

Peripheral Artery Disease (PAD) in T2D

Core Science Deck







- Introduction & Epidemiology of Peripheral artery disease (PAD)
- Symptoms and pathophysiology of PAD
- Association between PAD and T2D
- Diagnosis of PAD
- Management of PAD in people with diabetes
- GLP-1RAs in people with PAD and T2D
- Summary



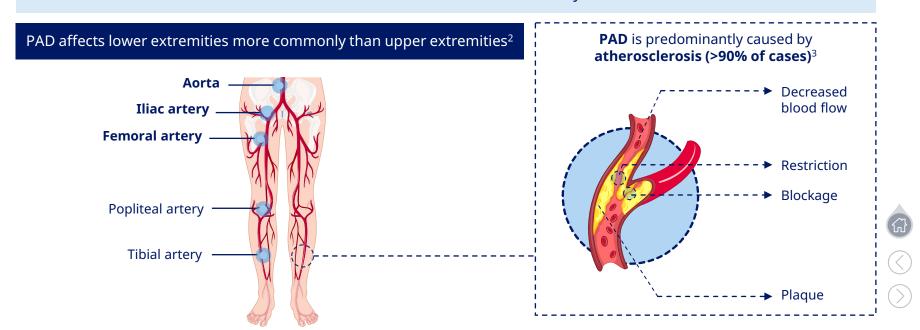




What is peripheral artery disease?

Peripheral artery disease is the narrowing

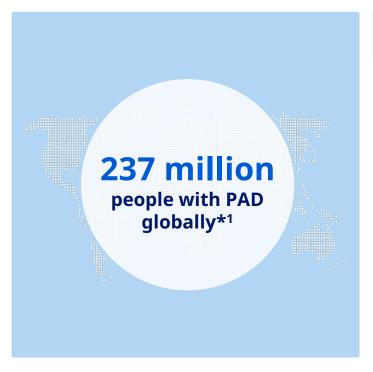
and/or occlusion of arteries outside of the heart and brain mainly attributed to atherosclerosis1



PAD, peripheral artery disease. ASCVD- Atherosclerotic cardiovascular disease, a leading cause of mortality and morbidity globally, is caused by the build-up of plaque in the arterial walls and comprise conditions such as Coronory Heart disease, Cerebrovascular disease, PAD, aortic atherosclerotic disease 5

1. Soyoye DO et al. World J Diabetes 2021;12:827–38; 2. Zemaitis M et al. StatPearls Publishing LLC. Available at: https://www.ncbi.nlm.nih.gov/books/NBK430745/ Accessed December 2023;3. Shu J, Santulli G. Atherosclerosis 2018;275:379–81.

Global prevalence of PAD is high







PAD affects **12–14%** of the general population²

 However, PAD remains underdiagnosed compared with other CVD conditions,³ making absolute numbers difficult to obtain



Prevalence increases with age, affecting **up to 20%** of those **aged ≥60 years** and **nearly 50%** of those aged **≥85 years** in USA⁴

- Incidence doubles with each advancing decade of life³
- 72% increase in prevalence from 1990 to 2019⁵



PAD affects women as often or more as men³

• IC is more common in men than in women, who are more likely to present as asymptomatic or with atypical symptoms³







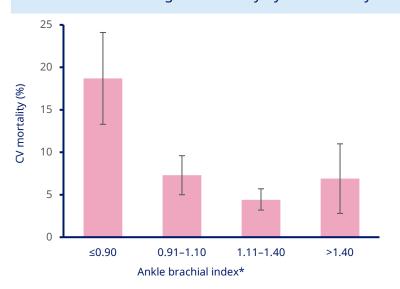
^{*}Estimated global prevalence of PAD in people aged ≥25 years in 2015, according to a systematic review of 118 articles from 33 countries. CVD, cardiovascular disease; IC, intermittent claudication; PAD, peripheral artery disease.

^{1.} Song P et al. Lancet Glob Health 2019;7:e1020-30; 2. Shammas NW. Vasc Health Risk Manag 2007;3:229-34; 3. Pabon M et al. Circ Res 2022;130:496-511;

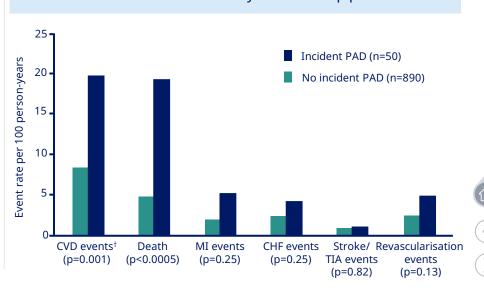
^{4.} Firnhaber JM, Powell CS. Am Fam Physician 2019;99:362-9; 5. Eid MA et al. J Vasc Surg 2023,77:1119-26.e1;

People with PAD have a higher risk of CV events vs those without PAD

Meta-analysis of male participants (N=24,955) from 16 studies showing CV mortality by PAD severity¹



Prospective cohort study of people with stable CAD ± incident PAD over a 7.2-year follow-up period²



PAD, peripheral artery disease; CV, Cardiovascular; *Ankle brachial index is a diagnostic tool for PAD where a lower value indicates a more severe disease state. Error bars represent 95% confidence intervals. 'The primary outcome (CVD events) included a composite of events: death, MI, CHF, stroke, TIA and coronary revascularisation. Secondary outcomes included those events assessed individually. CAD, coronary artery disease; CHF, congestive heart failure; CV, cardiovascular; CVD, cardiovascular disease; MI, myocardial infarction; PAD, peripheral arterial disease; TIA, transient ischaemic attack. 1. Ankle Brachial Index Collaboration. JAMA 2008;300:197–208; 2. Grenon SM et al. Vasc Med 2013;18:176–84.

PAD is a major economic burden associated with substantial healthcare costs



The total US PAD-related annualised healthcare cost in the symptomatic PAD patient population was **\$4,006** per patient¹



PAD prevalence increased by 13.1% in high-income countries and **28.7% in low- and middle-income** countries between 2000 and 2010²



The number of **annual CV events are higher for people with PAD compared with CAD**, leading to greater hospitalisation rates and higher direct costs¹





Total annual costs associated with vascular- related hospitalizations in PAD patients were estimated to be more than \$21 billion between 2004 and 2010.³





^{1. .} Bauersachs R et al. Cardiovasc Ther 2019;8295054. 2. Fowkes FG et al. Lancet 2013;382:1329–40; 3. Mahoney, Elizabeth M., et al. Circulation: Cardiovascular Quality and Outcomes 3.6 (2010): 642-651.

Symptoms and clinical manifestations

Spectrum of PAD symptoms

Increasing severity & risk of CV event or amputation



Asymptomatic PAD

No obvious symptoms¹

PAD can be detected with blood pressure and imaging tests^{1,2}

It has been estimated that 20–50% of people with PAD are asymptomatic¹



Intermittent claudication and/or atypical leg pain symptoms

Exercise-induced pain in the legs and/or buttocks that goes away with rest³

Atypical leg symptoms occur in 40–50% of cases;¹ may be confused with lower extremity arthritis or degenerative spinal disease⁴



CLI*

Exhibits the following:5

- 1. Ischemic rest pain or ulcer necrosis
- 2. Limited to chronic ischemia
- Limb-threatening ischemia usually requiring major amputation ≤6 months if blood flow does not improve
- Objective ischemia (ankle pressure ≤50-70 mmHg,[†] toe pressure ≤30-50 mmHg[†] and transcutaneous oxygen pressure ≤30-50 Torr)



ALI*

Sudden onset (<2 weeks), claudication deteriorates; severe hypoperfusion characterized by '6 Ps':5

- 1. Pain at rest
- 2. Pallor
- 3. Pulseless
- 4. 'Perishingly' cold
- 5. Paresthesia
- 6. Paralysis





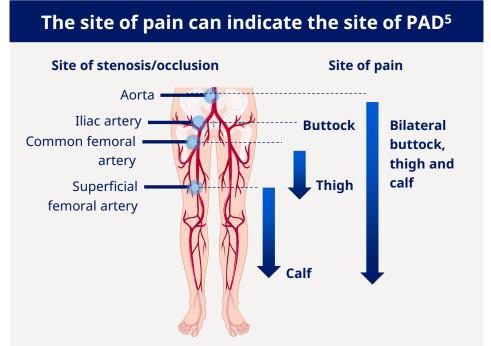




1. Olin JW, Sealove BA. Mayo Clin Proc 2010;85:678–92; 2. Aboyans V et al. Eur J Vasc Endovasc Surg 2018;55:305–68; .3. Criqui MH et al. Circulation 2021;144:e171–91; 4. Gerhard-Herman MD et al. Circulation 2017;135:e686–725; .5 Norgren L et al. Eur J Vasc Endovasc Surg 2007;33(Suppl 1):S1–75

Intermittent claudication is the most common symptom of PAD¹

- Early symptom experienced by 10–35% of people with PAD³
 - Characterized by pain in lower extremities (e.g. calf) that develops with walking or other exertion^{1,2,4}
 - Resolves within 10 minutes of rest⁴
- Prevalence of PAD is similar between men and women;1 however, incidence of IC is higher in men than women at all ages⁵











^{1.} Boyko EJ et al. Peripheral Arterial Disease, Foot Ulcers, Lower Extremity Amputations, and Diabetes. In: Cowie CC et al., editors. Diabetes in America. 3rd edition. 2018. Available at: https://www.ncbi.nlm.nih.gov/books/NBK567977/ January 2024; 2. Criqui MH et al. Circulation 2021;144:e171–91 3. Olin JW, Sealove BA. Mayo Clin Proc 2010;85:678–92; 4. Criqui MH, Aboyans V. Circ Res 2015;116:1509–26; 5. Kannel et al.. 1970 available at https://www.ahajournals.ora/doi/adf/10.1161/01.CIR.41.5.875

Risk factors for PAD



Modifiable risk factors



Diabetes^{1,2,4} Can double the risk of PAD





Hypertension^{1,2,4}
Association more evident for SBP than DBP¹



High homocysteine levels³



Smoking^{2,4}

Risk is higher in current vs former smokers



Dyslipidemia^{2,4}

Including high LDL-C and triglycerides



Physical inactivity¹

Inversely related to physical activity



Inflammation^{3,4}

As assessed by hs-CRP

★ Non-modifiable risk factors



Age^{2,4}

Risk increases with age



Race and ethnicity^{1,4}

Black individuals have greater risk for PAD vs White, Hispanic or Asian



Family history/genetic factors³

Greatest risk in those with family history of PAD. CVD or stroke



Other risk factors



Socioeconomic and/or psychosocial factors³



CKD and renal impairment^{2,3}

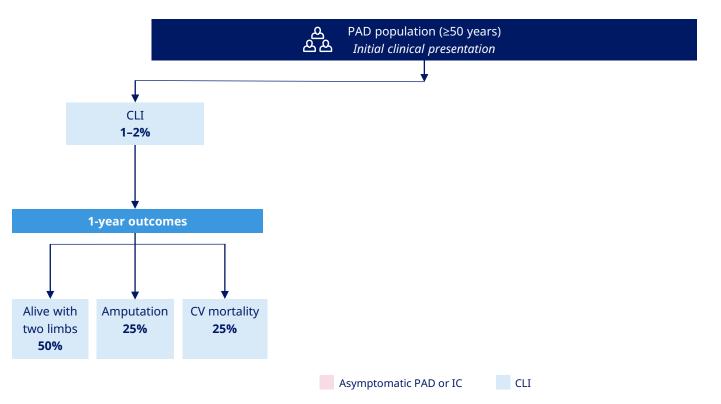


Heavy metals/ radiation/injury¹

PAD, peripheral artery disease CKD, chronic kidney disease; CVD, cardiovascular disease; DBP, diastolic blood pressure; hs-CRP, high sensitivity-C-reactive protein; LDL-C, low-density lipoprotein-cholesterol; PAD, peripheral arterial disease; SBP, systolic blood pressure

1. Criqui MH et al. Circulation 2021;144:e171–91; 2. Song P et al. Lancet Glob Health 2019;7:e1020–30; 3. Criqui MH, Aboyans V. Circ Res 2015;116:1509–26. 4. Olin JW, Sealove BA. Mayo Clin Proc 2010;85:678–92

Natural history of PAD: typical disease progression

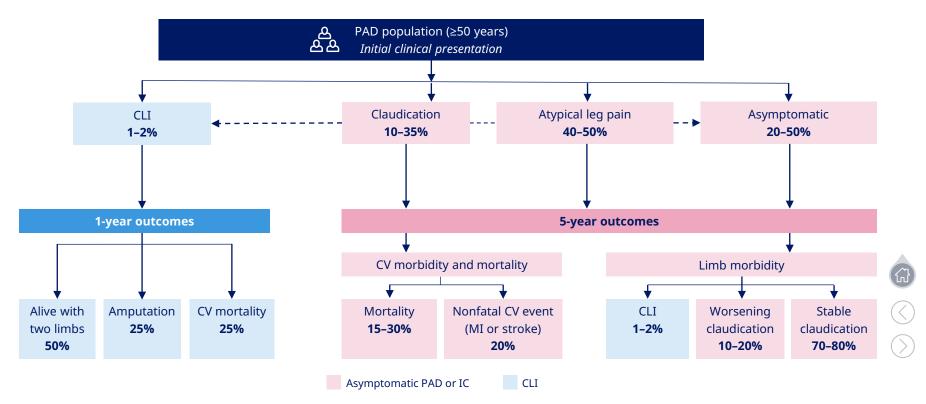








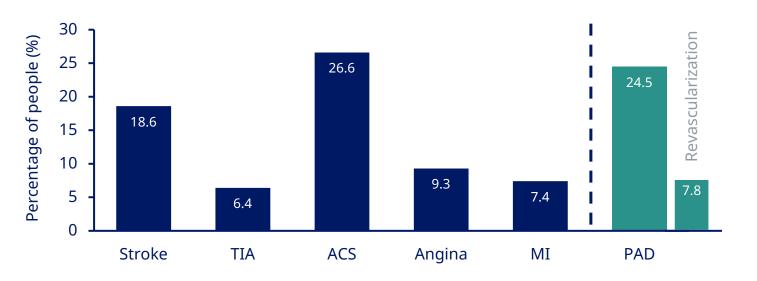
Natural history of PAD: typical disease progression



PAD, peripheral artery disease; CV, Cardiovascular; CLI/CLTI, critical limb ischaemia/chronic limb-threatening ischaemia; CV, cardiovascular; IC, intermittent claudication; MI, myocardial infarction; PAD, peripheral arterial disease. Olin JW, Sealove BA. Mayo Clin Proc 2010;85:678-92.

People with diabetes have an increased risk of ASCVD, including PAD

RWE database study of people with T2D in the USA (N=1,202,596), of which 45.2% had established ASCVD



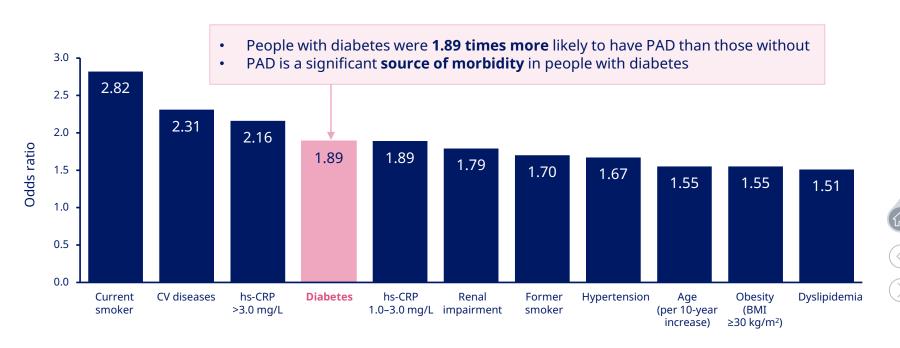






Diabetes is one of the main risk factors of PAD

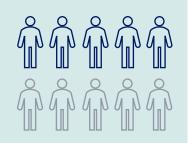
Worldwide estimates from a meta-analysis of 118 studies from low- and high-income countries



BMI, body mass index; CV, cardiovascular; hs-CRP, high-sensitivity C-reactive protein; PAD, peripheral artery disease. Song P et al. Lancet Glob Health 2019;7:e1020–30.

Prevalence of PAD is higher in people with T2D than in those without

The prevalence of PAD is up to twice as high in people with diabetes as in the general population, and the prevalence of concomitant PAD in people with diabetes is over 50% ^{1,2}



Glucose intolerance is associated with a >20% prevalence of an abnormal ankle brachial index* relative to 7% in those with normal glucose tolerance²



Approximately
50–70% of people with
chronic limbthreatening ischemia
have diabetes³



50-70%



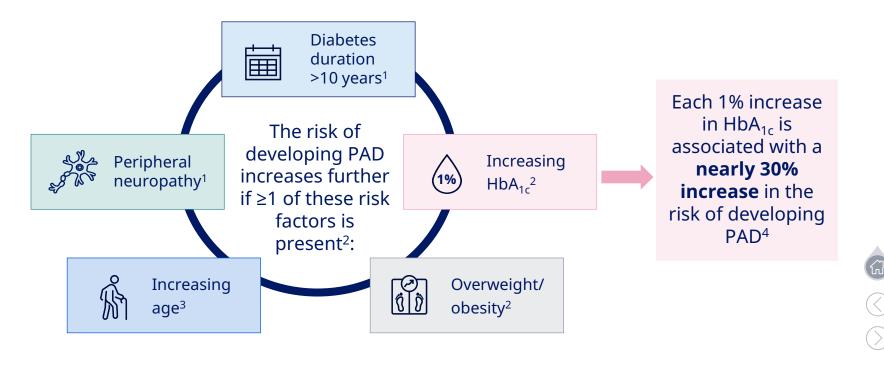




^{*}Ankle brachial index is a diagnostic tool for PAD where a lower value indicates a more severe disease state. CV, cardiovascular; PAD, peripheral artery disease.

^{1.} Soyoye DO et al. World J Diabetes 2021;12:827–38; 2. Thiruvoipati T et al. World J Diabetes 2015;6:961–69; 3. Marx N et al. Eur Heart J 2023:44:4043–140.

PAD Risk Factors within the T2D Population

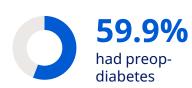




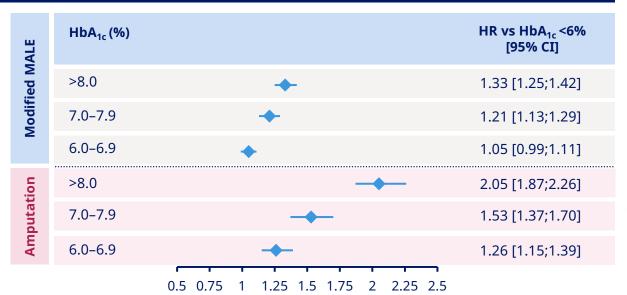
^{1.} Piliponiene L et al. Medicina 2021;57:1380; 2. American Heart Association. Peripheral Artery Disease and Diabetes, 2021. Available at: https://www.heart.org/en/health-topics/diabetes/diabetes/complications-and-risks/peripheral-artery-disease-diabetes Accessed January 2024; 3. Thiruvoipati T et al. World J Diabetes 2015;6:961–9. 4. Heikkinen et al. "Diabetes care for patients with peripheral arterial disease." European Journal of Vascular and Endovascular Surgery 33.5 (2007): 583-591.

Poor glycemic control associated with worse outcomes in PAD patients

Database study of people undergoing revascularization for PAD (N=26,799)







HR [95% CI]









There are several unmet needs for people with PAD and T2D

Under-diagnosis

People may be asymptomatic due to altered pain perception from comorbidities of diabetes, such as peripheral neuropathy¹⁻³

Screening tools may be underutilized¹

50% of people >50 years old were not aware that **diabetes is** a **risk factor** for PAD⁴

People with diabetes lack awareness of PAD symptoms¹

Poor prognosis

People with diabetes have worse PAD outcomes than those without (mortality and amputation rates)⁵ PAD+T2D patients are ~5x likely to have limb amputations vs PAD patients without T2D⁵ and have ~3-4x higher mortality and higher risk of stroke and heart failure compared to either condition alone⁵

Diabetes negatively affects neovascularization of the limb in PAD⁶

Limited treatment options

Data on efficacy of glucose-lowering treatments in PAD and T2D **are limited** and there is **need** for disease-modifying glucose-lowering **treatment options** for people with PAD and T2D^{7,8}

T2D is associated with **greater severity** and more diffuse PAD relative to nondiabetics. It also correlates **to greater risk of mortality** and **impaired quality of life**. ³







PAD, peripheral artery disease.

^{1.} Akalu Y, Birhan A. J Diabetes Res 2020:9419413; 2. Sunner SS et al. CJC Open 2021;3:936-49; 3. Thiruvoipati T et al. World J Diabetes 2015;6:961-9; 4. Hirsch AT et al. Circulation 2007;116:2086-94;

Recommended diagnostic testing for people with suspected PAD 2016 AHA/ACC,¹ 2023 ESC² and 2024 ESVS³ guidelines

ABI for diagnosing PAD

 Resting ABI is recommended to establish a PAD diagnosis in people where PAD is suspected¹

Physiological testing

- TBI should be measured to diagnose people when ABI is considered high or in people with clinical suspicion of PAD, despite a normal ABI*1,3
- People with exertional non-joint-related leg symptoms² and a normal ABI (>0.90 and ≤1.40)^{1,3} or a normal ABI at rest,³ should undergo exercise treadmill ABI testing to evaluate PAD¹⁻³

Imaging for anatomic assessment

- Duplex ultrasound, CTA or MRA of the lower extremities is useful to diagnose anatomic location and severity of stenosis when revascularization is considered^{1,2}
- Invasive angiography is useful for people with CLI when revascularization is considered¹







^{*}Defined as >1.3 or 1.4 across the literature. Information taken from the strongest class of recommendation and levels of evidence. ABI, ankle brachial index; ACC, American College of Cardiology; AHA, American Heart Association; CLI/CLTI, critical limb ischaemia/chronic limb-threatening ischaemia; CTA, computed tomography angiography; ESC, European Society of Cardiology; ESVS, European Society for Vascular Surgery; MRA, magnetic resonance angiography; PAD, peripheral artery disease. TBI, Toe-brachial index1. Gerhard-Herman MD et al. Circulation 2017;135:e686-725; 2. Marx N et al. Eur Heart | 2023:44:4043-140; 3. Nordanstia | et al. Eur | Vasc Endovasc Surg 2024:67:9-96.

ADA STANDARDS OF MEDICAL CARE IN DIABETES - 2024

Peripheral Artery Disease (PAD)

ADA recommends screening for asymptomatic PAD using ankle brachial index in people with diabetes at high risk for PAD, including any of the following:



age ≥ 50 years



diabetes with duration ≥ 10 years



comorbid microvascular disease



clinical evidence of foot complications



or any end-organ damage from diabetes.

Initial screening for PAD should include:

- Assessment of lower-extremity pulses, capillary refill time
- Rubor on dependency
- Pallor on elevation, and venous filling time
- Individuals with a history of leg fatigue, claudication, and rest pain relieved with dependency or decreased or absent pedal pulses should be referred for anklebrachial index with toe pressures and for further vascular assessment as appropriate







Early-stage PAD Management

Non-pharmacological measures for CV risk factor management in all people with PAD^{1,2}



Pharmacotherapy targeting individual risk factors	
Therapy	Options
Diabetes management for glycemic control that also reduce CV risk ^{2,3}	SGLT2i³ and GLP-1RAs³
Lipid-lowering drugs to improve CV outcomes ⁴	Statin therapy ⁵ , ezetimibe ⁵ and PCSK9 inhibitors ⁵
Antiplatelet therapy to manage aggregation of platelets and increased adhesions ^{4.} Antiplatelet and statin agents are customized to additional risk factors, such as whether the patient also has diabetes mellitus or hypertension ^{1.}	Aspirin ⁵ , clopidogrel ⁵ and ticagrelor ⁵
Anticoagulants to reduce risk of thrombi-dependant complications ⁴ should be considered for individuals with stable coronary and/or peripheral artery disease (PAD) and low bleeding risk to prevent major adverse limb and cardiovascular events ¹ .	Heparin ³ and rivaroxaban ⁵
Peripheral vasodilators/viscosity reductors to reduce symptoms of peripheral ischaemia and claudication ⁴	Cilostazol (although infrequently used owing to heart failure contraindications and subtle benefits) 4
Blood pressure management to reduce CV risk while maintaining limb perfusion ⁴	ACE inhibitors and angiotensin receptor blockers ⁴

ACE, angiotensin-converting enzyme; CV, cardiovascular; GLP-1RA, glucagon-like peptide-1 receptor agonist; PAD, peripheral artery disease; PCSK9, proprotein convertase subtilisin/kexin type 9; SGLT2i, sodium-glucose cotransporter 2 inhibitor.

^{1.} American Heart Association. Peripheral Artery Disease and Diabetes, 2021. Available at https://www.heart.org/en/health-topics/peripheral-artery-disease/prevention-and-treatment-of-pad: 2. Aboyans V et al. Eur Heart J 2018;39:763–816; 3. Nordanstig J et al. Eur J Vasc Endovasc Surg 2024;67:9-96.; 4. Abramson BL et al. Can J Cardiol 2022;38:56087; 5. Bevan GH, White Solaru KT. Arterioscler Thromb Vasc Biol 2020;40:541-53.









Undertreatment and poor outcomes are unmet needs in people with PAD and T2D

A 2021 Scientific Statement from the American Heart Association



Underutilization of evidence-based prevention measures

Despite guidelines recommending antiplatelet therapy, statins, antihypertensive agents, glycaemic control, smoking cessation and supervised exercise therapy, people with PAD remain undertreated and better adherence is needed



Improvement in diagnosis is needed to improve outcomes

Increasing awareness of PAD, including the diagnosis, clinical manifestation, and complications, is **critical to improving overall outcomes** among this population



Need for better control of symptoms

RCTs investigating the effects of medical therapies (in particular, glucose-lowering) on **functional capacity and walking distance**, in people with PAD and T2D are needed

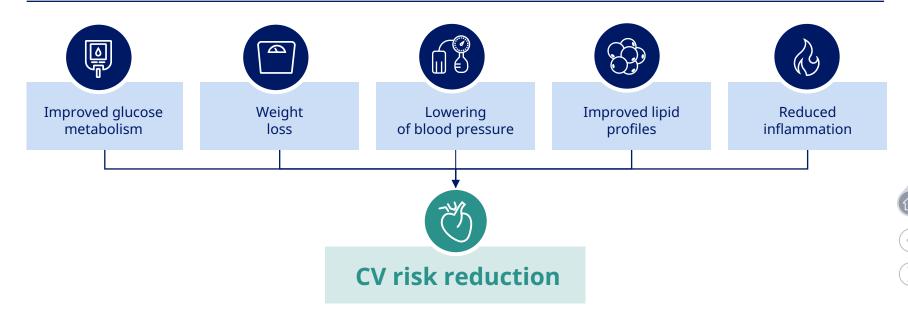






GLP-1RAs have several effects that support the rationale for use in people with PAD and T2D

Potential benefits

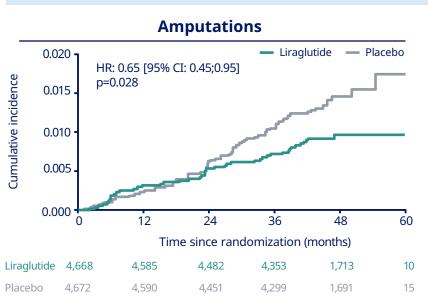


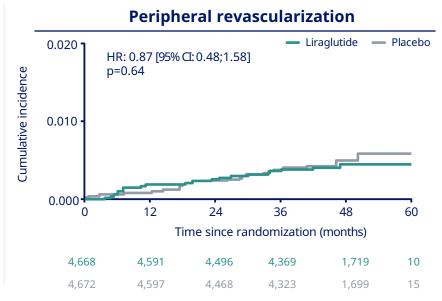


Liraglutide (1.8 mg) significantly reduced the risk of amputations vs placebo

Post hoc analyses of LEADER

Proportion of people with T2D and high CV risk experiencing amputation or peripheral revascularization

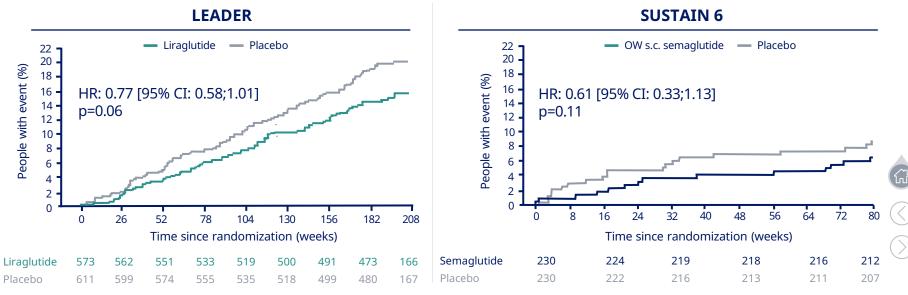




CI, confidence interval; CV, cardiovascular; HR, hazard ratio; PAD, peripheral artery disease. Dhatariya K et al. Diabetes Care 2018;41:2229–35.

Liraglutide (1.8 mg) and OW s.c. semaglutide (0.5 mg,1.0 mg) numerically reduced the risk of MACE in people with PAD and T2D vs placebo

Proportion of people with T2D and PAD at baseline experiencing MACE over time



Analysis from time to randomisation to first EAC-confirmed MACE. Data are without adjustment for baseline variables. All randomised people were included in the analyses from the time of randomisation until death or end of follow-up. CI, confidence interval; EAC, event adjudication committee; HR, hazard ratio; MACE, major adverse cardiovascular events; OW, once-weekly; PAD, peripheral artery disease; s.c., subcutaneous.

Verma S et al. Diabetes Obes Metab 2022;24:1288–99.

STRIDE: Effects of OW s.c semaglutide (1.0 mg) on Functional Capacity in Patients With T2D and Peripheral Artery Disease: Trial design

792 people with T2D

- Age ≥18 years
- T2D diagnosis ≥180 days prior to screening
- HbA_{1c} ≤10%
- PAD with intermittent claudication (Fontaine stage IIa) ≥3 months and:
 - o Pain-free walking distance >200 m
 - Maximum walking distance ≤600 m on a graded treadmill test
 - ABI ≤0.90 or TBI ≤0.70



Trial information

- Trial objective: to compare the effect of OW s.c. semaglutide on functional capacity in terms of maximum walking distance in people with PAD and T2D, vs placebo
- Randomized, phase 3b, double-blind, parallel-group trial

Primary endpoint

Change from baseline in maximum walking distance on a constant load treadmill test[†] at week 52

^{*}OW s.c. semaglutide dose escalation from starting dose of 0.25 mg; doubled every 4 weeks until trial maintenance dose achieved. †Treadmill at constant speed and incline (3.2 km/h, 12%). ABI, ankle brachial index; OW, once-weekly; PAD, peripheral artery disease; s.c., subcutaneous; SoC, standard of care; TBI, toe brachial index; VascuQol-6, Vascular Quality of Life Questionnaire-6.



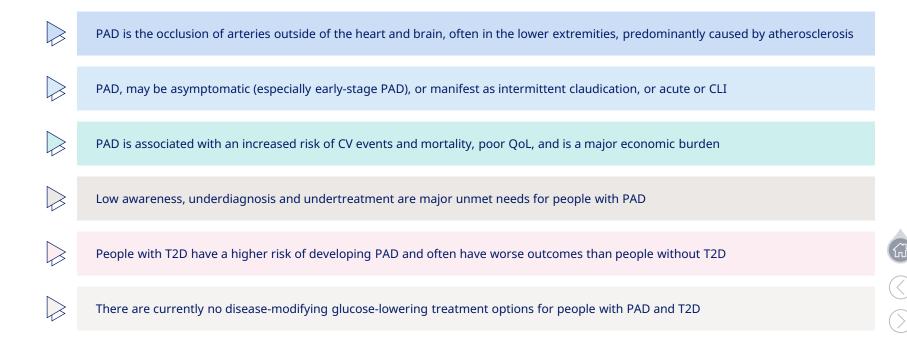




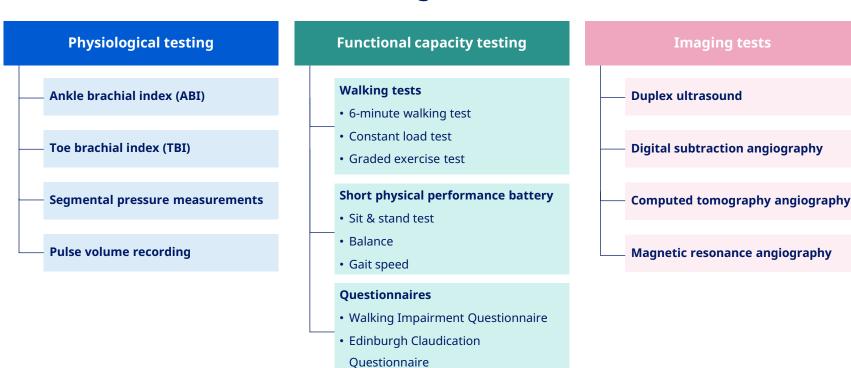




Summary



Different tests can be used to diagnose PAD^{1–4}



ABI, ankle brachial index; PAD, peripheral artery disease; TBI, toe brachial index.

^{1.} Gerhard-Herman MD et al. Circulation 2017;135:e686–725; 2. Aboyans V et al. Eur J Vasc Endovasc Surg 2018;55:305–68; 3. Criqui MH et al. Circulation 2021;144:e171–91. 4. American Heart Association. Peripheral Artery Disease and Diabetes, 2021. Available at https://www.heart.org/en/health-topics/peripheral-artery-disease/diagnosing-pad . 5. Yasuda, Tomohiro, et al. Scientific reports 7.1 (2017): 17425.



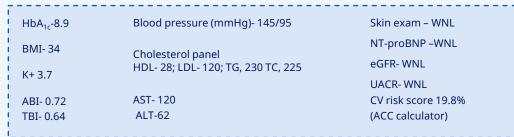


WALTER



- Male, 50
- Architect
- Enjoys walks in the park and reading
- Additionally, gained some weight in the past years and developed lower leg pain when walking.
- Comes to see a specialist to discuss HbA_{1c} that has increased over the past months
- Is very scared of disease progression and wants to do what he can for a healthy retirement
- Wants to feel control over the disease and ensure no further increase in CV risk

Patient characteristics





Medical history

- Metabolic syndrome >10 years
- T2D duration 4 years
- Metformin 1000mg BID
- Sitagliptin 100mg QD
- Simvastatin 40mg QHS
- Ramipril 5mg
- Dilated Eye Exam -4 yrs ago
- Foot Exam 4 yrs ago
- · No family history of CVD disease
- Non-smoker





