



*2018 Clinical Practice Guidelines for
the Prevention and Management of
Diabetes in Canada*

The Essentials

Presented Feb.25, 2019

2018 Diabetes Canada CPG – The Essentials

Faculty/Presenter Disclosure

- **Faculty:** Lengim Ingram, BscPharm, EPPh, CDE

- **Relationships with commercial interests:**

Consultant/Advisory Board Honorarium: Sanofi, Novo Nordisk

Speaker's Honorarium: Merck, Janssen, Astra Zeneca,

Grant or other payments: Abbott, Servier, BMI



Learning Objectives

- **By the end of this session, participants will be able to:**
 - Understand the **major changes** within the 2018 *Diabetes Canada clinical practice guidelines*
 - Apply the recommendations in clinical practice
 - Use the tools in the guideline to provide better diabetes care for patients

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Key Message

- Throughout the guidelines remains the importance of individualizing therapy for the person with diabetes



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Screening for and Diagnosing Diabetes



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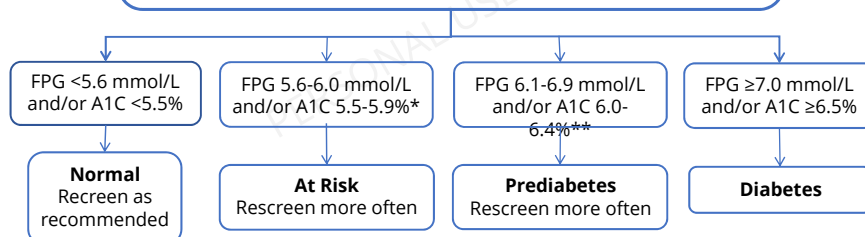
2018 Diabetes Canada CPG – Chapter 4. Screening for Diabetes in Adults

2018

Screening for type 2 diabetes in adults

Screen every **3 years** in individuals **≥40 years of age** or in individuals at high risk using a risk calculator.

Screen earlier and/or more frequently (every 6 to 12 months) in people with additional risk factors for diabetes or for those at very high risk using a risk calculator



If both FPG and A1C are available, but discordant, use the test that appears furthest to the right side of the algorithm.

*Consider 75-g OGTT if 1 risk factors; ** Consider 75-g OGTT

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2018 Diabetes Canada CPG – Chapter 3. Definition, Diagnosis & Classification of Diabetes, Prediabetes, Metabolic Syndrome

Diagnosis of Diabetes

FPG ≥ 7.0 mmol/L

Fasting = no caloric intake for at least 8 hours

or

A1C $\geq 6.5\%$ (in adults)

Using a standardized, validated assay in the absence of factors that affect the accuracy of the A1C and not for suspected type 1 diabetes

or

2hPG in a 75 g OGTT ≥ 11.1 mmol/L

or

Random PG ≥ 11.1 mmol/L

Random = any time of the day, without regard to the interval since the last meal

FPG, fasting plasma glucose; OGTT, oral glucose tolerance test; PG, plasma glucose

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Confirmatory test required

- In the **absence of symptomatic hyperglycemia**, if a single lab test result is in the diabetes range, a **repeat confirmatory lab test** (FPG, A1C, 2hPG in a 75 g OGTT) must be done on another day
- **Repeat the same test** (in a timely fashion) to confirm
- But a **random PG** in the diabetes range in an **asymptomatic** individual should be **confirmed with an alternate test**
- If **results of two different tests** are available and **both are above** the diagnostic thresholds, the diagnosis of diabetes is **confirmed**

2hPG, 2-hour plasma glucose; A1C, glycated hemoglobin; FPG, fasting plasma glucose; OGTT, oral glucose tolerance test; PG, plasma glucose.

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Screening for and Diagnosing Diabetes

 Print

Healthcare Provider Tool

▼ Screening and Diagnosis Calculator

Measure Fasting Plasma Glucose (FPG) and/or A1C and enter test results:

FPG: mmol/L A1C: %

Calculate Screening / Diagnosis Result

► Risk Factors for Type 2 Diabetes

► When to Screen for Diabetes

► Screening Algorithm

This is only to be used as a decision support tool and is subject to these terms.
For more information, please see terms of use.

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Approach to Management



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ABCDE³ of Diabetes Care

- ✓ **A** • A1C – optimal glycemic control (usually $\leq 7\%$)
- ✓ **B** • BP – optimal blood pressure control ($< 130/80$)
- ✓ **C** • Cholesterol – LDL < 2.0 mmol/L or $> 50\%$ reduction
- ✓ **D** • Drugs to protect the heart
 A – ACEi or ARB | S – Statin | A – ASA if indicated | SGLT2i/GLP-1 RA
 with demonstrated CV benefit if type 2 DM with CVD and A1C not at target
- ✓ **E** • Exercise / Healthy Eating
- ✓ **S** • Screening for complications
- ✓ **S** • Smoking cessation
- ✓ **S** • Self-management, stress and other barriers

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A1C Targets

2018

≤6.5	Adults with type 2 diabetes to reduce the risk of CKD and retinopathy if at low risk of hypoglycemia
≤7.0	MOST ADULTS WITH TYPE 1 OR TYPE 2 DIABETES
7.1 ↓ 8.5	7.1-8.0%: Functionally dependent* 7.1-8.5%: <ul style="list-style-type: none"> • Recurrent severe hypoglycemia and/or hypoglycemia unawareness • Limited life expectancy • Frail elderly and/or with dementia**
Avoid higher A1C to minimize risk of symptomatic hyperglycemia and acute and chronic complications	
End of life	A1C measurement not recommended. Avoid symptomatic hyperglycemia and any hypoglycemia

* Based on class of antihyperglycemic medication(s) utilized and person's characteristics

** see Diabetes in Older People chapter

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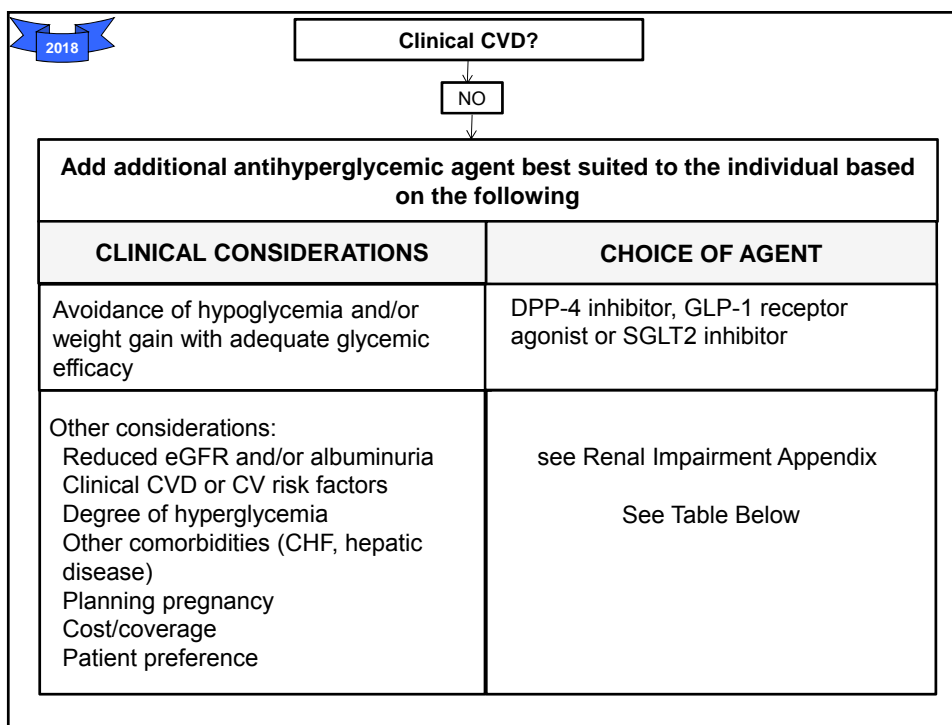
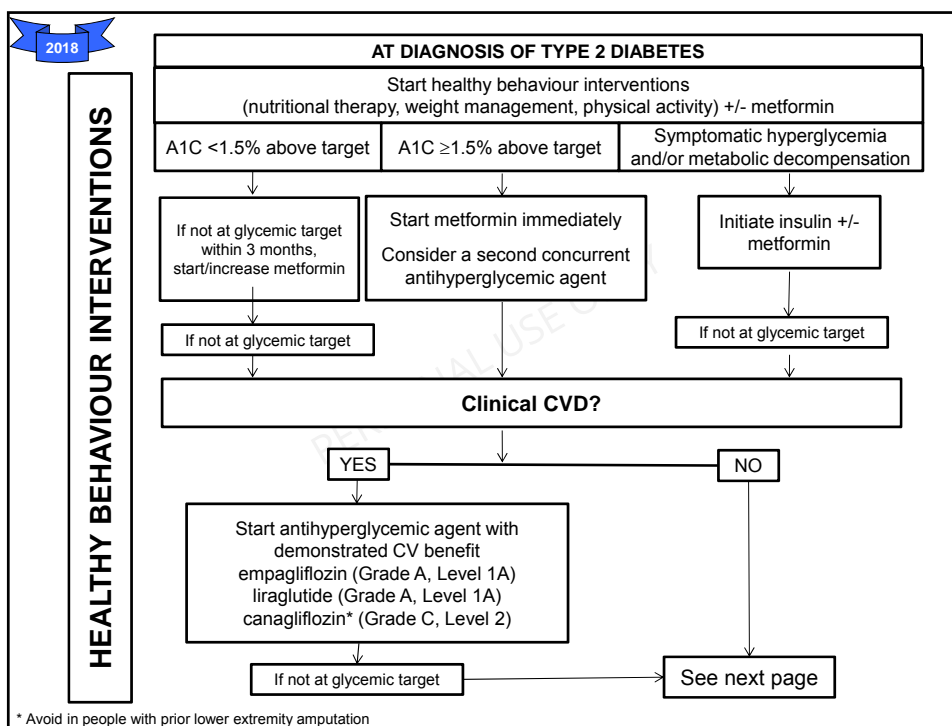
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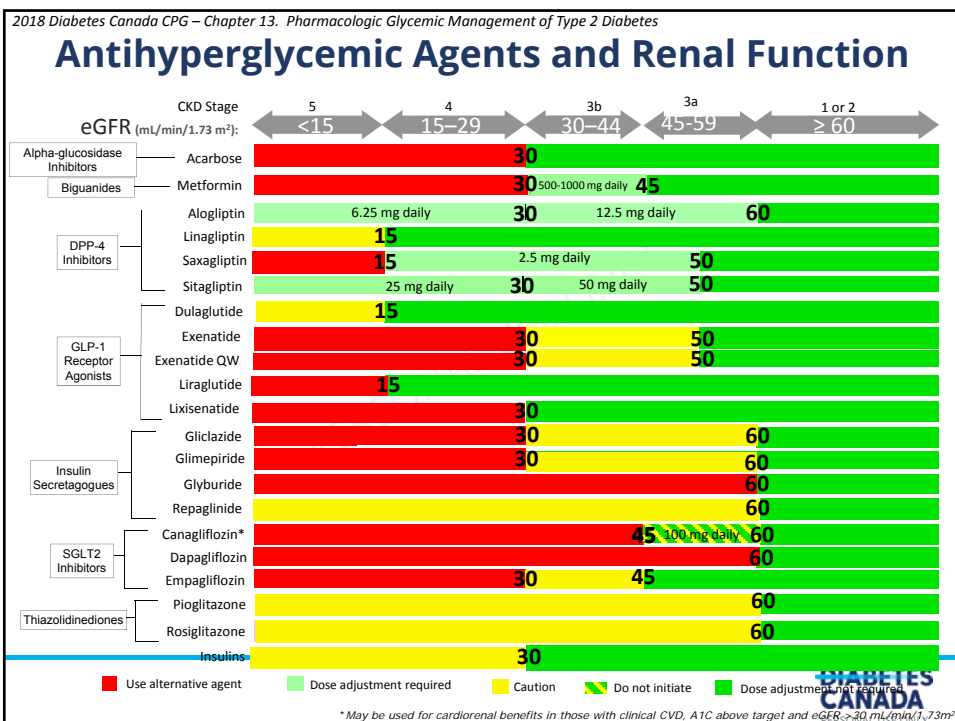
Individualizing your Patient's A1C Target

Print

For Patients with Type 1 and Type 2 Diabetes

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2018 Diabetes Canada CPG – Chapter 13. Pharmacologic Glycemic Management of Type 2 Diabetes

Add additional antihyperglycemic agent best suited to the individual by prioritizing patient characteristics (agents listed in alphabetical order by CV outcome data)

Class	Effect on CVD Outcomes	Hypo-glycemia	Weight	Relative A1C Lowering when added to metformin	Other therapeutic considerations	Cost
GLP-1R agonists	lira: Superiority in T2DM with clinical CVD exenatide LAR & lixi: Neutral	Rare	↓↓	↓↓ to ↓↓↓	GI side-effects, Gallstone disease Contraindicated with personal / family history of medullary thyroid cancer or MEN 2 Requires subcutaneous injection	\$\$\$\$
SGLT2 inhibitors	Cana & empa: Superiority in T2DM patients with clinical CVD	Rare	↓↓	↓↓ to ↓↓↓	Genital infections, UTI, hypotension, dose-related changes in LDL-C. Caution with renal dysfunction, loop diuretics, in the elderly. Dapagliflozin not to be used if bladder cancer. Rare diabetic ketoacidosis (may occur with no hyperglycemia). Increased risk of fractures and amputations with canagliflozin. Reduced progression of nephropathy & CHF hospitalizations with empagliflozin and canagliflozin in those with clinical CVD	\$\$\$
DPP-4 Inhibitors	alo, saxa, sita: Neutral	Rare	Neutral	↓↓	Caution with saxagliptin in heart failure Rare joint pain	\$\$\$
Insulin	glar: Neutral degludec: noninferior to glar	Yes	↑↑	↓↓↓↓	No dose ceiling. Flexible regimens Requires subcutaneous injection	\$- \$\$\$\$
Thiazolidinediones	Neutral	Rare	↑↑	↓↓	CHF, edema, fractures, rare bladder cancer (pioglitazone), cardiovascular controversy (rosiglitazone), 6-12 weeks for maximal effect	\$
α-glucosidase Inhibitor (acarbose)		Rare	Neutral	↓	GI side-effects common Requires 3 times daily dosing	\$
Insulin secretagogue: Meglitinide		Yes	↑	↓↓	More rapid BG-lowering response Reduced postprandial glycemia with meglitinides but usually requires 3 to 4 times daily dosing.	\$
Sulfonylurea		Yes	↑	↓↓	Gliclazide and glimepiride associated with less hypoglycemia than glyburide. Poor durability	\$
Weight loss agent (orlistat)		None	↓	↓	GI side effects Requires 3 times daily dosing	\$\$\$

↓

If not at glycemic targets

↓

Add another antihyperglycemic agent from a different class and/or add/intensify insulin regimen
Make timely adjustments to attain target A1C within 3-6 months

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Sulfonylurea		Yes	↑	↓↓	Gliclazide and glimepiride associated with less hypoglycemia than glyburide. Poor durability	\$
Weight loss agent (orlistat)		None	↓	↓	GI side effects Requires 3 times daily dosing	\$\$\$

Types of insulin			
Insulin type (trade name)	Onset	Peak	Duration
BOLUS (prandial or mealtime) insulins			
Rapid-acting insulin analogues (clear) • Insulin aspart (NovoRapid®) • Insulin glulisine (Apidra®) • Insulin lispro (Humalog®) U-100 U-200 • Faster-acting insulin aspart (Fiasp®)	9–20min 10–15min 10–15min 4min	1–1.5h 1–1.5h 1–2h 0.5–1.5h	3–5h 3.5–5h 3–4.75h 3–5h
Short-acting insulins (clear) • Insulin regular (Humulin®-R, Novolin® ge Toronto) • Insulin regular U-500 (Entuzity® (U-500))	30min 15min	2–3h 4–8h	6.5h 17–24h
BASAL insulins			
Intermediate-acting (cloudy) • Insulin neutral protamine Hagedorn (Humulin® N, Novolin® ge NPH)	1–3h	5–8h	Up to 18h
Long-acting insulin (clear) • Insulin detemir (Levemir®) • Insulin glargine U-100 (Lantus®) • Insulin glargine U-300 (Toujeo®) • Insulin glargine biosimilar (Basaglar®) • Insulin degludec U-100, U-200 (Tresiba®)	90min	Not applicable	U-100 glargine 24h, detemir 16–24h U-300 glargine >30h degludec 42h
PREMIXED insulins			
Premixed regular insulin –NPH (cloudy) • Humulin® 30/70 • Novolin® ge 30/70, 40/60, 50/50	A single vial or cartridge contains a fixed ratio of insulin (% of rapid-acting or short-acting insulin to % of intermediate-acting insulin)		
Premixed insulin analogues (cloudy) • Biphasic insulin aspart (NovoMix® 30) • Insulin lispro/lispro protamine (Humalog® Mix25 and Mix50)			

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Examples of Insulin Initiation and Titration Regimens in People With Type 2 Diabetes

All people starting insulin should be counseled about the recognition, prevention and treatment of hypoglycemia. Consider a change in type or timing of insulin administration if glycemic targets are not being reached.

Example A: Basal insulin (degludec U-100 or U-200, detemir, glargine U-100 or U-300, NPH) added to non-insulin antihyperglycemic agents

- Insulin should be titrated to achieve target fasting BG levels of **4.0 to 7.0 mmol/L** or individualized targets (e.g. **4.0 to 5.5 mmol/L** if A1C target **≤7.0% not achieved**; higher fasting BG targets may be considered in some people with diabetes where the goal of avoiding hypoglycemia is important, see Chapter 8. Targets for Glycemic Control, p. S42).
- Individuals can be taught self-titration, or titration may be done in conjunction with a health-care provider.
- Suggested starting dose is 10 units once daily at bedtime.
- Suggested titration is 1 unit per day until target is reached. (Degludec should be titrated by 2 units every 3 to 4 days or 4 units once a week).
- A lower starting dose, slower titration and higher targets may be considered for elderly or normal-weight subjects.
- In order to safely titrate insulin, people with diabetes must perform self-monitoring of blood glucose at least once a day fasting.
- Insulin dose should not be increased if the individual experiences 2 episodes of hypoglycemia (BG <4.0 mmol/L) in 1 week or any episode of nocturnal hypoglycemia.
- Non-insulin antihyperglycemic agents (especially insulin secretagogues) may need to be reduced if daytime hypoglycemia occurs.

Insulin Initiation and Titration Suggestions for Type 2 Diabetes

People starting insulin should be counseled about the prevention, recognition and treatment of hypoglycemia.

The following are suggestions for insulin initiation and titration. Clinical judgment must always be used as the suggestions may not apply to every patient.

Basal Insulin (only as an add-on to Antihyperglycemic Agents)

(Basaglar[®], Lantus[®], Levemir[®], Toujeo[®], Tresiba[®], Humulin[®] N, Novolin[®] ge NPH)

- Target fasting blood glucose (BG) of 4-7 mmol/L. The fasting BG target can be changed to 4-5.5 mmol/L if not achieving adequate overall glycemic control.
- Most patients will need 40-50 units at bedtime to achieve target but there is no maximum dose.
- Start at a low dose of 10 units at bedtime (may start at lower dose [0.1-0.2 units/kg] for lean patients <50 kg).
- If using Tresiba[®], the dose can be increased by 2-4 units every week until fasting BG target is achieved.
- Patient should gently self-titrate by increasing the dose by 1 unit every 1 night until fasting BG target is achieved.
- If fasting hypoglycemia occurs, the dose of bedtime basal should be reduced.
- Metformin and the secretagogue are usually maintained when basal insulin is added.

Dosing and Titration Example

Starting dose: 10 units at bedtime.
Increase dose by 1 unit every 1 night until fasting blood glucose has reached the target of 4-7 mmol/L.

guidelines.diabetes.ca/BloodGlucoseLowering/Insulin PrescriptionTool

- For current basal insulin users, maintain the basal dose and add bolus insulin with each meal at a dose equivalent to 10% of the basal dose. For example, if the patient is on 50 units of basal insulin, add 5 units of bolus insulin with each meal.
- For new insulin users starting a full Basal + Bolus regimen, calculate Total Daily Insulin dose (TDI) as 0.3 to 0.5 units/kg, then distribute as follows:
 - 40% of TDI dose as basal insulin (Lantus[®], Levemir[®], Toujeo[®], Humulin[®] N, Novolin[®] ge NPH) at bedtime.
 - 60% of TDI dose as prandial (bolus) insulin prior to each meal.
- Rapid-acting insulin analogues (Apidra[®], Fiasp[®], Humalog[®], Novolog[®]) should be given 0-10 minutes before eating.
- Short-acting insulin (Humulin[®] R, Novolin[®] ge Toronto) should be given 30 minutes before eating.
- An alternative distribution is 50% basal insulin (at bedtime) and 50% bolus insulin (distributed among the meals of the day).
- Adjust the dose of the basal insulin to achieve the target fasting BG level (usually 4-7 mmol/L).
- Adjust the dose of the bolus (prandial) insulin to achieve postprandial BG levels (usually 5-10 mmol/L) or pre-prandial BG levels for the subsequent meal (usually 4-7 mmol/L).

Basal insulin = 40% of TDI:

40% x 50 units

= Basal bedtime = 20 units

Bolus insulin = 60% of TDI:

60% x 50 units

= Bolus = 30 units

= 10 units with each meal

Premixed Insulin Before Breakfast and Before Dinner

(Humalog[®] Mix25[®], Humalog[®] Mix50[®], NovoMix[®] 30, Humulin[®] 30/70, Novolin[®] ge 30/70)

- Target fasting and pre-supper BG levels of 4-7 mmol/L.
- Most patients with type 2 diabetes will need 40-50 units twice a day to achieve target but there is no maximum dose.
- Start at a low dose of 5 to 10 units twice daily (before breakfast and before supper).
- Patient can gently self-titrate by increasing the breakfast dose by 1 unit every day until the pre-supper BG is at target.
- Patient can gently self-titrate by increasing the supper dose by 1 unit every day until the fasting BG target is at target.
- Beware of hypoglycemia post-breakfast or post-supper. Stop increasing dose if this occurs.
- Premixed analogue insulins (Humalog[®] Mix25[®], Humalog[®] Mix50[®], NovoMix[®] 30) should be given 0 to 10 minutes before eating.
- Premixed regular insulins (Humulin[®] 30/70, Novolin[®] ge 30/70) should be given 30 minutes before eating.
- Continue Metformin and consider stopping secretagogue.

Dosing and Titration Example

10 units ac breakfast, 10 units ac supper.

Increase breakfast dose by 1 unit every

1 day until pre-supper blood glucose

has reached the target of 4-7 mmol/L

(usual target).

Increase supper dose by 1 unit every

1 day until fasting blood glucose has

reached the target of 4-7 mmol/L

(usual target).

Selection of Pen Needle

- Forum for Injection Technique (FIT) Canada recommends that 4, 5, and 6mm needles are suitable for all people with diabetes regardless of BMI. In addition, there is no clinical reason for recommending needles longer than 8mm. Initial insulin therapy should start with the shorter needle length (Berard L, et al. FIT Forum for Injection Technique Canada. Recommendations for Best Practice in Injection Technique. October 2011).

CHOOSE AN INSULIN TYPE		CHOOSE A BRAND		DOSING SEE REVERSE FOR TIPS	
STEP 1: Choose Insulin Type					
BASAL Long-acting analogues (Clear)		<input type="checkbox"/> Basaglar® <input type="checkbox"/> Cartridge <input type="checkbox"/> Kwikpen® (prefilled)		<input type="checkbox"/> Levemir® <input type="checkbox"/> Cartridge <input type="checkbox"/> FlexTouch® (prefilled)	
<input type="checkbox"/> Tresiba® <input type="checkbox"/> FlexTouch® 100 U/ml (prefilled) <input type="checkbox"/> FlexTouch® 200 U/ml (prefilled)		<input type="checkbox"/> Lantus® <input type="checkbox"/> Cartridge <input type="checkbox"/> SoloSTAR® (prefilled)		<input type="checkbox"/> Toujeo® <input type="checkbox"/> SoloSTAR® (prefilled)	
Intermediate-acting (Cloudy)		<input type="checkbox"/> Humulin® N <input type="checkbox"/> Cartridge <input type="checkbox"/> Kwikpen® (prefilled)		<input type="checkbox"/> Novolin® ge NPH <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial	
PRANDIAL (BOLUS) Rapid-acting analogues (Clear)		<input type="checkbox"/> Humalog® <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial <input type="checkbox"/> Kwikpen® (prefilled)		<input type="checkbox"/> Fiasp® <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial <input type="checkbox"/> FlexTouch® (prefilled)	
<input type="checkbox"/> Humalog® 200 units/mL <input type="checkbox"/> Kwikpen® (prefilled)		<input type="checkbox"/> NovoRapid® <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial <input type="checkbox"/> FlexTouch® (prefilled)		<input type="checkbox"/> Apidra® <input type="checkbox"/> Cartridge <input type="checkbox"/> SoloSTAR® (prefilled)	
Short-acting (Clear) Give 30 minutes before meal.		<input type="checkbox"/> Humulin® R <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial		<input type="checkbox"/> Novolin® ge Toronto <input type="checkbox"/> Cartridge <input type="checkbox"/> Vial	
PREMIXED Premixed analogues (Cloudy)		<input type="checkbox"/> Humalog® Mix25® <input type="checkbox"/> Cartridge <input type="checkbox"/> Kwikpen® (prefilled)		<input type="checkbox"/> NovoMix® 30 <input type="checkbox"/> Cartridge	
<input type="checkbox"/> Humalog® Mix50® <input type="checkbox"/> Cartridge <input type="checkbox"/> Kwikpen® (prefilled)					
SELECT PEN DEVICE		CHECK OFF SUPPLIES		QUANTITY & REPEATS	
PEN DEVICE Required if insulin cartridges selected. Insulin pen should match the insulin brand.		<input type="checkbox"/> HumaPen® Savvio® <input type="checkbox"/> HumaPen LUXURA® HD		<input type="checkbox"/> NovoPen® 4 <input type="checkbox"/> NovoPen Echo® <input type="checkbox"/> NovoPen® 5	
<input type="checkbox"/> CliSTAR®					
OTHER SUPPLIES		<input type="checkbox"/> Pen needles (if using a pen): Check needle size (refer to back for information): <input type="checkbox"/> 4mm <input type="checkbox"/> 5mm <input type="checkbox"/> 6mm <input type="checkbox"/> 8mm OR			
<input type="checkbox"/> At discretion of pharmacist <input type="checkbox"/> Glucose test strips <input type="checkbox"/> Lancets <input type="checkbox"/> Insulin Syringe (if using vials) <input type="checkbox"/> Glucagon Kit (if applicable) <input type="checkbox"/> Ketone Strips (if applicable)					
QUANTITY and REPEATS		Supplies Mitre: _____ boxes Repeats x _____			
Signature: _____		Print Name: _____		Signature #: _____	
This tool was developed by the Ontario College of Family Physicians and the New Brunswick Diabetes Task Group and was re-produced with permission by Diabetes Canada. Diabetes Canada will keep this tool updated and available at guidelines.diabetes.ca . Updated December 2017 416584					

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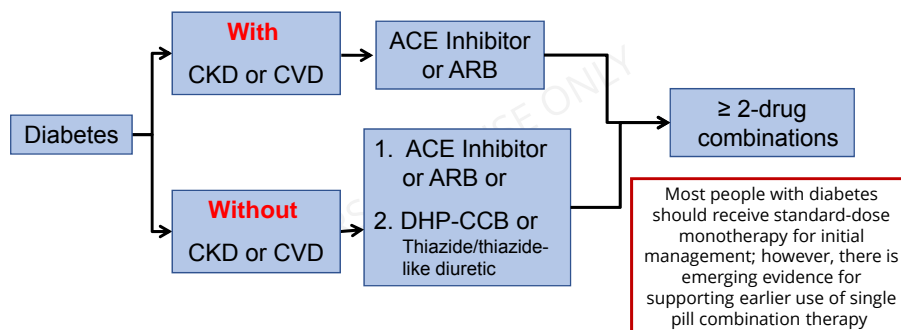
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- ✓ **D** • Drugs to protect the heart
 - A – ACEi or ARB | S – Statin | A – ASA if indicated | SGLT2i/GLP-1 RA
 - with demonstrated CV benefit if type 2 DM with CVD and A1C not at target
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2018 Diabetes Canada CPG – Chapter 26. Hypertension

Pharmacotherapy for Hypertension in Patients with Diabetes – Summary

Threshold $\geq 130/80$ mmHg and Target $< 130/80$ mmHgCheck serum **potassium** and **creatinine** at **baseline** and **within 1 to 2 weeks** of initiation of an **ACEi or ARB****Combinations** of agents that block the **RAAS** (ACEi, ARB, DRI) **should not be used****More than 3 drugs** may be needed to reach target values for people with diabetes

CKD, chronic kidney disease; CVD, cardiovascular disease; DHP-CCB, dihydropyridine calcium channel blocker; DRI, direct renin inhibitor



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Who Should Receive Statins?

(regardless of baseline LDL-C)

- **Cardiovascular disease** *or*
- **Age ≥ 40 yrs** *or*
- **Microvascular complications** *or*
- **DM > 15 yrs duration and age > 30 yr** *or*
- **Warrants therapy** based on the 2016 *Canadian Cardiovascular Society Guidelines for the Diagnosis and Treatment of Dyslipidemia*

Among women with childbearing potential, statins should only be used in the presence of proper preconception counselling & reliable contraception. Stop statins prior to conception.

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Antihyperglycemic therapy selection

In adults with type 2 diabetes with **clinical CVD** in whom glycemic targets are not achieved with existing antihyperglycemic medication(s) and with **eGFR >30 mL/min/1.73m²**, an antihyperglycemic agent with demonstrated **CV outcome benefit should be added** to reduce the risk of major CV events [Grade A, Level 1A for **empagliflozin**; Grade A, Level 1A for **liraglutide**; Grade C, Level 2 for **canagliflozin**]

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- ✓ **S** • Self-management, stress and other barriers

2018 Diabetes Canada CPG – Chapter 10. Physical Activity

Tools available at guidelines.diabetes.ca

guidelines.diabetes.ca/selfmanagementeducation/pa-tool

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Physical Activity Interactive Decision Tool

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Healthcare Provider Tool

Complete this tool to help assess your patient's physical ability and level of motivation to start and/or progress through a physical activity program.

When your patient is succeeding at being physically active you can encourage progression through an increase in the duration of physical activity followed by a progression in intensity, provided they are not limited by angina or other medical issues. The recommendations and programs from the use of this tool should be reassessed at regular intervals - every 6 - 12 months.

▼ STEP 1: Pre-Activity Screening [Reset](#)

Does your patient currently have symptoms of angina that would limit participation in physical activity ¹ (such as chest pain or severe pressure on physical exertion)?

☐ Yes ☐ No

▼ STEP 2: Physical Activity Level and Participation

Please complete step 1

▼ STEP 3: Physical Activity Program Recommendation

Please complete step 2

2018 Diabetes Canada CPG – Chapter 10. Physical Activity

Smarter Step Count Prescription

Health-care Provider Guidelines				
Suggest step count increments at each clinic visit to reach goal (an increase of 3,000 steps per day above the baseline step count within 1 year).				
Steps per day at baseline	Clinic Visit 1 (0 months)	Clinic Visit 2 (3 months)	Clinic Visit 3 (6 months)	Clinic Visit 4 (9 months)
<5,000	+500	+750	+750	+1,000
5,000–7,499	+750	+1,000	+1,250	
≥7,500	+1,000/+2,000	+1,000		

Start Date: _____

Patient's Name: _____

Baseline Step Count: _____ Recommended Step Count: _____

Clinic Visit Number: _____ Start Date for Step Increase: _____

Patient Step Count Prescription	
Today's Date: _____	
Patient's Name: _____	
• Please try to complete at least _____ steps per day until your next clinic visit. • Please record your step counts in your log sheets at the end of each day. • Please bring your log sheets to the next clinic visit which will be in about three months.	
Physician's Name: _____	
Physician's Signature: _____	

2018 Diabetes Canada CPG – Appendix 4

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Nutritional management of hyperglycemia in type 2 diabetes

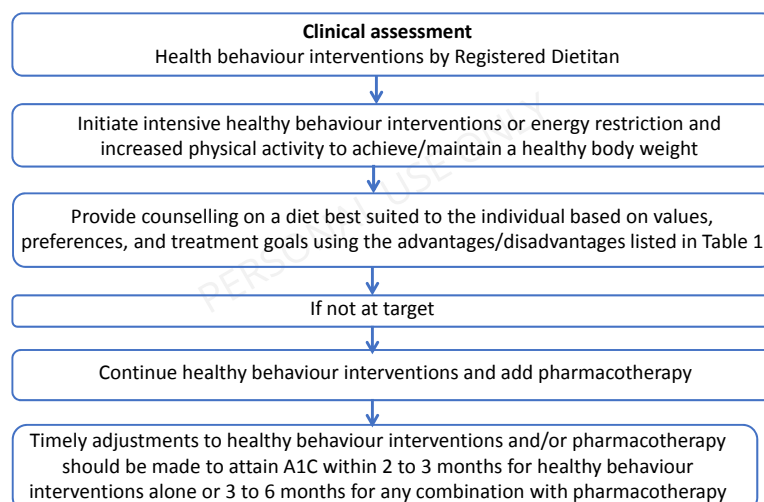


Table 1. Properties of dietary interventions

Properties of dietary interventions (listed in the order they are presented in the text)

Dietary interventions	A1C	CV benefit	Other advantages	Disadvantages
Macronutrient-based approaches				
Low-glycemic-index diets	↓	↓CVD	↓LDL-C, ↓CRP, ↓hypoglycemia, ↓diabetes Rx	None
High fibre diets	↓ (viscous fibre)	↓CVD	↓LDL-C, ↓non-HDL-C, ↓apo B (viscous fibre)	GI side effects (transient)
High MUFA diets	↔	↓CVD	↓Weight, ↓TG, ↓BP	None
Low-carbohydrate diets	↔	-	↓TG	↓Micronutrients, ↑renal load
High-protein diets	↓	-	↓TG, ↓BP, preserve lean mass	↓Micronutrients, ↑renal load
Mediterranean dietary pattern	↓	↓CVD	↓retinopathy, ↓BP, ↓CRP, ↑HDL-C	None
Alternate dietary patterns				
Vegetarian	↓ (144,249)	↓CHD (151)	↓Weight (147), ↓LDL-C (148)	↓vitamin B12
DASH	↓ (250)	↓CHD (159)	↓Weight, ↓LDL-C (157,250), ↓BP (28,157), ↓CRP	None
Portfolio	-	↓CVD (160,161)	↓LDL-C (160,161), ↓CRP, ↓BP	None
Nordic	-	-	↓LDL-C+↓non-HDL-C (167-169)	None
Popular weight loss diets				
Atkins	↔	-	↓Weight, ↓TG, ↑HDL-C, ↓CRP	↑LDL-C, ↓micronutrients, ↓adherence
Protein Power Plan	↓	-	↓Weight, ↓TG, ↑HDL-C	↓Micronutrients, ↓adherence, ↑renal load
Ornish	-	-	↓Weight, ↓LDL-C, ↓CRP	↔ FPG, ↓adherence
Weight Watchers	-	-	↓Weight, ↓LDL-C, ↑HDL-C, ↓CRP	↔ FPG, ↓adherence
Zone	-	-	↓Weight, ↓LDL-C, ↓TG, ↑HDL-C	↔ FPG, ↓adherence
Dietary patterns of specific foods				
Dietary pulses/legumes	↓ (174)	↓CVD	↓Weight (177), ↓LDL-C (175), ↓BP (176)	GI side effects (transient)
Fruit and vegetables	↓ (181,182)	↓CVD (78)	↓BP	None
Nuts	↓ (186)	↓CVD (142)	↓LDL-C (187,251), ↓TG, ↓FPG	Nut allergies (some individuals)
Whole grains	↓ (oats)	↓CHD (98)	↓LDL-C, FPG (oats, barley)	GI side effects (transient)
Dairy	↔	↓CVD (147,196)	↓BP, ↓TG (when replacing SSBs)	Lactose intolerance (some individuals)
Meal replacements	↓	-	↓Weight	Temporary intervention

* ↓ t = <1% decrease in A1C.

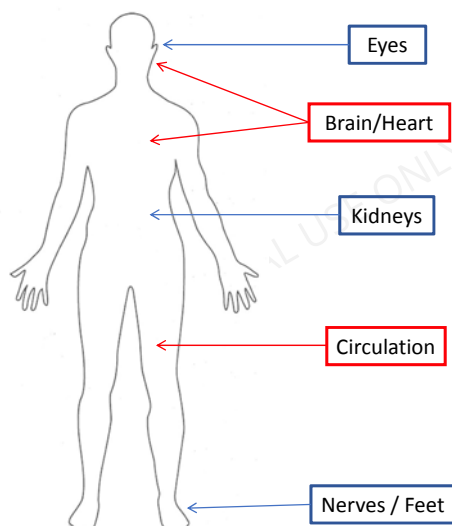
† Adjusted for medication changes.

A1C, glycated hemoglobin; apo B, apolipoprotein B; BMI, body mass index; BP, blood pressure; CHD, coronary heart disease; CHO, carbohydrate; CRP, C reactive protein; CV, cardiovascular; CVD, cardiovascular disease; DASH, dietary approaches to stop hypertension; FPG, fasting plasma glucose; GI, gastrointestinal; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; MUFA, monounsaturated fatty acid; SSBs, sugar-sweetened beverages; TG, total cholesterol; TG, triglycerides.

ABCDE³ of Diabetes Care

- ✓ **A** • A1C – optimal glycemic control (usually $\leq 7\%$)
- ✓ **B** • BP – optimal blood pressure control ($<130/80$)
- ✓ **C** • Cholesterol – LDL <2.0 mmol/L or $>50\%$ reduction
- ✓ **D** • Drugs to protect the heart
 A – ACEi or ARB | S – Statin | A – ASA if indicated | SGLT2i/GLP-1 RA
 with demonstrated CV benefit if type 2 DM with CVD and A1C not at target
- ✓ **E** • Exercise / Healthy Eating
- ✓ **S** • Screening for complications
- ✓ **S** • Smoking cessation
- ✓ **S** • Self-management, stress and other barriers

Screening for complications



Type of diabetes: <input type="checkbox"/> Type 1 <input type="checkbox"/> Type 2 <input type="checkbox"/> Other		Date Diagnosed:		Patient Name:	
Comorbidities: <input type="checkbox"/> Hypertension <input type="checkbox"/> Coronary artery disease <input type="checkbox"/> Stroke/TIA <input type="checkbox"/> Dyslipidemia <input type="checkbox"/> Peripheral arterial disease <input type="checkbox"/> Depression/Anxiety <input type="checkbox"/> CKD - stage _____ <input type="checkbox"/> Other(s): _____				Date of Birth: _____	
Weight (kg): _____ Height (cm): _____ BMI: _____ Waist circumference (cm): _____		Date: _____ Time: _____ Wt: _____ Ht: _____ BMI: _____ WC: _____		Date: _____ Time: _____ Wt: _____ Ht: _____ BMI: _____ WC: _____	
Nutrition Physical Activity (Average 150 minutes/week, Resistance 2-3x/week) <input type="checkbox"/> Non-smoker <input type="checkbox"/> Ex-smoker <input type="checkbox"/> Smoker				Smoking Status <input type="checkbox"/> Non-smoker <input type="checkbox"/> Ex-smoker <input type="checkbox"/> Smoker	
A1C (target: <7% or _____ %) (Individualized based on patient characteristics and antihyperglycemic medications) - see CPG, (q) months if at target and stable - (q) months if not at target				Test Date: _____ Result: _____	
Antihyperglycemic Medication(s) Drug Name(s)/Dose(s): _____				Test Date: _____ Result: _____	
Therapy Adherence/Concerns BG Record (Targets: premeal: 4-7 mmol/L or _____ mmol/L; 2hr postmeal: 5-10 mmol/L or _____ mmol/L) (Individualized based on ability to achieve A1C target + risk of hypoglycemia)				Test Date: _____ Result: _____	
Hypoglycemic Episodes (Frequency/pattern/driving risk)				Test Date: _____ Result: _____	
BP (Target: <130/80 mm Hg, 3 readings recommended)				Test Date: _____ Result: _____	
Antihypertensive(s) Drug Name(s)/Dose(s): _____				Test Date: _____ Result: _____	
CVD Symptoms (angina, decreased exercise tolerance, SOB, HF symptoms, claudication)				<input type="checkbox"/> None <input type="checkbox"/> Yes	
Resting ECG, every 3-5 yrs (If any: age >40 yrs, duration of diabetes >10 yrs + age >60 yrs; non-targeted imaging, abnormal ECG)				Date: _____ Report: _____	
Lipids (Primary target: LDL-C <2.6 mmol/L or >50% reduction in LDL-C or non-HDL-C <2.6 mmol/L or age >40 yrs)				LDL-C: _____ non-HDL-C: _____ test date: _____	
Lipid-lowering Therapy (Statins +/- 2nd line agent(s))				<input type="checkbox"/> Continue <input type="checkbox"/> Start <input type="checkbox"/> No - reason: _____	
ACE inhibitor/ARB Drug Name(s)/Dose(s): _____				<input type="checkbox"/> Continue <input type="checkbox"/> Start <input type="checkbox"/> No - reason: _____	
(If any: clinical CVD, age >40 yrs, age >60 yrs with an additional CV risk factor or end organ damage, albuminuria, microalbuminuria, CKD, atherosclerotic complications)				<input type="checkbox"/> Continue <input type="checkbox"/> Start <input type="checkbox"/> No - reason: _____	
Antihyperglycemic Agent with Demonstrated CV Outcome Benefit Drug Name(s)/Dose(s): _____				<input type="checkbox"/> Continue <input type="checkbox"/> Start <input type="checkbox"/> No - reason: _____	
(If type 2 DM with clinical CVD not at glycemic target - empagliflozin, liraglutide, canagliflozin)				<input type="checkbox"/> Not indicated <input type="checkbox"/> Yes	
Antidiabetic Agent(s) Drug Name(s)/Dose(s): _____				<input type="checkbox"/> Not indicated <input type="checkbox"/> Yes	
(If established CVD; consider if additional CV risk factors)				<input type="checkbox"/> Not indicated <input type="checkbox"/> Yes	
Urine ACR (normal <2 mg/mmol)				Test Date: _____ Result: _____	
Serum Creatinine/eGFR				Test Date: _____ Result: _____	
CKD				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Dilated Eye Exam (type 1 - annually; type 2 - q1-2 years or as recommended by vision care professional)				Date of last visit: _____ Reminded: _____	
Retinopathy				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Severity/Therapies				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Neuropathy Symptoms (e.g. pain, paresthesia, GI symptoms, sexual dysfunction)				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Diabetic Foot Exam (includes 10 g monofilament or 128 Hz tuning fork; structural abnormalities, skin changes, pulses)				Sensation: _____ Pulses: _____ Skin: _____ Other: _____	
(annually for screening; every visit if diabetic foot complications) See Appendices T1A, T1B and T2				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Neuropathy				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Screens for Depression, Anxiety, Other Stressors (consider use of PHQ-9, GAD-7)				Concerns: _____	
Influenza (annual)				<input type="checkbox"/> No <input type="checkbox"/> Yes Date: _____	
Pneumococcal (once repeat if >65 yrs)				<input type="checkbox"/> No <input type="checkbox"/> Yes Date: _____	
Patient Goals (e.g. coverage, accessibility, competing demands)				<input type="checkbox"/> Yes <input type="checkbox"/> No Reasons: _____	
Women: Contraception/monoclonation planning				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Driving Guidelines Reviewed				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Sick-day Management (advice holding metformin, SGLT2s, SLA2/ARB, diuretics, NSAIDs if inadequate fluid intake and IB)				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Referrals Made				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Changes to Medications or Other Management				<input type="checkbox"/> Yes <input type="checkbox"/> No	
Resources provided				<input type="checkbox"/> Yes <input type="checkbox"/> No	
RECALL (usually q3-4 months)				<input type="checkbox"/> Appointment given <input type="checkbox"/> Appointment given <input type="checkbox"/> Appointment given <input type="checkbox"/> Noted in recall system	
(If any: clinical CVD, age >40 yrs with an additional CV risk factor or end organ damage, albuminuria, microalbuminuria, CKD, atherosclerotic complications)				<input type="checkbox"/> Noted in recall system	

For additional diabetes management resources, visit www.guidelines.diabetes.ca.

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2018 Diabetes Canada CPG – Chapter 29. Chronic Kidney Disease in Diabetes

Chronic Kidney Disease (CKD) Checklist

- ✓ **SCREEN** with random urine albumin creatinine ratio (ACR) and serum creatinine for estimated glomerular filtration rate (eGFR) **at diagnosis** then **annually** (T2D)
- ✓ **DIAGNOSE** with repeat confirmed ACR ≥ 2.0 mg/mmol and/or eGFR <60 mL/min
- ✓ **DELAY** onset and/or progression with glycemic and blood pressure control and ACEi or ARB
- ✓ **PREVENT** complications with dose adjustment, “sick day management” counselling and referral when appropriate

ACEi, angiotensin converting enzyme inhibitor; ARB, angiotensin receptor blocker

Counsel all Patients About

Sick Day Medication List

Visit
guidelines.diabetes.ca
for patient
handout

Instructions for Healthcare Professionals:

If patients become ill and are unable to maintain adequate fluid intake, or have an acute decline in renal function (e.g. due to gastrointestinal upset or dehydration), they should be instructed to hold medications which will:

A) Increase risk for a decline in kidney function:

- Angiotensin-converting enzyme inhibitor
- Angiotensin receptor blockers
- Direct renin inhibitors
- Non-steroidal anti-inflammatory drugs
- Diuretics
- SGLT2 inhibitors

B) Have reduced clearance and increase risk for adverse effects:

- Metformin
- Sulfonylureas (gliclazide, glimepiride, glyburide)

S sulfonylureas
A ACE-inhibitors
D diuretics, direct renin inhibitors

M metformin
A angiotensin receptor blockers
N non-steroidal anti-inflammatory
S SGLT2 inhibitors

Please complete the following card and give it to your patient.

Patients should be instructed that increased frequency of self blood glucose monitoring will be required and adjustments to their doses of insulin or oral antihyperglycemic agents may be necessary.

Instructions for Patients

When you are ill, particularly if you become dehydrated (e.g. vomiting or diarrhea), some medicines could cause your kidney function to worsen or result in side effects.

If you become sick and are unable to drink enough fluid to keep hydrated, you should **STOP** the following medications:

- Blood pressure pills
- Water pills
- Metformin
- Diabetes pills
- Pain medications
- Non-steroidal anti-inflammatory drugs (see below)

Please be careful not to take non-steroidal anti-inflammatory drugs (which are commonly found in pain medications (e.g. Advil) and cold remedies).

Please check with your pharmacist before using over-the-counter medications and discuss all changes in medication with your healthcare professional.

Please increase the number of times you check your blood glucose levels. If they run too high or too low, contact your healthcare professional.

If you have any problems, you can call:

2018

Role of SGLT2 inhibitor in CKD

In adults with type 2 diabetes with **clinical CVD** in whom **glycemic targets are not achieved** with existing antihyperglycemic medication(s) and who have an eGFR >30 mL/min/1.73 m², an **SGLT2 inhibitor with proven renal benefit** may be considered to reduce progression of nephropathy [Grade B, Level 2 for empagliflozin; Grade C, Level 3 for canagliflozin]

2018 Diabetes Canada CPG – Chapter 32. Foot Care

Foot Care Checklist

- ✓ **EDUCATE** about proper foot care
- ✓ **EXAMINE** for structural, vascular, neuropathy problems **at diagnosis** then **annually**
- ✓ **DO** a 10 gram monofilament assessment
- ✓ **IDENTIFY** those at high risk of foot ulcers and educate, assess **more frequently**, consider footwear
- ✓ **REFER** persons with foot ulcers and other complications to those specialized in foot care

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2018 Diabetes Canada CPG – Chapter 33. Sexual Dysfunction & Hypogonadism in Men with Diabetes

Erectile Dysfunction (ED) Checklist

- ✓ **SCREEN** all adult men regularly with sexual function history
- ✓ **TREAT** erectile dysfunction with PDE-5 inhibitor as first-line therapy (if no contraindication)
- ✓ **INVESTIGATE** for hypogonadism if men with ED do not respond to PDE-5 inhibitor therapy

PDE-5, phosphodiesterase type 5

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ABCDE³ of Diabetes Care

- ✓ **A** • A1C – optimal glycemic control (usually $\leq 7\%$)
- ✓ **B** • BP – optimal blood pressure control ($< 130/80$)
- ✓ **C** • Cholesterol – LDL < 2.0 mmol/L or $> 50\%$ reduction
- ✓ **D** • Drugs to protect the heart
 A – ACEi or ARB | S – Statin | A – ASA if indicated | SGLT2i/GLP-1 RA
 with demonstrated CV benefit if type 2 DM with CVD and A1C not at target
- ✓ **E** • Exercise / Healthy Eating
- ✓ **S** • Screening for complications
- ✓ **S** • Smoking cessation
- ✓ **S** • Self-management, stress and other barriers

ABCDE³ of Diabetes Care

- ✓ **A** • A1C – optimal glycemic control (usually $\leq 7\%$)
- ✓ **B** • BP – optimal blood pressure control ($< 130/80$)
- ✓ **C** • Cholesterol – LDL < 2.0 mmol/L or $> 50\%$ reduction
- ✓ **D** • Drugs to protect the heart
 A – ACEi or ARB | S – Statin | A – ASA if indicated | SGLT2i/GLP-1 RA
 with demonstrated CV benefit if type 2 DM with CVD and A1C not at target
- ✓ **E** • Exercise / Healthy Eating
- ✓ **S** • Screening for complications
- ✓ **S** • Smoking cessation
- ✓ **S** • Self-management, stress and other barriers

2018 Diabetes Canada CPG – Chapter 18. Diabetes and Mental Health

Screening for depressive and anxious symptoms is important in patients with diabetes

Purpose	Screening Tools
Diabetes-specific	<ul style="list-style-type: none"> • Problem Areas in Diabetes (PAID) Scale • Diabetes Distress Scale (DDS)
Quality of Life	<ul style="list-style-type: none"> • WHO-5
Depression/Anxiety	<ul style="list-style-type: none"> • Hospital Anxiety and Depression Scale (HADS) • Patient Health Questionnaire (PHQ-9) • Beck Depression Inventory (BDI)

Websites with psychological / psychiatric scales:

www.phqscreeners.com
www.outcometracker.org/scales_library.php.

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Formulate an Action Plan to support Self-management

guidelines.diabetes.ca/self-management/education/self-management-actionplan

Managing My Diabetes – My Action Plan

Date: _____

The change I want to make happen is: _____

My goal for the next month is: _____

Action Plan:
 The specific steps I will take to reach my goal (what, when, where, how often): _____

Things that could make it difficult to achieve my goal: _____

My plan for overcoming these challenges are: _____

Support and resources I will need: _____

How **important** is it to me that I achieve my goal?
 (scale of 0 to 10, with 0 being not important at all and 10 being extremely important): _____

How **confident** am I that I can achieve my goal?
 (scale of 0 to 10, with 0 being not confident at all and 10 being extremely confident): _____

Follow-up date: _____

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Other considerations

2018 Diabetes Canada CPG – Chapter 21. Diabetes & Driving

Diabetes and Driving

- The **fitness** of people with diabetes to drive should be assessed on an **individual** basis
- All drivers with diabetes should undergo a **medical examination** at least **every two years** to assess fitness to drive.
- **People with diabetes** should play an **active role in assessing their fitness to drive**
- Should **not drive** when **BG <4.0 mmol/L** and should **wait at least 40 minutes** after treatment of hypoglycemia has increased their BG level to **at least 5.0 mmol/L**

Special Populations

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2018 Diabetes Canada CPG – Chapter 36. Diabetes and Pregnancy

Preconception Checklist for Women with Pre-existing Diabetes

2018

- ✓ Use **reliable birth control** until adequate glycemic control
- ✓ Attain a **preconception A1C of $\leq 7.0\%$** ($\leq 6.5\%$ if safe)
- ✓ May remain on **metformin + glyburide** until pregnancy, otherwise **switch to insulin**
- ✓ Assess for and manage any **diabetes complications**
- ✓ **Folic Acid 1 mg/d**: 3 months pre-conception to at least 12 weeks gestation
- ✓ **Discontinue** potential embryopathic meds:
 - ✓ **ACE-inhibitors / ARB** (prior to or upon detection of pregnancy in those with significant proteinuria)
 - ✓ **Statin** therapy

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2018 Diabetes Canada CPG – Chapter 37. Diabetes in Older People

Diabetes in the Elderly Checklist

2018

- ✓ **ASSESS** for level of functional dependency (frailty)
- ✓ **INDIVIDUALIZE** glycemic targets based on the above (A1C $\leq 8.5\%$ for frail elderly) but if otherwise healthy, use the same targets as younger people
- ✓ **AVOID** hypoglycemia in cognitive impairment
- ✓ **SELECT** or **ADJUST** antihyperglycemic therapy carefully
 - ✓ Caution with sulfonylureas or thiazolidinediones
 - ✓ DPP-4 inhibitors should be used over sulfonylureas
 - ✓ Basal analogues instead of NPH or human 30/70 insulin
- ✓ **GIVE** regular diets instead of “diabetic diets” or nutritional formulas in nursing homes

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2018 Diabetes Canada CPG – Chapter 38. Type 2 Diabetes and Indigenous Peoples

Type 2 Diabetes and Indigenous Peoples

2018

- ✓ Among the **highest-risk** populations
- ✓ **Prevention** strategies are **essential**
- ✓ Management **targets should be no different** from general population
- ✓ Focus on building a **therapeutic relationship**
- ✓ **Acknowledge** the legacy of colonization and its ongoing adverse effects on Indigenous health
- ✓ Use the **Educating for Equity (E4E) framework** to address social barriers and identify strategies for facilitating outcomes using a cultural approach

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Diabetes Canada is helping you provide patient-centred diabetes care and chronic disease management.

NEW 2018 Guidelines Available



Reduce the Risk of Diabetes Complications



Keep Patients Safe



Support Self-management

Interactive Tools

PHYSICAL ACTIVITY DECISION TOOL

SCREENING FOR AND DIAGNOSING DIABETES

SELF-MONITORING BLOOD GLUCOSE

REDUCING VASCULAR RISK

PHARMACOTHERAPY FOR TYPE 2 DIABETES

INDIVIDUALIZING YOUR PATIENT'S ATC TARGET


News & Events

Diabetes Canada recently released the 2018 Clinical Practice Guidelines For the Prevention and Management of Diabetes in Canada


The Canadian Diabetes Association has changed its name to Diabetes Canada! Find out more


The Importance of a Healthy Mind
Dealing with the Psychosocial Aspects of Diabetes


Or download the App




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Diabetes Canada Clinical Practice Guidelines

<http://guidelines.diabetes.ca> – for health-care
providers

1-800-BANTING (226-8464)

<http://diabetes.ca> – for people with diabetes

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