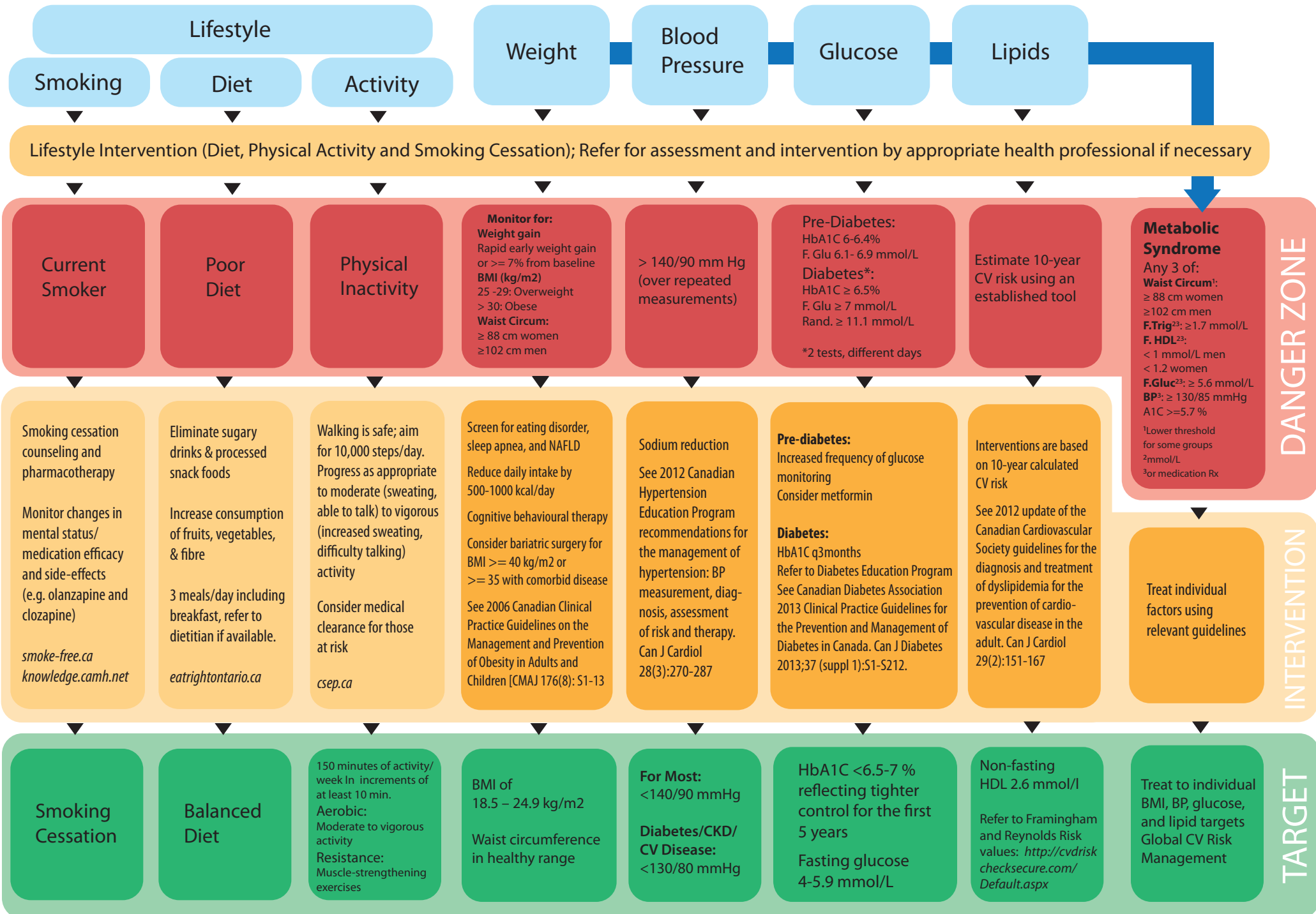


Cardiometabolic Risk Management

Antipsychotics (Mood Stabilizers & Antidepressants*)

*Despite less available evidence, similar principles apply to these medication classes



Monitoring: How Often and What to Do

Applies to patients prescribed antipsychotics and metabolically active mood stabilizers and antidepressants

Frequency: As a minimum review those prescribed a new agent at baseline and at least once after 3 months. Weight should be assessed monthly in the first 3 months of taking a new antipsychotic as rapid early weight gain may predict severe weight gain in the longer term. Subsequent review should take place annually unless an abnormality of physical health emerges, which should then prompt appropriate action and/or continuing review at least every 3 months.

	Baseline	4 weeks	8 weeks	12 weeks	Quarterly	Annually
Personal/FHx	X					X
Lifestyle Review ¹	X	X	X	X	X	X
Weight/WC	X	X	X	X	X	X
BP	X			X		X
FPG/HbA1C	X			X		X
Lipid Profile ²	X			X		X

History:

Ask about family history (diabetes, obesity, CVD in first degree relatives <60 yrs), gestational diabetes. Note ethnicity.

¹Smoking, diet, and physical activity ²If fasting lipid profile cannot be obtained, a non-fasting sample is satisfactory
Derived from consensus guidelines 2004, *J clin. psych* 65:2

Investigations

Fasting estimates of plasma glucose (FPG), HbA1c, and lipids (total cholesterol, non-HDL, HDL, triglycerides). If fasting samples are impractical, then non-fasting samples are satisfactory for most measurements except for LDL and triglycerides.

Review of Antipsychotic Medication

Choose lower metabolic liability medication first-line when possible. Response in first episode psychosis is robust independent of agent.

Changing or discontinuing antipsychotic requires careful clinical judgment, balancing metabolic benefits against relapse risk.

Ideally psychiatrist supervised. Should be a priority if there is:

- Rapid weight gain (e.g. 5kg <3 months) following antipsychotic initiation.
- Rapid development (<3 months) of abnormal lipids, BP, or glucose.

The psychiatrist should consider whether the antipsychotic drug regimen has played a causative role in these abnormalities and, if so, whether an alternative regimen could be expected to offer less adverse effect:

If clinical judgment and patient preference support continuing with the same treatment then ensure appropriate further monitoring and clinical considerations.

Avoid antipsychotic polypharmacy when possible; Avoid off-label use of antipsychotics.

Specific Adjunctive Pharmacological Interventions

For mitigation of excess weight gain associated with antipsychotic use, the strongest evidence is for off-label use of Metformin and Topiramate (Mayan, 2010). Consider Metformin first due to better tolerability profile unless there is a co-morbid binge-eating disorder (McElroy, SI, 2009). Please be advised that off-label use requires documented informed consent. Discontinue if no sign of efficacy (continuing weight gain if used for weight loss or stabilization) after 3 months at therapeutic dose.

Metformin:

Weight Gain & Primary Prevention of

Diabetes: Start 250 mg po BID, titrate every 1-2 weeks as clinically indicated and tolerated. Dose range is 750 mg - 2 Gm/day

Caution with renal or hepatic impairment. Avoid excessive alcohol use. Monitor for GI side effects. Monitor for B12 deficiency.

Topiramate:

Weight Gain: Start 12.5 mg po BID; titrate by 25-50 mg per day in divided doses every 1-2 weeks as tolerated to a maximum of 100 mg po BID.

Caution with renal or hepatic impairment. Avoid excessive alcohol use. Monitor for cognitive changes. Paresthesias are common but generally well-tolerated.

**DON'T JUST
SCREEN:
INTERVENE!**

FOR ALL PATIENTS IN THE DANGER ZONE

The primary care provider and psychiatrist will work together to ensure appropriate monitoring and interventions are provided and communicated to avoid over or under monitoring. The primary care provider will usually lead on supervising the provision of physical health interventions. The psychiatrist will usually lead on decisions to significantly change antipsychotic medicines.



With thanks to the On Track FEP program for their collaboration

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