 25 – 74 years, both sex
- Sample size : 5000 subjects
- Study period : July 2003 to June 2004
- Methods : STEPs Instrument for NCD Risk Factors (Core and Expanded version 1.3)

Diabetes Mellitus detectable by OGTT : Men and Women

Age	Men			Women		
	N	Diabetes n (%) SE	Others n (%) SE	N	Diabetes n (%) SE	Others n (%) SE
25-34	320	2 (0.87),0.68	318 (99.13),0.68	387	8 (2.28), 0.80	379 (97.72), 0.80
35-44	389	29 (8.40),1.34	360 (91.60),1.34	555	40 (7.14), 1.08	515 (92.86), 1.08
45-54	492	62 (15.20),1.94	430 (84.80),1.94	638	96 (15.81), 1.56	542 (84.19), 1.56
55-64	442	59 (14.72),2.61	383 (85.28),2.61	538	101 (19.79), 2.76	437 (80.21), 2.76
65-74	351	47 (15.03),1.40	304 (84.97),1.34	333	45 (15.96), 1.64	288 (84.04), 1.64
25-74	1994	199 (11.51),1.11	1795 (88.49),1.11	2451	290 (12.64), 1.07	2161 (87.36), 1.07

National Prevalence of Diabetes and Prediabetes in Myanmar (2014)

Tint-Swe Latt¹,

Ko-Ko Zaw², Ko Ko^{3,5}, Moh-Moh Hlaing², Mya Ohnmar²,

Ei-Sandar Oo⁴, Kyaw-Myint-Myat Thein⁵

¹Myanmar Diabetes Association, ²Department of Medical Research,

*³Department of Medical Services, ⁴Department of Health Professional
Development*

and Management, ⁵Diabetes Project, University of Medicine (2), Yangon

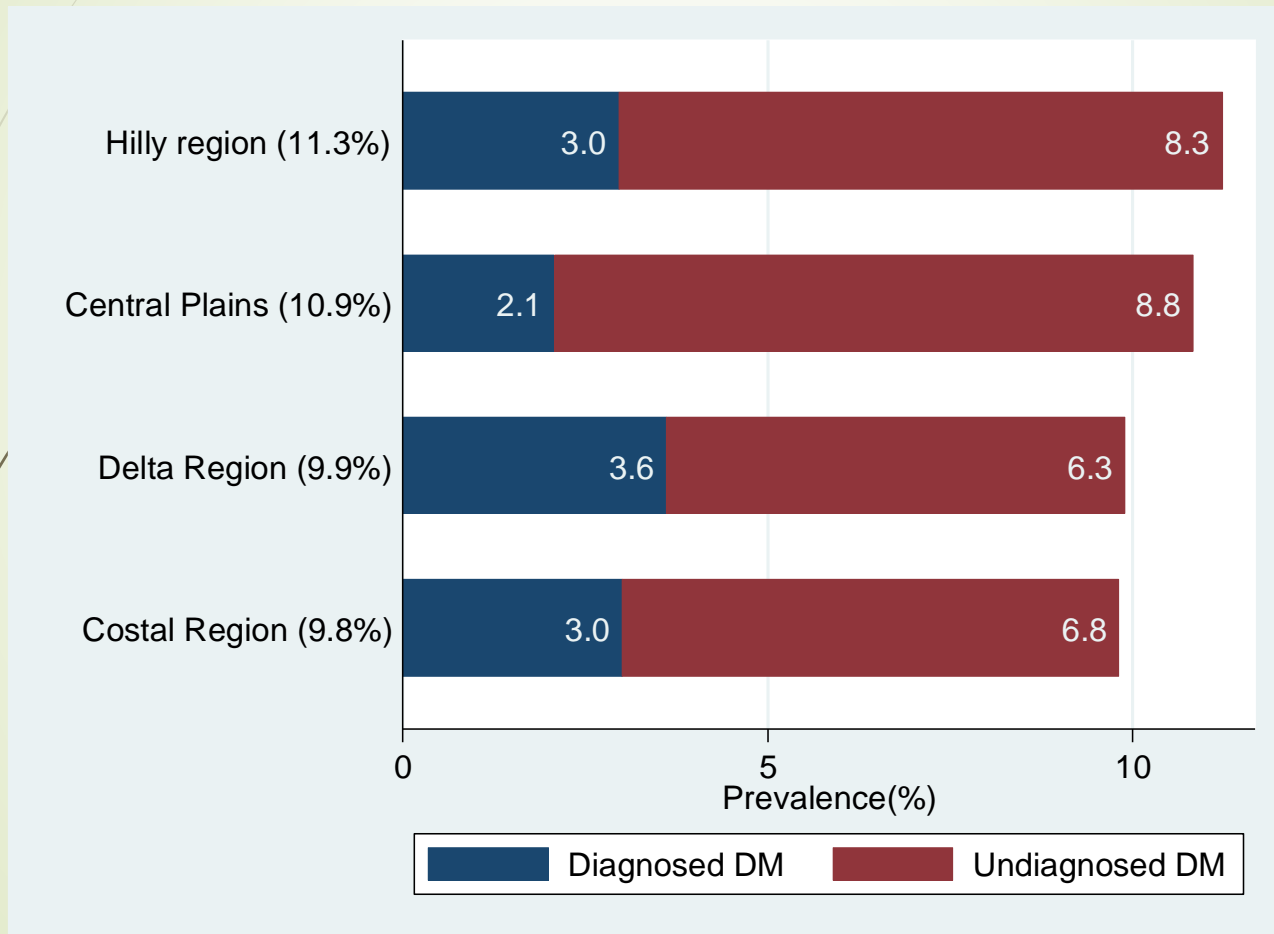
Prevalence of diabetes, Myanmar, 2014

Sex	Age in years	Pop.	Diagnosed diabetes		Undiagnosed diabetes		Total diabetes	
			%	95%CI	%	95%CI	%	95%CI
Men	25-34	616	0.1	[0.0,0.4]	3.5	[1.5,8.0]	3.6	[1.6,8.1]
	35-44	738	1.7	[0.9,3.1]	5.9	[3.7,9.2]	7.6	[5.0,11.2]
	45-54	832	4.8	[2.5,8.9]	7	[5.2,9.4]	11.8	[8.7,15.9]
	55-64	707	3.5	[2.0,6.0]	16	[10.4,23.6]	19.5	[13.4,27.4]
	25-64	2,893	2.1	[1.4,3.1]	7	[5.3,9.1]	9.1	[6.9,11.8]
Women	25-34	1,099	0.4	[0.2,1.0]	4.1	[2.7,6.2]	4.5	[3.1,6.6]
	35-44	1,559	2.3	[1.4,3.7]	8.7	[5.6,13.3]	11	[7.9,15.0]
	45-54	1,537	5.3	[3.6,7.7]	10	[8.1,13.1]	15.6	[12.6,19.1]
	55-64	1,236	11	[7.5,15.1]	13	[8.6,19.2]	23.7	[19.2,28.8]
	25-64	5,431	3.7	[2.6,5.1]	8.2	[6.0,11.1]	11.8	[9.6,14.6]
Both sexes	25-34	1,715	0.2	[0.1,0.5]	3.8	[2.4,6.0]	4.1	[2.6,6.2]
	35-44	2,297	2	[1.3,3.0]	7.3	[5.0,10.4]	9.3	[6.8,12.4]
	45-54	2,369	5	[3.6,7.0]	8.7	[7.2,10.4]	13.7	[11.3,16.6]
	55-64	1,943	7.1	[5.0,10.1]	15	[9.8,20.9]	21.6	[16.5,27.7]
	25-64	8,324	2.9	[2.2,3.8]	7.6	[5.7,9.9]	10.5	[8.3,13.1]

Urban Rural Difference in Diabetes Prevalence

	<u>Prevalence (%)</u>	<u>95% CI</u>
➡ Urban pop.	14	12 to 13
➡ Rural pop.	9	7 to 12

Diabetes prevalence by geographical regions

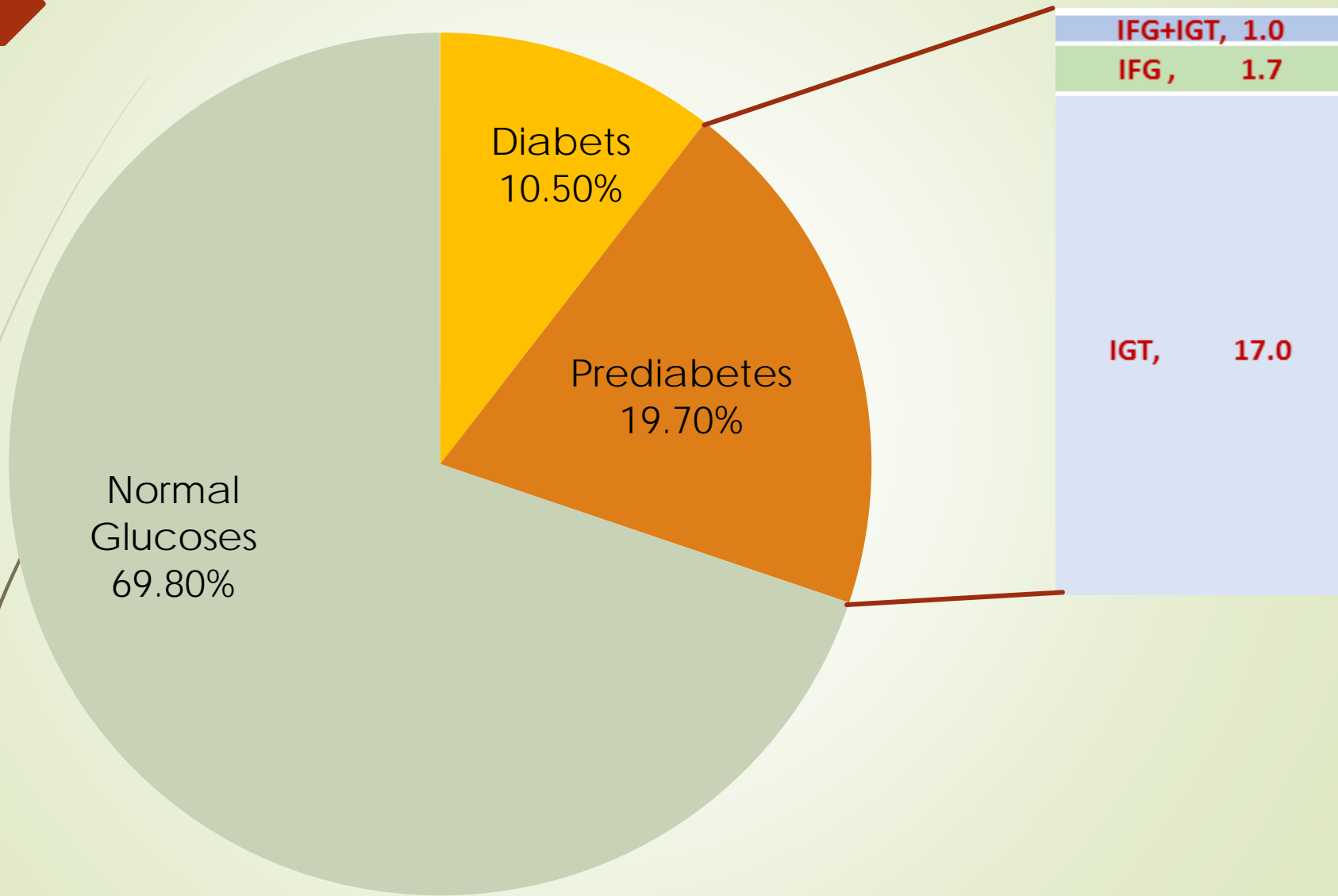


Relative Risk Ratios (RRR) for pre-diabetes and diabetes from multinomial logistic regression

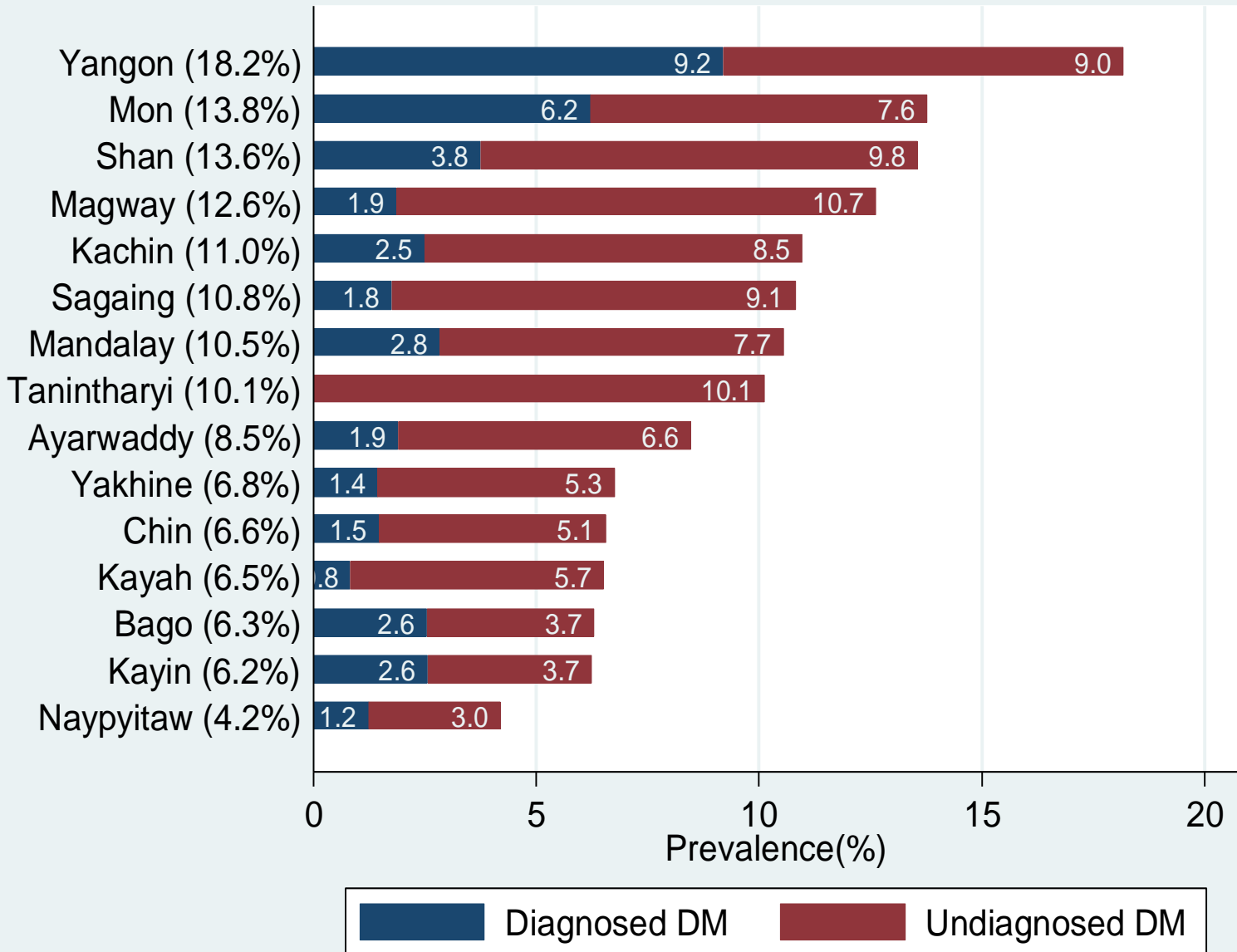
Variable	Prediabetes		Diabetes	
	RRR	95%CI	RRR	95%CI
Sex				
<i>Male (Ref.)</i>	1.0		1.0	
<i>Female</i>	1.5***	[1.2,1.8]	1.3	[1.0,1.8]
Age (in years)				
<i>25-34 (Ref.)</i>	1.0		1.0	
<i>35-44</i>	1.7**	[1.3,2.2]	2.5***	[1.6,4.0]
<i>45-54</i>	2.0***	[1.5,2.7]	3.4***	[2.1,5.3]
<i>55-64</i>	2.7***	[2.0,3.8]	6.4***	[3.9,10.3]
Urban-rural				
<i>Rural (Ref.)</i>	1.0		1.0	
<i>Urban</i>	1.1	[0.8,1.3]	1.5***	[1.3,2.2]
Central obesity				
<i>Absent (Ref.)</i>	1.0		1.0	
<i>Present</i>	1.4**	[1.1,1.8]	3.1**	[2.4,4.2]
Triglycerides(TG)				
<i>Low (Ref.)</i>	1.0		1.0	
<i>High</i>	1.3*	[1.0,1.6]	1.7***	[1.3,2.2]
Observations	6555			
Wald χ^2 (d.f=14)	327			
P value	<0.001			

* p<0.05, ** p<0.01, ***p<0.001

Prevalence of diabetes



Prevalence according to regions



Different level of prevention of Diabetes

1. Primordial prevention(the whole population)
2. Primary prevention (high risk persons)
3. Secondary prevention (diabetes without complications)
4. Tertiary prevention.(Diabetes with complications)

- All medical doctors should change their attitude toward prevention of diabetes
- It is mandatory for all health care providers
- It is prevention, not, treatment of diabetes

INTENSITY STRATIFIED BY BURDEN OF OBESITY AND RELATED COMPLICATIONS

Nutrition	<ul style="list-style-type: none"> • Maintain optimal weight • Calorie restriction • Plant-based diet; high polyunsaturated and monounsaturated fatty acids • Avoid <i>trans</i> fatty acids; limit saturated fatty acids 	+	<ul style="list-style-type: none"> • Structured counseling • Meal replacement 		
Physical Activity	<ul style="list-style-type: none"> • 150 min/week moderate exertion (eg. walking, stair climbing) • Strength training • Increase as tolerated 	+	<ul style="list-style-type: none"> • Structured program 	+	<ul style="list-style-type: none"> • Medical evaluation/clearance • Medical supervision
Sleep	<ul style="list-style-type: none"> • About 7 hours per night 	+	<ul style="list-style-type: none"> • Screen for obstructive sleep apnea 		
Behavioral Support	<ul style="list-style-type: none"> • Community engagement • Screen for mood disorders 	+	<ul style="list-style-type: none"> • Refer to mental healthcare professional • Behavioral therapy 		
Smoking Cessation	<ul style="list-style-type: none"> • No tobacco products 	+	<ul style="list-style-type: none"> • Structured programs 		

American Association of Clinical Endocrinologists Healthful Eating Recommendations for Patients with Diabetes Mellitus

Topic	Recommendation
General eating habits	<p>Regular meals and snacks; avoid fasting to lose weight</p> <p>Plant-based diet (high in fiber, low calories/glycemic index, and high in phytochemicals/antioxidants)</p> <p>Understand Nutrition Facts Label information</p> <p>Incorporate beliefs and culture into discussions</p> <p>Informal physician-patient discussions</p> <p>Use mild cooking techniques instead of high-heat cooking</p>

American Association of Clinical Endocrinologists Healthful Eating Recommendations for Patients with Diabetes Mellitus

Topic	Recommendation
Carbo- hydrate	<p>Explain the 3 types of carbohydrates: sugars, starch, and fiber and the effects on health for each type</p> <p>Specify healthful carbohydrates (fresh fruits and vegetables, pulses, whole grains); target 7-10 servings per day</p> <p>Lower-glycemic index foods may facilitate glycemic control (glycemic index score <55 out of 100: multigrain bread, pumpernickel bread, whole oats, legumes, apple, lentils, chickpeas, mango, yams, brown rice), but there is insufficient evidence to support a formal recommendation to educate patients that sugars have both positive and negative health effects</p>

American Association of Clinical Endocrinologists Healthful Eating Recommendations for Patients with Diabetes Mellitus

Topic	Recommendation
Fat	<p>Specify healthful fats (low mercury/contaminant-containing nuts, avocado, certain plant oils fish)</p> <p>Limit saturated fats (butter, fatty red meats, tropical plant oils, fast foods) and <i>trans</i> fat; no- or low-fat dairy products</p>
Protein	<p>Consume protein in foods preferably with low saturated fats (fish, egg whites, beans); there is no need to avoid animal protein</p> <p>Avoid or limit processed meats</p>

American Association of Clinical Endocrinologists Healthful Eating Recommendations for Patients with Diabetes Mellitus

Topic	Recommendation
Micronutrients	<p>Routine supplementation is not necessary</p> <p>Specifically, chromium, vanadium, magnesium, vitamins A, C, and E, and CoQ10 are not recommended for glycemic control</p> <p>Supplementation to avoid insufficiency or deficiency in at-risk patients</p> <p>A healthful eating meal plan can generally provide sufficient micronutrients</p>

Abbreviations: BEL, best evidence level; CPG, clinical practice guideline; EL, evidence level; MNRCT, meta-analysis of nonrandomized prospective or case-controlled trials; NE, no evidence (theory, opinion, consensus, review, or preclinical study); PCS, prospective cohort study; RCT, randomized controlled trial.

STEP 1

EVALUATION FOR COMPLICATIONS AND STAGING

CARDIOMETABOLIC DISEASE | BIOMECHANICAL COMPLICATIONS

NO COMPLICATIONS

BMI ≥ 25

COMPLICATIONS

BMI 25–26.9

BMI ≥ 27 : Stage Severity of Complications

MILD TO MODERATE

SEVERE

STEP 2

SELECT:

Therapeutic targets for improvement in complications

+

Treatment modality

+

Treatment intensity based on staging

Lifestyle Therapy:

Physician/RD counseling, web/remote program, structured multidisciplinary program

Medical Therapy (BMI ≥ 27):

Phentermine, orlistat, lorcaserin, phentermine/topiramate ER, naltrexone/bupropion, liraglutide 3 mg

Surgical Therapy (BMI ≥ 35):

Gastric banding, sleeve, or bypass

STEP 3

If therapeutic targets for complications not met, intensify lifestyle, medical, and/or surgical treatment modalities for greater weight loss.

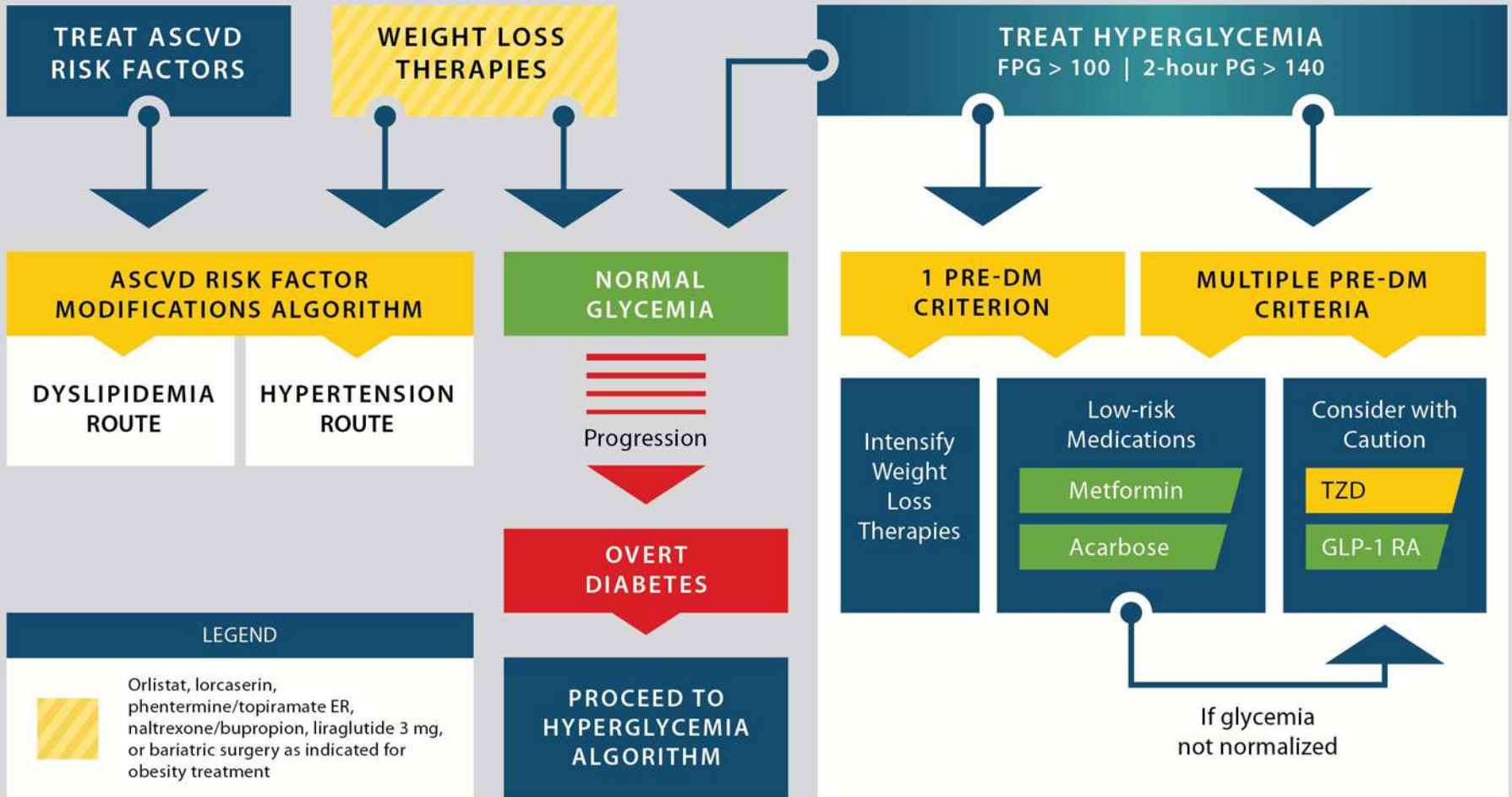


PREDIABETES ALGORITHM



IFG (100-125) | IGT (140-199) | METABOLIC SYNDROME (NCEP 2001)

LIFESTYLE THERAPY (Including Medically Assisted Weight Loss)





INDIVIDUALIZE GOALS

$A1C \leq 6.5\%$

For patients without
concurrent serious
illness and at low
hypoglycemic risk

$A1C > 6.5\%$

For patients with
concurrent serious
illness and at risk
for hypoglycemia

Table 5.1—Mean glucose levels for specified A1C levels (24,28)

A1C % (mmol/mol)	Mean plasma glucose*		Mean fasting glucose		Mean premeal glucose		Mean postmeal glucose		Mean bedtime glucose	
	mg/dL	mmol/L	mg/dL	mmol/L	mg/dL	mmol/L	mg/dL	mmol/L	mg/dL	mmol/L
6 (42)	126	7.0								
<6.5 (48)			122	6.8	118	6.5	144	8.0	136	7.5
6.5–6.99 (48–53)			142	7.9	139	7.7	164	9.1	153	8.5
7 (53)	154	8.6								
>7.0–7.49 (53–58)			152	8.4	152	8.4	176	9.8	177	9.8
7.5–7.99 (58–64)			167	9.3	155	8.6	189	10.5	175	9.7
8 (64)	183	10.2								
>8.0–8.5 (64–69)			178	9.9	179	9.9	206	11.4	222	12.3
9 (75)	212	11.8								
10 (86)	240	13.4								
11 (97)	269	14.9								
12 (108)	298	16.5								

A calculator for converting A1C results into eAG, in either mg/dL or mmol/L, is available at <http://professional.diabetes.org/eAG>.

*These estimates are based on ADAG data of ~2,700 glucose measurements over 3 months per A1C measurement in 507 adults with type 1, type 2, and no diabetes. The correlation between A1C and average glucose was 0.92 (28).

HbA1C and Estimated Average Glucose(eAVG)

HbA1C (%)	eAG (mg/dL)	eAG (mmol/l)
5	97	5.4
6	126	7.0
7	154	8.6
8	183	10.2
9	212	11.8
10	240	13.4
11	269	14.9
12	298	16.5

www.ngsp.org/Harmonizing HbA1C testing/accessed on 16th September, 2016

LIFESTYLE THERAPY (Including Medically Assisted Weight Loss)

Entry A1C < 7.5%

Entry A1C ≥ 7.5%

Entry A1C > 9.0%

MONOTHERAPY*

- ✓ Metformin
- ✓ GLP-1 RA
- ✓ SGLT-2i
- ✓ DPP-4i
- ⚠ TZD
- ✓ AGi
- ⚠ SU/GLN

If not at goal in 3 months proceed to Dual Therapy

DUAL THERAPY*

MET

or other 1st-line agent

+

- ✓ GLP-1 RA
- ✓ SGLT-2i
- ✓ DPP-4i
- ⚠ TZD
- ⚠ Basal Insulin
- ✓ Colesevelam
- ✓ Bromocriptine QR
- ✓ AGi
- ⚠ SU/GLN

If not at goal in 3 months proceed to Triple Therapy

TRIPLE THERAPY*

MET

or other 1st-line agent + 2nd-line agent

+

- ✓ GLP-1 RA
- ✓ SGLT-2i
- ⚠ TZD
- ⚠ Basal insulin
- ✓ DPP-4i
- ✓ Colesevelam
- ✓ Bromocriptine QR
- ✓ AGi
- ⚠ SU/GLN

If not at goal in 3 months proceed to or intensify insulin therapy

SYMPTOMS

NO YES

- | | |
|----------------|------------------------|
| DUAL Therapy | INSULIN ± Other Agents |
| OR | |
| TRIPLE Therapy | |

ADD OR INTENSIFY INSULIN

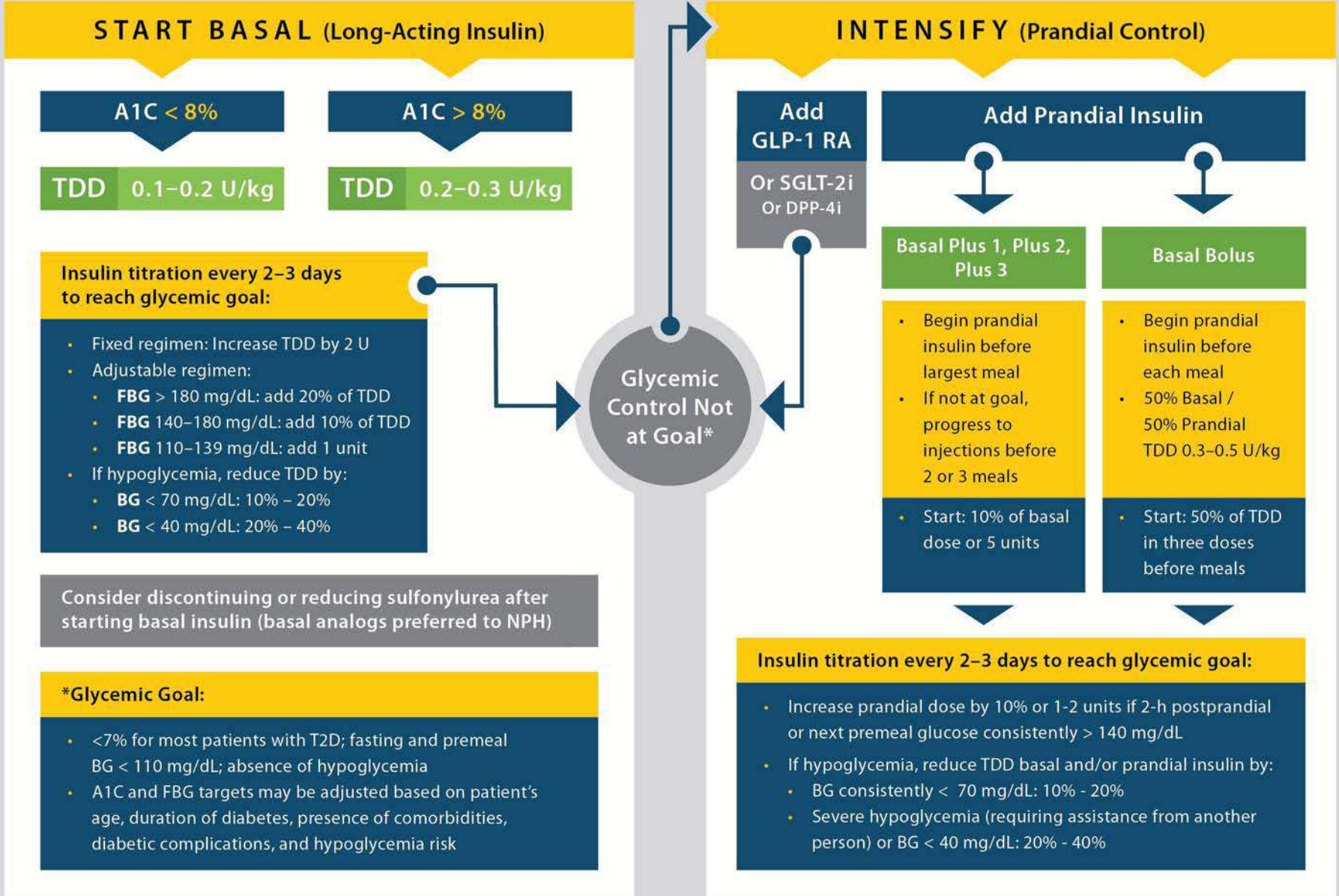
Refer to Insulin Algorithm

LEGEND

- ✓ Few adverse events and/or possible benefits
- ⚠ Use with caution

* Order of medications represents a suggested hierarchy of usage; length of line reflects strength of recommendation

PROGRESSION OF DISEASE



START BASAL (Long-Acting Insulin)

A1C < 8%

A1C > 8%

TDD 0.1–0.2 U/kg

TDD 0.2–0.3 U/kg

Insulin titration every 2–3 days to reach glycemic goal:

- Fixed regimen: Increase TDD by 2 U
- Adjustable regimen:
 - FBG > 180 mg/dL: add 20% of TDD
 - FBG 140–180 mg/dL: add 10% of TDD
 - FBG 110–139 mg/dL: add 1 unit
- If hypoglycemia, reduce TDD by:
 - BG < 70 mg/dL: 10% – 20%
 - BG < 40 mg/dL: 20% – 40%

Consider discontinuing or reducing sulfonylurea after starting basal insulin (basal analogs preferred to NPH)

*Glycemic Goal:

- <7% for most patients with T2D; fasting and premeal BG < 110 mg/dL; absence of hypoglycemia
- A1C and FBG targets may be adjusted based on patient’s age, duration of diabetes, presence of comorbidities, diabetic complications, and hypoglycemia risk

INTENSIFY (Prandial Control)

Add GLP-1 RA
Or SGLT-2i
Or DPP-4i

Add Prandial Insulin

Basal Plus 1, Plus 2, Plus 3

Basal Bolus

- Begin prandial insulin before largest meal
- If not at goal, progress to injections before 2 or 3 meals

Start: 10% of basal dose or 5 units

- Begin prandial insulin before each meal
- 50% Basal / 50% Prandial TDD 0.3–0.5 U/kg

Start: 50% of TDD in three doses before meals

Insulin titration every 2–3 days to reach glycemic goal:

- Increase prandial dose by 10% or 1–2 units if 2-h postprandial or next premeal glucose consistently > 140 mg/dL
- If hypoglycemia, reduce TDD basal and/or prandial insulin by:
 - BG consistently < 70 mg/dL: 10% - 20%
 - Severe hypoglycemia (requiring assistance from another person) or BG < 40 mg/dL: 20% - 40%

DYSLIPIDEMIA

HYPERTENSION

LIFESTYLE THERAPY (Including Medically Assisted Weight Loss)

LIPID PANEL: Assess ASCVD Risk

STATIN THERAPY

If TG > 500 mg/dL, fibrates, Rx-grade omega-3 fatty acids, niacin

If statin-intolerant

Try alternate statin, lower statin dose or frequency, or add nonstatin LDL-C-lowering therapies

Repeat lipid panel; assess adequacy, tolerance of therapy

Intensify therapies to attain goals according to risk levels

RISK LEVELS	HIGH	DM but no other major risk and/or age <40	VERY HIGH	DM + major ASCVD risk(s) (HTN, Fam Hx, low HDL-C, smoking) or ASCVD*
	DESIRABLE LEVELS		DESIRABLE LEVELS	
LDL-C (mg/dL)	<100		<70	
Non-HDL-C (mg/dL)	<130		<100	
TG (mg/dL)	<150		<150	
TC/HDL-C	<3.5		<3.0	
Apo B (mg/dL)	<90		<80	
LDL-P (nmol/L)	<1200		<1000	

IF NOT AT DESIRABLE LEVELS:

Intensify lifestyle therapy (weight loss, physical activity, dietary changes) and glycemic control; consider additional therapy

TO LOWER LDL-C:
TO LOWER Non-HDL-C, TG:
TO LOWER Apo B, LDL-P:
TO LOWER LDL-C in FH:**

Intensify statin, add ezetimibe, PCSK9i, colesovelam, or niacin
 Intensify statin and/or add Rx-grade OM3 fatty acid, fibrate, and/or niacin
 Intensify statin and/or add ezetimibe, PCSK9i, colesovelam, and/or niacin
 Statin + PCSK9i

Assess adequacy & tolerance of therapy with focused laboratory evaluations and patient follow-up

* EVEN MORE INTENSIVE THERAPY MIGHT BE WARRANTED ** FAMILIAL HYPERCHOLESTEROLEMIA

GOAL: SYSTOLIC <130, DIASTOLIC <80 mm Hg

ACEi or ARB

For initial blood pressure >150/100 mm Hg:
DUAL THERAPY

ACEi or ARB

- + Calcium Channel Blocker ✓
- + β-blocker ✓
- + Thiazide ✓

If not at goal (2–3 months)

Add calcium channel blocker, β-blocker or thiazide diuretic

If not at goal (2–3 months)

Add next agent from the above group, repeat

If not at goal (2–3 months)

Additional choices (α-blockers, central agents, vasodilators, aldosterone antagonist)

Achievement of target blood pressure is critical



PROFILES OF ANTIDIABETIC MEDICATIONS



	MET	GLP-1 RA	SGLT-2i	DPP-4i	AGi	TZD (moderate dose)	SU GLN	COLSVL	BCR-QR	INSULIN	PRAML
HYPO	Neutral	Neutral	Neutral	Neutral	Neutral	Neutral	Moderate/ Severe Mild	Neutral	Neutral	Moderate to Severe	Neutral
WEIGHT	Slight Loss	Loss	Loss	Neutral	Neutral	Gain	Gain	Neutral	Neutral	Gain	Loss
RENAL/ GU	Contra- indicated CKD Stage 3B,4,5	Exenatide Not Indicated CrCl < 30	Not Effective with eGFR < 45 Genital Mycotic Infections	Dose Adjustment Necessary (Except Linagliptin)	Neutral	Neutral	More Hypo Risk	Neutral	Neutral	More Hypo Risk	Neutral
GI Sx	Moderate	Moderate	Neutral	Neutral	Moderate	Neutral	Neutral	Mild	Moderate	Neutral	Moderate
CHF CARDIAC ASCVD	Neutral Benefit	Neutral	Possible Benefit	Neutral	Neutral	Moderate Neutral	Neutral ?	Neutral	Neutral Safe	Neutral	Neutral
BONE	Neutral	Neutral	Neutral	Neutral	Neutral	Moderate Fracture Risk	Neutral	Neutral	Neutral	Neutral	Neutral

■ Few adverse events or possible benefits
 ■ Use with caution
 ■ Likelihood of adverse effects
 ■ ? Uncertain effect



PRINCIPLES OF THE AAACE/ACE COMPREHENSIVE TYPE 2 DIABETES MANAGEMENT ALGORITHM



1. Lifestyle therapy, including medically supervised weight loss, is key to managing type 2 diabetes.
2. The A1C target must be individualized.
3. Glycemic control targets include fasting and postprandial glucoses.
4. The choice of therapies must be individualized on basis of patient characteristics, impact of net cost to patient, formulary restrictions, personal preferences, etc.
5. Minimizing risk of hypoglycemia is a priority.
6. Minimizing risk of weight gain is a priority.
7. Initial acquisition cost of medications is only a part of the total cost of care which includes monitoring requirements, risk of hypoglycemia, weight gain, safety, etc.
8. This algorithm stratifies choice of therapies based on initial A1C.
9. Combination therapy is usually required and should involve agents with complementary actions.
10. Comprehensive management includes lipid and blood pressure therapies and related comorbidities.
11. Therapy must be evaluated frequently until stable (e.g., every 3 months) and then less often.
12. The therapeutic regimen should be as simple as possible to optimize adherence.
13. This algorithm includes every FDA-approved class of medications for diabetes.

Approach to the management of hyperglycemia

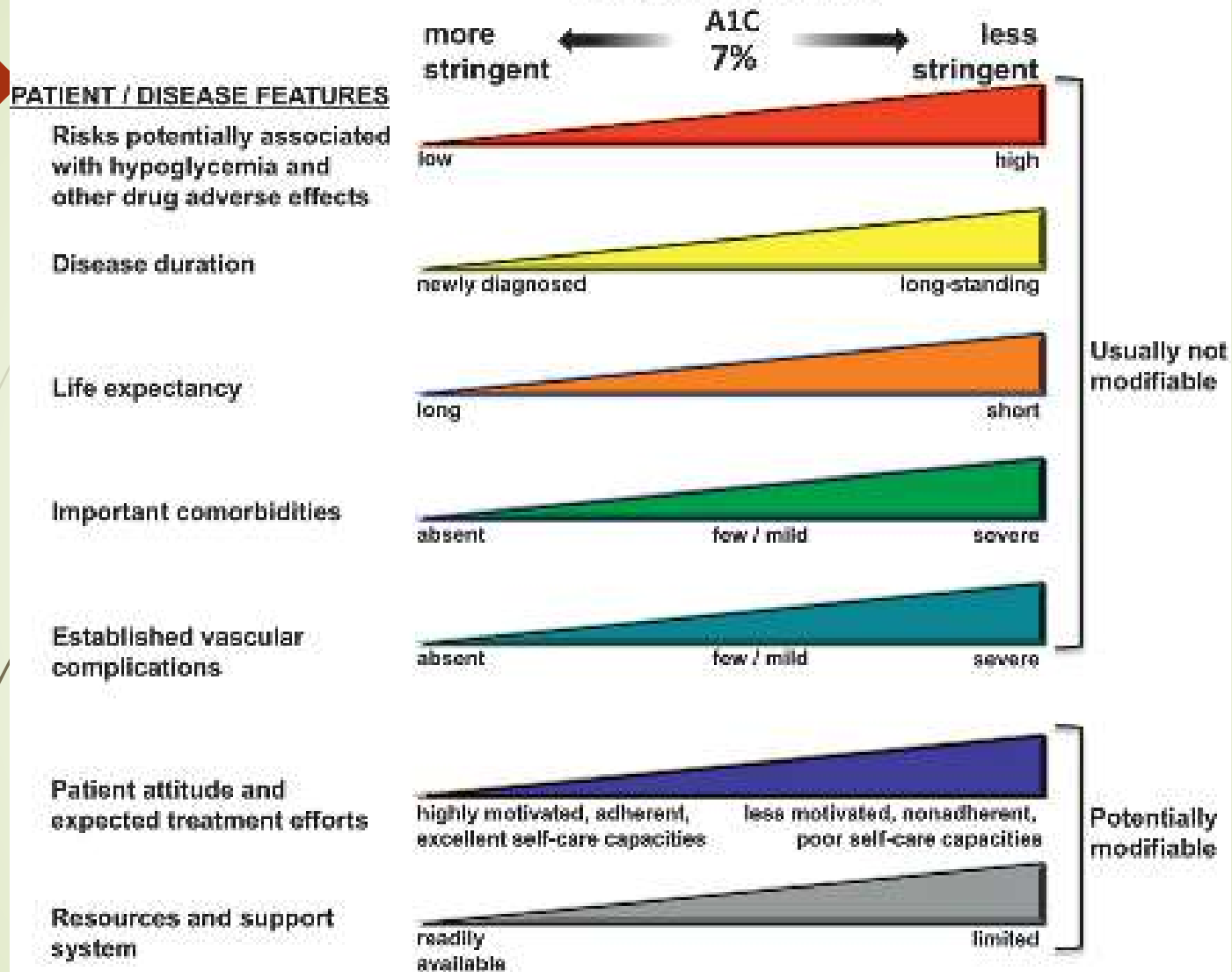


Figure —Depicted are patient and disease factors used to determine optimal A1C targets. Characteristics and predicaments toward the left justify more stringent efforts to lower A1C; those toward the right suggest less stringent efforts. Adapted with permission from Inzucchi et al. (53).

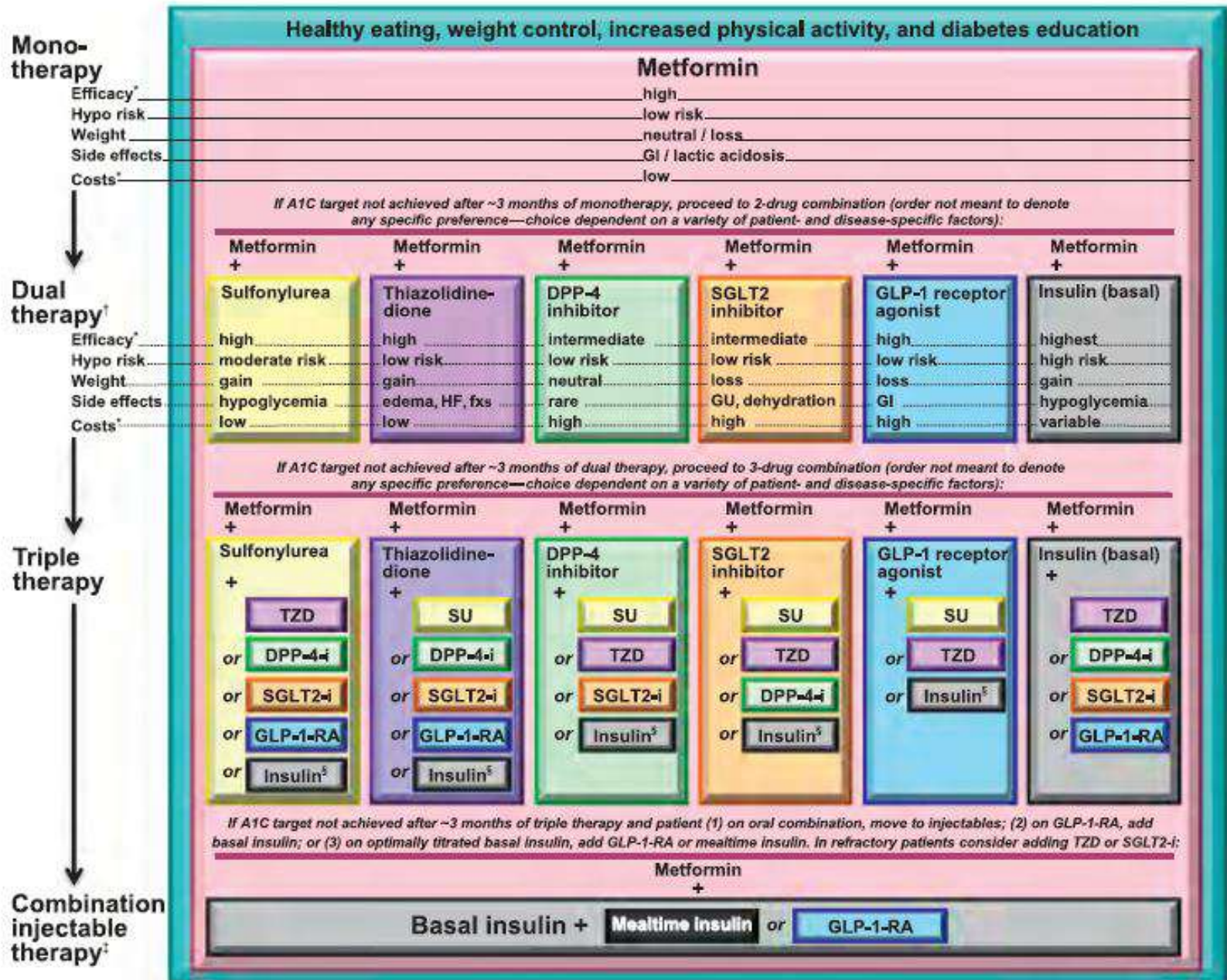


Figure — Antihyperglycemic therapy in type 2 diabetes: general recommendations .

Diabetes Prevention Programs with Lifestyle Intervention:

	N	Design	Diagnostic test	Mean age \pm SD	BMI, kg/m ² (SD)	Follow-up (yrs)
U.S.DPP 1996 – 2001	3234	RCT 27 sites	IGT & FPG >95 mg/dl	50.6 \pm 10.7	34 (6.7)	2.8
Finnish DPS 1993 – 2003	522	RCT 5 sites	IGT	55 \pm 7	31.3 (4.6)	3.2
Da Qing China 1986 - 1992	577	RCT, cluster 33 sites	IGT	45 \pm 9.1	25.8 (3.8)	4.51 – 4.62
Swedish Malmo 1974 - 1985	415	Nonrando- mized 1 site	IGT	45.9	26.6 (3.1) Intervention; 26.7 (4.0) Control	5.0
Indian DPP 2001 – 2002	531	RCT	IGT	35 – 55	25.8	2.5
Japan Prevention Trial	458 (males)	RCT	IGT	51.5	23.8	3.64

	Interventions	Goal : weight	Goal : diet	Exercise
U.S.DPP 1996 – 2001	<ul style="list-style-type: none"> Lifestyle Metformin Troglitazone 	7% weight loss	500 – 1,000 kcal/day <25% kcal from fat	150 minutes/week
Finnish DPS 1993 - 2003	Lifestyle	>5% weight loss	<30% kcal from fat <10% saturated fat >fiber /1,000 kcal	30 minutes moderate intensity daily
Da Qing China 1986 - 1992	<ul style="list-style-type: none"> Diet alone Exercise alone Diet and exercise 	BMI = 23 kg/m ²	25-30 kcal/kg 55-65% carbohydrate 10-15% protein 25-30% fat lower cal if BMI >25	Increase LTPA by 30 minutes of light or 20 minutes of moderate intensity
Swedish Malmo 1974 - 1985	Lifestyle	Not stated	Decreased simple carbohydrates and saturated fats, substitute PUFAs, increase complex carbohydrates Reduce calories if obese	Not stated
Indian DPP 2001 – 2002	Lifestyle Metformin	Not stated	Portion control, decrease fat, high fruit, vegetable fiber	
Japan Prevention Trial	Lifestyle	BMI < 24 kg/m ² for control < 22 kg/m ² for intervention	Portion control, decrease fat, high fruit, vegetable fiber	

	Tracking	Counseling	Risk reduction intervention vs. control
U.S.DPP 1996 – 2001	Goal setting, self-monitoring, food records, exercise logs, weight	16 core curriculum sessions, individual and group first 24 weeks, lifestyle coaches	(3 years) 58% lifestyle
Finnish DPS 1993 - 2003	Goal setting, self-monitoring, food records, exercise logs, weight	Quarterly visits with review of food records and 7 visits per year with nutritionist, 3 in first 6 weeks	(6 years) 58% lifestyle
Da Qing China 1986 - 1992	Goal setting, no self-monitoring or logging	Individual counseling quarterly and group counseling once weekly for 1 month and then quarterly	(6 years) 31% diet 46% exercise 42% diet and exercise
Swedish Malmo 1974 - 1985		Small-group or individual counseling monthly for 6 months, 60-minutes activity sessions twice/week under the glucose of a physiotherapist	(6 years) 63% lifestyle
Indian DPP 2001 – 2002		Individual counseling session at baseline and every 6 months	(3 years) 45.2% lifestyle
Japan Prevention Trial		Detailed instruction on lifestyle every 3-4 months	(4 years) 67.4% lifestyle

	Long-term mean follow-up	Follow-up	Diet and exercise	Outcomes
U.S.DPP 1996 – 2001	(7 years) 34% lifestyle 18% metformin	Every 2 months face-to-face group or individual, phone call in between	Twice-weekly session, including personal training, self-monitoring weight, fat, calories	(3 years) 28.9% controls 14.4% lifestyle 21.7% metformin
Finnish DPS 1993 - 2003	(7 years) 43%	Quarterly	Supervised exercise offered free	(4 years) 23% control 11% lifestyle
Da Qing China 1986 - 1992	(14 years) 14%	One group session quarterly		(6 years) 67.7% control 43.8% diet 41.1% exercise 46.0% diet and exercise
Swedish Malmö 1974 - 1985	N/A			(6 years) 28.6% control 10.6% lifestyle
Indian DPP 2001 – 2002	N/A	Every 6 months, phone contact after 2 weeks then monthly		(3 years) 55% control 39.3% lifestyle 40.5% metformin 39.5% lifestyle and metformin

TABLE 4. SUCCESS IN ACHIEVING THE GOALS OF THE INTERVENTION BY ONE YEAR, ACCORDING TO TREATMENT GROUP.*

GOAL	INTERVENTION GROUP	CONTROL GROUP	P VALUE†
	% of subjects		
Weight reduction >5%	43	13	0.001
Fat intake <30% of energy intake	47	26	0.001
Saturated-fat intake <10% of energy intake	26	11	0.001
Fiber intake \geq 15 g/1000 kcal	25	12	0.001
Exercise >4 hr/wk‡	86	71	0.001

*Nutrient intakes were calculated from three-day food records.

†P values were determined by the chi-square test for the difference between the groups.

‡Exercise frequency was reported by the subjects who chose one of the four categories described in Table 3. The goal identified here was a frequency in category 2 or higher.

Diabetes Prevention Trials with Pharmacotherapy: Design and Outcomes

	Design	RRR	Adverse effects
DPP	RCT, Metformin N = 3234	Outcome = 31% Long term = 18%	Mild diarrhea
IDPP	RCT, Metformin N = 532	Outcome = 25.4% Long term = N/A	Not reported
DREAM	RCT, Rosiglitazone Ramipril N = 5269	Outcome = No significant risk reduction Long term = N/A	No significant difference for Ramipril
ACT NOW	RCT, Pioglitazone N = 602	Outcome = 72% Long term = N/A	Edema (12.9 vs. 6.7%)
IDPP	RCT, Pioglitazone N = 401	Outcome = No significant risk reduction Long term = N/A	
STOP- NIDDM	RCT, Acarbose N = 1429	Outcome = 25% Long term = N/A	Flatulence (15.9 vs. 6.1%) Bloating (9.5 vs. 2.3%)

Diabetes Prevention Trials with Pharmacotherapy: Design and Outcomes

	Design	RRR	Adverse effects
Kawamori	RCT, Voglibose N = 1780	Outcome = 40.5% Long term = N/A	Mild gastro-intestinal symptoms
Glipizide	RCT, Glipizide N = 37	Outcome = 80% Long term = N/A	Hypoglycemia symptoms (41 vs. 32%)
NAVIGATOR	Nateglinide N = 9306	Outcome = No significant risk reduction Long term = N/A	Hypoglycemia
NAVIGATOR	Valsartan N = 9306	Outcome = No significant risk reduction Long term = N/A	Nasopharyn gitis, back pain, arthralgia hypotension
Tenenbum	Bezafibrate N = 339	Outcome = 59% Long term = N/A	

Diabetes Prevention Trials with Pharmacotherapy: Design and Outcomes

	Population	Design & Follow-up (years)	RRR	Adverse effects
XENDOS	N = 3277 Age 43 BMI 37.3 kg/m ²	RCT Orlistat FU = 4 years	Outcome = 37.3% Long term = N/A	Mild to moderate gastrointestinal events
Heymsfield	N = 675 Age 43.9 & 44.3 I/C BMI 35.6 kg/m ² , 35 kg/m ² I/C Obese	RCT Orlistat IGT	Outcome = 39% Long term = N/A	

Clinical Outcomes of Metabolic Surgery

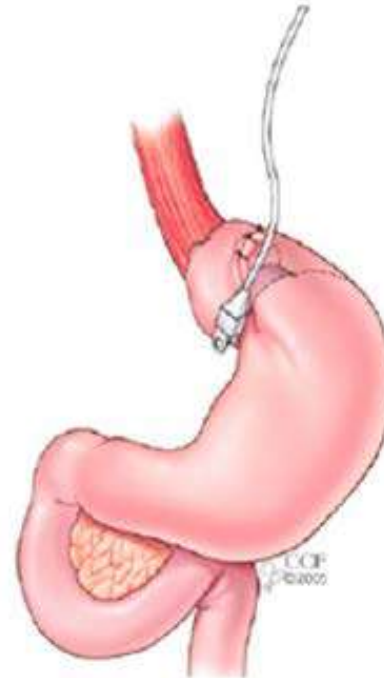
- Gastrointestinal procedures intended to yield long-term weight loss in patients with severe obesity, otherwise known as bariatric surgery.
- The original indications for bariatric surgery were based on BMI and were derived from the seminal National Institutes of Health (NIH) Consensus Conference in 1991, which considered surgery an option in patients with BMI ≥ 40 kg/m² or with BMI ≥ 35 kg/m² with significant obesity-related comorbidities.



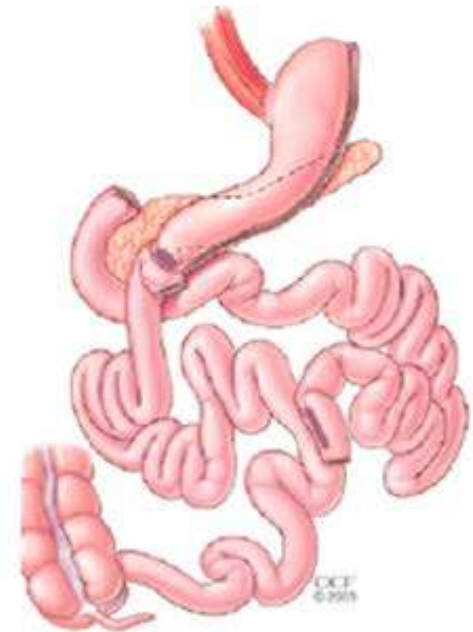
Sleeve Gastrectomy (SG)
Frequency 49%



Roux-en-Y Gastric Bypass (RYGB)
Frequency 43%



Laparoscopic Adjustable Gastric Banding (LAGB)
Frequency 6%



Biliopancreatic Diversion with Duodenal Switch (BPD+DS)
Frequency 2%

Fig (1): Common Metabolic Procedures and their frequency of use

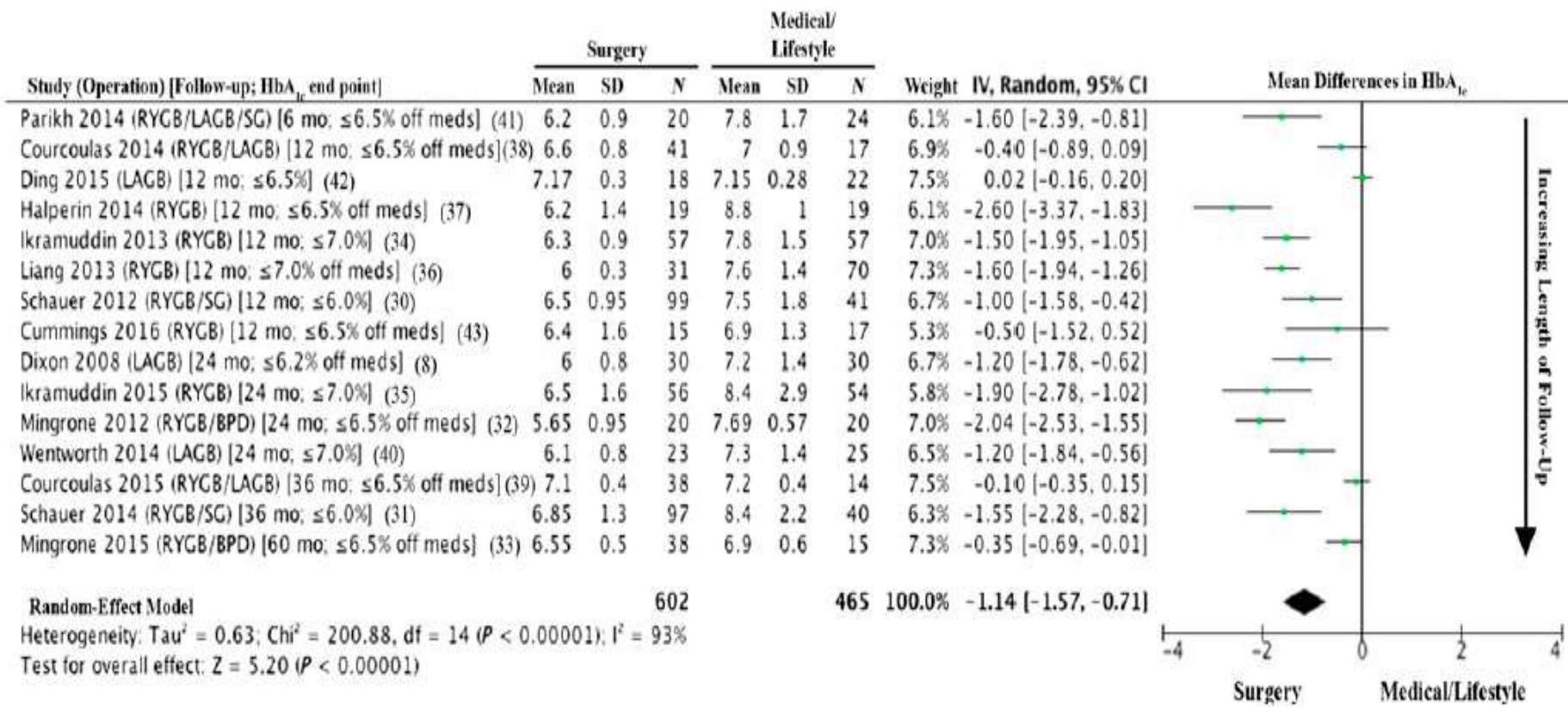


Fig (2): Forest plot of mean differences (MDs) of %HbA1c serum levels after bariatric/metabolic surgery compared with medical/lifestyle treatments in published RCTs.

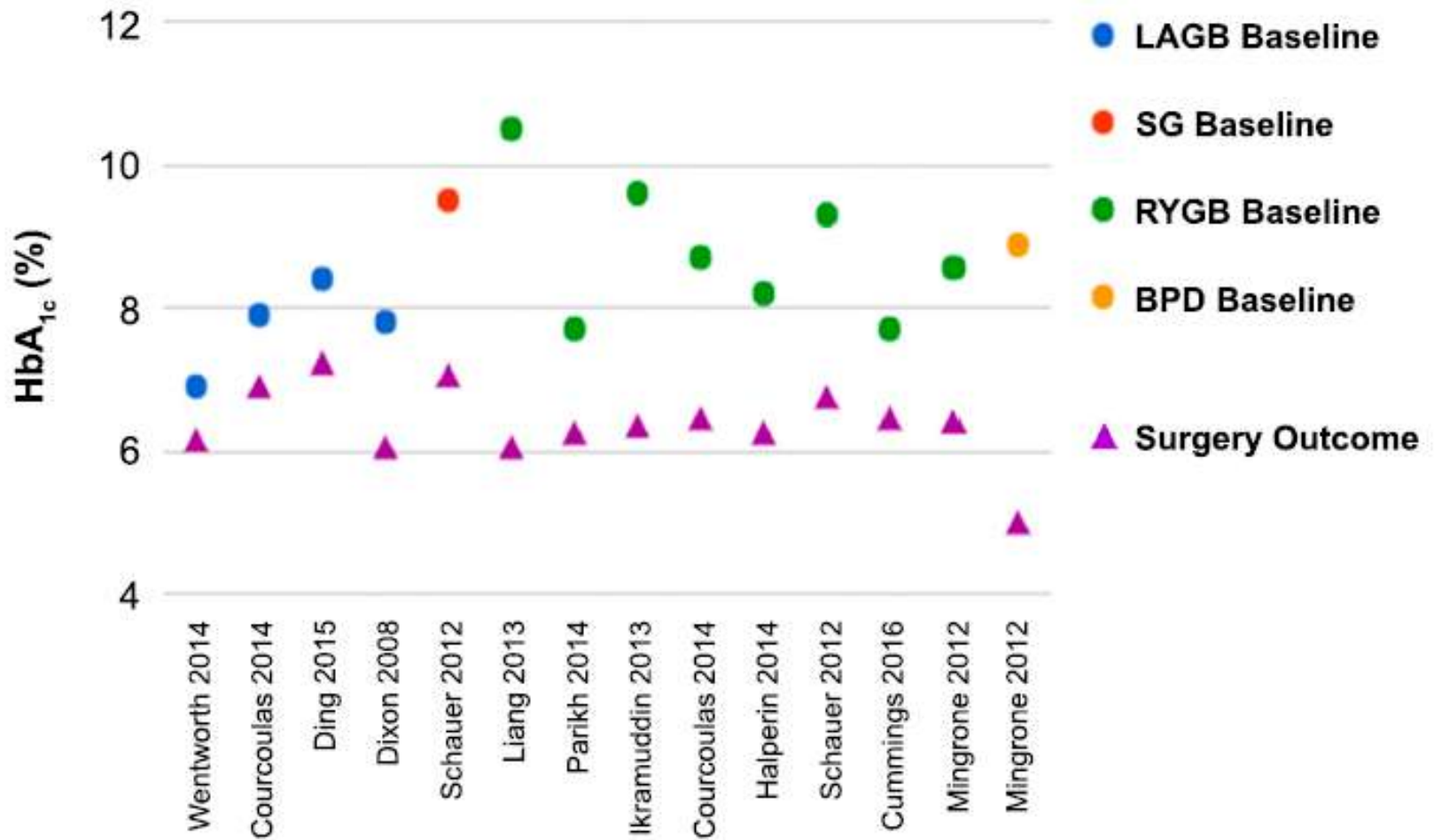
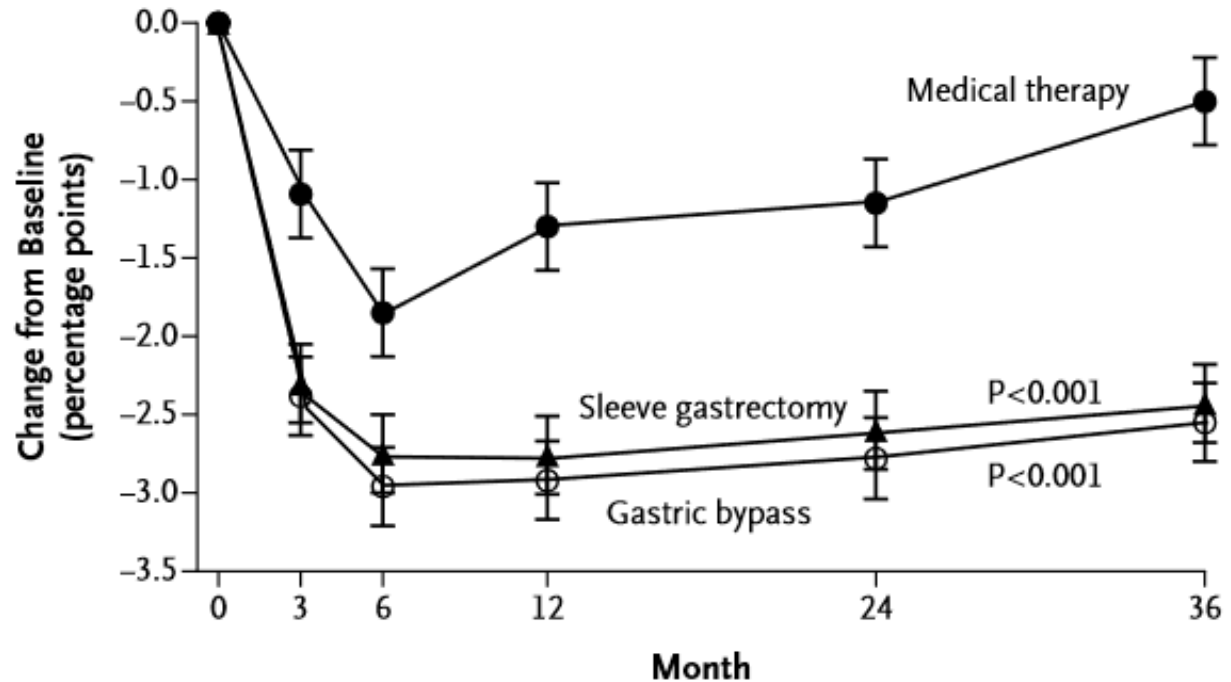


Fig (3) : Change in HbA_{1c} after LAGB, RYGB, SG, and BPD in 11 RCTs.

CHANGES IN HAEMOGLOBIN A₁C

A Glycated Hemoglobin



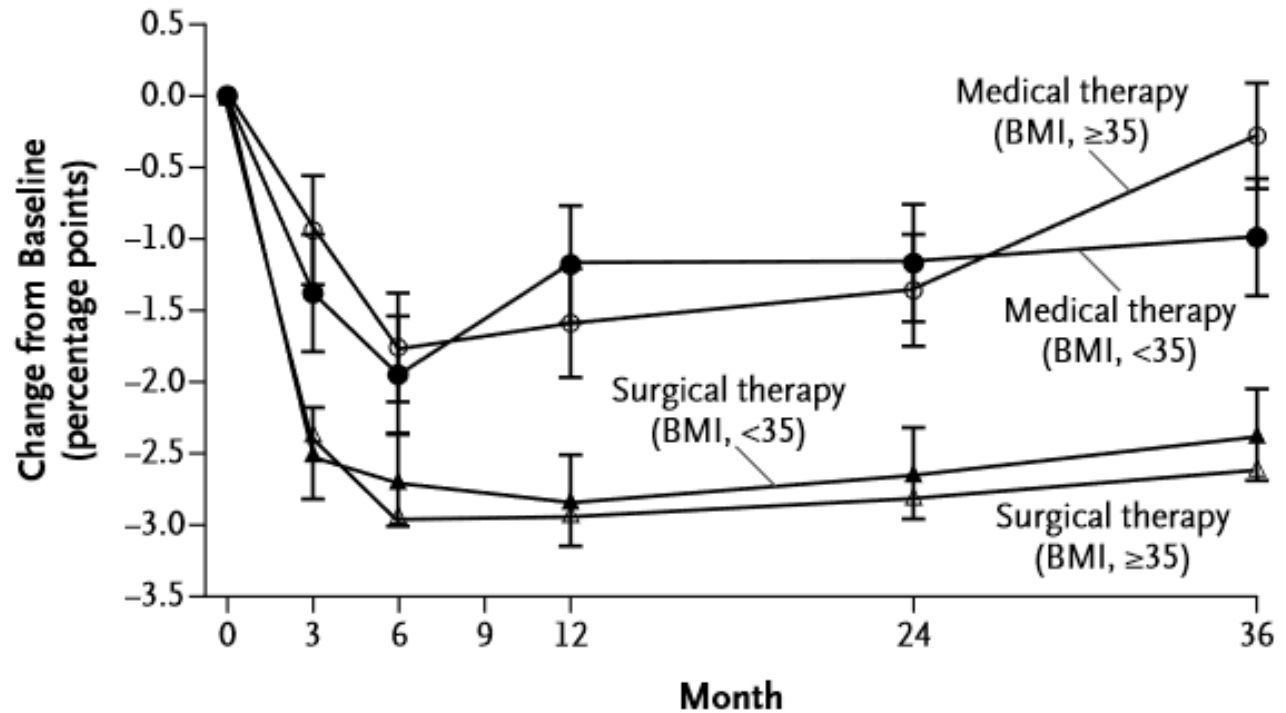
Value at Visit

Medical therapy	9.0 (8.5)	7.1 (6.8)	7.5 (6.9)	7.7 (7.3)	8.4 (7.6)
Sleeve gastrectomy	9.5 (8.9)	6.7 (6.4)	6.6 (6.4)	6.8 (6.8)	7.0 (6.6)
Gastric bypass	9.3 (9.2)	6.3 (6.2)	6.3 (6.1)	6.5 (6.4)	6.7 (6.6)

Fig (4): Change in HbA₁c after surgical vs. medical treatment of T2D in the studies

CHANGES IN HAEMOGLOBIN A₁C

B Glycated Hemoglobin According to Body-Mass Index



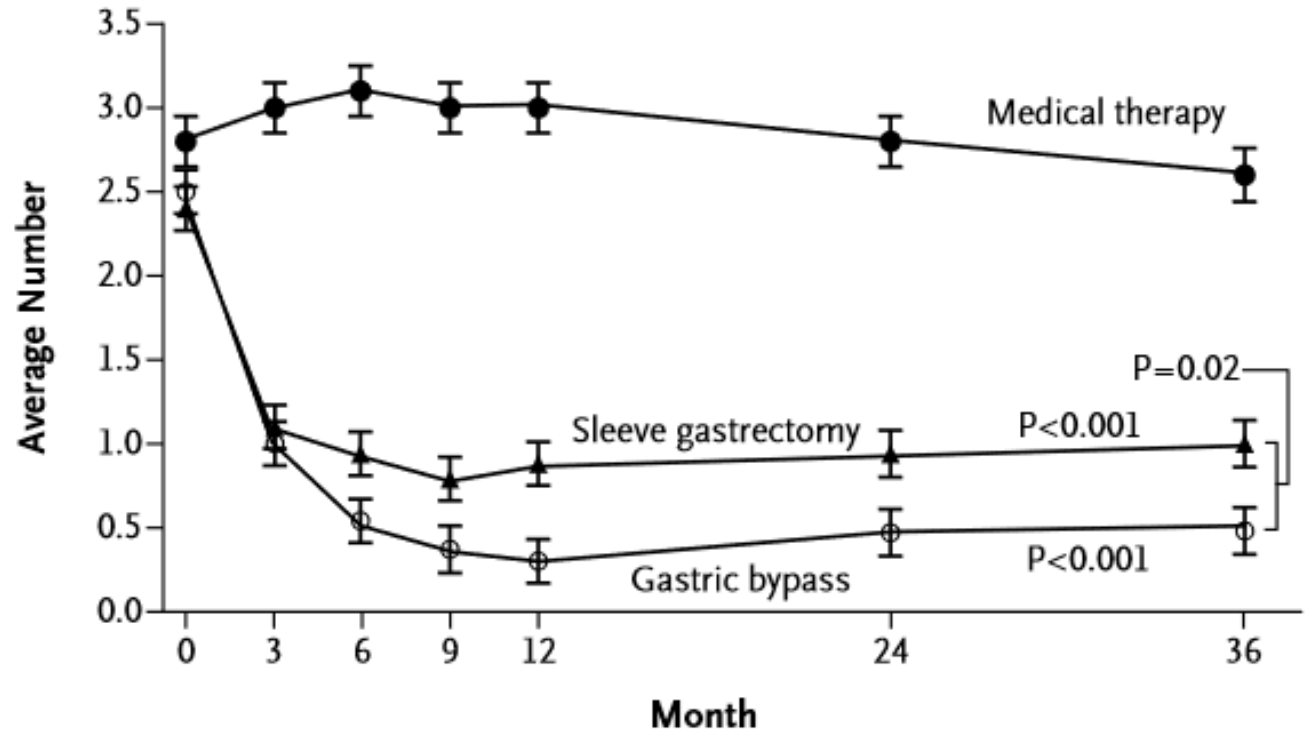
Value at Visit

Medical <35 BMI	9.1 (8.9)	7.2 (6.8)	7.9 (6.9)	8.0 (7.4)	8.1 (7.8)
Medical ≥35 BMI	8.8 (8.5)	7.1 (6.8)	7.2 (6.7)	7.4 (6.9)	8.5 (7.3)
Surgical <35 BMI	9.4 (9.1)	6.7 (6.9)	6.6 (6.6)	6.8 (6.8)	7.1 (6.7)
Surgical ≥35 BMI	9.3 (9.2)	6.4 (6.2)	6.4 (6.1)	6.6 (6.4)	6.7 (6.4)

Fig (4): Change in HbA₁c after surgical vs. medical treatment of T2D in the studies

CHANGES IN HAEMOGLOBIN A₁C

C Diabetes Medications



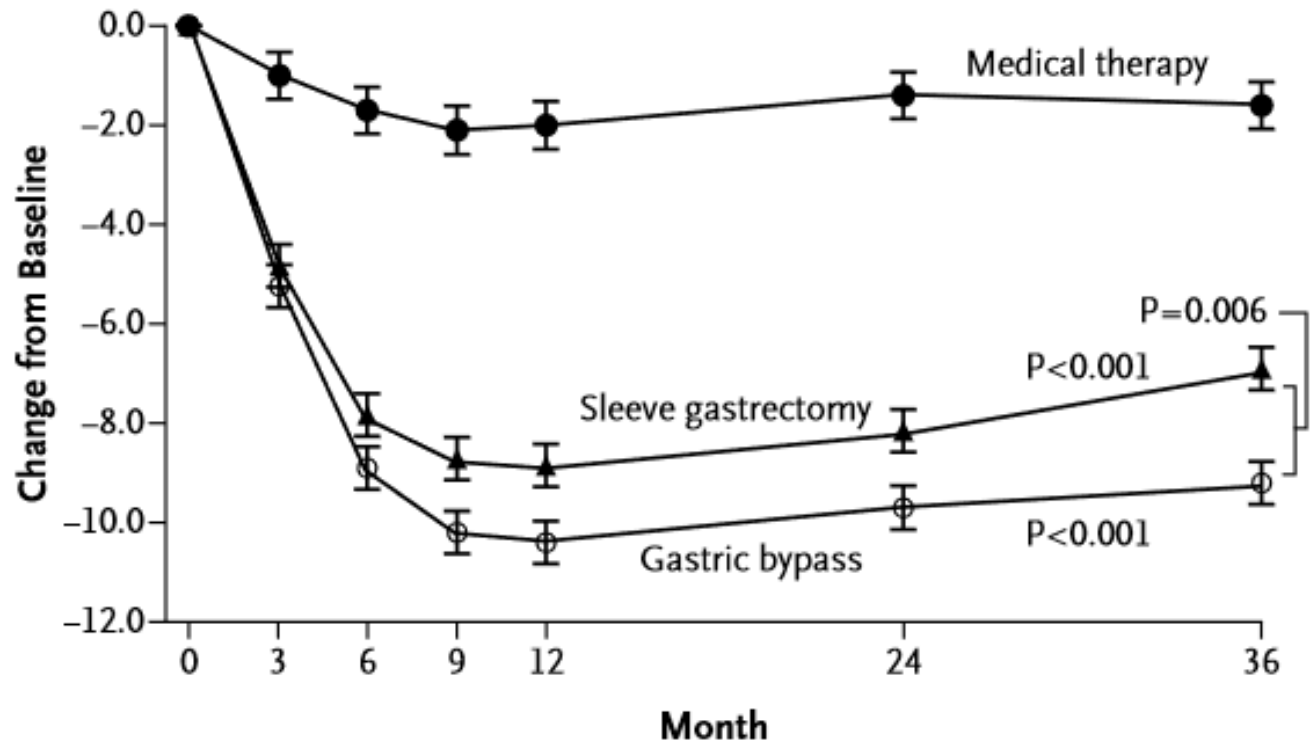
Value at Visit

Medical therapy	2.8	3.1	3.0	2.8	2.6
Sleeve gastrectomy	2.4	0.94	0.88	0.94	1.0
Gastric bypass	2.5	0.54	0.3	0.47	0.48

Fig (4): Change in HbA₁c after surgical vs. medical treatment of T2D in the studies

CHANGES IN HAEMOGLOBIN A₁C

D Body-Mass Index



Value at Visit

Medical therapy	36.4	34.6	34.2	35.0	34.8
Sleeve gastrectomy	36.1	28.3	27.1	27.9	29.2
Gastric bypass	37.1	28.2	26.7	27.3	27.9

Fig (4): Change in HbA₁c after surgical vs. medical treatment of T2D in the studies

A Weight Loss

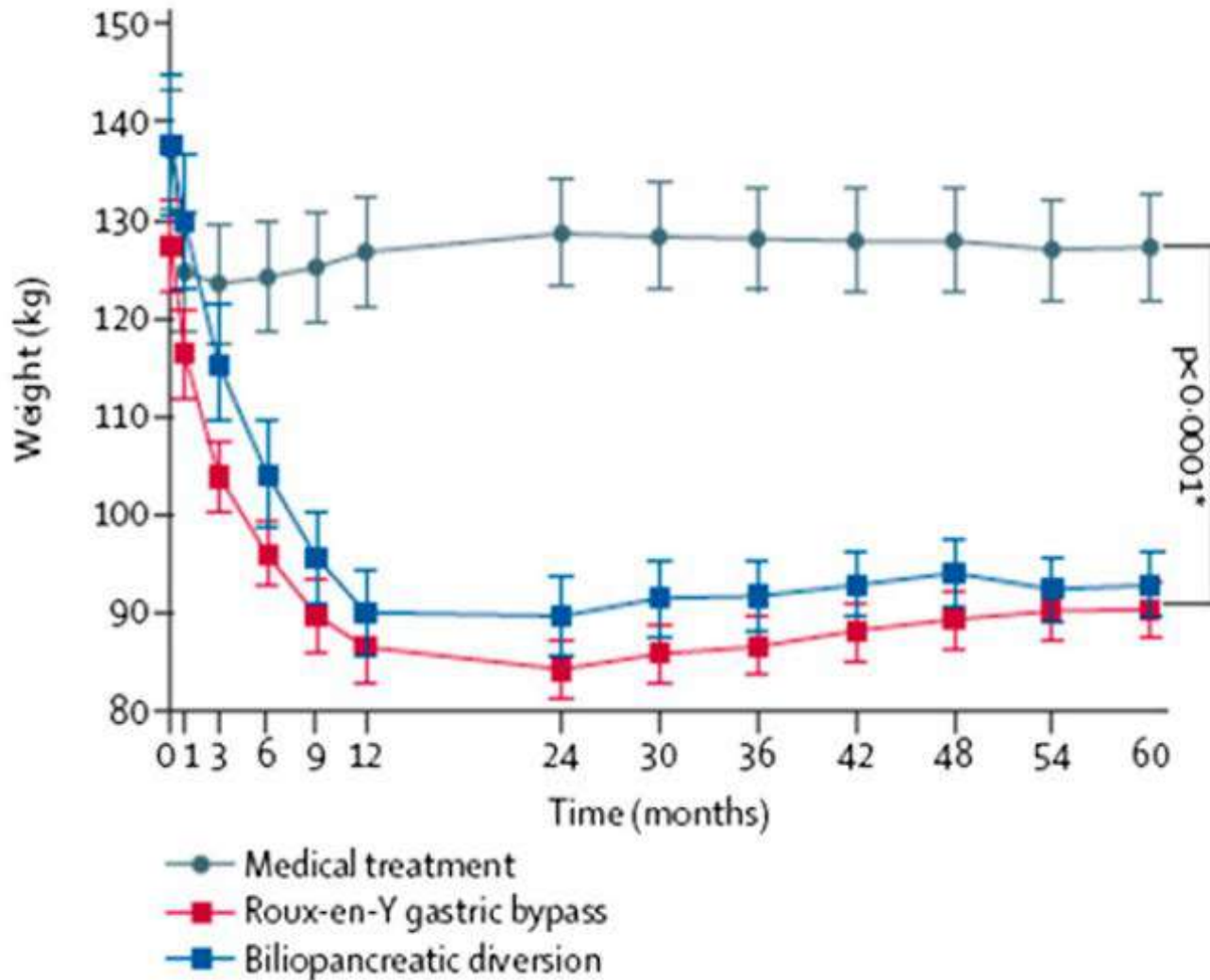


Fig (5): Other secondary end points favoring surgery over medical treatment.

B Change in HDL

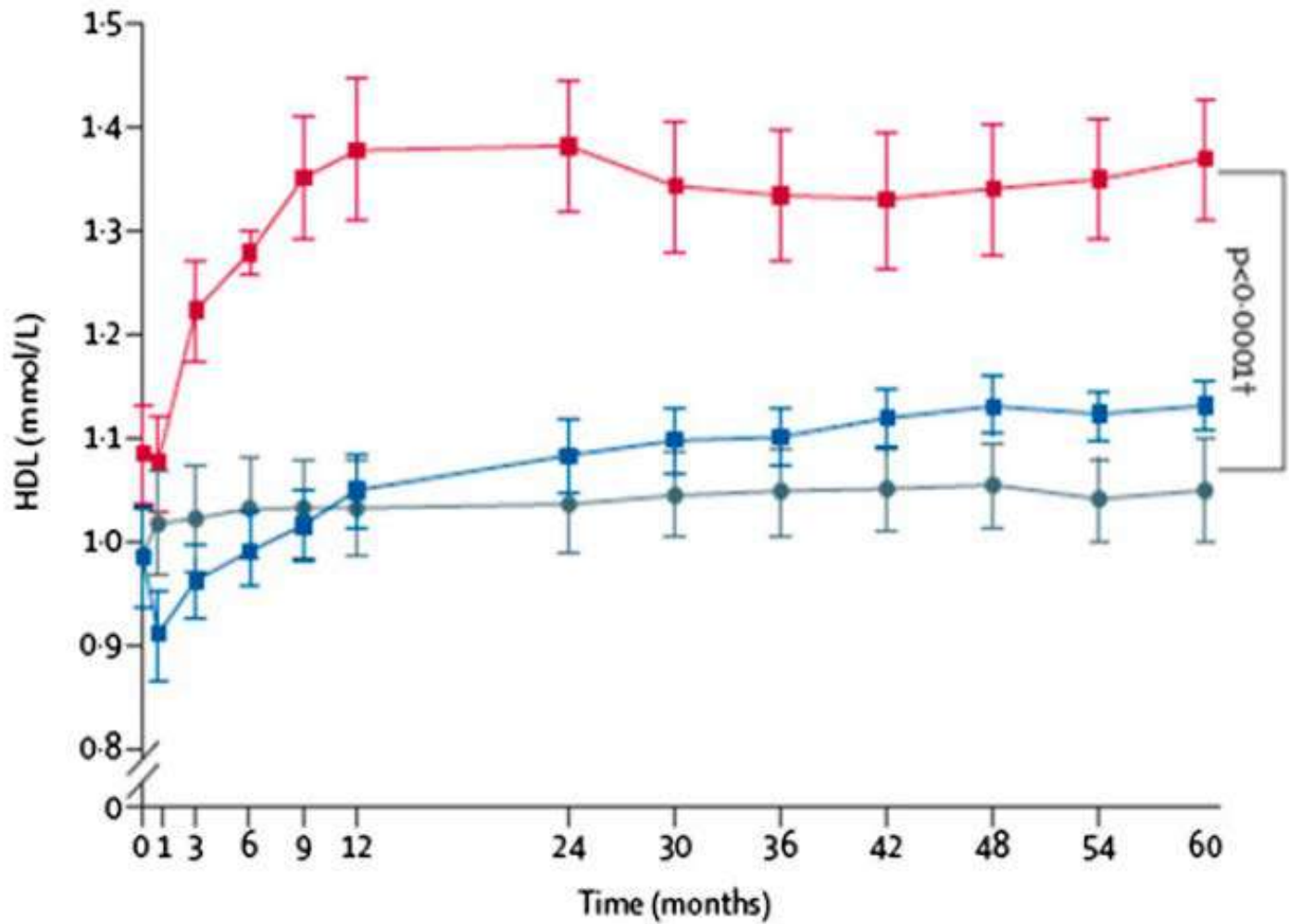
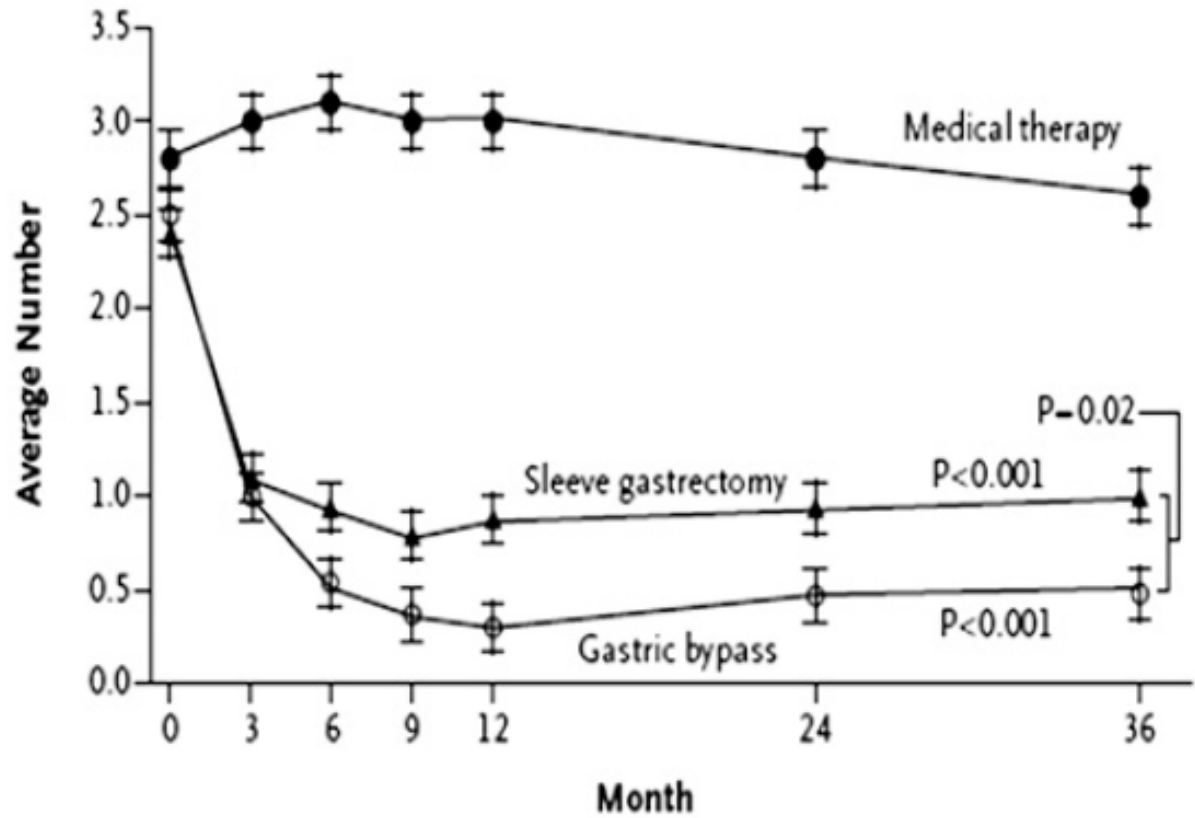


Fig (5): Other secondary end points favoring surgery over medical treatment.

C Change in Medications



Value at Visit

Medical therapy	2.8	3.1	3.0	2.8	2.6
Sleeve gastrectomy	2.4	0.94	0.88	0.94	1.0
Gastric bypass	2.5	0.54	0.3	0.47	0.48

Fig (5): Other secondary end points favoring surgery over medical treatment.

D Change in Quality of Life

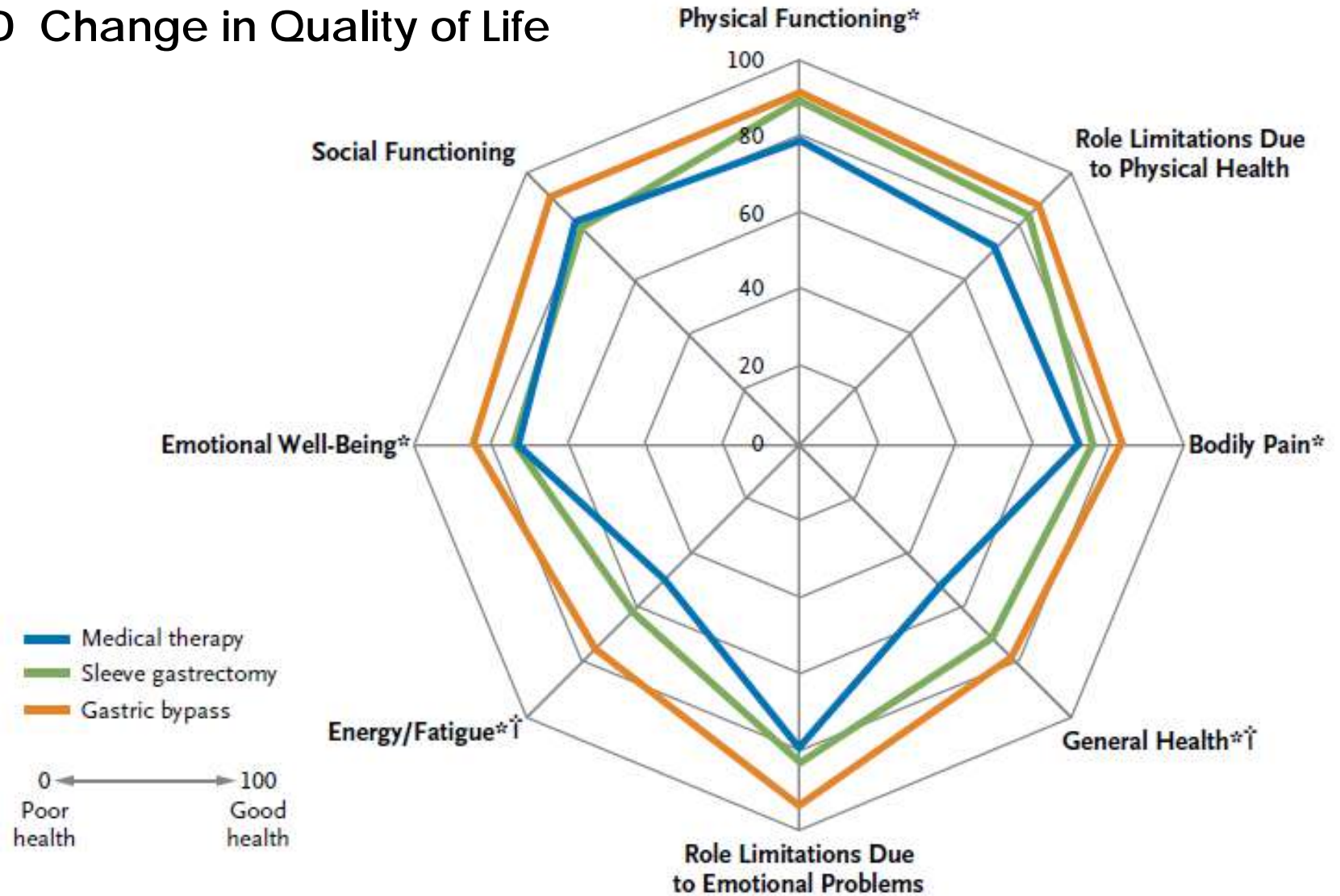


Fig (5): Other secondary end points favoring surgery over medical treatment.

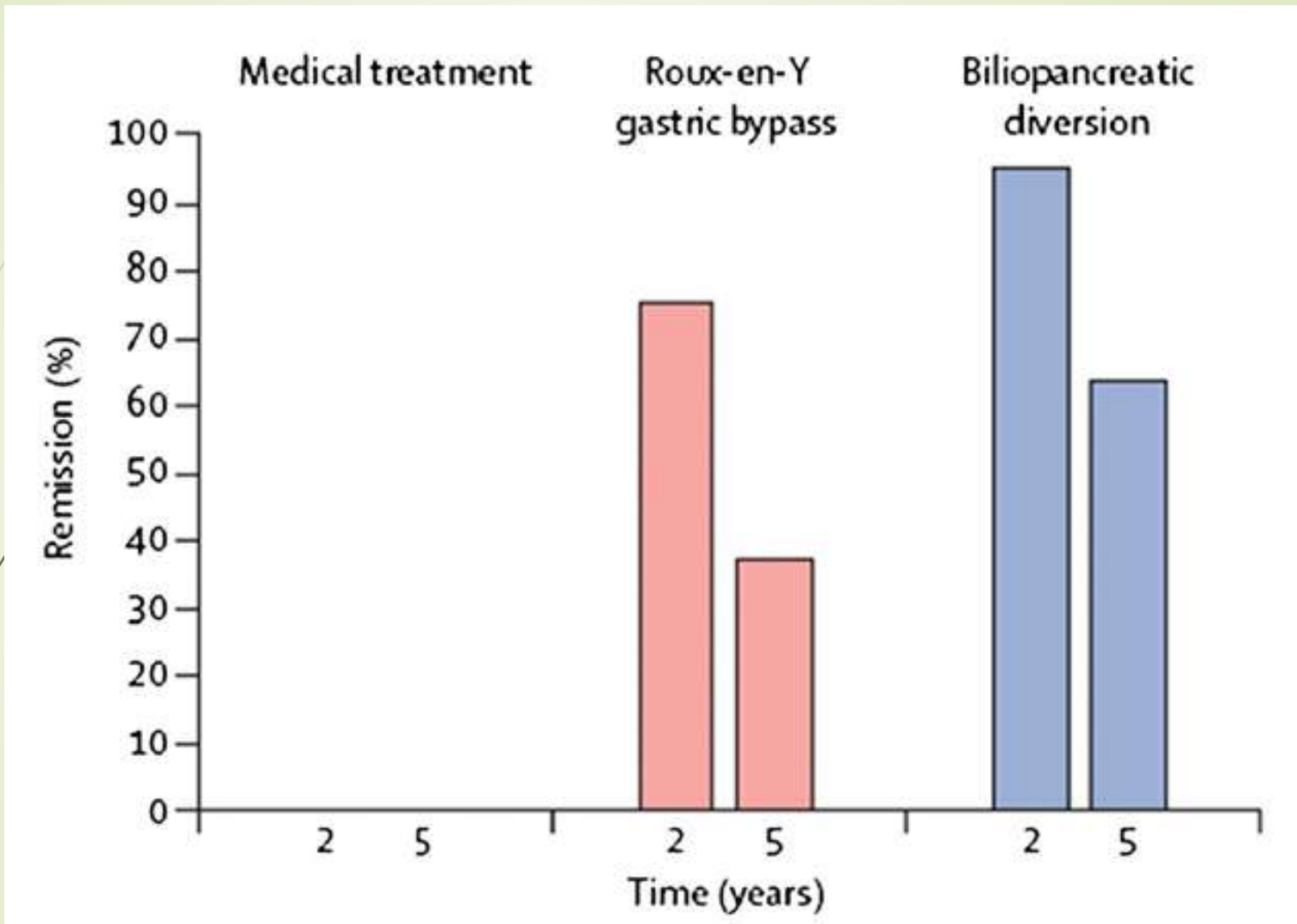


Fig (6): Remission at 5 years in the study

Complications of Metabolic Surgery

Complications	Frequency (%)
Sepsis from anastomotic leak	0.1–5.6
Hemorrhage	1–4
Cardiopulmonary events	<1
Thromboembolic disease	0.34
Death	0.1–0.3
Late complications for LAGB	
Band slippage	15
Leakage	2–5
Erosion	1–2
Late complications of bypass procedures	
Anastomotic strictures	1–5
Marginal ulcers	1–5
Bowel obstructions	0.5–2
Kidney stones	NK
Metabolic bone disease	NK
Alcohol use disorder	NK
Micronutrient and macronutrient deficiencies from RYGB	
2–3 years postoperative	
Iron deficiency	45–52
Vitamin B ₁₂ deficiency	8–37
Calcium deficiency	10
Vitamin D deficiency	51
Fat-soluble vitamin deficiencies (A, D, E, and K) and protein calorie malnutrition from BPD+DS procedures	1–5

NK, not known.

Early Action Type 2 Diabetes

The Four Pillars of Early Action are firmly rooted in the evidence base on what is most valuable in reducing the economic, societal and personal burden of type 2 diabetes:

Prevention

Taking steps to prevent people from developing type 2 diabetes from the outset



Early detection

Identifying those at high risk of developing type 2 diabetes and diagnosing them as early as possible



Early control

Ensuring that people with diabetes are given the treatment and support they need to achieve good control of their blood glucose levels as early as possible, to reduce the risk of complications



Early access to the right interventions

Ensuring that health systems are addressing the need for equitable, early access to the personalized education, lifestyle change programmes and treatments that people with type 2 diabetes need



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OUR VISION IS.....

A Nation with
minimal burden of health
due to Diabetes

OUR MISSION IS.....

Prevention of Diabetes and
its complications
in the country
through the delivery of
comprehensive
diabetes care



Myanmar Diabetes Association

