

Est.
1841

YORK
ST JOHN
UNIVERSITY

Mazzolai, Lucia ORCID:

<https://orcid.org/0000-0001-9650-3822>, Belch, Jill, Venermo, Maarit

ORCID: <https://orcid.org/0000-0001-8814-0988>, Aboyans, Victor

ORCID: <https://orcid.org/0000-0002-0322-9818>, Brodmann,

Marianne, Bura-Rivière, Alessandra, Debus, Sebastien, Espinola-

Klein, Christine ORCID: <https://orcid.org/0000-0001-9243-1761>,

Harwood, Amy E., Hawley, John A., Lanzi, Stefano ORCID:

<https://orcid.org/0000-0003-1089-6309>, Madarič, Juraj, Mahé,

Guillaume, Malatesta, Davide, Schlager, Oliver ORCID:

<https://orcid.org/0000-0001-6934-5355>, Schmidt-Trucksäss, Arno,

Seenan, Chris, Sillesen, Henrik, Tew, Garry ORCID:

<https://orcid.org/0000-0002-8610-0613> and Visonà, Adriana (2024)

Exercise therapy for chronic symptomatic peripheral artery disease.

Vasa, 53 (2). pp. 87-108.

Downloaded from: <http://ray.yorks.ac.uk/id/eprint/9734/>

The version presented here may differ from the published version or version of record. If you intend to cite from the work you are advised to consult the publisher's version:

<https://doi.org/10.1024/0301-1526/a001112>

Research at York St John (RaY) is an institutional repository. It supports the principles of open access by making the research outputs of the University available in digital form.

Copyright of the items stored in RaY reside with the authors and/or other copyright

owners. Users may access full text items free of charge, and may download a copy for

|

!

RaY

Research at the University of York St John

For more information please contact RaY at ray@yorks.ac.uk

1 **Title page - consensus document**

2

3 **Exercise therapy for chronic symptomatic peripheral artery disease: a clinical**
4 **consensus document of the ESC Working Group on Aorta & Peripheral Vascular**
5 **Diseases in collaboration with the European Society of Vascular Medicine, and**
6 **the European Society for Vascular Surgery**

7

8 **Lucia Mazzolai^{1*#}, Jill Belch^{2#}, Maarit Venermo^{3#}**, Victor Aboyans⁴, Marianne
9 Brodmann⁵, Alessandra Bura-Rivière⁶, Sebastien Debus⁷, Christine Espinola-Klein⁸,
10 Amy E. Harwood⁹, John A. Hawley¹⁰, Stefano Lanzi¹, Juraj Madarič¹¹, Guillaume
11 Mahé^{12,13}, Davide Malatesta¹⁴, Oliver Schlager¹⁵, Arno Schmidt-Trucksäss¹⁶, Chris
12 Seenan¹⁷, Henrik Sillesen¹⁸, Garry A. Tew¹⁹, Adriana Visonà²⁰

13

14 **# shared first authors**

15

16 ¹ Angiology Department, Lausanne University Hospital, University of Lausanne,
17 Switzerland

18 ² Institute of Cardiovascular Research, University of Dundee, Ninewells Hospital and
19 Medical School, Dundee, Scotland, United Kingdom

20 ³ Department of Vascular Surgery, Abdominal Center, Helsinki University Hospital,
21 Haartmaninkatu 4, 00029 Helsinki; University of Helsinki, Yliopistonkatu 4, 00100
22 Helsinki

23 ⁴ Department of Cardiology, Dupuytren-2 University Hospital, and EpiMaCT, INSERM
24 1094/IRD270, Limoges University, Limoges, France

- 1 ⁵ Division of Angiology, Department of Internal Medicine, Medical University, Graz,
2 Austria
- 3 ⁶ Department of Vascular Medicine, Toulouse University Hospital, France
- 4 ⁷ Department of Vascular Medicine, Vascular Surgery - Angiology - Endovascular
5 Therapy University of Hamburg-Eppendorf Hamburg Germany
- 6 ⁸ Center of Cardiology, Department of Cardiology III-Angiology, University Medical
7 Center of the Johannes Gutenberg-University Mainz, Mainz, Germany
- 8 ⁹ Department for Sport and Exercise Sciences, Manchester Metropolitan University,
9 Manchester, UK
- 10 ¹⁰ Exercise and Nutrition Research Programme, Mary MacKillop Institute for Health
11 Research, Australian Catholic University, Melbourne, VIC, Australia
- 12 ¹¹ Department of Angiology, Comenius University and National Institute of
13 Cardiovascular Diseases, Bratislava, Slovakia
- 14 ¹² Vascular Medicine Unit, Centre Hospitalier Universitaire de Rennes, 35033 Rennes,
15 France
- 16 ¹³ INSERM CIC 1414, Université de Rennes, 35033 Rennes, France
- 17 ¹⁴ Institute of Sport Sciences, University of Lausanne, Lausanne, Vaud, Switzerland
- 18 ¹⁵ Division of Angiology, Department of Medicine II, Medical University of Vienna,
19 Vienna, Austria
- 20 ¹⁶ Division of Sport and Exercise Medicine, Department of Sport, Exercise and Health,
21 University of Basel, Basel, Switzerland
- 22 ¹⁷ School of Health and Life Sciences, Glasgow Caledonian University, Glasgow, UK
- 23 ¹⁸ Department of Vascular Surgery, Rigshospitalet, Blegdamsvej 9, Copenhagen, DK,
24 2100, Denmark, and Department of Clinical medicine, University of Copenhagen

1 ¹⁹ Institute for Health and Care Improvement, York St John University, York, YO31
2 7EX, UK

3 ²⁰ Angiology Unit, Ospedale Castelfranco Veneto, Castelfranco Veneto, Italy

4

5 ***Corresponding author**

6 Prof. Lucia Mazzolai

7 Division of Angiology, Heart and Vessel Department

8 Lausanne University Hospital

9 Switzerland - Ch. de Mont-Paisible 18

10 1011 Lausanne, Switzerland

11 Tél. +41 021 314 07 68

12 lucia.mazzolai@chuv.ch

1 **Abstract**

2 All guidelines worldwide strongly recommend exercise as a pillar of the management
3 of patients affected by lower extremity peripheral artery disease (PAD). Exercise
4 therapy in this setting presents different modalities, and a structured programme
5 provides optimal results. This clinical consensus paper is intended for clinicians to
6 promote and assist for the set-up of comprehensive exercise programmes to best
7 advice in patients with symptomatic chronic PAD. Different exercise training protocols
8 specific for patients with PAD are presented. Data on patient assessment and outcome
9 measures are narratively described based on the current best evidence. The document
10 ends by highlighting disparities in access to supervised exercise programmes across
11 Europe, and the series of gaps for evidence requiring further research.

Graphical abstract

Included patients

- Women and men with symptomatic chronic peripheral artery disease
- Patients undergoing revascularisation

Initial exercise training

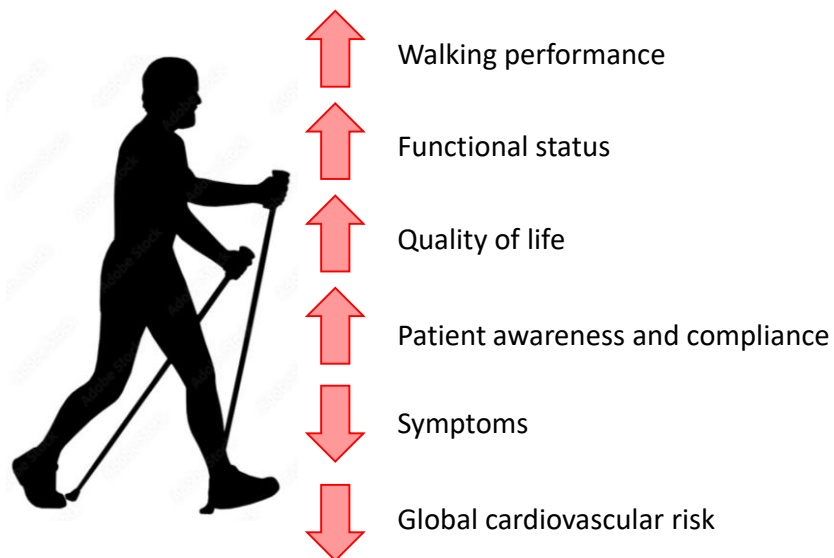
- Supervised exercise or home-based exercise training programmes
- *Training frequency*: at least 3 times per week
- *Training modality*: intermittent bouts of walking alternating with periods of rest are the first option. When walking is not an option, alternatives modalities (resistance training, arm-cranking, cycling, combinations of exercise) could also be considered.
- *Claudication pain intensity*: Based on strong evidence, patients should exercise to moderate-high claudication pain. No or low-pain approaches also shown improvement in walking ability, but the level of evidence is low. A flexible approach to pain intensity prescription is required, considering the patient's needs and preferences, and what might achieve high adherence.
- *Exercise intensity*: begin with a "lead-in period" of low-to-moderate intensity followed by, if tolerated, a gradual progression to vigorous exercise intensity.
- *Session duration*: at least 30 min
- *Programme duration*: at least 12 weeks
- Programmes should include advice and education about peripheral artery disease, cardiovascular risk factors, and lifestyle aiming for longer-term behavior change

Assessments prior and following exercise therapy

- Complete medical history, physical examination, and screening for contraindications
- Functional assessment
- Quality of life assessment
- Vascular assessments

Chronic exercise training

- Following initial exercise training (supervised or home-based), patients are encouraged to sustain lifelong and high levels of regular physical activity



1 **Introduction**

2 Physical activity, including regular exercise, is one of the pillars of cardiovascular (CV)
3 health and a major component of management of patients with most CV diseases
4 (CVD). In 2020, the European Society of Cardiology (ESC) issued a guideline
5 document addressing the main aspects of exercise therapy and sports practice for
6 cardiac diseases ¹.

7 In this consensus document, the acronym PAD will be used to indicate lower extremity
8 peripheral artery disease. PAD is one of the most prevalent clinical presentations of
9 atherosclerotic disease, affecting approximately 237 million people worldwide ². The
10 first symptoms of PAD are usually related to walking impairment, and the 2017
11 ESC/European Society for Vascular Surgery (ESVS) guidelines on the management
12 of PAD underscore the importance of exercise therapy, preferably supervised, for the
13 management of patients with intermittent claudication (IC) ³. Similarly, the 2019 PAD
14 guidelines of the European Society of Vascular Medicine (ESVM) encourage
15 structured exercise for symptomatic PAD patients ⁴. However, none of the
16 aforementioned documents provided in-depth guidance for exercise therapy in this
17 specific setting.

18 To address this gap, the ESC Working Group on Aorta & Peripheral Vascular Disease,
19 the ESVM, and the ESVS joined in a collaborative effort aiming to provide a roadmap
20 and guidance for the set-up and implementation of exercise therapy programmes for
21 patients with PAD.

1 **Consensus statements**

2 ▪ **1. In patients with PAD and exercise-induced limb symptoms due to**
3 **vascular origin, supervised exercise programmes should be the first line**
4 **treatment modalities.**

5
6 ▪ **2. In patients with PAD undergoing revascularisation, supervised exercise**
7 **programmes should be included as adjuvant therapy.**

8
9 ▪ **3. Supervised exercise programmes should ideally be coordinated by**
10 **vascular physicians, and sessions should be ideally supervised by**
11 **clinical exercise physiologists or physiotherapists.**

12
13 ▪ **4. Prior to exercise training initiation, complete medical history and**
14 **examination, and screening for contraindications should be investigated.**

15
16 ▪ **5. Measures of walking ability, functional status, and quality of life should**
17 **be assessed at the beginning and end of the programme to determine the**
18 **patient's response to exercise training. Clinical outcomes and patient**
19 **experience should also be documented.**

20
21 ▪ **6. Walking training (overground, pole striding, treadmill) should be**
22 **proposed as first line exercise modality. When walking is not an option,**
23 **alternative training modalities (resistance and strength training, arm-**
24 **cranking, cycling, combinations of exercise) should be performed.**

25

- 1 ▪ **7. The training frequency should be at least three times per week.**
- 2
- 3 ▪ **8. The training session duration should last a minimum of 30 min.**
- 4
- 5 ▪ **9. The training programme duration should last a minimum of 3 months.**
- 6
- 7 ▪ **10. Both claudication pain (A) and exercise intensity (B, based on common**
- 8 **training intensity measures such as heart rate or the rate of perceived**
- 9 **exertion (RPE) on Borg’s scale) should be evaluated during training**
- 10 **sessions:**
- 11 **A) The current consensus is that patients should exercise to moderate-**
- 12 **high claudication pain based on strong evidence. However, some**
- 13 **trials have recently demonstrated improvement in walking ability**
- 14 **using a low, or no pain approach. As claudication pain is a commonly**
- 15 **cited barrier to exercise, universally prescribing high-pain exercise**
- 16 **may lead to poor uptake of, and adherence to, exercise training**
- 17 **programmes. A more flexible approach to exercise prescription may**
- 18 **therefore be required, considering the patient’s needs and**
- 19 **preferences, and what might achieve a high level of (long term)**
- 20 **adherence.**
- 21 **B) Following a “lead-in period” of low-to-moderate exercise intensity, a**
- 22 **gradual progression to vigorous/high exercise intensity may be**
- 23 **proposed if well tolerated by the patient.**
- 24

- 1 ▪ **11. If supervised exercise is not available or feasible, a structured**
2 **community- or home-based exercise programme that includes behaviour**
3 **change techniques should be proposed.**
- 4
- 5 ▪ **12. Supervised exercise programmes should include structured education**
6 **and counselling on cardiovascular disease and PAD risk factor reduction.**
7 **Smoking cessation should be a cornerstone of risk factor counselling.**
- 8
- 9 ▪ **13. Following initial exercise training (supervised or home-based),**
10 **patients are encouraged to sustain lifelong and high levels of regular**
11 **physical activity.**

1 **Pathophysiology of intermittent claudication and functional impairment**

2 IC is characterised by exertional leg pain limiting walking ability⁵⁻⁷. PAD induces a wide
3 range of exercise-related symptoms experienced by nearly half of the PAD population
4⁸. The classical IC symptomology was first defined as calf pain, discomfort or fatigue
5 appearing during exercise and forcing the patient to stop⁹. Typically, IC is relieved
6 within 2-5 min after discontinuation of exertion⁹. Apart from this typical symptom, it is
7 now admitted that some patients with PAD may present atypical exercise-induced limb
8 symptoms¹⁰. These may be localised in lower limb muscles other than calves, may be
9 present at rest, may be described by patients as “burning”, “compressive” feeling, or
10 just “fatigue” without pain and may mimic limb pain due to spinal stenosis. Exercise-
11 induced limb symptoms in PAD are caused by a metabolic mismatch between oxygen
12 demand and supply⁵. The mismatch is linked to the reduction of the arterial lumen by
13 the atherosclerosis process, but it also induces cellular and metabolic disorders that
14 contribute to the functional impairment¹¹. Mechanisms of exercise-induced symptoms
15 are multifactorial among which nociceptive pain¹², nerve dysfunction¹³ and skeletal
16 muscle abnormalities¹¹ are suggested.

17 Potential mechanistic drivers of exertional limb symptoms in addition to arterial
18 obstruction and reduced perfusion include inflammation, vascular dysfunction, reduced
19 microvascular flow, impaired angiogenesis, and altered skeletal muscle function¹⁴⁻¹⁶
20 (Figure 1). A healthy vascular endothelium produces several vasodilator substances,
21 including nitric oxide (NO), which has pluripotent vascular benefits such as platelet
22 inhibition, smooth muscle cell proliferation inhibition, leukocyte adhesion prevention,
23 and angiogenesis induction. Diminished NO bioactivity in the lower limbs prevents
24 increased blood flow with exercise¹¹. Vascular dysfunction may also exacerbate the
25 vasoconstrictive effects of catecholamines and limit flow-mediated dilation¹⁷⁻²⁰.

1 Inadequate angiogenesis and collateral vessel formation may potentiate limb ischemia
2 and serve as a mechanism driving functional impairment ²¹. Skeletal muscle ischemia
3 may drive local inflammation, exacerbating symptoms and altering muscle metabolism
4 ²²⁻²⁴.

5 Patients with PAD present impaired walking endurance ²⁵, slower walking velocity ²⁶⁻
6 ²⁸, gait abnormalities ^{26,27,29-31}, poorer muscle strength ³², and poorer balance ^{33,34}
7 compared to individuals without PAD. They may also reduce their walking activity and
8 total activity to avoid leg symptoms ³⁵, and studies have shown a functional decline
9 occurring over time ^{25,28,36}.

10

11 **Vascular and functional assessment in PAD**

12 ***Vascular assessment***

13 General assessment of CV risk factors should be performed prior to exercise training
14 rehabilitation to improve preventive measures and reach preventive goals. Ankle-
15 Brachial Index (ABI) should be assessed before starting a training programme to detect
16 and diagnose PAD and assess disease severity (Figure 2) ³. The measurement of ABI
17 after exercise is also important to further detect ankle pressure drop, as some patients
18 may have leg symptoms on exercise while ABI can be ≥ 0.91 at rest. A post-exercise
19 ankle systolic blood pressure drop $>30\text{mmHg}$ or a post-exercise ABI decrease $>20\%$
20 should be considered for PAD diagnosis ³⁷. In patients with media calcinosis (for
21 example in patients with diabetes or chronic kidney disease) measurement of ABI
22 might not be possible because the arteries cannot be compressed by the cuff. In these
23 cases, toe brachial index (TBI) can be used as alternative assessment (the
24 pathological threshold usually retained is <0.70) ³.

25 ***Walking distance assessment***

1 Walking distance is considered an important clinical outcome both for patients and
2 clinicians. Standardised exercise testing should be used for assessment of functional
3 impairment in patients with PAD (Figure 2).

4 *Treadmill assessment.* Treadmill testing should be performed with patients familiarised
5 to the treadmill and under reproducible conditions (i.e. avoiding exercise and alcohol
6 prior to assessment). Patients should be asked to walk until maximal levels of pain,
7 lightly holding or not holding onto the treadmill. If the tests are stopped for reasons
8 other than leg pain, then this should be recorded. Patients are asked to indicate the
9 claudication pain score they reached during walking, especially the point at which pain
10 begins, and recovery based on a five-point scale (0 = no pain, 1 = onset of pain, 2 =
11 mild pain, 3 = moderate pain, 4 = severe/maximal pain)³⁸. Common treadmill protocols
12 include constant-load (single-stage) or graded exercise testing^{39,40}. The latter is
13 performed at constant speed varying the slope of the treadmill. Established graded
14 protocols include the Gardner/Skinner (3.2 km/h and a 2% increase in slope every 2
15 minutes) or the Hiatt protocol (3.2 km/h and an increase in slope of 3.5% every 3
16 minutes). Constant-load treadmill tests are performed at a fixed speed of 2 to 4 km/h
17 and fixed gradient of 10 to 12%. Constant-load protocols have poorer reliability both
18 for pain-free walking distance (PFWD) and maximal walking distance (MWD)
19 compared with graded protocols (coefficient of variance 30 and 45%, respectively)^{41,42}.
20 Treadmill tests have limitations including learning effect during repeated evaluations.
21 Also, some patients are unable or are unwilling to perform a treadmill test, mainly due
22 to balance impairment or limited walking abilities.

23 *Six-minute Walk Test.* The six-minute walk test (6MWT) is performed along a flat
24 corridor with a length of 30m with turning points marked by a cone. Patients are asked
25 to walk self-paced for the full duration and may stop and rest at any point in the test⁴³.

1 The total distance walked is measured and reported as the six-minute walking distance
2 (6MWD) ⁴³. Any encouragement given/phrases used should be the same for every test
3 performed to ensure test-retest reliability ⁴³. Further, there may be a learning effect so
4 it is recommended that the best out of two walks is recorded or the first test discounted
5 ⁴⁴. Although treadmill-based exercise tests can establish maximum walking capacity,
6 there may be a poor correlation between treadmill outcomes, habitual walking, and
7 self-reported walking distance ⁴⁵. On the other hand, compared to treadmill test, the
8 6MWT has been shown to better represent daily life walking in patients with PAD ⁴⁶.
9 The 6MWT is a well-validated and low-cost test. It has good reliability, with a correlation
10 coefficient of 0.90 (p<0.001) and a coefficient of variation of 8.9% with testing
11 performed one to two weeks apart ⁴⁷. Changes in the 6MWT can be used to predict
12 mortality and mobility loss in patients with PAD ^{7,48}. The minimal detectable changes
13 (i.e. the statistical detectability of change beyond measurement error) in the 6MWT are
14 represented by a change >46 meters ⁴⁹. The minimal clinically important difference (i.e.
15 the clinical relevance or importance of the observed change from the patient's
16 perspective) in the 6MWT in patients with PAD is represented by an improvement of 8
17 ⁵⁰ or 9 meters ⁵¹ for small changes, and 20 ⁵⁰ or 38 meters ⁵¹ for large changes.

18

19 *Connected Devices.* A measure of “real-life” walking performances may be performed
20 by use of global positioning systems (GPS) or commercially available devices such as
21 activity trackers, smart watches and phones ⁵². Research has shown that GPS
22 recorders have good accuracy and reliability when compared to known distances
23 walked ^{53,54}, and measurement of step counts with mobile phones has been shown to
24 be highly reliable even at low walking speeds ⁵⁵. Further, GPS recorded walking
25 distances correlate well with treadmill walking distances ⁵⁶. Patients should be able to

1 note the initial onset of claudication pain and the maximal walking distance either in
2 total or between bouts of walking using the GPS system.

3

4 ***Muscle strength assessment***

5 The presence of PAD is associated with impaired lower extremity muscle strength and
6 function ⁵⁷, which is associated with high prevalence of frailty and sarcopenia ⁵⁸.
7 Muscle strength and function should therefore be assessed before and after
8 supervised exercise training (SET, Figure 2). There is heterogeneity in how muscle
9 strength and function are assessed. Muscle isokinetic strength and endurance can be
10 assessed via isokinetic dynamometry, which is a chair device that patients sit on and
11 the specific joint is tested in an appropriate position with the dynamometer attached to
12 the limb. Patients push against the dynamometer as it provides resistance to maintain
13 a set speed. Isokinetic dynamometry has demonstrated good reliability at the ankle
14 (reliability coefficients ranging from 0.77 to 0.96) ⁵⁹. Testing can be done in various
15 joints, including ankle, knee, and hip, in various planes such as extension and flexion.
16 As isokinetic dynamometry assessment includes specialised equipment it may not be
17 practical or convenient to assess patients using this device. As an alternative, the short
18 physical performance battery (SPPB) which includes a 4-metre walk test, a sit-to-stand
19 chair test, and a standing balance test, should be used ⁶⁰. A recent study showed that
20 the sit-to-stand is a validated test to estimate muscle power in patients with
21 symptomatic PAD ⁶¹. Interestingly, muscle power assessed by the sit-to-stand test was
22 related to overall functional performance prior and following SET ⁶¹.

23

24

25

1 ***Self-reported functional impairment and quality of life assessment***

2 In addition to objective assessment of functional impairment, a subjective (self-
3 reported) evaluation of walking abilities and health-related quality of life (HRQoL)
4 should be incorporated to have a complete assessment of the functional status of the
5 patient (Figure 2) ⁶²⁻⁶⁴. Following exercise interventions, assessing HRQoL is usually
6 used to determine if an objective improvement in functional performance is also
7 perceived by the patients in their daily life. Table 1 reports the most used subjective
8 tools used for walking ability and HRQoL assessment in patients with PAD. Trials used
9 a wide variety of questionnaires of patient reported outcomes measurements (PROMs)
10 ⁶²⁻⁶⁴. The most used are the short-form health 36 (SF-36), a generic questionnaire
11 including physical and mental items related to health), and the Walking Impairment
12 Questionnaire (WIQ), a PAD-specific questionnaire focusing on PAD and functional
13 limitations. Studies have shown that HRQoL burden is greater in magnitude in patients
14 with both PAD and CVD than with CVD alone ⁶⁵. In the PARTNERS study, the SF-36
15 Physical Component Summary of the combined PAD-other-CVD group was 46.3 ± 1.2
16 compared with 55.5 ± 1.1 in the other-CVD group ⁶⁵. Cross-sectional studies show that
17 in patients with PAD the degree of difficulty in walking distance and stair climbing are
18 significantly related to HRQoL ⁶⁶. The ESVS VASCUNET and the International
19 Consortium of Vascular Registries consensus statement recommended the Vascular
20 Quality of Life Questionnaire-6 (VasCu-QoL6) as a primary assessment of PROMs in
21 patients with symptomatic PAD ⁶².

22 Greater amounts of physical activity are associated with higher ratings of both
23 perceived health and HRQoL, correlating with objective health outcomes and life
24 expectancy ⁶⁷. One of the most important factors linked to both subjective and objective
25 health, across both cognitive and physical domains, is physical activity ⁶⁸.

1 **Exercise therapy in patients with PAD**

2 ***Screening prior to exercise training participation***

3 All patients should be medically screened before SET programme initiation (Figure 2).
4 It is suggested to include a complete medical history and examination ³⁸. Patients with
5 contraindications to exercise training (Table 2) should be excluded from SET until the
6 relevant condition stabilises or is successfully treated. For patients with current or prior
7 symptomatic cardiac disease (Table 3), we recommend that they are referred for
8 cardiology work-up, including an exercise test to assess for evidence of exercise-
9 induced coronary ischaemia, to identify if additional treatment for cardiac disease is
10 required before proceeding with SET. Comorbidities (such as neurological and
11 orthopedic diseases leading to gait abnormalities) should be documented and
12 considered for how they may limit SET programme participation feasibility. After SET
13 programme initiation, patients should continue to be closely monitored for changes in
14 health status (e.g., any symptom or situation which may suspect undiagnosed/incident
15 cardiac condition, ischemic limb pain at rest, toe or foot wounds) that might necessitate
16 interruption of the programme, at least temporarily.

17

18 ***Supervised exercise training***

19 SET is considered among first-line therapies for patients with chronic and symptomatic
20 PAD (Figure 2) ^{3,64,69,70}. SET is safe and is usually conducted in the hospital setting ⁷¹.
21 Over the past 60 years, many trials have reported the effectiveness of SET on walking
22 distances in these patients ^{72,73}. The most recent Cochrane meta-analysis showed that
23 SET improves PFWD (82 m; 95% IC: 72 – 92) and MWD (120 m; 95% IC 51 – 190) ⁷⁴.
24 Similar findings were observed in another meta-analysis [PFWD: 128 m (95% IC: 92 –
25 165); MWD: 180 m (95% IC: 130 – 238)] ⁷⁵. Although less well investigated or usually

1 reported as a secondary outcome, SET also improved functional status, gait pattern,
2 self-reported walking ability and quality of life ^{64,74,76-82}. It is interesting to note that
3 cardiac rehabilitation programmes also increase walking distance, HRQoL, and
4 physical activity in patients with symptomatic PAD, suggesting that other types of
5 rehabilitation than SET may also be useful ⁸³. Finally, some vasoactive drugs such as
6 cilostazol (phosphodiesterase type 3 inhibitor), pentoxifylline (xanthine derivative),
7 bosentan, sildenafil and others are claimed to increase walking capacity in patients
8 with PAD ⁸⁴⁻⁸⁷. However, the objective documentation of their effect is very limited to
9 draw extensive conclusions ^{84,88}. More studies are needed to confirm additive effect of
10 drug therapies to supervised exercise.

11

12 *Training modalities.* There are different types of exercise training for patients with PAD,
13 but the common aim is to improve walking capacity and reduce symptoms. In addition,
14 exercise should aim to improve balance and muscle strength to promote independence
15 and a reduced risk of falling in the long-term ³³. Treadmill and overground walking are
16 the most common and recommended training modalities in patients with IC (Figure 2)
17 ^{64,70}. However, due to severe exercise-induced ischemia, low pain tolerance, the risk
18 of falling and/or other co-morbidities, some patients are unwilling or unable to perform
19 walking sessions. In addition to walking training, there are several other forms of
20 training that are used, although much less frequently, in the rehabilitation of patients
21 with PAD. A recent meta-analysis reported that other non-walking training modes are
22 also effective as traditional walking training in improving walking performance, whereas
23 there was no clear evidence for changes in quality of life following exercise
24 interventions. However, the authors concluded that the certainty of this evidence was
25 judged to be low ⁸⁹. Different training modes include strength training of large muscle

1 groups ^{90,91}, cycling ⁹², pole striding ^{93,94}, multimodal training ^{76,77,95-98} and training with
2 an arm-crank ergometer ^{99,100}. The beneficial effect of these training modalities can
3 usually be described as large and even reach those of typical walking training ¹⁰¹.
4 However, the PFWD and the MWD have the tendency to be higher with walking training
5 than with strength training when all studies are considered ⁸⁹. In contrast, self-reported
6 ability to climb stairs (assessed by the Walking Impairment Questionnaire) is more
7 improved following strength training (29.2% vs. 43.8% after 6 months) compared to
8 walking training on the treadmill (39.6% vs. 43.8% after 6 months) ¹⁰². Therefore, when
9 walking is not an option, alternative training modalities might also be effective. These
10 training modalities also elicit lower or no pain during exertion compared to walking,
11 which might lead to higher rates of adherence.

12

13 *Training frequency.* Based on a previous meta-analysis, and shared by most of the
14 studies and guidelines, the training frequency associated with greater improvements
15 in walking distance is at least 3 times per week ^{103,104}.

16

17 *Training duration.* Identifying an optimal training duration is difficult to elucidate, mainly
18 due to differences in training modalities, frequencies, and intensities among studies.
19 Current guidelines reported that optimal training duration ranges between 12 and 24
20 weeks ^{64,70,103}. The optimal training session duration has not been widely investigated.
21 Additionally, in most studies, the total session duration is usually reported without
22 specifying the actual time spent exercising. The literature shows that exercise sessions
23 lasting 30 to 60 min were the most effective to improve walking performance ^{103,104}.

24

1 *Training intensity.* In most studies, no clear distinction is made between symptom
2 intensity (claudication pain scale) and exercise training intensity [based on heart rate
3 (HR), oxygen uptake ($\dot{V}O_2$) or rate of perceived exertion (RPE) on Borg's scale: 6: "very
4 very light"; 20: "maximal effort"] to monitor the exercise therapy. The Borg scale is a
5 subjective assessment tool used to measure an individual's perceived exertion or effort
6 during physical activity. The scale assigns a numerical rating ranging from 6 to 20 to
7 indicate the intensity of exertion experienced by the person ¹⁰⁵.

8 First, the majority of trials used claudication pain severity to provide guidance during
9 the training sessions. In PAD research, the claudication pain scale, an ordinal scale
10 from 0 (no pain) to 4 (severe/maximal pain), is the most commonly used tool. A
11 distinction is made between walking training with and without muscle pain caused by
12 ischemia. With regards to claudication pain intensity, international guidelines are
13 heterogeneous ^{38,64,70}. The UK NICE guideline encourages patients to exercise to the
14 point of maximal pain , the American Heart Association guideline recommends
15 moderate to moderate/severe claudication pain as tolerated ⁶⁴, while an international
16 consensus as well as the Australian guideline does not specify pain intensity for
17 exercise dosage ¹⁰⁶. Based on strong evidence ^{64,73-75,104}, the current consensus is that
18 patients should exercise to moderate-high claudication pain to improve walking
19 performance. Also, one-year home-based walking training performed at high-intensity
20 pain has been found to be more effective than walking training performed at low-
21 intensity for improving walking and functional performance in patients with PAD ^{107,108}.

22 These findings indicate that claudication pain intensity may be a key factor for walking
23 improvement in these individuals. In contrast, others have reported that improvements
24 in walking performance may be obtained with less severe claudication pain during
25 exertion ¹⁰¹. According with recent findings, walking training with pain is not clearly

1 superior to walking training without pain regarding changes in walking distances ¹⁰⁹⁻¹¹².

2 It may be assumed that walking training with moderate, low, or no pain is associated
3 with higher compliance and possibly long-term maintenance of training or change in
4 activity behaviour ¹¹². This indicates that a more flexible approach to exercise
5 prescription may therefore be required, considering the patient's needs and
6 preferences, and what might achieve a high level of (long term) adherence. Larger
7 studies with a higher number of cases and longer duration, taking compliance into
8 account, are needed for a conclusive statement ¹¹³.

9 Second, the optimal no/low pain-based exercise training intensity is understudied in
10 this population. Indeed, it is interesting to note that the claudication pain severity does
11 not necessarily rely on common measures of exercise intensity ^{78,114}. For example,
12 when performing vigorous-intensity exercise, some patients may experience
13 moderate-to-severe claudication pain, whereas others, low levels of claudication only.
14 Assuming that exercise intensity is a cornerstone determinant of physiological
15 response to training ¹¹⁵, monitoring claudication pain only is limiting and prevents
16 accurate comparison of exercise effectiveness in patients with PAD. This may also
17 explain the large variability in the magnitude of improvements following exercise
18 interventions ^{64,103}. Fassora et al. ⁷⁸ recently reported that both training modality and
19 exercise intensity should be considered when looking for the best results in walking
20 performance and cardiorespiratory fitness. Notably, these results showed that walking
21 at vigorous intensity ($\%HR_{peak}$: 77-95, $\% \dot{V}O_{2peak}$: 64-90, RPE: ≥ 14 ¹¹⁵) induced the
22 greatest improvement in MWD, while cycling and other non-walking modalities
23 performed at vigorous intensity elicited the greatest improvements in cardiorespiratory
24 fitness ⁷⁸. These findings suggest that both walking and cardiorespiratory capacities
25 are desirable outcomes but that they need different exercise therapy programmes ⁷⁸.

1 It is however important to note that training programmes should start with a lead-in
2 period performed at low-to-moderate exercise intensity and, if tolerated, gradually
3 progressed to vigorous exercise intensity. This approach may allow to determine the
4 patient's exercise response and tolerance, reducing the risk of complications.

5 The monitoring of the exercise intensity during a resistance training program is
6 mediated by the percentage of the one repetition maximum (1RM) ¹¹⁶. The
7 determination of the 1RM plays a key role to objectively set an individualised
8 resistance-based program ¹¹⁶. Compared to a direct assessment of the 1RM, the
9 multiple RM assessment (such as 10RM, the maximum weight a person can lift for 10
10 repetitions) is considered to be a safe and well tolerated approach to evaluate muscle
11 strength for a given muscle group in patients with cardiovascular diseases ¹¹⁶.

12 Following the multiple RM test, different prediction equations are available to estimate
13 the 1RM ¹¹⁷. As also used in the cardiac rehabilitation, a target exercise intensity of 30-
14 70% of 1RM for the upper body, and 40–80% of 1RM for the lower body should be
15 considered ¹¹⁷. Exercise intensity should be progressively increased to determine the
16 patient's exercise response and exercise tolerance. It has been shown that resistance
17 training improves walking performance and muscular strength in patients with PAD ¹¹⁸.

18 Notably, high intensity (i.e. 80% 1RM) induces the best improvements in walking
19 performance when compared to low-to-moderate (i.e. <50% 1RM) strength training
20 intensity in these patients ^{90,118}.

21 Table 4 summarises the main exercise prescription recommendations with some
22 practical applications.

23
24
25

1 ***Home-based exercise training***

2 In comparison with patients not undergoing exercise training, a home-based training
3 (HBT) strategy resulted in a non-significant increase of MWD in a recent meta-analysis
4 (mean difference: 136 m; 95% CI: -2 to 273 m; $p = 0.05$) ¹¹⁹. When comparing HBT
5 with basic exercise advice, no improvement of MWD was observed in patients
6 following a HBT strategy (mean difference: 39 m; 95% CI: -123.1 to 201.1 m; $p = 0.64$)
7 ¹¹⁹. Regarding PFWD, HBT led to a greater increase than exercise advice did (mean
8 difference: 64.5 m; 95% CI: 14.1 to 114.8 m; $p = 0.01$) ¹¹⁹. In comparison with HBT,
9 SET was more effective in improving MWD (mean difference: 139 m; 95% CI: 45 to
10 232 m; $p = 0.004$) and PFWD (mean difference: 84 m; 95% CI: 25 to 143 m; $p = 0.005$)
11 ¹¹⁹.

12 Considering the effect of monitoring in HBT, no difference in the change of MWD and
13 PFWD were observed between monitored HBT and SET (mean difference in MWD: 8
14 m; 95% CI: -81 to 97 m; $p = 0.86$; mean difference in PFWD: 43 m; 95% CI: -29 to 114
15 m; $p = 0.24$) ¹¹⁹. The equality in training efficacy of monitored HBT and SET
16 emphasises the role of monitoring in HBT programmes. Apart from regular on-site
17 visits or phone calls, activity diaries or log books have been used for HBT monitoring
18 ¹¹⁹. Additional tools for self-monitoring, such as wrist-worn activity trackers with
19 smartwatch-like functions or smartphone accelerometer applications have been
20 assessed, however, it still needs to be clarified, which modality is most appropriate ⁵⁵.
21 The effect of training on patients' daily physical activity was assessed by several
22 studies implementing pedometer- and accelerometer-measurements. A network meta-
23 analysis demonstrated improvements of daily physical activity in HBT to a similar
24 extent as it was observed in patients undergoing SET ¹²⁰.

1 Focusing on quality of life, most studies reported improvements in patients undergoing
2 HBT ¹¹⁹. In comparison with SET, improvements of individual SF-36 measures (pain
3 and social functioning) and Walking Impairment Questionnaire measures (distance)
4 were less pronounced in patients undergoing HBT ¹¹⁹. In addition, HBT improves
5 measures of self-efficacy for walking, satisfaction with functioning, pain acceptance
6 and social functioning in patients with claudication ¹²¹. Follow-up data of patients who
7 had undergone HBT suggest sustained improvements in measures of quality of life,
8 functional and walking capacity after termination of the active training intervention
9 ^{122,123}.

10 Safety of HBT was analysed in a systematic review including 27 studies, which
11 reported a cardiac event rate of 1 per 49,270 and a non-cardiac event rate of one per
12 147,810 ¹²⁴. Event rates of HBT were lower than event rates reported for SET (HBT
13 vs. SET: cardiac 1:49,270 vs. 1:13,788; non-cardiac: 1:147,810 vs. 1:41,363) ¹²⁴.
14 Regarding overall mortality, retrospective data suggest a reduction of long-term
15 mortality in patients undergoing HBT ¹²⁵. Comparing HBT with SET, overall mortality
16 rates do not differ between patients undergoing HBT and patients following a SET
17 programme ¹²⁶. The results of the reported meta-analyses and reviews should be
18 viewed with caution according to a moderate to low quality of evidence ^{119,126,127}. Due
19 to the limited availability and utilisation of SET programmes, HBT programmes can be
20 used as a valid alternative training modality for patients with IC ¹²⁸⁻¹³¹.

21 Data on sex-specific differences in the efficacy of HBT are inconsistent ^{132,133}. In
22 females, the efficacy of HBT appears to be more strongly related to the individual
23 training intensity than in males ¹³⁴. Regarding co-morbidities, HBT seems to be less
24 effective in patients with diabetes with respect to the potential increase in walking
25 capacity ¹³⁵. In elderly patients, HBT potentially improves quality of life to a similar

1 extent as revascularisation does ¹³⁶. Considering the frequency of HBT training, 3
2 weekly sessions was the most commonly training strategy (range: 3 weekly sessions
3 to daily sessions) ¹¹⁹. For initiation, patients should start with a duration of 20 minutes
4 per session, progressively increasing the duration to 60 minutes per session. HBT can
5 be performed outside, around a track or in a hallway at a self-selected pace ^{51,137}.

7 ***Long-term adherence to exercise therapy***

8 In clinical practice, long-term adherence to therapy is a major problem. Participating in
9 SET programmes may help patients to acquire awareness of the disease and learn the
10 importance of exercise and how to practice it. SET programmes can be regarded as a
11 transition phase to improve self-management and may serve as a bridge for those
12 patients that need it to other forms of exercise approach such as community or home-
13 based exercise. Telemedical monitoring through step counting with pedometers or
14 activity monitors proved to be effective ^{138,139}, as did supervised structured walking
15 exercise to improve pain-free and maximal walking distance ¹¹⁹. In addition to
16 monitoring, factors such as education, self-efficacy, goal setting, feedback, and a
17 training plan were critical to successful outcomes ¹¹⁹. This should be used more
18 frequently in clinical practice to increase long-term adherence but needs to be
19 demonstrated in long-term studies.

21 **Mechanisms of response to exercise in PAD**

22 Exercise represents a major challenge to whole-body homeostasis provoking
23 widespread perturbations in numerous cells, tissues, and organs that are caused by or
24 are in response to the increased bio-energetic activity of the contracting skeletal
25 musculature ¹⁴⁰. The exercise training-induced increase in functional capacity and the

1 concomitant amelioration of diverse maladaptive responses that ultimately reduce
2 claudication symptoms in patients with PAD, are underpinned by several inter-
3 dependent physiological, metabolic, and mechanical mechanisms. After several
4 months of exercise training there is extensive remodelling of the vascular system, and
5 although direct sampling of the vasculature in humans *in vivo* is limited, the trained
6 musculature provides a valid proxy, being the primary tissue involved in training
7 adaptation ¹⁴⁰. The dynamic biochemical and mechanical environment around blood
8 vessels arising from the forces provoked during skeletal muscle contractile activity (i.e.,
9 shear stress and passive stretch), as well as signals stimulated by the increases in
10 muscle energetic demand (i.e., increases in AMP concentration, reduced oxygen
11 delivery) activate several intracellular signalling pathways responsible for promoting a
12 regulatory network governing the transcriptional control of mitochondrial biogenesis
13 and respiratory function along with enhanced expression of pro-angiogenic factors ¹⁴¹
14 (Figure 3).

15 Over time, this results in the initiation of capillary growth and a proliferation in the
16 number of arterioles. Such structural remodelling is driven by a complex and often-
17 redundant sequence of events that include NO, and prostaglandins. Indeed,
18 mechanical, neural, and humoral factors, including those released from contracting
19 skeletal muscle, have all been implicated in the remodelling response, with the
20 vascular endothelial growth factor (VEGF) signalling pathway and downstream targets
21 ultimately driving skeletal muscle capillary expansion ¹⁴¹. Muscle activity increases
22 VEGF in the muscle interstitium and subsequently acts on the VEGF receptors,
23 VEGFR-1 and VEGFR-2 on the capillary endothelium, activating multiple downstream
24 pathways via signalling intermediates such as mitogen activated protein kinases
25 (MAPK), phosphatidylinositol-3-Kinase ¹⁴². The time-course of remodelling varies and

1 is largely a function of the blood vessel size, and while many of these adaptations are
2 restricted to the vascular beds of the trained muscles, improved endothelial function
3 appears to be a whole-body response to exercise training, even in individuals with
4 PAD.

5 VEGF expression is partially regulated by the hypoxia-inducible factor-1 α (HIF-1 α) but
6 recently the peroxisome proliferator-activated receptor gamma coactivator-1 α (PGC-
7 1 α) has emerged as an important candidate in the exercise-induced angiogenic
8 response. PGC-1 α regulates the coordinated expression of mitochondrial proteins
9 encoded in the nuclear and mitochondrial genomes and is rapidly induced after
10 exercise. This protein has been called the “master regulator” of mitochondrial
11 biogenesis, and controls various aspects of muscle oxidative phenotype, while
12 transducing and integrating physiological signals governing metabolism,
13 differentiation, and cell growth, and suppressing a broad inflammatory response ¹⁴³.
14 Thus, the PGC-1 coactivators serve as a central component of the transcriptional
15 regulatory circuitry that coordinates the energy-generating functions of mitochondria in
16 accordance with the metabolic demands imposed by exercise training undertaken by
17 patients with PAD.

18

19 **Exercise and revascularisation**

20 Current guidelines recommend SET programmes as an initial treatment modality for
21 patients with IC ^{3,144}. Revascularisation is recommended for patients with IC when they
22 do not respond to initial exercise and medical therapies ¹⁴⁵. However, the role of
23 revascularisation as an initial treatment option alone or as an upstream adjunct to SET
24 in patients with IC remains controversial.

1 Several trials have compared endovascular therapies with or without SET versus SET
2 alone as an initial treatment strategy for patients with PAD with IC and reported
3 inconsistent results ¹⁴⁶⁻¹⁴⁹.

4 The relevant aspect of exercise training may be the reduction of the inflammatory
5 process in patients with PAD. In a recent trial, reactive oxygen species (ROS) formation
6 was measured using the luminol analogue L-012 for patients with IC, randomised
7 either to home-based training alone or in addition to endovascular therapy (EVT) ¹⁵⁰.
8 Follow-up was performed after 3 months. ROS production after NOX2 (NAPDH
9 oxidase 2) stimulation showed a significant reduction in both groups at follow-up (EVT
10 group: $p = 0.002$, exercise group: $p = 0.019$), with a higher relative reduction in ROS
11 in the EVT group than in the exercise group ($p = 0.014$).

12 The data regarding the benefit of SET alone or in combination with EVT or EVT alone
13 are rare. A robust evaluation of existing data comes from a meta-analysis comparing
14 the different treatment approaches ¹⁵¹. A total of 987 patients from 7 randomized
15 control trials (constituting 9 total comparison arms) with a median follow-up duration of
16 12.4 months (range 10 to 18 months) were enrolled. Of these, 530 patients were
17 randomized to EVT versus SET alone, and 457 patients to EVT plus SET versus SET
18 alone ¹⁵¹. For the effect of EVT alone versus SET alone (5 comparison arms) a random
19 effects model showed no significant difference in the MWD (standardised mean
20 difference (SMD): -0.11 (95% CI: -0.59 to 0.36); $p = 0.64$) on follow-up between the 2
21 groups, neither for the PFWD, need for revascularisation or amputation. On pooled
22 analysis, the ABI was significantly higher among participants that underwent EVT
23 alone as compared with SET only (SMD: 0.64 ; 95% CI: 0.38 to 0.90 , $p < 0.0001$;
24 weighted mean difference (WMD): 0.15 ; 95% CI: 0.10 to 0.19 , $p < 0.0001$).

1 On pooled analysis using random effects models, EVT plus SET (4 comparison arms)
2 was associated with significantly higher MWD on follow-up compared with SET alone
3 (SMD: 0.79; 95% CI: 0.18 to 1.39, $p = 0.01$), as well as significantly higher ABI on
4 follow-up compared with SET only (SMD: 0.62; 95% CI: 0.33 to 0.91; WMD: 0.14; 95%
5 CI: 0.10 to 0.17, $P < 0.0001$).

6 The combination of EVT plus SET was also associated with a significantly lower risk
7 of revascularisation or amputation on follow-up (3.5% vs. 17.3%, OR: 0.19; 95% CI:
8 0.09 to 0.40, $P < 0.0001$). The corresponding number needed to treat was 8 patients
9 (95% CI: 6 to 12). PFWD was reported in 2 studies with no difference between the 2
10 groups in random effects pooled analysis ¹⁵¹. However, EVT alone is not associated
11 with better outcomes than SET ^{151,152}. Among patients with stable PAD and IC,
12 compared with SET alone, endovascular revascularisation in combination with SET is
13 associated with improved outcomes.

14 Exercise training after surgical revascularisation also improves outcomes compared to
15 revascularisation without exercise training. Although much less investigated, few
16 publications exist on the impact of exercise on the outcome after surgical
17 revascularisation of symptomatic PAD. One small RCT compared patients after bypass
18 surgery ($n=14$) ¹⁵³. Group I had standard preoperative and postoperative care and the
19 intervention group (group II) had SET 4-10 weeks postoperatively. MWD, mean
20 increase in ABI and improvement in WIQ were significantly better in group II. In another
21 recent study, patients who underwent above knee femoropopliteal bypass were divided
22 into two groups: those who continued regular exercise after bypass operation with
23 those who discontinued exercise after surgery ¹⁵⁴. After propensity score matching, 5-
24 year primary and secondary patency (PP: 97% vs. 61%, $p = 0.0041$; SP: 100% vs.
25 69%, $p = 0.0021$), and freedom from major adverse cardiovascular events (61% vs.

1 24%, $p = 0.0071$) were significantly better in patients who continued exercise. One
2 systematic review included all RCTs with either surgical or endovascular
3 revascularisation to evaluate the evidence on the efficacy of lower limb
4 revascularisation combined with supervised exercise training in patients with PAD ¹⁵⁵.
5 Eight trials with 726 patients showed that combined therapy led to greater
6 improvements in PFWD and MWD compared with revascularisation or supervised
7 training alone. In 2 out of 8 studies, revascularisation was surgical and in 6 studies it
8 was endovascular.

9

10 **Effect of exercise on health-related quality of life and cognitive function**

11 Poor HRQoL is associated with higher rate of mortality in patients with PAD ¹⁵⁶.
12 Randomised controlled trials have shown that exercise training versus usual medical
13 care in patients with PAD not only improves the perceived walking distance and speed,
14 but also the functional status as measured by specific impairment questionnaires, as
15 the WIQ. When compared to controls, patients who complete any form of exercise
16 training significantly improve their WIQ speed (mean difference: 9.60; 95% CI: 6.98 to
17 12.23, $p \leq 0.001$); WIQ distance (mean difference: 7.41; 95% CI: 4.49 to 10.33, $p \leq$
18 0.001) and WIQ stair-climbing (mean difference: 5.07; 95% CI: 3.16 to 6.99, $p \leq 0.001$)
19 ⁸⁰. In addition, more general HRQoL evaluation scores (Short-Form Physical
20 Component Summary) also showed significant improvement following exercise
21 therapy (mean difference: 1.24; 95% CI: 0.48 to 2.01) ⁸⁰. Most of the studies showed
22 that 3- ¹⁵⁷⁻¹⁵⁹, or 6/12-month ^{94,102,160} exercise training improves patient's perception of
23 physical HRQoL, with lesser effects on mental HRQoL. However, in the current
24 literature, findings are inconsistent ^{74,80,161} and other studies did not find the same
25 effects ¹⁶²⁻¹⁶⁴. It is interesting to note that the improvement in general HRQoL scores

1 (as SF-36) were mainly predicted by physical functional markers, such as the distance
2 covered during a 6MWT (6MWD) and the history of stumbling ¹⁶⁵. These data indicate
3 that greater improvements in physical function following exercise therapy are expected
4 to have greater improvements in self-perceived HRQoL ¹⁶⁵. It has recently been
5 showed that improvements in 6MWD following SET are predictive of augmentations in
6 general HRQoL in patients with PAD ⁹⁶. Interestingly, changes in treadmill
7 performance, which are less representative of functional walking ⁴⁶, were not related
8 to improvements in HRQoL ⁹⁶.

9 Regular physical activity is also known to improve cognitive functioning and brain
10 health across the lifespan ¹⁶⁶. Cross-sectional and experimental studies show that
11 greater amounts of physical activity are linked to better cognitive function in adults,
12 with the best performances for exercise programmes that are structured,
13 individualised, higher intensity, longer duration, and multicomponent ¹⁶⁷. These results
14 support a dose-dependent neuroprotective relationship between physical exercise and
15 cognitive performance. Physical exercise interventions aimed at improving brain health
16 through neuroprotective mechanisms show promise for preserving cognitive
17 performance ¹⁶⁷. Scientific evidence based on functional and neuroimaging approach
18 has demonstrated that this relation could be mediated by improved brain integrity,
19 including adaptations in cerebral blood flow, volume and white matter integrity ¹⁶⁸.

20

21 **Patient education**

22 All patients with PAD should be offered oral and written information about their disease
23 so they can share decision-making and understand what they can do to help manage
24 their condition. The role of exercise should be clearly explained, and patients should
25 be supported to exercise regularly (assuming no contraindications). The impact of

1 patient education regarding exercise is probably dependent on several factors,
2 including the specific information that is provided, the timing and mode of delivery, and
3 the nature of any interventions that are delivered concomitantly (e.g., SET). Patient
4 education in the form of brief exercise advice, when delivered in isolation, confers little
5 benefit and results in minimal improvement in individuals' walking distances ¹⁶⁹.
6 Structured education programmes, on the other hand, may have greater potential to
7 improve exercise behaviour and walking distances by building the knowledge and skills
8 of patients to enable them to successfully self-manage their condition ¹⁷⁰. Key
9 programme features include: a structured evidence-based curriculum that includes
10 content on the nature of the condition and the role of exercise; delivery by trained
11 educators; and embedded quality assurance processes ¹⁷⁰.

12 A systematic review by Abaraogu et al. ¹⁷⁰ identified six studies (1,087 participants)
13 that had investigated the effects of structured education for patients with PAD and IC.
14 The interventions varied widely, but all included education sessions, exercise
15 prescription, and behaviour change techniques. Four trials reported improvements in
16 walking ability in intervention versus control comparisons ¹⁷⁰. Effects on physical
17 activity and quality of life were mixed. Overall, the evidence was inconclusive and more
18 rigorous trials are needed that include a clear and complete description of the
19 education intervention. Participant feedback from three studies highlights intervention
20 features that may be important for improving physical activity: providing information
21 about PAD/IC and exercise; providing encouragement and support with self-
22 monitoring; and having group interaction while allowing space for individual discussion
23 ¹⁷⁰.

24 Three other trials have tested exercise programmes that had an educational
25 component in patients with PAD ¹⁷¹⁻¹⁷³. The GOALS trial ¹⁷² randomized 194

1 participants either to a group-mediated cognitive behavioural intervention or an
2 attention control group. The intervention consisted of group meetings with a facilitator
3 once weekly for 6 months. Discussion topics included effective behaviour change
4 methods, self-monitoring, exercising in cold weather, managing leg pain during
5 exercise, and overcoming other obstacles to exercise adherence. At the 6-month
6 follow-up, the intervention group achieved a 53.5 meters greater increase in 6MWD
7 compared with the control group. Next, the HONOR trial ¹⁷³ tested the efficacy of
8 telephone coaching combined with a wearable activity monitor and showed no
9 improvement in 6MWD at the 9-month follow-up. Finally, the MOSAIC trial explored
10 the effect of a physiotherapist-delivered motivational interviewing intervention in 190
11 patients with PAD and IC ¹⁷¹. A statistically significant mean difference of 16.7 m in
12 6MWD was observed at 3 months follow-up compared with usual care control ¹⁷¹. The
13 contrasting results of these trials indicate that exercise programmes that include
14 education are more likely to be successful if they include periodic visits to a medical
15 centre to meet with a coach or include tailored behaviour change components.

16

17 **Sex and exercise**

18 Prevalence of PAD in women is similar to men at all ages ^{174,175}. However, women are
19 more likely to have asymptomatic PAD and less likely to report IC ¹⁷⁶. Decreased
20 detection and subsequent intervention may then result in a higher proportion of women
21 with severe disease and chronic limb-threatening ischemia. Further, women who
22 undergo revascularisation tend to be older and have more severe PAD compared to
23 men, and these factors can affect outcomes of procedures adversely ¹⁷⁷. Contradictory
24 results exist on women with PAD and mortality rates ¹⁷⁸⁻¹⁸⁰. Population studies suggest
25 a trend towards higher mortality rates in women with lower ABI ¹⁷⁹.

1 Exercise performance has been used to suggest that women decline faster in terms of
2 functional ability once PAD is established. However, this difference may in fact merely
3 be due to the smaller muscles in the calves of women ¹⁸¹. McDermott et al. ¹⁸² showed
4 that at 4 years of follow-up, women were more likely to become unable to walk for 6
5 min continuously than men, more likely to develop mobility disability, had faster
6 declines in walking velocity, and the distance achieved in the 6MWT was less.
7 However, these apparent sex differences in functional decline were attenuated after
8 additional adjustment for baseline calf muscle area, and so may be attributable to
9 smaller baseline calf muscle area in women. Interestingly poorer leg strength is
10 associated with increased mortality in men, but not in women, with PAD ¹⁸¹.

11 The data on the efficacy of exercise rehabilitation in women with PAD compared to
12 men are scarce. What is known, however, is that women with IC seem to have a poorer
13 response to exercise rehabilitation, smaller changes in PFWD and MWD following
14 three months of exercise than men (Δ 280 meters for men vs Δ 220 meters for women;
15 $p = 0.04$) ¹⁸³. This is particularly so in those with diabetes ¹³². Reduced blood volume
16 expansion and slower oxygen kinetics occur in the calf musculature during exercise in
17 women with PAD with IC ¹⁸⁴. Further, recent data showed that this poor response to
18 exercise in women with IC and diabetes was not related to where the intervention was
19 performed, being impaired both in a supervised exercise class and a home exercise
20 setting ¹³². This poorer response to exercise was also demonstrated in the EXITPAD
21 study, which showed that women with IC, independent of confounding factors including
22 diabetes, benefit less from supervised exercise and have significantly lower MWD after
23 12 months. Higher level of metabolic syndrome presents in postmenopausal women
24 compared with similarly aged men, may contribute to this ¹⁸³. On the contrary, it has
25 recently been shown that multimodal SET (combining strengthening of lower limbs and

1 Nordic walking) significantly improves walking performance (treadmill and overground)
2 in women and men, with no difference between groups ^{98,185}. Although not significant,
3 it is interesting to note that women had greater improvements (i.e., delta) than men ⁹⁸.
4 The clinical implication is that women with IC may respond less well to current exercise
5 interventions and either need a greater 'dose' of exercise, or another intervention
6 separate or in combination with exercise, to obtain similar improvements in IC as that
7 seen in men with exercise alone.

8

9 **Situation in Europe**

10 Despite of the large body of evidence highlighting benefits, SET is underused, and its
11 availability and adherence is low ^{128-130,186-192}. To note, the rate of clinicians referred a
12 patient for SET is very low ¹²⁸. The reasons and barriers for not participating in SET
13 programmes are lack of facilities, feeling worse, costs, time, lack of motivation, and
14 comorbidities ^{128,130,187}.

15 The situation with SET in Europe varies from country to country. A recent European
16 survey showed that supervised exercise programmes exist in Austria, Belgium, Czech
17 Republic, France, Germany, Italy, Sweden, Switzerland, and United Kingdom ¹⁹³.
18 However, SET is reimbursed by the health insurance only in Austria, Belgium, France,
19 Germany, Sweden, and Switzerland ¹⁹³. In the United Kingdom, SET programmes are
20 funded by the National Health Service. In contrast, SET is not reimbursed in Czech
21 Republic, Italy, and it even does not exist for patients with PAD in Denmark, Greece,
22 Ireland, Poland, Serbia, Slovakia, Slovenia, or Ukraine ¹⁹³. Similarly, the structured
23 home-based exercise programme is not routinely present in European countries ¹⁹³.
24 Importantly, there is heterogeneity in form of SET in most of individual countries, with
25 existence of individual programmes or practice of each hospital or community ¹⁹³. They

1 differ in respect of frequency, length and duration of training, type of exercise, as well
2 as by supervising professional ¹⁹³. Mostly, the SET is coordinated by
3 angiologist/vascular physician, but sessions are predominantly supervised by clinical
4 exercise physiologists or physiotherapists. SET for patients with PAD is sometimes
5 offered in cardiac rehabilitation centres. Training programme duration is mostly 12
6 weeks or less, with session duration 30-60 min. Most often used training modalities
7 are combination of walking and resistance training or walking training alone ¹⁹³.
8 To standardise SET programmes and provision across Europe, the following steps are
9 required: 1) a more widespread availability of SET programmes and standardised
10 outcomes to assess their effectiveness; 2) a more defined harmonisation of SET
11 characteristics (establish process of referral, supervision, coordination, selection of
12 patients, SET protocols); 3) health insurance reimbursement of costs; and 4) action to
13 improve the public knowledge about the benefits of SET ¹⁹³.

14

15 **Gaps in evidence and further studies**

16 Awareness and access to supervised exercise programmes should be a field of further
17 studies. Additionally, there are still many areas of insufficient or inconsistent evidence
18 in the treatment of claudication with exercise therapy. We do not know the optimal
19 therapy in terms of duration of the single walking session or intensity of training. We
20 have few studies on the impact of no, or low pain-based exercise and the data on sex
21 differences are inconsistent. The combination of walking exercise with non-walking
22 training has not been yet established. Also, we need more evidence to better
23 understand the potential role of wearable monitoring during exercise interventions, and
24 to evaluate on the efficacy of supportive interventions that can be used together with
25 exercise therapy. For example, the effect of different hydration strategies used during

1 exercise training needs more evidence. In a non-randomised study, Parodi et al.
2 reported mean increase in treadmill walking from 100 meters to 535 meters in 131
3 patients, who were treated with hydration, determined as drinking at least 2000 mL of
4 water during 24 hours for a period of 6 months and to ingest albumin and salt (3.5
5 g/day) ¹⁹⁴.

6 Moreover, data on the interference of exercise training, as well as of individual training
7 modalities, with medical treatment in patients with IC is scarce: one historic RCT
8 suggested an augmentation of the beneficial effect of exercise training by antiplatelet
9 therapy ¹⁹⁵. Another more recent RCT suggested an additive effect of cilostazol on top
10 of exercise treatment on absolute claudication distance ¹⁹⁶. However, it needs to be
11 taken into account that both studies had very small sample sizes. Therefore, larger
12 prospective trials are needed to further elucidate the interaction between exercise
13 training and medication in PAD.

14 Another area of future research should be exploration of the best modalities to
15 transition patients from supervised exercise programmes to everyday life while
16 maintaining the beneficial effects. Finally, we need more research on how to measure
17 success in exercise training in an accurate and reproducible way.

1 **APPOINTED REVIEWERS**

2 Vinko Boc (Department of Vascular Diseases, University Medical Centre Ljubljana,
3 1000 Ljubljana, Slovenia); Tristan Mirault (Université Paris Cité, Inserm, PARCC, F-
4 75015, Paris, France; Service de médecine vasculaire, Hopital Européen G.
5 Pompidou, Paris, France); Frederico Bastos Gonçalves (Department of Angiology and
6 Vascular Surgery, Hospital de Santa Marta, Centro Hospitalar de Lisboa Central,
7 Lisbon, Portugal); Christian-Alexander Behrendt (Department of Vascular Medicine,
8 University Heart and Vascular Centre UKE Hamburg, University Medical Centre
9 Hamburg-Eppendorf, Hamburg, Germany).

1 **References**

- 2 1. Pelliccia A, Sharma S, Gati S, Back M, Borjesson M, Caselli S, et al. 2020 ESC
3 Guidelines on sports cardiology and exercise in patients with cardiovascular
4 disease. *Eur Heart J* 2021;42:17-96. 10.1093/eurheartj/ehaa605
- 5 2. Song P, Rudan D, Zhu Y, Fowkes FJI, Rahimi K, Fowkes FGR, et al. Global,
6 regional, and national prevalence and risk factors for peripheral artery disease in
7 2015: an updated systematic review and analysis. *Lancet Glob Health*
8 2019;7:e1020-e30. 10.1016/S2214-109X(19)30255-4
- 9 3. Aboyans V, Ricco JB, Bartelink MEL, Bjorck M, Brodmann M, Cohnert T, et al.
10 2017 ESC Guidelines on the Diagnosis and Treatment of Peripheral Arterial
11 Diseases, in collaboration with the European Society for Vascular Surgery
12 (ESVS): Document covering atherosclerotic disease of extracranial carotid and
13 vertebral, mesenteric, renal, upper and lower extremity arteries Endorsed by: the
14 European Stroke Organization (ESO) The Task Force for the Diagnosis and
15 Treatment of Peripheral Arterial Diseases of the European Society of Cardiology
16 (ESC) and of the European Society for Vascular Surgery (ESVS). *Eur Heart J*
17 2018;39:763-816. 10.1093/eurheartj/ehx095
- 18 4. Frank U, Nikol S, Belch J, Boc V, Brodmann M, Carpentier PH, et al. ESVM
19 Guideline on peripheral arterial disease. *Vasa* 2019;48:1-79. 10.1024/0301-
20 1526/a000834
- 21 5. McDermott MM. Lower extremity manifestations of peripheral artery disease: the
22 pathophysiologic and functional implications of leg ischemia. *Circ Res*
23 2015;116:1540-50. 10.1161/CIRCRESAHA.114.303517

- 1 6. McDermott MM, Greenland P, Liu K, Guralnik JM, Celic L, Criqui MH, et al. The
2 ankle brachial index is associated with leg function and physical activity: the
3 Walking and Leg Circulation Study. *Ann Intern Med* 2002;136:873-83.
- 4 7. McDermott MM, Liu K, Ferrucci L, Tian L, Guralnik JM, Liao Y, et al. Decline in
5 functional performance predicts later increased mobility loss and mortality in
6 peripheral arterial disease. *J Am Coll Cardiol* 2011;57:962-70.
7 10.1016/j.jacc.2010.09.053
- 8 8. Weitz JI, Byrne J, Clagett GP, Farkouh ME, Porter JM, Sackett DL, et al.
9 Diagnosis and treatment of chronic arterial insufficiency of the lower extremities:
10 a critical review. *Circulation* 1996;94:3026-49. 10.1161/01.cir.94.11.3026
- 11 9. Rose GA. The diagnosis of ischaemic heart pain and intermittent claudication in
12 field surveys. *Bull World Health Organ* 1962;27:645-58.
- 13 10. McDermott MM, Mehta S, Greenland P. Exertional leg symptoms other than
14 intermittent claudication are common in peripheral arterial disease. *Arch Intern*
15 *Med* 1999;159:387-92. 10.1001/archinte.159.4.387
- 16 11. Hiatt WR, Armstrong EJ, Larson CJ, Brass EP. Pathogenesis of the limb
17 manifestations and exercise limitations in peripheral artery disease. *Circ Res*
18 2015;116:1527-39. 10.1161/CIRCRESAHA.116.303566
- 19 12. Seretny M, Colvin LA. Pain management in patients with vascular disease. *Br J*
20 *Anaesth* 2016;117 Suppl 2:ii95-ii106. 10.1093/bja/aew212
- 21 13. Tew GA, Ouedraogo N, Nicolas G, Leftheriotis G, Copeland RJ, Abraham P.
22 Impaired somatosensation in patients with isolated proximal-without-distal
23 exercise-related lower-limb ischemia. *Clin J Pain* 2012;28:404-9.
24 10.1097/AJP.0b013e3182340c01

- 1 14. Hammad TA, Strefling JA, Zellers PR, Reed GW, Venkatachalam S, Lowry AM,
2 et al. The Effect of Post-Exercise Ankle-Brachial Index on Lower Extremity
3 Revascularization. *JACC Cardiovasc Interv* 2015;8:1238-44.
4 10.1016/j.jcin.2015.04.021
- 5 15. McDermott MM, Dayanidhi S, Kosmac K, Saini S, Slysz J, Leeuwenburgh C, et
6 al. Walking Exercise Therapy Effects on Lower Extremity Skeletal Muscle in
7 Peripheral Artery Disease. *Circ Res* 2021;128:1851-67.
8 10.1161/CIRCRESAHA.121.318242
- 9 16. Sheikh MA, Bhatt DL, Li J, Lin S, Bartholomew JR. Usefulness of postexercise
10 ankle-brachial index to predict all-cause mortality. *Am J Cardiol* 2011;107:778-
11 82. 10.1016/j.amjcard.2010.10.060
- 12 17. Flammer AJ, Anderson T, Celermajer DS, Creager MA, Deanfield J, Ganz P, et
13 al. The assessment of endothelial function: from research into clinical practice.
14 *Circulation* 2012;126:753-67. 10.1161/CIRCULATIONAHA.112.093245
- 15 18. Gokce N, Vita JA, Bader DS, Sherman DL, Hunter LM, Holbrook M, et al. Effect
16 of exercise on upper and lower extremity endothelial function in patients with
17 coronary artery disease. *Am J Cardiol* 2002;90:124-7. 10.1016/s0002-
18 9149(02)02433-5
- 19 19. Meredith IT, Currie KE, Anderson TJ, Roddy MA, Ganz P, Creager MA.
20 Postischemic vasodilation in human forearm is dependent on endothelium-
21 derived nitric oxide. *Am J Physiol* 1996;270:H1435-40.
22 10.1152/ajpheart.1996.270.4.H1435
- 23 20. Vita JA, Hamburg NM. Does endothelial dysfunction contribute to the clinical
24 status of patients with peripheral arterial disease? *Can J Cardiol* 2010;26 Suppl
25 A:45A-50A. 10.1016/s0828-282x(10)71062-x

- 1 21. Robbins JL, Jones WS, Duscha BD, Allen JD, Kraus WE, Regensteiner JG, et al.
2 Relationship between leg muscle capillary density and peak hyperemic blood flow
3 with endurance capacity in peripheral artery disease. *J Appl Physiol* (1985)
4 2011;111:81-6. 10.1152/jappphysiol.00141.2011
- 5 22. Beckman JA, Preis O, Ridker PM, Gerhard-Herman M. Comparison of usefulness
6 of inflammatory markers in patients with versus without peripheral arterial disease
7 in predicting adverse cardiovascular outcomes (myocardial infarction, stroke, and
8 death). *Am J Cardiol* 2005;96:1374-8. 10.1016/j.amjcard.2005.07.041
- 9 23. Tzoulaki I, Murray GD, Lee AJ, Rumley A, Lowe GD, Fowkes FG. C-reactive
10 protein, interleukin-6, and soluble adhesion molecules as predictors of
11 progressive peripheral atherosclerosis in the general population: Edinburgh
12 Artery Study. *Circulation* 2005;112:976-83.
13 10.1161/CIRCULATIONAHA.104.513085
- 14 24. Vidula H, Tian L, Liu K, Criqui MH, Ferrucci L, Pearce WH, et al. Biomarkers of
15 inflammation and thrombosis as predictors of near-term mortality in patients with
16 peripheral arterial disease: a cohort study. *Ann Intern Med* 2008;148:85-93.
17 10.7326/0003-4819-148-2-200801150-00003
- 18 25. McDermott MM, Ferrucci L, Liu K, Guralnik JM, Tian L, Liao Y, et al. Leg symptom
19 categories and rates of mobility decline in peripheral arterial disease. *J Am*
20 *Geriatr Soc* 2010;58:1256-62. 10.1111/j.1532-5415.2010.02941.x
- 21 26. Gardner AW, Forrester L, Smith GV. Altered gait profile in subjects with peripheral
22 arterial disease. *Vasc Med* 2001;6:31-4.
- 23 27. Gommans LNM, Smid AT, Scheltinga MRM, Cancrinus E, Brooijmans FAM,
24 Meijer K, et al. Prolonged stance phase during walking in intermittent
25 claudication. *J Vasc Surg* 2017;66:515-22. 10.1016/j.jvs.2017.02.033

- 1 28. McDermott MM, Liu K, Greenland P, Guralnik JM, Criqui MH, Chan C, et al.
2 Functional decline in peripheral arterial disease: associations with the ankle
3 brachial index and leg symptoms. *JAMA* 2004;292:453-61.
4 10.1001/jama.292.4.453
- 5 29. Gardner AW, Montgomery PS, Ritti-Dias RM, Forrester L. The effect of
6 claudication pain on temporal and spatial gait measures during self-paced
7 ambulation. *Vasc Med* 2010;15:21-6. 10.1177/1358863X09106836
- 8 30. Koutakis P, Johanning JM, Haynatzki GR, Myers SA, Stergiou N, Longo GM, et
9 al. Abnormal joint powers before and after the onset of claudication symptoms. *J*
10 *Vasc Surg* 2010;52:340-7. 10.1016/j.jvs.2010.03.005
- 11 31. Koutakis P, Pipinos, II, Myers SA, Stergiou N, Lynch TG, Johanning JM. Joint
12 torques and powers are reduced during ambulation for both limbs in patients with
13 unilateral claudication. *J Vasc Surg* 2010;51:80-8. 10.1016/j.jvs.2009.07.117
- 14 32. Schieber MN, Hasenkamp RM, Pipinos, II, Johanning JM, Stergiou N,
15 DeSpiegelaere HK, et al. Muscle strength and control characteristics are altered
16 by peripheral artery disease. *J Vasc Surg* 2017;66:178-86 e12.
17 10.1016/j.jvs.2017.01.051
- 18 33. Gardner AW, Montgomery PS. Impaired balance and higher prevalence of falls
19 in subjects with intermittent claudication. *J Gerontol A Biol Sci Med Sci*
20 2001;56:M454-8.
- 21 34. Gohil RA, Mockford KA, Mazari F, Khan J, Vanicek N, Chetter IC, et al. Balance
22 impairment, physical ability, and its link with disease severity in patients with
23 intermittent claudication. *Ann Vasc Surg* 2013;27:68-74.
24 10.1016/j.avsg.2012.05.005

- 1 35. Chaudru S, Jehannin P, de Mullenheim PY, Klein H, Jaquinandi V, Mahe G, et
2 al. Using wearable monitors to assess daily walking limitations induced by
3 ischemic pain in peripheral artery disease. *Scand J Med Sci Sports*
4 2019;29:1813-26. 10.1111/sms.13511
- 5 36. McDermott MM, Guralnik JM, Tian L, Liu K, Ferrucci L, Liao Y, et al. Associations
6 of borderline and low normal ankle-brachial index values with functional decline
7 at 5-year follow-up: the WALCS (Walking and Leg Circulation Study). *J Am Coll*
8 *Cardiol* 2009;53:1056-62. 10.1016/j.jacc.2008.09.063
- 9 37. Aboyans V, Criqui MH, Abraham P, Allison MA, Creager MA, Diehm C, et al.
10 Measurement and interpretation of the ankle-brachial index: a scientific statement
11 from the American Heart Association. *Circulation* 2012;126:2890-909.
12 10.1161/CIR.0b013e318276fbc
- 13 38. Treat-Jacobson D, McDermott MM, Beckman JA, Burt MA, Creager MA, Ehrman
14 JK, et al. Implementation of Supervised Exercise Therapy for Patients With
15 Symptomatic Peripheral Artery Disease: A Science Advisory From the American
16 Heart Association. *Circulation* 2019;140:e700-e10.
17 10.1161/CIR.0000000000000727
- 18 39. Gardner AW, Skinner JS, Cantwell BW, Smith LK. Progressive vs single-stage
19 treadmill tests for evaluation of claudication. *Med Sci Sports Exerc* 1991;23:402-
20 8.
- 21 40. Hiatt WR, Hirsch AT, Regensteiner JG, Brass EP. Clinical trials for claudication.
22 Assessment of exercise performance, functional status, and clinical end points.
23 Vascular Clinical Trialists. *Circulation* 1995;92:614-21. 10.1161/01.cir.92.3.614

- 1 41. Hiatt WR, Rogers RK, Brass EP. The treadmill is a better functional test than the
2 6-minute walk test in therapeutic trials of patients with peripheral artery disease.
3 *Circulation* 2014;130:69-78. 10.1161/CIRCULATIONAHA.113.007003
- 4 42. Nicolai SP, Viechtbauer W, Kruidenier LM, Candel MJ, Prins MH, Teijink JA.
5 Reliability of treadmill testing in peripheral arterial disease: a meta-regression
6 analysis. *J Vasc Surg* 2009;50:322-9. 10.1016/j.jvs.2009.01.042
- 7 43. ATS Committee on Proficiency Standards for Clinical Pulmonary Function
8 Laboratories. ATS statement: guidelines for the six-minute walk test. *Am J Respir
9 Crit Care Med* 2002;166:111-7. 10.1164/ajrccm.166.1.at1102
- 10 44. Chandra D, Kulkarni HS, Sciurba F. Learning from the learning effect in the six-
11 minute-walk test. *Am J Respir Crit Care Med* 2012;185:684; author reply -5.
12 10.1164/ajrccm.185.6.684
- 13 45. Tew G, Copeland R, Le Faucheur A, Gernigon M, Nawaz S, Abraham P.
14 Feasibility and validity of self-reported walking capacity in patients with
15 intermittent claudication. *J Vasc Surg* 2013;57:1227-34.
16 10.1016/j.jvs.2012.02.073
- 17 46. McDermott MM, Guralnik JM, Criqui MH, Liu K, Kibbe MR, Ferrucci L. Six-minute
18 walk is a better outcome measure than treadmill walking tests in therapeutic trials
19 of patients with peripheral artery disease. *Circulation* 2014;130:61-8.
20 10.1161/CIRCULATIONAHA.114.007002
- 21 47. McDermott MM, Ades PA, Dyer A, Guralnik JM, Kibbe M, Criqui MH. Corridor-
22 based functional performance measures correlate better with physical activity
23 during daily life than treadmill measures in persons with peripheral arterial
24 disease. *J Vasc Surg* 2008;48:1231-7, 7 e1. 10.1016/j.jvs.2008.06.050

- 1 48. McDermott MM, Guralnik JM, Tian L, Ferrucci L, Liu K, Liao Y, et al. Baseline
2 functional performance predicts the rate of mobility loss in persons with peripheral
3 arterial disease. *J Am Coll Cardiol* 2007;50:974-82. 10.1016/j.jacc.2007.05.030
- 4 49. Sandberg A, Cider A, Jivegard L, Nordanstig J, Wittboldt S, Back M. Test-retest
5 reliability, agreement, and minimal detectable change in the 6-minute walk test in
6 patients with intermittent claudication. *J Vasc Surg* 2020;71:197-203.
7 10.1016/j.jvs.2019.02.056
- 8 50. McDermott MM, Tian L, Criqui MH, Ferrucci L, Conte MS, Zhao L, et al.
9 Meaningful change in 6-minute walk in people with peripheral artery disease. *J*
10 *Vasc Surg* 2021;73:267-76 e1. 10.1016/j.jvs.2020.03.052
- 11 51. Gardner AW, Montgomery PS, Wang M. Minimal clinically important differences
12 in treadmill, 6-minute walk, and patient-based outcomes following supervised and
13 home-based exercise in peripheral artery disease. *Vasc Med* 2018;23:349-57.
14 10.1177/1358863X18762599
- 15 52. de Mullenheim PY, Chaudru S, Mahe G, Prioux J, Le Faucheur A. Clinical Interest
16 of Ambulatory Assessment of Physical Activity and Walking Capacity in
17 Peripheral Artery Disease. *Scand J Med Sci Sports* 2016;26:716-30.
18 10.1111/sms.12512
- 19 53. Abraham P, Noury-Desvaux B, Gernigon M, Mahe G, Sauvaget T, Leftheriotis G,
20 et al. The inter- and intra-unit variability of a low-cost GPS data logger/receiver to
21 study human outdoor walking in view of health and clinical studies. *PLoS One*
22 2012;7:e31338. 10.1371/journal.pone.0031338
- 23 54. Taoum A, Chaudru S, PY DEM, Congnard F, Emily M, Noury-Desvaux B, et al.
24 Comparison of Activity Monitors Accuracy in Assessing Intermittent Outdoor

- 1 Walking. *Med Sci Sports Exerc* 2021;53:1303-14.
2 10.1249/MSS.0000000000002587
- 3 55. Hochsmann C, Knaier R, Eymann J, Hintermann J, Infanger D, Schmidt-
4 Trucksass A. Validity of activity trackers, smartphones, and phone applications
5 to measure steps in various walking conditions. *Scand J Med Sci Sports*
6 2018;28:1818-27. 10.1111/sms.13074
- 7 56. Le Faucheur A, Abraham P, Jaquinandi V, Bouye P, Saumet JL, Noury-Desvaux
8 B. Measurement of walking distance and speed in patients with peripheral arterial
9 disease: a novel method using a global positioning system. *Circulation*
10 2008;117:897-904. 10.1161/CIRCULATIONAHA.107.725994
- 11 57. McDermott MM, Tian L, Ferrucci L, Liu K, Guralnik JM, Liao Y, et al. Associations
12 between lower extremity ischemia, upper and lower extremity strength, and
13 functional impairment with peripheral arterial disease. *J Am Geriatr Soc*
14 2008;56:724-9. 10.1111/j.1532-5415.2008.01633.x
- 15 58. Pizzimenti M, Meyer A, Charles AL, Giannini M, Chakfe N, Lejay A, et al.
16 Sarcopenia and peripheral arterial disease: a systematic review. *J Cachexia*
17 *Sarcopenia Muscle* 2020;11:866-86. 10.1002/jcsm.12587
- 18 59. Ritti-Dias RM, Basyches M, Camara L, Puech-Leao P, Battistella L, Wolosker N.
19 Test-retest reliability of isokinetic strength and endurance tests in patients with
20 intermittent claudication. *Vasc Med* 2010;15:275-8. 10.1177/1358863X10371415
- 21 60. Guralnik JM, Simonsick EM, Ferrucci L, Glynn RJ, Berkman LF, Blazer DG, et al.
22 A short physical performance battery assessing lower extremity function:
23 association with self-reported disability and prediction of mortality and nursing
24 home admission. *J Gerontol* 1994;49:M85-94.

- 1 61. Lanzi S, Pousaz A, Calanca L, Mazzolai L. Sit to Stand Muscle Power Is Related
2 to Functional Performance at Baseline and After Supervised Exercise Training in
3 Patients with Lower Extremity Peripheral Artery Disease. *Eur J Vasc Endovasc*
4 *Surg* 2023;65:521-7. 10.1016/j.ejvs.2022.12.029
- 5 62. Arndt H, Nordanstig J, Bertges DJ, Budtz-Lilly J, Venermo M, Espada CL, et al.
6 A Delphi Consensus on Patient Reported Outcomes for Registries and Trials
7 Including Patients with Intermittent Claudication: Recommendations and
8 Reporting Standard. *Eur J Vasc Endovasc Surg* 2022;64:526-33.
9 10.1016/j.ejvs.2022.08.011
- 10 63. Raja A, Spertus J, Yeh RW, Secemsky EA. Assessing health-related quality of
11 life among patients with peripheral artery disease: A review of the literature and
12 focus on patient-reported outcome measures. *Vasc Med* 2021;26:317-25.
13 10.1177/1358863X20977016
- 14 64. Treat-Jacobson D, McDermott MM, Bronas UG, Campia U, Collins TC, Criqui
15 MH, et al. Optimal Exercise Programs for Patients With Peripheral Artery
16 Disease: A Scientific Statement From the American Heart Association.
17 *Circulation* 2019;139:e10-e33. 10.1161/CIR.0000000000000623
- 18 65. Regensteiner JG, Hiatt WR, Coll JR, Criqui MH, Treat-Jacobson D, McDermott
19 MM, et al. The impact of peripheral arterial disease on health-related quality of
20 life in the Peripheral Arterial Disease Awareness, Risk, and Treatment: New
21 Resources for Survival (PARTNERS) Program. *Vasc Med* 2008;13:15-24.
22 10.1177/1358863X07084911
- 23 66. Kim M, Kim Y, Ryu GW, Choi M. Functional Status and Health-Related Quality of
24 Life in Patients with Peripheral Artery Disease: A Cross-Sectional Study. *Int J*
25 *Environ Res Public Health* 2021;18:10.3390/ijerph182010941

- 1 67. Franco OH, de Laet C, Peeters A, Jonker J, Mackenbach J, Nusselder W. Effects
2 of physical activity on life expectancy with cardiovascular disease. *Arch Intern*
3 *Med* 2005;165:2355-60. 10.1001/archinte.165.20.2355
- 4 68. Dostalova R, Stillman C, Erickson KI, Slepicka P, Mudrak J. The Relationship
5 between Physical Activity, Self-Perceived Health, and Cognitive Function in Older
6 Adults. *Brain Sci* 2021;11:10.3390/brainsci11040492
- 7 69. Frank U, Nikol S, Belch J. 5 Conservative treatment for PAD - Risk factor
8 management. *Vasa* 2019;48:1-12. 10.1024/0301-1526/a000835
- 9 70. Harwood A, Pymer S, Ingle L, Doherty P, Chetter I, Parmenter B, et al. Exercise
10 training for intermittent claudication: a narrative review and summary of
11 guidelines for practitioners. *BMJ Open Sport & Exercise Medicine*
12 2020;6:e000897. doi:10.1136/bmjsem-2020-000897
- 13 71. Gommans LN, Fokkenrood HJ, van Dalen HC, Scheltinga MR, Teijink JA, Peters
14 RJ. Safety of supervised exercise therapy in patients with intermittent
15 claudication. *J Vasc Surg* 2015;61:512-8 e2. 10.1016/j.jvs.2014.08.070
- 16 72. Bronas UG, Regensteiner JG. Connecting the past to the present: A historical
17 review of exercise training for peripheral artery disease. *Vasc Med* 2022;27:174-
18 85. 10.1177/1358863X211073620
- 19 73. Penin-Grandes S, Lopez-Ortiz S, Maroto-Izquierdo S, Menendez H, Pinto-Fraga
20 J, Martin-Hernandez J, et al. Winners do what they fear: exercise and peripheral
21 arterial disease-an umbrella review. *Eur J Prev Cardiol*
22 2023;10.1093/eurjpc/zwad261
- 23 74. Lane R, Harwood A, Watson L, Leng GC. Exercise for intermittent claudication.
24 *Cochrane Database Syst Rev* 2017;12:CD000990.
25 10.1002/14651858.CD000990.pub4

- 1 75. Fakhry F, van de Luijngaarden KM, Bax L, den Hoed PT, Hunink MG, Rouwet EV,
2 et al. Supervised walking therapy in patients with intermittent claudication. *J Vasc*
3 *Surg* 2012;56:1132-42. 10.1016/j.jvs.2012.04.046
- 4 76. Lanzi S, Boichat J, Calanca L, Aubertin P, Malatesta D, Mazzolai L. Gait changes
5 after supervised exercise training in patients with symptomatic lower extremity
6 peripheral artery disease. *Vasc Med* 2021;26:259-66.
7 10.1177/1358863X20984831
- 8 77. Lanzi S, Boichat J, Calanca L, Mazzolai L, Malatesta D. Supervised Exercise
9 Training Improves 6 min Walking Distance and Modifies Gait Pattern during Pain-
10 Free Walking Condition in Patients with Symptomatic Lower Extremity Peripheral
11 Artery Disease. *Sensors (Basel)* 2021;21:7989. 10.3390/s21237989
- 12 78. Fassora M, Calanca L, Jaques C, Mazzolai L, Kayser B, Lanzi S. Intensity-
13 dependent effects of exercise therapy on walking performance and aerobic
14 fitness in symptomatic patients with lower-extremity peripheral artery disease: A
15 systematic review and meta-analysis. *Vasc Med* 2022;27:158-70.
16 10.1177/1358863X211034577
- 17 79. Lanzi S, Pousaz A, Calanca L, Mazzolai L. Time-course evolution of functional
18 performance during a 3-month supervised exercise training program in patients
19 with symptomatic peripheral artery disease. *Vasc Med*
20 2023;1358863X231191908. 10.1177/1358863X231191908
- 21 80. Parmenter BJ, Dieberg G, Phipps G, Smart NA. Exercise training for health-
22 related quality of life in peripheral artery disease: a systematic review and meta-
23 analysis. *Vasc Med* 2015;20:30-40. 10.1177/1358863X14559092

- 1 81. Parmenter BJ, Dieberg G, Smart NA. Exercise training for management of
2 peripheral arterial disease: a systematic review and meta-analysis. *Sports Med*
3 2015;45:231-44. 10.1007/s40279-014-0261-z
- 4 82. Schieber MN, Pipinos, II, Johanning JM, Casale GP, Williams MA,
5 DeSpiegelaere HK, et al. Supervised walking exercise therapy improves gait
6 biomechanics in patients with peripheral artery disease. *J Vasc Surg*
7 2019;71:575-83. 10.1016/j.jvs.2019.05.044
- 8 83. Siercke M, Jorgensen LP, Missel M, Thygesen LC, Moller SP, Sillesen H, et al.
9 Cardiovascular Rehabilitation Increases Walking Distance in Patients With
10 Intermittent Claudication. Results of the CIPIC Rehab Study: A Randomised
11 Controlled Trial. *Eur J Vasc Endovasc Surg* 2021;62:768-76.
12 10.1016/j.ejvs.2021.04.004
- 13 84. Brown T, Forster RB, Cleanthis M, Mikhailidis DP, Stansby G, Stewart M.
14 Cilostazol for intermittent claudication. *Cochrane Database Syst Rev*
15 2021;6:CD003748. 10.1002/14651858.CD003748.pub5
- 16 85. De Haro J, Bleda S, Varela C, Esparza L, Acin F, Bosentan Population-Based
17 Randomized Trial for C, et al. Effect of Bosentan on Claudication Distance and
18 Endothelium-Dependent Vasodilation in Hispanic Patients With Peripheral
19 Arterial Disease. *Am J Cardiol* 2016;117:295-301.
20 10.1016/j.amjcard.2015.10.032
- 21 86. Omarjee L, Le Pabic E, Custaud MA, Fontaine C, Locher C, Renault A, et al.
22 Effects of sildenafil on maximum walking time in patients with arterial claudication:
23 The ARTERIOFIL study. *Vascul Pharmacol* 2019;118-119:106563.
24 10.1016/j.vph.2019.05.003

- 1 87. Suzuki J, Shimamura M, Suda H, Wakayama K, Kumagai H, Ikeda Y, et al.
2 Current therapies and investigational drugs for peripheral arterial disease.
3 *Hypertens Res* 2016;39:183-91. 10.1038/hr.2015.134
- 4 88. Broderick C, Forster R, Abdel-Hadi M, Salhiyyah K. Pentoxifylline for intermittent
5 claudication. *Cochrane Database Syst Rev* 2020;10:CD005262.
6 10.1002/14651858.CD005262.pub4
- 7 89. Jansen SC, Abaraogu UO, Lauret GJ, Fakhry F, Fokkenrood HJ, Teijink JA.
8 Modes of exercise training for intermittent claudication. *Cochrane Database Syst*
9 *Rev* 2020;8:CD009638. 10.1002/14651858.CD009638.pub3
- 10 90. Parmenter BJ, Raymond J, Dinnen P, Lusby RJ, Fiatarone Singh MA. High-
11 intensity progressive resistance training improves flat-ground walking in older
12 adults with symptomatic peripheral arterial disease. *J Am Geriatr Soc*
13 2013;61:1964-70. 10.1111/jgs.12500
- 14 91. Ritti-Dias RM, Wolosker N, de Moraes Forjaz CL, Carvalho CR, Cucato GG, Leao
15 PP, et al. Strength training increases walking tolerance in intermittent claudication
16 patients: randomized trial. *J Vasc Surg* 2010;51:89-95. 10.1016/j.jvs.2009.07.118
- 17 92. Sanderson B, Askew C, Stewart I, Walker P, Gibbs H, Green S. Short-term
18 effects of cycle and treadmill training on exercise tolerance in peripheral arterial
19 disease. *J Vasc Surg* 2006;44:119-27. 10.1016/j.jvs.2006.03.037
- 20 93. Collins EG, Edwin Langbein W, Orebaugh C, Bammert C, Hanson K, Reda D, et
21 al. PoleStriding exercise and vitamin E for management of peripheral vascular
22 disease. *Med Sci Sports Exerc* 2003;35:384-93.
23 10.1249/01.MSS.0000053658.82687.FF

- 1 94. Collins EG, Langbein WE, Orebaugh C, Bammert C, Hanson K, Reda D, et al.
2 Cardiovascular training effect associated with polestriding exercise in patients
3 with peripheral arterial disease. *J Cardiovasc Nurs* 2005;20:177-85.
- 4 95. Calanca L, Lanzi S, Ney B, Berchtold A, Mazzolai L. Multimodal Supervised
5 Exercise Significantly Improves Walking Performances Without Changing
6 Hemodynamic Parameters in Patients With Symptomatic Lower Extremity
7 Peripheral Artery Disease. *Vasc Endovascular Surg* 2020;54:605-11.
8 10.1177/1538574420940090
- 9 96. Lanzi S, Calanca L, Berchtold A, Mazzolai L. Improvement in 6-Minute Walking
10 Distance after Supervised Exercise Training Is Related to Changes in Quality of
11 Life in Patients with Lower Extremity Peripheral Artery Disease. *J Clin Med*
12 2021;10:3330. 10.3390/jcm10153330
- 13 97. Lanzi S, Calanca L, Borgeat Kaeser A, Mazzolai L. Walking performances and
14 muscle oxygen desaturation are increased after supervised exercise training in
15 Takayasu arteritis: a case report and a review of the literature. *Eur Heart J Case*
16 *Rep* 2018;2:yty123. 10.1093/ehjcr/yty123
- 17 98. Ney B, Lanzi S, Calanca L, Mazzolai L. Multimodal Supervised Exercise Training
18 Is Effective in Improving Long Term Walking Performance in Patients with
19 Symptomatic Lower Extremity Peripheral Artery Disease. *J Clin Med*
20 2021;10:2057. 10.3390/jcm10102057
- 21 99. Tew G, Nawaz S, Zwierska I, Saxton JM. Limb-specific and cross-transfer effects
22 of arm-crank exercise training in patients with symptomatic peripheral arterial
23 disease. *Clin Sci (Lond)* 2009;117:405-13. 10.1042/CS20080688

- 1 100. Treat-Jacobson D, Bronas UG, Leon AS. Efficacy of arm-ergometry versus
2 treadmill exercise training to improve walking distance in patients with
3 claudication. *Vasc Med* 2009;14:203-13. 10.1177/1358863X08101858
- 4 101. Parmenter BJ, Raymond J, Dinnen P, Singh MA. A systematic review of
5 randomized controlled trials: Walking versus alternative exercise prescription as
6 treatment for intermittent claudication. *Atherosclerosis* 2011;218:1-12.
7 10.1016/j.atherosclerosis.2011.04.024
- 8 102. McDermott MM, Ades P, Guralnik JM, Dyer A, Ferrucci L, Liu K, et al. Treadmill
9 exercise and resistance training in patients with peripheral arterial disease with
10 and without intermittent claudication: a randomized controlled trial. *JAMA*
11 2009;301:165-74. 10.1001/jama.2008.962
- 12 103. Bulmer AC, Coombes JS. Optimising exercise training in peripheral arterial
13 disease. *Sports Med* 2004;34:983-1003. 10.2165/00007256-200434140-00004
- 14 104. Gardner AW, Poehlman ET. Exercise rehabilitation programs for the treatment of
15 claudication pain. A meta-analysis. *JAMA* 1995;274:975-80.
- 16 105. Borg GA. Psychophysical bases of perceived exertion. *Med Sci Sports Exerc*
17 1982;14:377-81.
- 18 106. Au TB, Golledge J, Walker PJ, Haigh K, Nelson M. Peripheral arterial disease -
19 diagnosis and management in general practice. *Aust Fam Physician*
20 2013;42:397-400.
- 21 107. Hammond MM, Spring B, Rejeski WJ, Sufit R, Criqui MH, Tian L, et al. Effects of
22 Walking Exercise at a Pace With Versus Without Ischemic Leg Symptoms on
23 Functional Performance Measures in People With Lower Extremity Peripheral
24 Artery Disease: The LITE Randomized Clinical Trial. *J Am Heart Assoc*
25 2022;11:e025063. 10.1161/JAHA.121.025063

- 1 108. McDermott MM, Spring B, Tian L, Treat-Jacobson D, Ferrucci L, Lloyd-Jones D,
2 et al. Effect of Low-Intensity vs High-Intensity Home-Based Walking Exercise on
3 Walk Distance in Patients With Peripheral Artery Disease: The LITE Randomized
4 Clinical Trial. *JAMA* 2021;325:1266-76. 10.1001/jama.2021.2536
- 5 109. Mika P, Konik A, Januszek R, Petriczek T, Mika A, Nowobilski R, et al.
6 Comparison of two treadmill training programs on walking ability and endothelial
7 function in intermittent claudication. *Int J Cardiol* 2013;168:838-42.
8 10.1016/j.ijcard.2012.10.003
- 9 110. Novakovic M, Krevel B, Rajkovic U, Vizintin Cuderman T, Jansa Trontelj K, Fras
10 Z, et al. Moderate-pain versus pain-free exercise, walking capacity, and
11 cardiovascular health in patients with peripheral artery disease. *J Vasc Surg*
12 2019;70:148-56. 10.1016/j.jvs.2018.10.109
- 13 111. Perks J, Zaccardi F, Paterson C, Houghton JSM, Nickinson ATO, Pepper CJ, et
14 al. Effect of high-pain versus low-pain structured exercise on walking ability in
15 people with intermittent claudication: meta-analysis. *Br J Surg* 2022;109:686-94.
16 10.1093/bjs/znac134
- 17 112. Seed SA, Harwood AE, Sinclair J, Pymer S, Caldow E, Ingle L, et al. A Systematic
18 Review of Exercise Prescription in Patients with Intermittent Claudication: Does
19 Pain Matter? *Ann Vasc Surg* 2021;77:315-23. 10.1016/j.avsg.2021.06.025
- 20 113. Birkett ST, Sinclair J, Seed SA, Pymer S, Caldow E, Ingle L, et al. Effects of
21 exercise prescribed at different levels of claudication pain on walking
22 performance in patients with intermittent claudication: a protocol for a randomised
23 controlled trial. *Ther Adv Cardiovasc Dis* 2022;16:17539447221108817.
24 10.1177/17539447221108817

- 1 114. Lanzi S, Mazzolai L. Commentary to Seed et al. 'What is the correct level of
2 claudication pain to prescribe? Universal inconsistency within guidelines, a
3 painful issue'. *Vascular* 2023;17085381231160931.
4 10.1177/17085381231160931
- 5 115. Garber CE, Blissmer B, Deschenes MR, Franklin BA, Lamonte MJ, Lee IM, et al.
6 American College of Sports Medicine position stand. Quantity and quality of
7 exercise for developing and maintaining cardiorespiratory, musculoskeletal, and
8 neuromotor fitness in apparently healthy adults: guidance for prescribing
9 exercise. *Med Sci Sports Exerc* 2011;43:1334-59.
10 10.1249/MSS.0b013e318213fefb
- 11 116. Hansen D, Abreu A, Ambrosetti M, Cornelissen V, Gevaert A, Kemps H, et al.
12 Exercise intensity assessment and prescription in cardiovascular rehabilitation
13 and beyond: why and how: a position statement from the Secondary Prevention
14 and Rehabilitation Section of the European Association of Preventive Cardiology.
15 *Eur J Prev Cardiol* 2022;29:230-45. 10.1093/eurjpc/zwab007
- 16 117. Wood TM, Maddalozzo GF, Harter RA. Accuracy of Seven Equations for
17 Predicting 1-RM Performance of Apparently Healthy, Sedentary Older Adults.
18 *Measurement in Physical Education and Exercise Science* 2002;6:67-94.
19 10.1207/S15327841MPEE0602_1
- 20 118. Parmenter BJ, Mavros Y, Ritti Dias R, King S, Fiatarone Singh M. Resistance
21 training as a treatment for older persons with peripheral artery disease: a
22 systematic review and meta-analysis. *Br J Sports Med* 2019;54:452-61.
23 10.1136/bjsports-2018-100205
- 24 119. Pymer S, Ibeggazene S, Palmer J, Tew GA, Ingle L, Smith GE, et al. An updated
25 systematic review and meta-analysis of home-based exercise programs for

- 1 individuals with intermittent claudication. *J Vasc Surg* 2021;74:2076-85 e20.
2 10.1016/j.jvs.2021.03.063
- 3 120. van den Houten MML, Hageman D, Gommans LNM, Kleijnen J, Scheltinga MRM,
4 Teijink JAW. The Effect of Supervised Exercise, Home Based Exercise and
5 Endovascular Revascularisation on Physical Activity in Patients With Intermittent
6 Claudication: A Network Meta-analysis. *Eur J Vasc Endovasc Surg* 2019;58:383-
7 92. 10.1016/j.ejvs.2018.12.023
- 8 121. Rejeski WJ, Spring B, Domanchuk K, Tao H, Tian L, Zhao L, et al. A group-
9 mediated, home-based physical activity intervention for patients with peripheral
10 artery disease: effects on social and psychological function. *J Transl Med*
11 2014;12:29. 10.1186/1479-5876-12-29
- 12 122. Fakhry F, Spronk S, de Ridder M, den Hoed PT, Hunink MG. Long-term effects
13 of structured home-based exercise program on functional capacity and quality of
14 life in patients with intermittent claudication. *Arch Phys Med Rehabil*
15 2011;92:1066-73. 10.1016/j.apmr.2011.02.007
- 16 123. McDermott MM, Guralnik JM, Criqui MH, Ferrucci L, Zhao L, Liu K, et al. Home-
17 based walking exercise in peripheral artery disease: 12-month follow-up of the
18 GOALS randomized trial. *J Am Heart Assoc* 2014;3:e000711.
19 10.1161/JAHA.113.000711
- 20 124. Waddell A, Seed S, Broom DR, McGregor G, Birkett ST, Harwood AE. Safety of
21 home-based exercise for people with intermittent claudication: A systematic
22 review. *Vasc Med* 2022;27:186-92. 10.1177/1358863X211060388
- 23 125. Lamberti N, Lopez-Soto PJ, Guerzoni F, Napoli N, Gasbarro V, Zamboni P, et al.
24 Changes in exercise capacity and risk of all-cause mortality in patients with

- 1 peripheral artery disease: a 10-year retrospective cohort study. *Intern Emerg Med*
2 2020;15:289-98. 10.1007/s11739-019-02176-3
- 3 126. Hageman D, Fokkenrood HJ, Gommans LN, van den Houten MM, Teijink JA.
4 Supervised exercise therapy versus home-based exercise therapy versus
5 walking advice for intermittent claudication. *Cochrane Database Syst Rev*
6 2018;4:CD005263. 10.1002/14651858.CD005263.pub4
- 7 127. Al-Jundi W, Madbak K, Beard JD, Nawaz S, Tew GA. Systematic review of home-
8 based exercise programmes for individuals with intermittent claudication. *Eur J*
9 *Vasc Endovasc Surg* 2013;46:690-706. 10.1016/j.ejvs.2013.09.004
- 10 128. Dua A, Gologorsky R, Savage D, Rens N, Gandhi N, Brooke B, et al. National
11 assessment of availability, awareness, and utilization of supervised exercise
12 therapy for peripheral artery disease patients with intermittent claudication. *J*
13 *Vasc Surg* 2020;71:1702-7. 10.1016/j.jvs.2019.08.238
- 14 129. Haque A. Few UK vascular centres offer a fully NICE-compliant supervised
15 exercise programme: a national audit. *Ann R Coll Surg Engl*
16 2021;10.1308/rcsann.2021.0126
- 17 130. Harwood AE, Pymer S, Ibeggazene S, Ingle L, Caldow E, Birkett ST. Provision
18 of exercise services in patients with peripheral artery disease in the United
19 Kingdom. *Vascular* 2021;17085381211035259. 10.1177/17085381211035259
- 20 131. Makris GC, Lattimer CR, Lavidia A, Geroulakos G. Availability of supervised
21 exercise programs and the role of structured home-based exercise in peripheral
22 arterial disease. *Eur J Vasc Endovasc Surg* 2012;44:569-75; discussion 76.
23 10.1016/j.ejvs.2012.09.009

- 1 132. Gardner AW, Parker DE, Montgomery PS, Blevins SM. Diabetic women are poor
2 responders to exercise rehabilitation in the treatment of claudication. *J Vasc Surg*
3 2014;59:1036-43. 10.1016/j.jvs.2013.10.058
- 4 133. Manfredini R, Lamberti N, Manfredini F, Straudi S, Fabbian F, Rodriguez Borrego
5 MA, et al. Gender Differences in Outcomes Following a Pain-Free, Home-Based
6 Exercise Program for Claudication. *J Womens Health (Larchmt)* 2019;28:1313-
7 21. 10.1089/jwh.2018.7113
- 8 134. Gardner AW, Parker DE, Montgomery PS. Sex-specific predictors of improved
9 walking with step-monitored, home-based exercise in peripheral artery disease.
10 *Vasc Med* 2015;20:424-31. 10.1177/1358863X15596237
- 11 135. Collins TC, Lunos S, Carlson T, Henderson K, Lightbourne M, Nelson B, et al.
12 Effects of a home-based walking intervention on mobility and quality of life in
13 people with diabetes and peripheral arterial disease: a randomized controlled
14 trial. *Diabetes Care* 2011;34:2174-9. 10.2337/dc10-2399
- 15 136. Lamberti N, Malagoni AM, Ficarra V, Basaglia N, Manfredini R, Zamboni P, et al.
16 Structured Home-Based Exercise Versus Invasive Treatment: A Mission
17 Impossible? A Pilot Randomized Study in Elderly Patients With Intermittent
18 Claudication. *Angiology* 2016;67:772-80. 10.1177/0003319715618481
- 19 137. McDermott MM, Polonsky TS. Home-Based Exercise: A Therapeutic Option for
20 Peripheral Artery Disease. *Circulation* 2016;134:1127-9.
21 10.1161/CIRCULATIONAHA.116.023691
- 22 138. Chan C, Sounderajah V, Normahani P, Acharya A, Markar SR, Darzi A, et al.
23 Wearable Activity Monitors in Home Based Exercise Therapy for Patients with
24 Intermittent Claudication: A Systematic Review. *Eur J Vasc Endovasc Surg*
25 2021;61:676-87. 10.1016/j.ejvs.2020.11.044

- 1 139. Kim M, Kim C, Kim E, Choi M. Effectiveness of Mobile Health-Based Exercise
2 Interventions for Patients with Peripheral Artery Disease: Systematic Review and
3 Meta-Analysis. *JMIR Mhealth Uhealth* 2021;9:e24080. 10.2196/24080
- 4 140. Hawley JA, Hargreaves M, Joyner MJ, Zierath JR. Integrative biology of exercise.
5 *Cell* 2014;159:738-49. 10.1016/j.cell.2014.10.029
- 6 141. Hoier B, Hellsten Y. Exercise-induced capillary growth in human skeletal muscle
7 and the dynamics of VEGF. *Microcirculation* 2014;21:301-14.
8 10.1111/micc.12117
- 9 142. Egginton S. Invited review: activity-induced angiogenesis. *Pflugers Arch*
10 2009;457:963-77. 10.1007/s00424-008-0563-9
- 11 143. Handschin C, Spiegelman BM. The role of exercise and PGC1alpha in
12 inflammation and chronic disease. *Nature* 2008;454:463-9. 10.1038/nature07206
- 13 144. Gerhard-Herman MD, Gornik HL, Barrett C, Barshes NR, Corriere MA, Drachman
14 DE, et al. 2016 AHA/ACC Guideline on the Management of Patients With Lower
15 Extremity Peripheral Artery Disease: Executive Summary: A Report of the
16 American College of Cardiology/American Heart Association Task Force on
17 Clinical Practice Guidelines. *J Am Coll Cardiol* 2017;69:1465-508.
18 10.1016/j.jacc.2016.11.008
- 19 145. Heiss C, Olinic DM, Belch JJF, Brodmann M, Mazzolai L, Stanek A, et al.
20 Management of chronic peripheral artery disease patients with indication for
21 endovascular revascularization. *Vasa* 2022;10.1024/0301-1526/a000998
- 22 146. Anderson JL, Halperin JL, Albert NM, Bozkurt B, Brindis RG, Curtis LH, et al.
23 Management of patients with peripheral artery disease (compilation of 2005 and
24 2011 ACCF/AHA guideline recommendations): a report of the American College

- 1 of Cardiology Foundation/American Heart Association Task Force on Practice
2 Guidelines. *Circulation* 2013;127:1425-43. 10.1161/CIR.0b013e31828b82aa
- 3 147. Koelemay MJW, van Reijen NS, van Dieren S, Frans FA, Vermeulen EJJ,
4 Buscher H, et al. Editor's Choice - Randomised Clinical Trial of Supervised
5 Exercise Therapy vs. Endovascular Revascularisation for Intermittent
6 Claudication Caused by Iliac Artery Obstruction: The SUPER study. *Eur J Vasc*
7 *Endovasc Surg* 2022;63:421-9. 10.1016/j.ejvs.2021.09.042
- 8 148. Murphy TP, Cutlip DE, Regensteiner JG, Mohler ER, Cohen DJ, Reynolds MR,
9 et al. Supervised exercise versus primary stenting for claudication resulting from
10 aortoiliac peripheral artery disease: six-month outcomes from the claudication:
11 exercise versus endoluminal revascularization (CLEVER) study. *Circulation*
12 2012;125:130-9. 10.1161/CIRCULATIONAHA.111.075770
- 13 149. Murphy TP, Cutlip DE, Regensteiner JG, Mohler ER, 3rd, Cohen DJ, Reynolds
14 MR, et al. Supervised exercise, stent revascularization, or medical therapy for
15 claudication due to aortoiliac peripheral artery disease: the CLEVER study. *J Am*
16 *Coll Cardiol* 2015;65:999-1009. 10.1016/j.jacc.2014.12.043
- 17 150. Koppe-Schmeisser F, Schwaderlapp M, Schmeisser J, Dopheide JF, Munzel T,
18 Daiber A, et al. Influence of Peripheral Transluminal Angioplasty Alongside
19 Exercise Training on Oxidative Stress and Inflammation in Patients with
20 Peripheral Arterial Disease. *J Clin Med* 2021;10:10.3390/jcm10245851
- 21 151. Pandey A, Banerjee S, Ngo C, Mody P, Marso SP, Brilakis ES, et al. Comparative
22 Efficacy of Endovascular Revascularization Versus Supervised Exercise Training
23 in Patients With Intermittent Claudication: Meta-Analysis of Randomized
24 Controlled Trials. *JACC Cardiovasc Interv* 2017;10:712-24.
25 10.1016/j.jcin.2017.01.027

- 1 152. Fakhry F, Fokkenrood HJ, Spronk S, Teijink JA, Rouwet EV, Hunink MGM.
2 Endovascular revascularisation versus conservative management for intermittent
3 claudication. *Cochrane Database Syst Rev* 2018;3:CD010512.
4 10.1002/14651858.CD010512.pub2
- 5 153. Badger SA, Soong CV, O'Donnell ME, Boreham CA, McGuigan KE. Benefits of
6 a supervised exercise program after lower limb bypass surgery. *Vasc*
7 *Endovascular Surg* 2007;41:27-32. 10.1177/1538574406296209
- 8 154. Kobayashi T, Hamamoto M, Okazaki T, Honma T, Iba K, Takakuwa T, et al.
9 Effectiveness of continuous unsupervised exercise therapy after above-knee
10 femoropopliteal bypass. *Vascular* 2021;29:387-95. 10.1177/1708538120957488
- 11 155. Meneses AL, Ritti-Dias RM, Parmenter B, Golledge J, Askew CD. Combined
12 Lower Limb Revascularisation and Supervised Exercise Training for Patients with
13 Peripheral Arterial Disease: A Systematic Review of Randomised Controlled
14 Trials. *Sports Med* 2017;47:987-1002. 10.1007/s40279-016-0635-5
- 15 156. Issa SM, Hoeks SE, Scholte op Reimer WJ, Van Gestel YR, Lenzen MJ,
16 Verhagen HJ, et al. Health-related quality of life predicts long-term survival in
17 patients with peripheral artery disease. *Vasc Med* 2010;15:163-9.
18 10.1177/1358863X10364208
- 19 157. Gardner AW, Parker DE, Montgomery PS, Scott KJ, Blevins SM. Efficacy of
20 quantified home-based exercise and supervised exercise in patients with
21 intermittent claudication: a randomized controlled trial. *Circulation* 2011;123:491-
22 8. 10.1161/CIRCULATIONAHA.110.963066
- 23 158. Patterson RB, Pinto B, Marcus B, Colucci A, Braun T, Roberts M. Value of a
24 supervised exercise program for the therapy of arterial claudication. *J Vasc Surg*
25 1997;25:312-8. 10.1016/s0741-5214(97)70352-5

- 1 159. Tsai JC, Chan P, Wang CH, Jeng C, Hsieh MH, Kao PF, et al. The effects of
2 exercise training on walking function and perception of health status in elderly
3 patients with peripheral arterial occlusive disease. *J Intern Med* 2002;252:448-
4 55. 10.1046/j.1365-2796.2002.01055.x
- 5 160. Nicolai SP, Teijink JA, Prins MH, Exercise Therapy in Peripheral Arterial Disease
6 Study G. Multicenter randomized clinical trial of supervised exercise therapy with
7 or without feedback versus walking advice for intermittent claudication. *J Vasc*
8 *Surg* 2010;52:348-55. 10.1016/j.jvs.2010.02.022
- 9 161. Guidon M, McGee H. Exercise-based interventions and health-related quality of
10 life in intermittent claudication: a 20-year (1989-2008) review. *Eur J Cardiovasc*
11 *Prev Rehabil* 2010;17:140-54. 10.1097/HJR.0b013e3283377f08
- 12 162. Guidon M, McGee H. One-year effect of a supervised exercise programme on
13 functional capacity and quality of life in peripheral arterial disease. *Disabil Rehabil*
14 2013;35:397-404. 10.3109/09638288.2012.694963
- 15 163. Kakkos SK, Geroulakos G, Nicolaides AN. Improvement of the walking ability in
16 intermittent claudication due to superficial femoral artery occlusion with
17 supervised exercise and pneumatic foot and calf compression: a randomised
18 controlled trial. *Eur J Vasc Endovasc Surg* 2005;30:164-75.
19 10.1016/j.ejvs.2005.03.011
- 20 164. Savage P, Ricci MA, Lynn M, Gardner A, Knight S, Brochu M, et al. Effects of
21 home versus supervised exercise for patients with intermittent claudication. *J*
22 *Cardiopulm Rehabil* 2001;21:152-7. 10.1097/00008483-200105000-00006
- 23 165. Gardner AW, Montgomery PS, Wang M, Xu C. Predictors of health-related quality
24 of life in patients with symptomatic peripheral artery disease. *J Vasc Surg*
25 2018;68:1126-34. 10.1016/j.jvs.2017.12.074

- 1 166. Erickson KI, Hillman C, Stillman CM, Ballard RM, Bloodgood B, Conroy DE, et al.
2 Physical Activity, Cognition, and Brain Outcomes: A Review of the 2018 Physical
3 Activity Guidelines. *Med Sci Sports Exerc* 2019;51:1242-51.
4 10.1249/MSS.0000000000001936
- 5 167. Kirk-Sanchez NJ, McGough EL. Physical exercise and cognitive performance in
6 the elderly: current perspectives. *Clin Interv Aging* 2014;9:51-62.
7 10.2147/CIA.S39506
- 8 168. Gomez-Pinilla F, Hillman C. The influence of exercise on cognitive abilities.
9 *Compr Physiol* 2013;3:403-28. 10.1002/cphy.c110063
- 10 169. Gommans LN, Saarloos R, Scheltinga MR, Houterman S, de Bie RA, Fokkenrood
11 HJ, et al. Editor's choice--The effect of supervision on walking distance in patients
12 with intermittent claudication: a meta-analysis. *Eur J Vasc Endovasc Surg*
13 2014;48:169-84. 10.1016/j.ejvs.2014.04.019
- 14 170. Abaraogu UO, Dall PM, Seenan CA. The Effect of Structured Patient Education
15 on Physical Activity in Patients with Peripheral Arterial Disease and Intermittent
16 Claudication: A Systematic Review. *Eur J Vasc Endovasc Surg* 2017;54:58-68.
17 10.1016/j.ejvs.2017.04.003
- 18 171. Bearne LM, Volkmer B, Peacock J, Sekhon M, Fisher G, Galea Holmes MN, et
19 al. Effect of a Home-Based, Walking Exercise Behavior Change Intervention vs
20 Usual Care on Walking in Adults With Peripheral Artery Disease: The MOSAIC
21 Randomized Clinical Trial. *JAMA* 2022;327:1344-55. 10.1001/jama.2022.3391
- 22 172. McDermott MM, Liu K, Guralnik JM, Criqui MH, Spring B, Tian L, et al. Home-
23 based walking exercise intervention in peripheral artery disease: a randomized
24 clinical trial. *JAMA* 2013;310:57-65. 10.1001/jama.2013.7231

- 1 173. McDermott MM, Spring B, Berger JS, Treat-Jacobson D, Conte MS, Creager MA,
2 et al. Effect of a Home-Based Exercise Intervention of Wearable Technology and
3 Telephone Coaching on Walking Performance in Peripheral Artery Disease: The
4 HONOR Randomized Clinical Trial. *JAMA* 2018;319:1665-76.
5 10.1001/jama.2018.3275
- 6 174. Behrendt CA, Thomalla G, Rimmele DL, Petersen EL, Twerenbold R, Debus ES,
7 et al. Prevalence of peripheral arterial disease, abdominal aortic aneurysm, and
8 risk factors in the Hamburg City Health Study: A Cross-Sectional Analysis. *Eur J*
9 *Vasc Endovasc Surg* 2023;10.1016/j.ejvs.2023.01.002
- 10 175. Pabon M, Cheng S, Altin SE, Sethi SS, Nelson MD, Moreau KL, et al. Sex
11 Differences in Peripheral Artery Disease. *Circ Res* 2022;130:496-511.
12 10.1161/CIRCRESAHA.121.320702
- 13 176. Behrendt CA, Sigvant B, Kuchenbecker J, Grima MJ, Schermerhorn M, Thomson
14 IA, et al. Editor's Choice - International Variations and Sex Disparities in the
15 Treatment of Peripheral Arterial Occlusive Disease: A Report from VASCUNET
16 and the International Consortium of Vascular Registries. *Eur J Vasc Endovasc*
17 *Surg* 2020;60:873-80. 10.1016/j.ejvs.2020.08.027
- 18 177. Detriche G, Guedon A, Mohamedi N, Sellami O, Cheng C, Galloula A, et al.
19 Women Specific Characteristics and 1-Year Outcome Among Patients
20 Hospitalized for Peripheral Artery Disease: A Monocentric Cohort Analysis in a
21 Tertiary Center. *Front Cardiovasc Med* 2022;9:824466.
22 10.3389/fcvm.2022.824466
- 23 178. Heidemann F, Kuchenbecker J, Peters F, Kotov A, Marschall U, L'Hoest H, et al.
24 A health insurance claims analysis on the effect of female sex on long-term
25 outcomes after peripheral endovascular interventions for symptomatic peripheral

- 1 arterial occlusive disease. *J Vasc Surg* 2021;74:780-7 e7.
2 10.1016/j.jvs.2021.01.066
- 3 179. Hirsch AT, Allison MA, Gomes AS, Corriere MA, Duval S, Ershow AG, et al. A
4 call to action: women and peripheral artery disease: a scientific statement from
5 the American Heart Association. *Circulation* 2012;125:1449-72.
6 10.1161/CIR.0b013e31824c39ba
- 7 180. Kotov A, Heidemann F, Kuchenbecker J, Peters F, Marschall U, Acar L, et al. Sex
8 Disparities in Long Term Outcomes After Open Surgery for Chronic Limb
9 Threatening Ischaemia: A Propensity Score Matched Analysis of Health
10 Insurance Claims. *Eur J Vasc Endovasc Surg* 2021;61:423-9.
11 10.1016/j.ejvs.2020.11.006
- 12 181. Singh N, Liu K, Tian L, Criqui MH, Guralnik JM, Ferrucci L, et al. Leg strength
13 predicts mortality in men but not in women with peripheral arterial disease. *J Vasc*
14 *Surg* 2010;52:624-31. 10.1016/j.jvs.2010.03.066
- 15 182. McDermott MM, Ferrucci L, Liu K, Guralnik JM, Tian L, Kibbe M, et al. Women
16 with peripheral arterial disease experience faster functional decline than men with
17 peripheral arterial disease. *J Am Coll Cardiol* 2011;57:707-14.
18 10.1016/j.jacc.2010.09.042
- 19 183. Gommans LN, Scheltinga MR, van Sambeek MR, Maas AH, Bendermacher BL,
20 Teijink JA. Gender differences following supervised exercise therapy in patients
21 with intermittent claudication. *J Vasc Surg* 2015;62:681-8.
22 10.1016/j.jvs.2015.03.076
- 23 184. Regensteiner JG, Bauer TA, Reusch JE, Brandenburg SL, Sippel JM, Vogelsong
24 AM, et al. Abnormal oxygen uptake kinetic responses in women with type II

- 1 diabetes mellitus. *J Appl Physiol* (1985) 1998;85:310-7.
2 10.1152/jappl.1998.85.1.310
- 3 185. Lanzi S, Pousaz A, Calanca L, Mazzolai L. Sex-based differences in supervised
4 exercise therapy outcomes for symptomatic peripheral artery disease. *Vasc Med*
5 2023;1358863X221149454. 10.1177/1358863X221149454
- 6 186. Cetlin MD, Polonsky T, Ho K, Zhang D, Tian L, Zhao L, et al. Barriers to
7 participation in supervised exercise therapy reported by people with peripheral
8 artery disease. *J Vasc Surg* 2023;77:506-14. 10.1016/j.jvs.2022.09.014
- 9 187. Gupta T, Manning P, Kolte D, Smolderen KG, Stone N, Henry JG, et al. Exercise
10 therapy referral and participation in patients with peripheral artery disease:
11 Insights from the PORTRAIT registry. *Vasc Med* 2021;26:654-6.
12 10.1177/1358863X211033649
- 13 188. Harwood A, Smith G, Broadbent E, Cayton T, Carradice D, Chetter I. Access to
14 supervised exercise services for peripheral vascular disease patients. *Bull R Coll*
15 *Surgeons Engl* 2017;99:207–11.
- 16 189. Harwood AE, Smith GE, Cayton T, Broadbent E, Chetter IC. A Systematic Review
17 of the Uptake and Adherence Rates to Supervised Exercise Programs in Patients
18 with Intermittent Claudication. *Ann Vasc Surg* 2016;34:280-9.
19 10.1016/j.avsg.2016.02.009
- 20 190. Li Y, Rother U, Rosenberg Y, Hinterseher I, Uhl C, Mylonas S, et al. A prospective
21 survey study on the education and awareness about walking exercise amongst
22 inpatients with symptomatic peripheral arterial disease in Germany. *Vasa*
23 2023;52:218-23. 10.1024/0301-1526/a001057

- 1 191. Rother U, Dorr G, Malyar N, Muller OJ, Steinbauer M, Ito W, et al. How German
2 vascular surgeons and angiologists judge walking exercise for patients with PAD.
3 *Vasa* 2023;52:224-9. 10.1024/0301-1526/a001071
- 4 192. Saxon JT, Safley DM, Mena-Hurtado C, Heyligers J, Fitridge R, Shishehbor M,
5 et al. Adherence to Guideline-Recommended Therapy-Including Supervised
6 Exercise Therapy Referral-Across Peripheral Artery Disease Specialty Clinics:
7 Insights From the International PORTRAIT Registry. *J Am Heart Assoc*
8 2020;9:e012541. 10.1161/JAHA.119.012541
- 9 193. Lanzi S, Belch J, Brodmann M, Madaric J, Bura-Riviere A, Visona A, et al.
10 Supervised exercise training in patients with lower extremity peripheral artery
11 disease. *Vasa* 2022;51:267-74. 10.1024/0301-1526/a001024
- 12 194. Parodi JC, Fernandez S, Moscovich F, Pulmaria C. Hydration may reverse most
13 symptoms of lower extremity intermittent claudication or rest pain. *J Vasc Surg*
14 2020;72:1459-63. 10.1016/j.jvs.2020.05.066
- 15 195. Mannarino E, Pasqualini L, Innocente S, Scricciolo V, Rignanese A, Ciuffetti G.
16 Physical training and antiplatelet treatment in stage II peripheral arterial occlusive
17 disease: alone or combined? *Angiology* 1991;42:513-21.
18 10.1177/000331979104200701
- 19 196. Hobbs SD, Marshall T, Fegan C, Adam DJ, Bradbury AW. The effect of
20 supervised exercise and cilostazol on coagulation and fibrinolysis in intermittent
21 claudication: a randomized controlled trial. *J Vasc Surg* 2007;45:65-70;
22 discussion 10.1016/j.jvs.2006.08.084

1 **Figures titles and abbreviations**

2 **Structured Graphical Abstract.** Graphical summary of the exercise training
3 approaches in patients with peripheral artery disease.

4

5 **Figure 1.** Pathophysiology of limb symptoms in peripheral artery disease.

6

7 **Figure 2.** Algorithm of chronic symptomatic patients with PAD with indication for
8 exercise treatment. *PAD = peripheral artery disease; SPPB = short physical
9 performance battery; BMT = best medical treatment (including pharmacological and
10 non-pharmacological (lifestyle changes, exercise) approach); DUS = Duplex
11 ultrasound; SF-36 = short-form health 36 questionnaire; WIQ = Walking Impairment
12 Questionnaire; Vascu-QoL6 = Vascular Quality of Life Questionnaire-6.*

13

14 **Figure 3.** Dynamic exercise training induces extensive remodeling of the vascular
15 system. *Skeletal muscle contraction is associated with several physiological, metabolic
16 and mechanical mechanisms that when repeated over several weeks and months,
17 result in mitochondrial biogenesis, angiogenesis, and increases in the functional
18 capacity of individuals with peripheral arterial disease. AMPK = AMP-activated protein
19 kinase; PGC-1 α = peroxisome proliferator-activated receptor gamma coactivator-1 α ;
20 HIF-1 α = hypoxia inducible factor 1-alpha; ERR α = Estrogen-related receptor alpha;
21 VEGF = Vascular endothelial growth factor; NO = nitric oxide; ROS = reactive oxygen
22 species; PGI₂ = prostacyclin; CRP = C-reactive protein; IL-6 = interleukin-6; sICAM-1
23 = soluble intercellular adhesion molecule-1; sVCAM-1 = circulating vascular cell
24 adhesion molecule-1.*