Indigenous Services Services aux

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Name:									□ Type			/pe 1	Age a	at diagnosis	:	
Date of bir	rth:			□ Ma	ale		□ Fema	ale	Medica	l histo	ory:					
Band #:																
Healthcare	e Card #:							4	Allergi	es:						
				Ris	k Fact	ors a	& Co-mor	bidities	s (chec	k off al	ll applic	able)				
□ Hyperte	ension	□ Chror	nic kidr					ronary	_				heral	artery disea	ase	)
<ul> <li>Dyslipio</li> </ul>		□ Polyc	ystic o	varian	syndr	ome		ectile d					al illn	ess - dx:		
□ Retinop	oathy	□ Subst	ance r					bacco						or quit-date	э:	)
			DATE	Ro	outine i	Diab	etes Ass	essmer	it (eve	ry 3 to	6 mon	ths)				
BP		l	DATE												$\dashv$	
Di .	<	<130/80 fo	r most													
Wt (kg) / B	BMI / WC (															
	1110 400	BMI 18.														
A1C	WC <102	cm ♂; <88	scm ¥													
	0% for mos	st, or														
Blood Glu																
	nquire abou st, premea															
	ır postmeal															
Foot chec																
screening)	Any rodr	ace one	3 2ro2													
b	lister, sign	ness, oper of infection														
Self-mana																
client																
Nutrition		Inoto rofo	rrole)													
	n control/C	note refe														
	Healthy	dietary pa														
Physical A		- >450														
R	Aerobi esistance 3	c ≥150mir 3 sessions														
Smoking (			,													
		(if appli	cable)													
Psychoso Assess for	Cıal r diabetes-⊦	ralatad dis	etrace													
	on, anxiety,															
			etc.													
Medication		<b>v</b> e any char	200(c)													
Inquire abo																
, , ,		□ No change			□ No change		□ No change		ge	□ No change		ge □ No change		□ No change		
See progre	ess notes															
		(√ if appli	cable)												_	
		Sian	ature													
	Screenii			Compl	ication	ıs (a	nnually o	r as inc	dicated	l, but l	abworl	k must be	orde	ered by NP o	r M	ID)
Nephropat	thy				Neur	•	-						Reti	nopathy	٧	accinations
Date	AC	R	eG	FR			ssment (a	ccording	to BPG	is)			Ann	ual eye	F	lu (annual):
					Date:		betes foo	 t accac	emant f	form			exar	•		ate:
							frequenc				erv 6 r	nonths	Date	ı:		
							3 months						Date	•	0	ate:
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For vers	lor prets-	tion:			Date:		gets: If ind	Finding	gs:	I DL C	2 40 0	mmel/l	6-14	monite-i		Pland
For vascul  ☐ Statins if	-		> 30 va	are	-		reduction							-monitoring	Oī	B1000
and >15 yea					Date		dication	LDL	HDL	TG	(non-	(Apo		ual meter-to-	lab	comparison
disease											HDL- C)	В)	_	9:		•
□ ACEi/ARI organ dama;				nd							٥,					
CVD Asse		ovasculai (	aioca5 <del>C</del>										Review client techr (have client demonsti			
ECG:													meter)			IC WILLI OWIT
Stress ECG														e:		
Other:						Ì		1								

Care	Objective	Target
Self-monitoring	Ensure patient can use glucose meter, interpret SMBG results and modify	Premeal (mmol/L) = $4.0-7.0$ for most patients
of blood glucose	treatment as needed. Develop an SMBG schedule with patient and review	2-hour postmeal (mmol/L) = $5.0-10.0$ for most
(SMBG)	records. Inquire about hypoglycemia at each visit: estimate of cause,	patients
	frequency, symptoms, recognition, severity of treatment and driving	5.0–8.0 if not achieving A1C target
Blood glucose	Measure A1C every 3 months for most adults.	A1C ≤7.0% for most patients. Individualized
control	Consider testing at least every 6 months in adults during periods of	based on life expectancy, functional
	treatment and lifestyle stability when glycemic targets are being	dependency, extensive coronary artery disease
	consistently achieved.	at high risk of ischemia, multiple comorbidities,
		recurrent severe hypoglycemia, hypoglycemia
		unawareness, longstanding diabetes unable to
		achieve A1C ≤7% despite best efforts
		(including intensifying insulin)
Hypoglycemia	Enquire about hypoglycemia at each visit. Discuss recognition and treatment	Avoidance of hypoglycemia especially in the
	of hypoglycemia and risk/benefit of hypoglycemia and pharmacologic	elderly, those with hypoglycemia unawareness,
	management.	and those with criteria for less stringent control.
Blood glucose meter	Compare meter results with laboratory measurements at least annually, and	Simultaneous fasting glucose/meter lab
accuracy	when indicators of glycemic control do not match meter.	comparison within 20%
Hypertension	Measure BP at diagnosis of diabetes and at every diabetes clinic visit.	<130/80 mm Hg
Waist circumference	Measure as an indicator of abdominal fat.	Central obesity defined as:
		WC M ≥102cm; F ≥88cm (North America)
Body mass index	Calculate BMI (mass in kg/height in m <sup>2)</sup>	Healthy body weight target: BMI: 18.5–24.9
(BMI)		kg/m <sup>2</sup>
Nutrition	Encourage nutritional therapy (by a Registered Dietician) as an integral part of	Meet nutritional needs following <i>Canada's</i>
	treatment and self-management.	Food Guide
Physical activity	Discuss and encourage aerobic and resistance exercise. Evaluate those with	Aerobic: ≥150 minutes/week
	possible CAD or microvascular complications undertaking exercise	Resistance: 3 sessions/week
	substantially more vigorous than brisk walking.	
Smoking	Encourage patient to stop at each visit; provide support as needed	Smoking cessation
Chronic kidney	Identification of CKD requires screening for <b>proteinuria</b> using random urine	Normal ACR <2.0 mg/mmol
disease (CKD)	ACR (2 out of 3 samples over 3 months) and assessment of renal function	Normal eGFR >60mL/min
, , ,	using a serum creatinine converted to <b>eGFR</b> . <b>Type 1 diabetes</b> - screen at 5	Give patients with CKD "sick day" medication
	years duration and then annually if no CKD. <b>Type 2 diabetes</b> - Screen at	list that outlines which meds to hold during
	diagnosis and then yearly if no CKD.	episodes of acute illness
Retinopathy	<b>Type 1 diabetes</b> - Screen age ≥15 and 5 years duration, then rescreen yearly.	Early detection and treatment
rtomopumy	Type 2 diabetes - Screen at diagnosis then every1–2 years if no retinopathy	
	present. The interval for follow-up assessment should be tailored to the	
	severity of the retinopathy. Screening should be conducted by an experienced	
	eye care professional.	
Neuropathy/Foot	<b>Type 1 diabetes</b> - Screen after 5 years postpubertal duration then annually, or	Early detection and treatment
Examination	more frequently if high risk.	If peripheral neuropathy present: require foot
	<b>Type 2 diabetes</b> - Screen at diagnosis then annually, or more frequently if	care education, specialized footwear, smoking
Using a tool such as	high risk. Screen for neuropathy assessing: history of current or previous foot	cessation, and intensive glycemic control.
Inlow's 60-second	ulcers; sensation with 10-g monofilament or 128-Hz tuning fork at dorsum of	If ulcer present: manage by multidisciplinary
Diabetic Foot	great toe; structural and biomechanical abnormalities; circulation; and self-	team with expertise
Screen	care behaviour and knowledge.	team with emperiuse
Coronary Vascular	Conduct CCD risk assessment periodically: CV history, lifestyle, duration	Vascular protection: first priority in
Disease (CCD)	of diabetes, sexual function, abdominal obesity, lipid profile, BP, reduced	prevention of diabetes complications is
Discuse (CCD)	pulses, bruits, glycemic control, retinopathy, eGFR, ACR.	reduction of CV risk by vascular protection
	Baseline ECG and every 3-5 years if: age > 40 years; > 30 years and	through a comprehensive multifaceted
	duration of diabetes > 15 years; end organ damage; \ge 1 CVD risk factors.	approach.
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	Low dose ASA therapy in those with established CVD.	All people with DM: optimize BP, glycemic
		control and lifestyle.
		Statin if: CVD, age ≥40 years OR
		macrovascular disease OR microvascular
		disease OR long duration of DM (DM >15
		years and age >30 years). ACEi or ARB if:
		CVD, age \ge 55 years with additional CV risk
		factor OR EOD OR microvascular
		complications. Lipid targets for those who need therapy:
Developidan-i-	Footing on non-footing limid levels (TO HDI TO aslandated I DI 6/1	LI LING PARGERS FOR THOSE WHO BEED THERSING
Dyslipidemia	Fasting or non-fasting lipid levels (TC, HDL, TG, calculated LDL, &/or	
Dyslipidemia	apoB or non-HDL) at diagnosis, then repeat every 1-3 years based on CV	Primary target: LDL <2.0 mmol/L or >50%
Dyslipidemia	<b>apoB or non-HDL</b> ) at diagnosis, then repeat every 1-3 years based on CV risk. Repeat testing in 3-6 months after treatment initiation to verify if	Primary target: LDL <2.0 mmol/L or >50% reduction
Dyslipidemia	apoB or non-HDL) at diagnosis, then repeat every 1-3 years based on CV	Primary target: LDL <2.0 mmol/L or >50%

encourage self-management and use of interdisciplinary team approach to attain care objectives; 3) schedule diabetes-focused visits; and 4) use diabetes patient care flow sheets and systematic recall for visits.