Glucose Control: non-insulin therapies*

The Society for Endocrinology, Metabolism and Diabetes of South Africa Type 2 Diabetes Guidelines Expert Committee.


About

This chapter summarises information for each of the non-insulin drug classes that are used for blood glucose control. Each summary is accompanied by a table of recommendations to guide the clinical use of these medications. For the sake of completeness, and for those that are interested, we have included a more detailed review of each drug as an appendix to each summary. These can be found in the Appendix section of the guidelines. The treatment recommendations for each drug have been incorporated into the treatment algorithm in Chapter 11. The following abbreviations are used in this chapter:

- DPP-4: dipeptidyl peptidase-4
- GLP-1: glucagon-like peptide-1
- SGLT2- sodium-glucose linked transporter-2

9.1: Drug Summary – Metformin

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to appendix 9.1 for full text and references)

<table>
<thead>
<tr>
<th>Mechanism of Action</th>
<th>Metformin targets the liver, skeletal muscle and the gut to reduce hepatic glucose output, increase skeletal muscle glucose uptake, increase GLP-1 levels and reduce glucose absorption.</th>
</tr>
</thead>
</table>
| Glycaemic efficacy and indications | Mean HbA1C reductions  
- As monotherapy vs. placebo: -1.1%  
- As add-on therapy to other non-insulin agents: -0.9%  
- Ass add-on to insulin: -0.6% |
| Cardiovascular outcome trials | Proven superiority vs. diet alone in obese patients - reduced all-cause mortality, diabetes related mortality and myocardial infarction, but not peripheral vascular disease or microvascular disease.  
Proven superiority vs. SU and insulin in obese patients: reduced all cause mortality and stroke, but not myocardial infarction, diabetes-related deaths, peripheral vascular disease or microvascular disease.  
Safety when added to SU’s (glibenclamide and chlorpropamide) is unclear. |
| Hypoglycaemia | No severe hypoglycaemia as monotherapy. Some patients may have symptoms of hypoglycaemia. Can potentiate the hypoglycaemic effect of insulin or insulin secretagogues. |
| Weight | Weight neutral or causes modest weight loss (-1.2kg). No weight loss in non-diabetic individuals. |
| Non-glycaemic benefits | Improves lipid profile.  
Reduces cancer rates in population studies.  
May improve outcomes in mild to moderate heart failure; Improves laboratory measures of inflammation, coagulation, oxidative stress, endothelial function and tumour suppression; cancer rates are lower. |
Glucose Control: non-insulin therapies

Side Effects and Precautions

Gastrointestinal (GI) side-effects are common, are not dose-dependent, and occur in 20-30% of patients (diarrhoea, nausea, vomiting, cramping, bloating and flatulence). Up 10% will discontinue therapy due to GI side effects. Switching to an extended release formulation improves GI tolerability and adherence.

Lactic Acidosis is rare with current usage (0.04 cases per 1000 patient years) and not different to non-metformin users. Metformin should be discontinued at the time of, or before an iodinated contrast imaging procedure or general anaesthesia in the following categories of patients: those with an eGFR < 60 ml/minute/1.73 m²; those with a history of liver disease, alcoholism, or heart failure; those who will receive intra-arterial iodinated contrast. Re-evaluate eGFR 48 hours after the procedure and restart metformin if renal function is stable and the patient is eating normally.

Can cause low levels of serum vitamin B12 in 7-30% of long-term users, but is rarely associated with the clinical features of vitamin B12 deficiency. The exact mechanism and significance is unknown. There is no recommendation to screen for vitamin B12 deficiency routinely. Investigate and manage vitamin B12 deficiency according to standard clinical practice, with a high index of suspicion in patients who are vegetarian or anaemic, or have peripheral neuropathy.

Dosing and prescribing

Metformin (standard release): Dose range is 500 mg/day to 3 000 mg/day in two or three divided doses with meals. Optimum glycaemic efficacy is achieved with 2 000 mg/day; few patients have additional glycaemic benefit with higher doses. The optimum dose for cardiovascular benefit in obese patients is 2 550mg/day.

Metformin extended-release: Dose range is 500 mg/day to 2 000 mg/day as a single dose with the evening meal. The 2 000mg dose can be split to 1000mg twice daily without losing efficacy.

Start with 500 mg/day of standard metformin tablets, and increase the dose by 500 mg every one to two weeks to minimise side effects. If GI side effects occur reduce the dose and re-titrate slowly. If GI disturbances persist try switching to the extended-release formulation.

Renal dosing

<table>
<thead>
<tr>
<th>eGFR ≥60 ml/min</th>
<th>45-60 ml/min</th>
<th>30-45 ml/min</th>
<th>&lt; 30 ml/min</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard dosing.</td>
<td>Standard dosing.</td>
<td>Maximum dose: 1000 mg/day</td>
<td>Metformin is contraindicated</td>
</tr>
<tr>
<td>Monitor eGFR annually</td>
<td>Monitor eGFR 3 to 6 monthly</td>
<td>Monitor eGFR 3 to 6 monthly</td>
<td></td>
</tr>
</tbody>
</table>

SEMDSA 2017 Recommendations for metformin

- Initiate standard-release metformin therapy in all newly diagnosed obese patients with type 2 diabetes. (A)
- Initiate standard-release metformin therapy in all newly diagnosed non-obese patients with type 2 diabetes. (C)
- Dosing: Start with 500 mg once daily and up-titrante the dose slowly every 10 to 14 days until glycaemic targets are met or side effects occur. Few patients will achieve and maintain glycaemic targets with 500 mg once daily. Most patients will require 1000 – 2550 mg per day in two or three divided doses. The optimum dose for cardiovascular benefit in obese patients is 2550 mg/day (850 mg TDS). (B)

If gastrointestinal (GI) adverse events are limiting, try temporarily reducing or discontinuing the drug, and re-titrante when the GI disturbances resolve. The GI side-effects with metformin extended-release is not different to the standard release when used as initial therapy; however patients who switch due to the extended release may have improved tolerability. If GI disturbances remain intolerable with standard metformin tablets, try switching to a metformin extended release (XR) formulation and titrate the dose every 10-14 days again.

The extended release formulation should be dosed once daily with the evening meal at a dose not exceeding 2000 mg/day. The 2000 mg dose can be taken as 1000 mg twice a day without disadvantages if the patient so prefers. Patients not achieving their glycaemic target with 2000 mg of the extended release may benefit from switching to a higher dose of the standard release metformin. (B)

Monitor renal function (eGFR) in all patients at least annually. Do not exceed 1000 mg/day if the eGFR is 30-45 ml/min/1.73m². Stop metformin therapy if the eGFR is < 30 ml/min/1.73m². (B)

The significance of low serum vitamin B12 levels associated with long-term metformin use is not known. Measure and treat whenever clinically appropriate. (B)

Profile of the patient in whom metformin may not be a preferred option:
- Patients with irritable bowel syndrome or other chronic gastrointestinal disorders
- Normal weight individuals who do not wish to lose weight
- Patients at high risk for lactic acidosis (severe heart, lung, liver, renal or peripheral vascular disease)
- There is a history of metformin intolerance. (C)

Do not persist with any chosen treatment if the HbA1c has not decreased by > 0.5% after six months
9.2: Drug Summary – Gliclazide modified-release

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to appendix 9.2 for full text and references)

| Mode of action | Gliclazide modified-release is a sulphonylurea (SU) and insulin secretagogue that binds selectively to the sulphonylurea receptor-1A (SUR1A) binding-site of KATP channels on pancreatic beta cells resulting in membrane depolarization, calcium influx and exocytosis of stored insulin. |
| Glycaemic efficacy and indications | When used as monotherapy or as add-on to other non-insulin glucose lowering drugs the mean HbA1C reduction is -1% with sustained efficacy beyond 2 years. The ADVANCE study, using a gliclazide modified-release intensive treatment strategy achieved an HbA1c target of 6.5% (vs. 7.3% for conventional treatment) and demonstrated a significant reduction in microvascular outcomes (and therefore also the combined microvascular and macrovascular outcomes), driven mainly by the reduction in the incidence of nephropathy. |
| Cardiovascular outcome trials | There have been no dedicated cardiovascular safety studies with any sulphonylurea. Meta-analyses of observational and randomized controlled trials consistently demonstrate that gliclazide has a better cardiovascular safety profile than glibenclamide and glimepiride. |
| Hypoglycaemia | Gliclazide modified-release increases the risk of hypoglycaemia when used as monotherapy or combination therapy, but this is significantly lower when compared to glibenclamide and glimepiride. The rates of hypoglycaemia increase with lower glycaemic targets. |
| Weight | Weight change ranges from 0 to +1.5 kg. Mean weight gain in a meta-analysis of RCTs was 0.5kg. |
| Non-glycaemic benefits | Gliclazide modified-release increases peripheral insulin sensitivity, decreases hepatic glucose production, inhibits platelet aggregation and adhesion, increases t-PA activity and has antioxidant effects. There are also reports of possibly a lower cancer risk compared to other SUs and insulin. |
| Side Effects and Precautions | Side-effects apart from hypoglycaemia and weight gain are rare. Cross-reactivity with sulphonamide antibiotic allergy is uncommon and is not a contra-indication. Do not use with advanced liver disease because of hepatic metabolism. |
| Dosing and prescribing | • The usual starting dose for gliclazide modified-release is 30 to 60 mg administered once daily with the morning meal. • Consider starting with the higher (60 mg) dose when the HbA1c target is more than 0.5% from the prevailing HbA1c level, or if the patient is has symptomatic hyperglycaemia. • The dose can be escalated by 30 to 60mg every one to four weeks, guided by fasting glucose levels. • The maximum dose is 120 mg administered once daily with the morning meal. • If mild hypoglycaemia occurs unexpectedly reduce the dose by 30 to 60 mg. • A single episode of severe hypoglycaemia or recurrent episodes of mild hypoglycaemia would be a strong indication to switch to an alternative glucose lowering drug. • The 60 mg tablets are scored and can be broken to improve cost effectiveness. |
| Renal dosing | Current guidelines recommend that gliclazide modified-release can be used at all stages of chronic kidney disease using standard dosing guidelines. Caution is advised however, when the eGFR is < 30 ml/min/1.73m², and these patients with CKD stage 3 or worse should be managed with specialist supervision. |
| Cost of cheapest formulation at maximum dose | R93.00 for 120 mg (Single exit price as at March 2017). |

SEMDSA 2017 Recommendations for sulphonylureas

The sulphonylurea of choice should be gliclazide modified-release because:
• It has equivalent efficacy compared to other sulphonylureas.
• It is consistently associated with lower rates of hypoglycaemia and better cardiovascular and renal safety relative to other sulphonylureas.
• It has proven benefits for long-term microvascular disease outcomes.

Glibenclamide must not be used at primary care level.

Consider gliclazide modified-release as initial monotherapy when metformin is not tolerated or is contraindicated.

Consider gliclazide modified-release as add-on (dual therapy) to metformin (or other initial drug therapy) in most patients not achieving or maintaining their glycaemic targets.

If not already in use, consider gliclazide modified-release as a third glucose lowering drug.
To convert treatment from another sulphonylurea to gliclazide modified-release, use the following dose conversion:

- Glibenclamide 5 mg ≈ Gliclazide modified-release 30 mg
- Glimepiride 1-2 mg ≈ Gliclazide modified-release 30 mg

Only continue gliclazide modified-release beyond stage 3 chronic kidney disease (when the eGFR is less 30 ml/min/m^2) with specialist supervision.

Circumstances where gliclazide MR may be preferred to other treatment options:

- Gliclazide MR should be the preferred second drug for the majority of patients with type 2 diabetes.
- At diagnosis when rapid control of hyperglycaemic symptoms is required.

Circumstances where gliclazide MR may not be the preferred option:

- The individualised glycaemic target is ≤ 6.5% (as the risk of hypoglycaemia may be unacceptably high with this target).
- There is a history of severe hypoglycaemia or hypoglycaemia unawareness.
- There is a history of recurrent hypoglycaemia (any degree) despite dose adjustments.
- The risk of hypoglycaemia is high and/or its consequences are severe.
- The patient has advanced liver disease.

Do not persist with any chosen treatment if the HbA1C has not decreased by > 0.5% after six months

9.3: Drug Summary – Pioglitazone

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to appendix 9.3 for optional full text and references)

| Mode of action | Agonist of nuclear receptors called peroxisome proliferator-activated receptor-gamma (PPARγ). Leads to increased transcription of proteins that augment the post-receptor actions of insulin resulting in improved insulin sensitivity and β-cell function. |
| Glycaemic efficacy and indications | Good efficacy as monotherapy, dual therapy and triple therapy (± 1% HbA1C reduction); similar efficacy to metformin, SU or GLP-1RA. |
| Cardiovascular outcome trials | PROactive study (secondary prevention) showed reductions in secondary endpoints (composite of all-cause mortality, non-fatal myocardial infarction, and stroke) by 16%. Meta-analysis of RCTs also reported a significant 18% relative risk reduction in the composite outcome (death, myocardial infarction, or stroke). Increases heart failure hospitalisations because of fluid retention, but not mortality. |
| Hypoglycaemia | Does not cause hypoglycaemia except when combined with insulin or insulin secretagogues. |
| Weight | Causes dose-dependent weight gain (~2-4 kg) due to fluid retention and adipocyte differentiation. Weight gain correlates with therapeutic response. |
| Non-glycaemic benefits | Increases HDL-C; reduces triglycerides, reduces LDL atherogenicity, CRP and microalbuminuria; increases PAI-1 and adiponectin; reduces carotid intima media thickness and atheroma volume. Reduces hepatic fibrosis in non-alcoholic steatohepatitis (NASH); improves ovulation and metabolic abnormalities in polycystic ovary syndrome. |
| Side Effects and Precautions | Can cause fluid retention, oedema, and may worsen or precipitate congestive heart failure. Associated with an increase in distal long-bone fractures in women and men. Possible small increase in bladder cancer. |
| Dosing and prescribing | Start with 15 once daily in the morning; increase to 30 mg after one to three months if necessary. Most patients will derive optimum benefit at this dose; do not exceed 30 mg in the primary care setting. Maximum registered dose is 45 mg daily. |
| Renal dosing | No dose adjustment is necessary, but do not use if renal disease is causing fluid retention, or when the eGFR is <30 ml/min/1.73m^2. |
| Cost of cheapest formulation | R117.00 for 30 mg. |
SEMDSA 2017 Recommendations for pioglitazone

Consider pioglitazone as initial monotherapy when metformin is contraindicated or not tolerated.

Consider pioglitazone as add-on to metformin or other initial drug therapy, in selected patients not achieving or maintaining their glycaemic targets.

Consider pioglitazone as a third non-insulin glucose lowering drug in selected patients not achieving or maintaining their glycaemic targets on an existing oral two-drug regimen.

Circumstances where pioglitazone may be preferred to other treatment options:
- Gliclazide MR is contraindicated or not tolerated.
- Non-alcoholic steatohepatitis is present.
- The patient has features of severe insulin resistance.
- There is a history of previous myocardial infarction, previous stroke or chronic kidney disease stage-3 (pioglitazone offers probable benefit for secondary prevention).

Circumstances where pioglitazone may not be the preferred option:
- Age > 75 years old (risk of congestive heart failure (CHF), fracture and bladder cancer).
- History of congestive heart failure.
- History of osteoporosis.
- History of bladder cancer, or haematuria that has not been investigated.
- Stage-4 or worse chronic kidney disease (risk of fluid retention).
- Patients on insulin therapy (higher risk of fluid retention and CHF).
- Elevated liver enzymes (>2x ULN), which is not due to NASH.

Do not persist with any chosen treatment if the HbA1c has not decreased by > 0.5% after six months

9.4: Drug summary - DPP-4-inhibitors

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to Appendix 9.4 for review and references)

**Mode of action and pharmacology**
DPP-4 inhibitors (gliptins) are capable of inhibiting the degradation of endogenous GLP-1, thereby therapeutically raising circulating GLP-1 levels.

**Glycaemic efficacy and indications**
Can be used as monotherapy in selected patients where there is intolerance to metformin (where there is a high risk for hypoglycaemia). Can be used as dual or triple therapy when added to metformin / sulphonylurea / SGLT 2 inhibitor / insulin. HbA1c reduction when used as monotherapy is between 0.5 and 1.1 %.

**Macrovascular and Mortality Outcomes**
Cardiovascular outcome safety trials for all 3 DPP-4 inhibitors have been neutral for major adverse cardiovascular events. Saxagliptin was associated with increased rates of hospitalisation for heart failure.

**Hypoglycaemia**
Hypoglycaemia rates are not different to placebo except when DPP 4 inhibitors are combined with insulin or insulin secretagogues.

**Non-glycaemic benefits**
Weight-neutral.

**Side Effects and Precautions**
Small absolute risk for pancreatitis.
Increased risk of hospitalisation for heart failure with saxagliptin.

**Contraindications**
Acute, chronic or recurrent pancreatitis or high risk for pancreatitis. Pancreatic cancer.
All are contraindicated in severe liver disease. Use saxagliptin and vildagliptin with caution in moderate liver disease. Heart failure (saxagliptin).

<table>
<thead>
<tr>
<th>eGFR</th>
<th>Saxagliptin</th>
<th>Sitagliptin</th>
<th>Vildagliptin</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥50 ml/min</td>
<td>5 mg daily</td>
<td>100 mg daily</td>
<td>50 mg twice a day</td>
</tr>
<tr>
<td>30-50 ml/min</td>
<td>2.5 mg daily</td>
<td>50 mg daily</td>
<td>50 mg daily</td>
</tr>
<tr>
<td>&lt;30 ml/min</td>
<td>2.5 mg daily</td>
<td>25 mg daily</td>
<td>50 mg daily</td>
</tr>
</tbody>
</table>

**Cost at maximum dose**
Moderate (R260 – 340 per month - single exit pricing as at March 2017)
SEMDSA 2017 Recommendations for DPP-4 inhibitors

Consider a DPP-4 inhibitor as initial monotherapy when metformin is contraindicated or not tolerated.  

Consider a DPP-4 inhibitor as add-on to metformin or other initial drug therapy, in selected patients not achieving or maintaining their glycaemic targets.  

Consider a DPP-4 inhibitor as the third glucose lowering drug in selected patients not achieving or maintaining their glycaemic targets on an existing oral two-drug regimen.  

Combination DPP-4 inhibitor and insulin therapy should be initiated at specialist level.  

Be aware of dose adjustments for chronic kidney disease.  

Circumstances where a DPP-4 inhibitor may be preferred to other treatment options:  
- As the 2nd add-on drug when gliclazide MR is contraindicated or not tolerated.  
- As the 3rd add on drug for most patients if HbA1c targets are potentially achievable.  
- Older patients with multiple comorbidities.  
- Patients with stage-4 chronic kidney disease (can be used without risk of hypoglycaemia).  
- If a fixed-dose combination tablet will improve adherence, compliance and/or cost-effectiveness.

Circumstances where a DPP-4 inhibitor may not be the preferred option:  
- Very high HbA1c and the glycemic target is not likely to be achieved with a DPP-4 inhibitor.  
- History of pancreatitis or pancreatic tumour.  
- History of heart failure or high risk of heart failure (saxagliptin).  
- Liver disease: moderate (do not use saxagliptin or vildagliptin) or severe (do not any DPP-4 inhibitor).

Do not persist with any chosen treatment if the HbA1c has not decreased by > 0.5% after six months

9.5: Drug summary - GLP-1 receptor agonists

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to Appendix 9.5 for drug review and references)

| Mode of action and pharmacology | Reduces HbA1c by 0.9 to 1.2 %. Monotherapy: registered (liraglutide) but not recommended for primary healthcare. Dual therapy: registered but not recommended for primary healthcare. Triple therapy: Can be combined with any 2 of metformin / sulphonylurea / TZD. Do not combine with DPP-4 inhibitor or SGLT2 inhibitor. Exenatide can be added to basal insulin with or without oral agents. |
| Glycaemic efficacy and indications | Lixisenatide was neutral in the ELIXA trial. Liraglutide and semaglutide have each demonstrated reductions in a composite endpoint of major adverse cardiovascular events by 13% and 26% in the LEADER and SUSTAIN-6 trials respectively (vs. placebo), when used in patients with established cardiovascular disease. There are no cardiovascular outcomes trials with exenatide. |
| Macrovascular and Mortality Outcomes | Hypoglycaemia rates are not different to placebo except when GLP-1 receptor agonist are combined with insulin or insulin secretagogues. |
| Hypoglycaemia | Weight reduction of between 1 and 3 kg for exenatide and liraglutide. Both exenatide and liraglutide reduce SBP and DBP by 1 to 5 mmHg. Reduction in liver fat content (liraglutide). |
| Non-glycaemia benefits | Nausea and vomiting in up to 25%; Discontinuation rate 5-20%. Pancreatitis. Skin reactions. |
| Side Effects and Precautions | History of pancreatitis, or at high risk for pancreatitis (e.g. untreated gallstone disease, recent or planned ERCP, excessive alcohol use). |
| Contraindications | History of pancreatic tumour. History of medullary thyroid cancer (MTC) or multiple endocrine neoplasia (MEN) syndrome type 2 |
| **Dosing** | **eGFR**
| | ≥30 ml/min | Exenatide Initial dose is 5 ug BD for the 1st month; then 10 ug BD |
| | <35 ml/min | Liraglutide Initiate 0.6 mg daily Maximum dose of 1.8 mg daily |
| Cost | High (R620.00 to R2145.00 - single exit price March 2017) |
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### SEMDSA 2017 Recommendations for GLP-1 receptor agonists (GLP-1RA)

<table>
<thead>
<tr>
<th>Recommendation</th>
<th>Consider a GLP-1RA injectable as the third glucose lowering drug (triple therapy) in overweight and obese patients when glycaemic targets are not being achieved or maintained.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Consider adding a GLP-1RA to existing basal insulin therapy (with oral therapies) as an alternative to intensifying the insulin regimen, especially when weight gain and/or hypoglycaemia is a limiting factor.</td>
</tr>
<tr>
<td></td>
<td>Escalate the dose of GLP-1RA slowly to minimise side-effects.</td>
</tr>
<tr>
<td></td>
<td>Circumstances where a GLP-1RA may be preferred to other treatment options:</td>
</tr>
</tbody>
</table>
|                | - Overweight and obese patients  
|                | - Weight gain or hypoglycaemia has been, or is likely to be problematic with other treatment options.  
|                | - HbA1c is very high (GLP-1 RA and insulin are the most effective glucose lowering drugs for most patients).  
|                | - Patients with established cardiovascular disease (liraglutide benefit); to be managed at specialist care level. |
|                | Circumstances where a GLP-1RA may not be the preferred option: |
|                | - Patients with chronic gastrointestinal disorders.  
|                | - Patients with a history of pancreatitis or pancreatic tumour. |

Do not persist with any chosen treatment if the HbA1c has not decreased by > 0.5% after six months

### 9.6: Drug Summary - SGLT2 inhibitors

SEMDSA Type 2 Diabetes Guideline Expert Committee

(Refer to Appendix 9.6 for review and references)

| MOA | By inhibiting SGLT2, gliflozins reduces renal glucose reabsorption, leading to increased urinary excretion of excess glucose and a reduction in plasma glucose concentrations in a non-insulin dependent manner. |
| Glycaemic efficacy and indications | As monotherapy: 0.6% to -1.2%; non-inferior to metformin.  
| | As add on to metformin: -0.5% to -1.0%; non-inferior to other classes |
| | As add on to metformin + SU: -0.7% to -0.9%; non-inferior to DPP-4 inhibitors  
| | As add on to insulin therapy: -0.5% to -0.8% |

| Microvascular Outcomes | No primary outcome studies. |
| Macrovascular and Mortality Outcomes | Empagliflozin therapy was associated with a reduction in all-cause death, cardiovascular-death and hospitalisation for heart failure in patients with established cardiovascular disease. There has been no signal of adverse cardiovascular outcomes in systematic meta-analyses for the other SGLT2 inhibitors. |
| Hypoglycaemia | Hypoglycaemia rates are not different to placebo except when SGLT2 inhibitors are combined with insulin or insulin secretagogues. |
| Non-glycaemic benefits | Mean weight loss of 1.6 to 2.5 kg; not different to GLP-1RAs (meta-analysis) Systolic and diastolic blood pressure reduction (-4.0 and -1.5 mmHg) respectively. |
| Side effects and precautions | Mycotic genital infections are common (and more so in women); do not use in patients with a history of recurrent genital infections.  
| | The risk of urinary tract infections may be increased but can be minimised by advising adequate hydration and fastidious bathroom hygiene; do not prescribe SGLT2 inhibitors in patients with a history of recurrent UTI. |
| | Dehydration and hypotension can occur particularly in susceptible patients (treated with loop diuretics, have advanced cardiac disease or older than 65 years). Always ensure and emphasise adequate hydration when prescribing SGLT2 inhibitors. |
| | eGFR usually declines within the first weeks of initiating therapy and then gradually returns toward baseline. Monitor eGFR and discontinue SGLT2 inhibitors if the decline is ≥ 30%.  
| | Diabetic ketoacidosis may occur at mildly elevated levels of blood glucose particularly in insulin treated patients. Warn patients of the symptoms of DKA and advise them to seek medical attention immediately.  
| | The risk of fractures and lower-limb (especially toe) amputations is increased with canagliflozin. Therefore do not use canagliflozin, and exercise caution with the other drugs in this class, in patients who are at high risk for these conditions.  
| | Do not prescribe dapagliflozin to patients with bladder cancer or in combination with pioglitazone. |

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### Dosing

<table>
<thead>
<tr>
<th>eGFR</th>
<th>Dapagliflozin</th>
<th>Empagliflozin</th>
<th>Canagliflozin</th>
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</thead>
<tbody>
<tr>
<td>≥60 ml/min</td>
<td>5 mg or 10 mg</td>
<td>10 mg or 25 mg</td>
<td>100 mg or 300 mg</td>
</tr>
<tr>
<td>45-60 ml/min</td>
<td>Contraindicated</td>
<td>Continue 25 mg but do not initiate therapy.</td>
<td>Continue 100 mg but do not initiate therapy.</td>
</tr>
<tr>
<td>&lt;45 ml/min</td>
<td>Contraindicated</td>
<td>Contraindicated</td>
<td>Contraindicated</td>
</tr>
</tbody>
</table>

### Cost
Unknown; not registered in South Africa as at March 2017.

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**SEMDSA 2017 Recommendations for SGLT2 inhibitors**

- Do not use SGLT2 inhibitors as initial monotherapy.  
  
- Consider a SGLT2 inhibitor as add-on (dual therapy) to metformin (or other initial drug therapy) in selected patients not achieving or maintaining their glycaemic targets.  
  
- Consider a SGLT2 inhibitor as the 3rd glucose lowering drug in selected patients not achieving or maintaining their glycaemic targets on an existing oral two-drug regimen.

Circumstances where an SGLT2 inhibitor may be preferred to other treatment options:
- Overweight and obese patients.
- Weight gain or hypoglycaemia has been, or is likely to be problematic with other treatment options.
- Patients with established cardiovascular disease (empagliflozin benefit); to be managed at specialist care level.

Circumstances where an SGLT2 inhibitor may not be the preferred option:
- Patients with recurrent mycotic genital infections or urinary tract infections.
- Patients at risk for dehydration and hypotension.
- Patients at high risk for stroke, fracture (canagliflozin), amputation (canagliflozin), bladder cancer (dapagliflozin) or ketoacidosis (refer to drug review).

Do not initiate SGLT2 inhibitors when the eGFR is < 60 ml/min/m².

Stop all SGLT2 inhibitors when the eGFR is < 45 ml/min/m².

Do not persist with any chosen treatment if the HbA1C has not decreased by > 0.5% after six months.

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**9.7: Alpha-glucosidase inhibitors (acarbose)**

SEMDSA Type 2 Diabetes Guideline Expert Committee

**Mechanism of Action**
Prevents the conversion of complex carbohydrates into monosaccharides within the intestine thereby decreasing the ability to absorb monosaccharides.

**Glycaemic efficacy and indications**
HbA1C reduction when used as monotherapy of between 0.6 and 0.8%.  

**Microvascular Outcomes**
Reduction of Microvascular Complications are inferred from the benefit of improved glycaemic control.

**Macrovascular and Mortality Outcomes**
Acarbose significantly reduced the risk of cardiovascular events by 49% in patients with impaired glucose tolerance. No data for patients with Diabetes.

**Hypoglycaemia**
Hypoglycaemia rates are no different to placebo except when combined with Insulin or Insulin Secretagogues.

**Non glycaemic benefits**
Weight neutral.

**Side Effects and Precautions**
Transient elevation of hepatic transaminases.

<table>
<thead>
<tr>
<th>eGFR</th>
<th>Acarbose</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥30 ml/min</td>
<td>Start with 50 mg once daily with meals, and increase by 50 mg every two weeks if tolerated. Maximum dose is 100 mg x3 daily.</td>
</tr>
<tr>
<td>&lt;30 ml/min</td>
<td>Not recommended</td>
</tr>
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</table>

**Cost**
High (R407.00).