

An updated systematic review and meta-analysis of homebased exercise programs for individuals with intermittent claudication

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1 Review

- 2 An Updated Systematic Review and Meta-Analysis of
- 3 Home-based Exercise Programmes for Individuals with
- 4 Intermittent Claudication.
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- 31 **Running head:** Home-based exercise for intermittent claudication
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ARTICLE HIGHLIGHTS
Type of Research: Systematic Review and Meta-
Analysis
Key Findings: Supervised exercise programmes
are superior to structured home-based exercise
programmes for patients with intermittent
claudication ($p = .004$). However, when
monitoring was used via pedometers or activity
monitors, home-based exercise programmes were
equivalent to supervised exercise programmes (p
=.86).
, ,
Take home Message: When supervised exercise
programmes are unavailable, home-based exercise
programmes can be used. However, they must be
appropriately structured and monitored to be
effective.
Table of Contents Summary
In this meta-analysis, supervised exercise
programmes were superior to structured home-
based exercise programmes. However, home-
based programmes with monitoring methods were
equivalent. When supervised exercise
programmes are unavailable, home-based exercise
programmes can be used. However, they must be
appropriately structured and monitored to be
effective.

67 Abstract:

68 **Objectives:** Supervised exercise programmes (SEP) are effective for improving walking 69 distance in patients with intermittent claudication (IC) but provision and uptake rates are 70 sub-optimal. Access to such programmes has also been halted by the Coronavirus 71 pandemic. The aim of this review is to provide a comprehensive overview of the evidence 72 for home-based exercise programmes (HEP).

Data Sources: Medline, EMBASE, CINAHL, PEDro and Cochrane CENTRAL were
searched for terms relating to HEP and IC.

Review Methods: This review was conducted in according with the published protocol and PRISMA guidance. Randomised and non-randomised trials that compared a HEP to SEP, basic exercise advice or no exercise controls for IC were included. A narrative synthesis was provided for all studies and meta-analyses conducted using data from randomised trials. The primary outcome was maximal walking distance. Sub-group analyses were performed to consider the effect of monitoring. Risk of bias was assessed using the Cochrane tool and quality of evidence via GRADE.

82 **Results:** 23 studies with 1907 participants were included. Considering the narrative

83 review, HEPs were inferior to SEPs which was reflected in the meta-analysis (MD 139m,

84 95% CI 45 to 232m, p = .004, very-low-quality evidence). Monitoring was an important

component, as HEPs adopting this were equivalent to SEPs (MD: 8m, 95% CI -81 to 97,

p = .86; moderate-quality evidence). For HEPs versus basic exercise advice, narrative

87 review suggested HEPs can be superior, though not always significantly so. For HEPs

88 versus no exercise controls, narrative review and meta-analysis suggested HEPs were

potentially superior (MD: 136m, -2-273m p = .05, very-low-quality evidence).

90 Monitoring was also a key element in these comparisons.

91 Other elements such as appropriate frequency ($\geq 3x$ a week), intensity (to moderate-

92 maximum pain), duration (20 progressing to 60 minutes) and type (walking) of exercise

93 were important, as was education, self-regulation, goal setting, feedback and action

94 planning.

- 95 Conclusion: When SEPs are unavailable, HEPs are recommended. However, to elicit
- 96 maximum benefit they should be structured, incorporating all elements of our evidence-
- 97 based recommendations.

98 PROSPERO registration number: CRD42018091248

100 Introduction

101 Peripheral arterial disease (PAD) is categorised by stenotic or occlusive atherosclerotic 102 lesions in the arteries that supply the legs, limiting blood flow¹. Global estimates suggest that PAD affects 237 million people². The classic symptom of PAD is intermittent 103 104 claudication (IC); a reproducible ambulatory lower limb muscle pain, relieved by rest, caused by a muscle oxygen supply and demand imbalance^{3,4}. IC can impede daily 105 106 activities, functional capacity and quality of life (QoL) and carries an increased mortality 107 risk³⁻⁷. First-line treatment for IC includes exercise therapy, ideally in the form of a 108 supervised exercise programme $(SEP)^{8,9}$, with substantial evidence that SEPs significantly 109 improve walking distance ¹⁰⁻¹².

110 Despite this, only $\sim 30\%$ of patients with IC are eligible and willing to join a SEP and the 111 majority of vascular units in the United Kingdom and United States do not have access to one, suggesting they are under-utilised and under-valued¹³⁻¹⁵. Patient-cited barriers include 112 113 a lack of time and transport, whilst provider-cited barriers include a lack of funding, facilities or expertise^{14,16}. Consequently, there has been an increased interest in home-based 114 115 exercise programmes (HEP), with more recent evaluations including technological advancements such as wearable technology¹⁷⁻¹⁹. It is likely that interest in HEP provision 116 has been increased further by the Coronavirus disease 2019 (COVID-19) pandemic, which 117 118 meant that for some time, SEP access was not available, and this may still be the case in 119 some countries.

120

A systematic review in 2013 demonstrated that there was low quality, preliminary evidence
 that HEPs can provide improvements in walking capacity and QoL²⁰. The review

123 concluded that more robust trials were required. Other reviews have attempted to consider the contemporary evidence base for HEPs^{21,22}, However, significant limitations included 124 125 summating the evidence at the same time-points rather than the planned primary endpoint 126 of each trial, including asymptomatic patients and combining exercise advice with no 127 exercise controls, which limits their applicability Therefore, we aimed to update the 128 aforementioned 2013 systematic review and provide a comprehensive overview of the 129 evidence for HEPs versus SEPs, basic exercise advice or no exercise controls for improving walking distance in patients with IC. We also aimed to provide guidance for the most 130 effective HEP elements which can aid healthcare professionals in the design and 131 implementation of an evidence-based structured HEP for those with IC. 132

134 Methods

This systematic review was conducted in accordance with the PRISMA guidelines²³ and was prospectively registered on PROSPERO (CRD42018091248). Furthermore, our protocol outlining the full methodology, including search strategy, data management, outcome measures and the methods for assessing the risk of bias and rating the quality of evidence is published elsewhere²⁴.

Briefly, we included prospective non-randomised and randomised controlled trials (RCT's)
that considered the effect of a HEP versus a comparator arm (SEP, basic exercise advice

142 or no exercise control) on walking distance, QoL and/or physical activity for patients with

143 IC. Searches were performed from database inception and completed in March 2020.

144

145 Data analysis and synthesis

146 Both RCT's and non-RCT's were included and a summary of findings table produced for 147 each comparison including all studies. Where possible, a meta-analysis of RCT's was 148 performed. Where data was not provided to allow entry into a meta-analysis, study authors 149 were contacted, and relevant data requested. Meta-analysis was performed using Review 150 Manager 5 (RevMan 2014), to produce forest plots with an overall effect estimate of mean 151 difference and associated 95% confidence intervals. Random effects models were used for 152 all meta-analyses to consider heterogeneity as interventions and outcomes differed between trials²⁵. For meta-analyses, post-intervention mean and standard deviation was used unless 153 154 only change scores were given. We have summated the results at the planned primary

- assessment point of each trial, rather than at designated time-points (e.g. six weeks) as this
 is the point at which the intervention is designed to have greatest effect²².
- 157

158 A head-to-head analysis of the effectiveness of HEPs versus each comparator arm was 159 conducted and sub-group analyses were performed based on the presence or absence of 160 monitoring. Monitoring included either self-monitoring, using devices such as pedometers, 161 or remote monitoring, using activity monitors. Other pre-specified sub-group analyses 162 were not performed due to insufficient data. Furthermore, the robustness of the analyses 163 was determined via sensitivity analysis. For this, we removed RCT's with a higher risk of bias assessment and repeated the analysis²⁶. Further sensitivity analyses were also 164 performed using change scores from baseline (where reported) instead of final 165 measurement scores as has been recommended²⁷. When certain studies reported only final 166 167 measurement scores, these were used in conjunction with the change scores that were 168 reported for the purpose of sensitivity analyses. All sensitivity analyses are presented in 169 the supplementary material.

We also considered the components of effective HEP interventions, such as the frequency, intensity, time and type of exercise and the use of monitoring or dietary and lifestyle advice or psychological components. Effective HEP interventions were identified as those that induced a significantly greater change (p<0.05) for at least one outcome, when compared with the basic exercise advice or no exercise control comparator groups. For trials comparing a SEP and a HEP, without a no exercise control or basic exercise advice comparator group, the HEP intervention was considered effective if it induced a significant 177 positive change from baseline (p < 0.05). The effective individual components were then 178 identified as those that were evident (and similar) within the majority of these HEPs. 179

180 Results

181 Search Results

The search yielded a total of 4,411 results. Twenty-six articles^{17-19,28-50}, reporting 23 studies, were included in this review, with 18 contributing to meta-analyses (Figure 1). Nine articles included in the previous review were excluded due to lack of an appropriate comparator arm and the inclusion of patients with atypical leg pain. Seventeen additional articles were identified. The definition of HEPs was heterogenous with a number of studies referring to it as 'walking advice' or 'unsupervised exercise' when they were structured and included specific prescriptions.

189

190 Included trials

Of the included trials, three were non-randomised and compared HEPs with SEPs³³⁻³⁵. The remaining trials were RCT's, with nine comparing HEPs with SEPs^{28,30,36,38,41-43,45,47}, three comparing HEPs with basic exercise advice^{31,32,48}, two comparing HEPs with both these groups^{18,46} and six comparing HEPs to no exercise controls^{17,19,29,39,49,50}.

195

196 The total number of recruited patients was 1907. All studies used walking as the mode of 197 exercise. The frequency of training was varied, with three sessions per week being the 198 minimum prescription to a maximum prescription of three times per day. Duration of exercise was either prescribed as minutes per session or number of steps per day. Exercise
intensity was not always specified but was often based on reaching a mild or near-maximal
level of claudication pain. HEP duration and length of follow-up ranged from six weeks to
12 months.

All but one study³² reported treadmill and/or six-minute walk test (6-MWT) MWD, whilst 203 seven did not report PFWD^{17,29,30,32,39,46,50}. There was a lack of consistency between studies 204 205 with regards to how walking distances were reported; either in minutes or metres, or how 206 they were measured; with 15 using a graded treadmill test, five a constant load treadmill 207 test and two the 6-MWT. Three studies also reported both treadmill and 6-MWT MWD. 208 One study, from 1966, was included, but not used in meta-analyses because the treadmill 209 test was not standardised between patients. Generic and disease specific QoL was measured 210 in 14 studies via the Walking Impairment Questionnaire (WIQ), the Medical Outcomes 211 Study short form 36 (SF-36), 20 (SF-20), or 12 (SF-12), the Intermittent Claudication 212 Questionnaire (ICQ), the World Health Organisation quality of life questionnaire, the 213 Vascular Quality of Life Questionnaire and the Euroqol-5D.

214

215 Quality assessment and Risk of Bias

All outcomes were rated via GRADE as very low, low or moderate quality (supplementarytables I-III). The most common reason for rating down was imprecision, based on wide
confidence intervals and/or small sample sizes

219

Risk of bias summary is shown in Figure 2. All studies were rated as high risk forperformance bias due to the nature of the interventions. Across other domains, there was

little evidence of a high risk of bias (other than for selective outcome reporting). However,
there was often inadequate information to imply a low risk, resulting in several domains
being rated as 'unclear'.

225

226 HEP vs. SEP

227 Supplementary-table IV outlines the narrative findings of all studies that compared HEPs with SEPs^{18,28,30,33-36,38,41-43,45-47}. Overall, these studies show that for MWD there were 228 229 statistically significant improvements in half of the HEP groups, and in all of the SEP 230 groups. For between-group analyses, there were significantly greater improvements 231 following SEP in nine of the 14 studies. For PFWD, there were statistically significant 232 improvements in half of the HEP groups and in 11 of the 14 SEP groups, with four of the 233 SEP groups demonstrating significantly greater improvements than the HEP groups. For 234 three studies that adopted monitoring for the HEP via pedometers or step-monitors, there were no differences between groups for improvements in PFWD^{18,34,36}. For MWD, one 235 study reported no differences between groups³⁶, another reported a significantly greater 236 improvement in the SEP group¹⁸ and the final study noted a significant improvement in the 237 SEP group but not the HEP group $(p = .06)^{34}$. The latter study also reported that individual 238 239 increases were 'much higher' in the SEP group, though the difference in improvements 240 between groups was 5% and it was not compared statistically.

For QoL outcomes, there were improvements in the WIQ and the physical functioning domain and physical component summary score of the SF-36 with improvements largely similar between groups.

245 Meta-analysis for MWD from eight studies including 334 participants showed an overall 246 improvement favouring SEPs (MD 139m, 95% CI 45 to 232m, p = .004, very-low-quality 247 evidence; Figure 2A). PFWD, including seven studies and 306 participants also favoured 248 SEPs (MD 84m, 95% CI 25 to 143m, p = .005, very-low-quality evidence; Figure 2B). 249 However, these differences were no longer significant in the sub-group analyses including 250 only trials which included monitoring (moderate-quality evidence; Figure 2). 6-MWD was 251 not significantly different between groups (very-low-quality evidence). 252 The SF-36 measures of pain (p = .006, low-quality evidence) and social functioning (p = .006, low-quality evidence) and social 253 .04, low-quality evidence) significantly favoured SEPs. The WIQ domain of distance also

significantly favoured SEPs (p = .01, very-low-quality evidence). The remaining QoL measures showed no significant mean difference between groups, which was also the case for daily steps (very-low to moderate-quality evidence). (very-low to moderate-quality evidence).

258

259 *HEP vs. basic exercise advice*

Supplementary-table V outlines the narrative findings of the five studies that compared HEPs with basic exercise advice^{31,32,36,46,48}. Three studies reported change from baseline with two noting significant improvements in MWD and PFWD for the HEP groups. Two studies, which included monitoring, demonstrated significantly greater improvements in MWD for the HEP group compared to basic exercise advice.

265

For QoL, there were statistically significant improvements in the WIQ and the physical functioning domain of the SF-36, with the improvements in the WIQ being significantly greater than the basic exercise advice group in one study. For two of the three studies that reported physical activity measures, there were significantly greater improvements in daily steps and maximum 20-, 30- and 60-minute cadence for the HEP group in comparison to the basic exercise advice group^{32,36}.

272

273 Meta-analysis for MWD from four studies including 137 participants showed no significant 274 difference between groups (MD 39.0m, 95% CI -123.1 to 201.1m, p = .64, very-low-275 quality evidence; Figure 3A). For sub-group analysis, findings were not altered for studies adopting monitoring. However, monitoring appeared important as there was a trend (p =276 277 .05) for HEPs without it to be inferior to basic walking advice (very-low-quality evidence, 278 Figure 3A). For PFWD, including 3 studies and 109 participants, there was a significant 279 between group difference, favouring HEPs (MD 64.5m, 95% CI 14.1 to 114.8m, p = .01, 280 very-low-quality evidence; Figure 3B). Two of the three studies in this analysis adopted 281 monitoring, precluding sub-group analysis. There was also a significant between group difference for the ICQ, favouring HEPs (p = <.01, low-quality-evidence). There were no 282 283 significant mean differences for daily steps or the WIQ (very-low-quality evidence).

284 HEP vs. no exercise controls

Supplementary table VI outlines the narrative findings of all 6 studies that compared HEPs with no exercise controls^{17, 19,29,39,49,50}. Three studies provided statistical comparisons and there were significant improvements in MWD and PFWD for the HEP groups, which were generally, significantly greater than the control groups. Two studies provided statistical comparisons for the 6-MWD with one demonstrating significant improvements in the HEP group, whilst the other showed no significant difference compared to baseline or control. 291

For QoL outcomes, there were improvements in the WIQ though they were not analysed statistically. The SF-12 and SF-36 outcomes were variable between studies.

294 For two studies that reported physical activity measures, only one provided statistical comparison and reported no significant improvements in either group^{19,48}. For the three 295 296 studies that adopted monitoring via an activity monitor or pedometer, two reported 297 significant improvements in MWD for the HEP group and one also reported a greater improvement compared to the control group¹⁹. Meta-analysis including three studies and 298 299 100 participants revealed a mean difference in MWD of 136m, favouring HEPs, though it 300 was not significant (95% CI -2 to 273m, p = .05, very-low-quality evidence; figure 4). 301 There were insufficient studies to perform a meta-analysis of PFWD or sub-group analysis 302 for MWD. There were no significant mean differences for daily steps, 6-MWD, the WIQ 303 or the physical and mental component summaries of the SF-12/36 (moderate to very-low-304 quality evidence).

305

306 *HEP adherence*

307 HEP adherence was poorly reported, stated in only seven studies^{18,19,29,30,32,33,36} and 308 assessed via self-reported methods in four^{29,30,32,33}. Three studies were able to receive 309 quantified adherence information via their remote monitoring methods^{18, 19, 36}.

Four studies reported an adherence of $>80\%^{18,29,32,36}$, and the lowest reported was 67%.

311 The HEP prescribed on the basis of step count, reported poor adherence to the prescribed

312 steps, but did not report adherence to frequency of exercise¹⁹.

313

314 Discussion

The aim of this systematic review and meta-analysis was to provide an up-to-date comprehensive overview of the evidence for HEPs versus SEPs, basic exercise advice and no exercise controls for patients with IC. Comparable to a recent review⁵¹, the overall findings indicated that HEPs are inferior to SEPs for improvements in PFWD and MWD. However, HEPs may be more effective than basic exercise advice, and certainly more effective than no exercise at all. One novel finding is that for all comparisons, monitoring appeared to be an important contributing factor to an effective HEP.

322

323 The apparent superiority of SEPs compared to HEPs, could be due to differences in the 324 exercise dose between the two programme types. SEPs are, within reason, clearly defined 325 as structured exercise with recommended frequency, intensity, time and type (FITT) principles^{8,52-54}. HEPs are much less established, have varied utilisation and suffer greater 326 327 heterogeneity, especially in older studies. Indeed, three studies included SEPs that had (up to 40 minute) longer individual sessions than the HEP^{28,38,43}, whilst two SEP groups were 328 also told to complete the HEP in conjunction with the SEP^{34,38}, meaning they received at 329 330 least one extra exercise session per week, compared to the HEP only group. Conversely, three HEPs prescribed daily walking^{33,38,41}, up to a maximum of three times a day, versus 331 332 a frequency of two to three times a week in the SEP group. This HEP prescription may be 333 too intense and discourage engagement, especially given the reduced functional capacity evident in these patients¹. As such, heterogeneity may be greater for HEPs than it is for 334 335 SEPs, especially with regards to dose, contributing to their inferiority.

337 Additionally, the terminology used to describe HEPs may also be a contributing factor. 338 HEP descriptions included 'exercise advice' or 'unsupervised exercise', which for patients 339 can either be too vague, or even perceived as optional (in the case of exercise advice). It is 340 therefore important that patients are made aware that exercise therapy, including HEPs 341 when appropriate, constitutes part of their treatment regime and should be adhered to, as 342 well as being provided in a way that is structured and multifaceted, rather than simple 343 advice. This problem is compounded by recent guidelines which identify that home-based 344 walking is a useful alternative to SEPs, but refer only to simple 'unsupervised' or 'nonsupervised' exercise with no specific recommendations⁹. 345

346

347 Evidence from our sub-group analyses suggests that HEPs may not always be inferior to 348 SEPs. Specifically, HEPs adopting remote or self-monitoring, via pedometers and/or 349 activity monitors were equivalent to SEPs, or at least reduced their superiority by half for 350 improvements in MWD. Furthermore, the results also suggest that HEPs without 351 monitoring may be inferior to basic exercise advice. One possible explanation for the 352 apparent benefit of monitoring is that it can provide a form of remote supervision, with 353 four of the seven monitoring studies having the facility to regularly feedback data to the study team, potentially improving adherence^{17-19,36}. For SEPs, the intensity of supervision 354 is associated with the level of improvement in walking distance⁵¹. It would therefore be 355 356 reasonable to assume that this remote supervision will be more effective than little or no 357 supervision (or monitoring) at all. However, based on the findings of three studies included in this review^{17,18,36}, for remote monitoring to be most effective, and to add specificity to 358

feedback, the device should only be worn during exercise sessions, rather than at all timesduring the day.

361

362 In addition to remote monitoring, self-monitoring, with the use of pedometers and exercise 363 diaries, also appeared effective. This is not surprising given that pedometer use is 364 associated with a reactive effect, with the greatest reactivity seen in those who are asked to 365 record their daily step count in an activity diary⁵⁵. This process of recording daily step 366 count may increase awareness of activity levels, leading to effective goal-setting and 367 greater confidence for walking. Monitoring via exercise diaries (without step-monitors) or 368 telephone calls is ineffective. Clearly, given the variety of possible monitoring, 369 standardisation is required. However, we recommend pedometers in conjunction with an 370 exercise diary as the minimum.

371

372 In addition to monitoring, a number of HEP components were identified in studies which, in isolation, appeared to provide similar benefits to SEPs^{18,34,36,47}, or superior benefits to 373 basic exercise advice or no exercise controls^{19,29,36,48}. As such, we have created an example 374 375 supported home-based exercise programme (SHEP), outlined in table I. Our programme is 376 structured and includes a detailed prescription based on the FITT principle, and 377 incorporates support including regular feedback (ideally in real-time), goal setting and 378 patient education with appropriate theoretical underpinning. These elements also 379 demonstrated good patient adherence, have recently been highlighted as important from the PAD patient perspective⁵⁶ and provide a holistic, patient-centred approach. 380

Only one study has combined these components into a deliverable structured HEP⁴⁸, 382 383 though it was not an adequately powered RCT, meaning it is currently untested. Future, 384 larger, longer-term studies adopting this SHEP structure in a way that is accessible and 385 pragmatic to patients, such as via telehealth (alongside other monitoring), which has shown promise in other clinical populations^{57,58}, are required. These studies should report the 386 intervention in full to aid replication in clinical practice⁵⁹. In addition, they should also 387 388 report clinical and cost-effectiveness and the patient eligibility, recruitment, adherence and 389 completion rates. This important information is required to build an appropriate evidence 390 base for the effectiveness of a standardised, structured SHEP, whilst identifying if it is an 391 acceptable alternative to SEPs.

392

However, in the absence of such an evidence base, HEPs should currently only be considered when SEPs are unavailable or impractical. HEPs should also be considered in exceptional circumstances, such as the COVID-19 pandemic, which suspended SEP availability and practicality. Under these normal and exceptional circumstances, we recommend that a structured SHEP, based on the components outlined in table I, is likely most effective, and should be provided to engage more patients in appropriate lifestyle and exercise behaviour change.

Such a programme could also be recommended to aid continued engagement for those who
do complete a SEP, as currently, there is limited provision of long-term exercise
recommendations.

403

404 Limitations

A number of studies provided inadequate data to allow for meta-analysis, meaning the meta-analyses provided herein do not encompass the full evidence base. In addition, a number of meta-analysable outcomes were restricted by moderate to very-low-quality evidence, small sample sizes and a lack of robustness to sensitivity analyses, meaning their interpretation is limited. Finally, due to the limited number of studies included in the metaanalysis, publication bias could not be excluded via funnel plot.

411 Conclusion

412 HEPs still appear inferior to SEPs. However, with remote- and self-monitoring this 413 inferiority is markedly reduced. Compared to basic exercise advice, HEPs generally 414 provided a benefit, though this was not always significantly greater. However, HEPs did 415 appear to demonstrate superiority compared to no exercise controls for improvements in 416 MWD, though with very-low-quality evidence. As such, evidence for HEPs suggests they 417 should only be recommended when SEPs are unavailable or impractical. When HEPs are 418 appropriate, they should be structured and personalised, taking into account the specific 419 FITT (and other) principles, provided in the recommendations outlined above. Larger, 420 longer-term studies combining all of these elements into one accessible, pragmatic SHEP, 421 potentially via telehealth, should provide the future direction of HEP-based research for 422 patients with IC.

423

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433

434 Conflict of Interest Statement

435 The authors declare that there is no conflict of interest.

437 References:

438	1.	Hiatt WR. Medical treatment of peripheral arterial disease and claudication. N
439		Engl J Med 2001;344(21):1608-21.
440	2.	Song P, Rudan D, Zhu Y, Fowkes FJ, Rahimi K, Fowkes FGR, et al. Global, regional,
441		and national prevalence and risk factors for peripheral artery disease in 2015: an
442		updated systematic review and analysis. The Lancet Global Health
443		2019;7(8):e1020-e1030.
444	3.	Harwood A-E, Broadbent E, Totty JP, Smith GE, Chetter IC. "Intermittent
445		claudication a real pain in the calf"—Patient experience of diagnosis and
446		treatment with a supervised exercise program. Journal of Vascular Nursing
447		2017;35(3):131-135.
448	4.	Meru AV, Mittra S, Thyagarajan B, Chugh A. Intermittent claudication: an
449		overview. Atherosclerosis 2006;187(2):221-237.
450	5.	Golomb BA, Dang TT, Criqui MH. Peripheral arterial disease. Circulation
451		2006;114(7):688-699.
452	6.	Pell J. Impact of intermittent claudication on quality of life. European Journal of
453		Vascular and Endovascular Surgery 1995;9(4):469-472.
454	7.	Criqui MH, Langer RD, Fronek A, Feigelson HS, Klauber MR, McCann TJ, et al.
455		Mortality over a period of 10 years in patients with peripheral arterial disease.
456		New England Journal of Medicine 1992;326(6):381-386.
457	8.	NICE. Peripheral arterial disease: diagnosis and management. Clinical guidance
458		147.; 2012.
459	9.	Aboyans V, Ricco J-B, Bartelink M-LE, Björck M, Brodmann M, Cohnert T, et al.
460		2017 ESC Guidelines on the Diagnosis and Treatment of Peripheral Arterial
461		Diseases, in collaboration with the European Society for Vascular Surgery (ESVS)
462		Document covering atherosclerotic disease of extracranial carotid and vertebral,
463		mesenteric, renal, upper and lower extremity arteries Endorsed by: the
464		European Stroke Organization (ESO) The Task Force for the Diagnosis and
465		Treatment of Peripheral Arterial Diseases of the European Society of Cardiology
466		(ESC) and of the European Society for Vascular Surgery (ESVS). European heart
467		journal 2017;39(9):763-816.
468	10.	Lane R, Harwood A, Watson L, Leng GC. Exercise for intermittent claudication.
469		The Cochrane database of systematic reviews 2017;12:CD000990.
470	11.	Gardner AW, Montgomery PS, Flinn WR, Katzel LI. The effect of exercise intensity
471		on the response to exercise rehabilitation in patients with intermittent
472		claudication. Journal of vascular surgery 2005;42(4):702-9.
473	12.	Lauret GJ, Fakhry F, Fokkenrood HJ, Hunink MG, Teijink JA, Spronk S. Modes of
474		exercise training for intermittent claudication. Cochrane Database Syst Rev
475		2014(7):CD009638.
476	13.	Harwood A-E, Smith GE, Cayton T, Broadbent E, Chetter IC. A systematic review
477		of the uptake and adherence rates to supervised exercise programs in patients
478		with intermittent claudication. Annals of vascular surgery 2016;34:280-289.

479	1.4	Harwood A. Smith C. Broadhant F. Cautan T. Carradian D. Chattar I. According
479	14.	Harwood A, Smith G, Broadbent E, Cayton T, Carradice D, Chetter I. Access to
		supervised exercise services for peripheral vascular disease patients. The Bulletin
481	45	of the Royal College of Surgeons of England 2017;99(6):207-211.
482	15.	Dua A, Gologorsky R, Savage D, Rens N, Gandhi N, Brooke B, et al. National
483		assessment of availability, awareness, and utilization of supervised exercise
484		therapy for peripheral artery disease patients with intermittent claudication.
485		Journal of vascular surgery 2019.
486	16.	Harwood AE, Hitchman LH, Ingle L, Doherty P, Chetter IC. Preferred exercise
487		modalities in patients with intermittent claudication. Journal of Vascular Nursing
488		2018.
489	17.	McDermott MM, Spring B, Berger JS, Treat-Jacobson D, Conte MS, Creager MA,
490		et al. Effect of a home-based exercise intervention of wearable technology and
491		telephone coaching on walking performance in peripheral artery disease: the
492		HONOR randomized clinical trial. Jama 2018;319(16):1665-1676.
493	18.	Gardner AW, Parker DE, Montgomery PS, Blevins SM. Step-monitored home
494		exercise improves ambulation, vascular function, and inflammation in
495		symptomatic patients with peripheral artery disease: a randomized controlled
496		trial. Journal of the American Heart Association 2014;3(5):e001107.
497	19.	Duscha BD, Piner LW, Patel MP, Crawford LE, Jones WS, Patel MR, et al. Effects of
498		a 12-week mHealth program on FunctionalCapacity and physical activity in
499		patients with peripheralArtery disease. The American journal of cardiology
500		2018;122(5):879-884.
501	20.	Al-Jundi W, Madbak K, Beard JD, Nawaz S, Tew GA. Systematic review of home-
502		based exercise programmes for individuals with intermittent claudication.
503		European journal of vascular and endovascular surgery : the official journal of
504		the European Society for Vascular Surgery 2013;46(6):690-706.
505	21.	Hageman D, Fokkenrood H, Gommans L, van den Houten M, Teijink J. Supervised
506		exercise therapy versus home-based exercise therapy versus walking advice for
507		intermittent claudication. The Cochrane database of systematic reviews
508		2018;4:CD005263.
509	22.	Golledge J, Singh T, Alahakoon C, Pinchbeck J, Yip L, Moxon J, et al. Meta-analysis
510		of clinical trials examining the benefit of structured home exercise in patients
511		with peripheral artery disease. British Journal of Surgery 2019;106(4):319-331.
512	23.	Liberati A, Altman DG, Tetzlaff J, Mulrow C, Gøtzsche PC, Ioannidis JP, et al. The
513		PRISMA statement for reporting systematic reviews and meta-analyses of
514		studies that evaluate health care interventions: explanation and elaboration.
515		PLoS medicine 2009;6(7):e1000100.
516	24.	Pymer S, Tew GA, Palmer J, Ingle L, Smith GE, Chetter IC, et al. Home-based
517	2	exercise programmes for individuals with intermittent claudication: A protocol
518		for an updated systematic review and meta-analysis. SAGE open medicine
519		2018;6:2050312118818295.
520	25.	Ryan R. Heterogeneity and subgroup analyses in Cochrane Consumers and
520	23.	Communication Group
522	review	vs: Planning the analysis at protocol stage. <u>http://cccrg.cochrane.org</u> . In; 2016.
522		vs. Flamming the analysis at protocol stage. <u>http://ccorg.cocmanc.org</u> . m, 2010.

523	26.	Bown M, Sutton A. Quality control in systematic reviews and meta-analyses.
524		European Journal of Vascular and Endovascular Surgery 2010;40(5):669-677.
525	27.	Fu R, Holmer HK. Change score or followup score? An empirical evaluation of the
526		impact of choice of mean difference estimates. 2015.
527	28.	Allen JD, Stabler T, Kenjale A, Ham KL, Robbins JL, Duscha BD, et al. Plasma nitrite
528		flux predicts exercise performance in peripheral arterial disease after 3 months
529		of exercise training. Free Radical Biology and Medicine 2010;49(6):1138-1144.
530	29.	Brenner IK, Brown CA, Hains SJ, Tranmer J, Zelt DT, Brown PM. Low-Intensity
531		Exercise Training Increases Heart Rate Variability in Patients With Peripheral
532		Artery Disease. Biological research for nursing 2020;22(1):24-33.
533	30.	Cheetham D, Burgess L, Ellis M, Williams A, Greenhalgh R, Davies A. Does
534		supervised exercise offer adjuvant benefit over exercise advice alone for the
535		treatment of intermittent claudication? A randomised trial. European journal of
536		vascular and endovascular surgery 2004;27(1):17-23.
537	31.	Christman SK. Intervention to slow progression of peripheral arterial disease.
538		2003.
539	32.	Cunningham M, Swanson V, O'Caroll R, Holdsworth R. Randomized clinical trial
540		of a brief psychological intervention to increase walking in patients with
541		intermittent claudication. British journal of surgery 2012;99(1):49-56.
542	33.	Degischer S, Labs K-H, Hochstrasser J, Aschwanden M, Tschoepl M, Jaeger KA.
543		Physical training for intermittent claudication: a comparison of structured
544		rehabilitation versus home-based training. Vascular Medicine 2002;7(2):109-115.
545	34.	Dopheide JF, Geissler P, Rubrech J, Trumpp A, Zeller GC, Daiber A, et al. Influence
546		of exercise training on proangiogenic TIE-2 monocytes and circulating angiogenic
547		cells in patients with peripheral arterial disease. Clinical Research in Cardiology
548		2016;105(8):666-676.
549	35.	Fakhry F, Spronk S, de Ridder M, den Hoed PT, Hunink MM. Long-term effects of
550		structured home-based exercise program on functional capacity and quality of
551		life in patients with intermittent claudication. Archives of physical medicine and
552		rehabilitation 2011;92(7):1066-1073.
553	36.	Gardner AW, Parker DE, Montgomery PS, Scott KJ, Blevins SM. Efficacy of
554		quantified home-based exercise and supervised exercise in patients with
555		intermittent claudication: a randomized controlled trial. Circulation
556		2011;123(5):491-8.
557	37.	Imfeld S, Singer L, Degischer S, Aschwanden M, Thalhammer C, Jaeger K. Quality
558		of life improvement after hospital-based rehabilitation or home-based physical
559		training in intermittent claudication. VASA Zeitschrift fur Gefasskrankheiten
560		2006;35(3):178-184.
561	38.	Kakkos S, Geroulakos G, Nicolaides A. Improvement of the walking ability in
562		intermittent claudication due to superficial femoral artery occlusion with
563		supervised exercise and pneumatic foot and calf compression: a randomised
564		controlled trial. European journal of vascular and endovascular surgery
565		2005;30(2):164-175.

566	39.	McDermott MM, Liu K, Guralnik JM, Criqui MH, Spring B, Tian L, et al. Home-
567		based walking exercise intervention in peripheral artery disease: a randomized
568		clinical trial. Jama 2013;310(1):57-65.
569	40.	McDermott MM, Guralnik JM, Criqui MH, Ferrucci L, Zhao L, Liu K, et al. Home-
570		based walking exercise in peripheral artery disease: 12-month follow-up of the
571		GOALS randomized trial. Journal of the American Heart Association
572		2014;3(3):e000711.
573	41.	Nicolaï SP, Teijink JA, Prins MH. Multicenter randomized clinical trial of
574		supervised exercise therapy with or without feedback versus walking advice for
575		intermittent claudication. Journal of vascular surgery 2010;52(2):348-355.
576	42.	Parmenter BJ, Raymond J, Dinnen P, Lusby RJ, Fiatarone Singh MA. High-Intensity
577		Progressive Resistance Training Improves Flat-Ground Walking in Older Adults
578		with Symptomatic Peripheral Arterial Disease. Journal of the American Geriatrics
579		Society 2013;61(11):1964-1970.
580	43.	Patterson RB, Pinto B, Marcus B, Colucci A, Braun T, Roberts M. Value of a
581		supervised exercise program for the therapy of arterial claudication. Journal of
582		Vascular Surgery 1997;25(2):312-319.
583	44.	Pinto BM, Marcus BH, Patterson RB, Roberts M, Colucci A, Braun C. On-Site
584		Versus Home Exercise Programs: Psychological Benefits for Individuals With
585		Arterial Claudication. Journal of Aging & Physical Activity 1997;5(4).
586	45.	Regensteiner JG, Meyer TJ, Krupski WC, Cranford LS, Hiatt WR. Hospital vs home-
587		based exercise rehabilitation for patients with peripheral arterial occlusive
588		disease. Angiology 1997;48(4):291-300.
589	46.	Sandercock GR, Hodges LD, Das SK, Brodie DA. The impact of short term
590		supervised and home-based walking programmes on heart rate variability in
591		patients with peripheral arterial disease. Journal of sports science & medicine
592		2007;6(4):471.
593	47.	Savage P, Ricci MA, Lynn M, Gardner A, Knight S, Brochu M, et al. Effects of home
594		versus supervised exercise for patients with intermittent claudication. Journal of
595		Cardiopulmonary Rehabilitation and Prevention 2001;21(3):152-157.
596	48.	Tew GA, Humphreys L, Crank H, Hewitt C, Nawaz S, Al-Jundi W, et al. The
597		development and pilot randomised controlled trial of a group education
598		programme for promoting walking in people with intermittent claudication.
599		Vascular Medicine 2015;20(4):348-357.
600	49.	Larsen A, Lassen N. Effect of daily muscular exercise in patients with intermittent
601		claudication. The Lancet 1966;288(7473):1093-1095.
602	50.	Galea-Holmes M, Weinman J, Bearne L. A randomized controlled feasibility trial
603		of a home-based walking behavior-change intervention for people with
604		intermittent claudication. Journal of vascular nursing: official publication of the
605		Society for Peripheral Vascular Nursing 2019;37(2):135-143.
606	51.	Gommans L, Saarloos R, Scheltinga M, Houterman S, de Bie R, Fokkenrood H, et
607		al. The effect of supervision on walking distance in patients with intermittent
608		claudication: a meta-analysis. Journal of Vascular Surgery 2014;60(2):535-536.

609	52.	Gerhard-Herman MD, Gornik HL, Barrett C, Barshes NR, Corriere MA, Drachman
610		DE, et al. 2016 AHA/ACC guideline on the management of patients with lower
611		extremity peripheral artery disease: executive summary: a report of the
612		American College of Cardiology/American Heart Association Task Force on
613		Clinical Practice Guidelines. Journal of the American College of Cardiology
614		2017;69(11):1465-1508.
615	53.	Tew GA, Harwood AE, Ingle L, Chetter I, Doherty PJ. The BASES Expert Statement
616		on Exercise Training for People with Intermittent Claudication due to Peripheral
617		Arterial Disease. The Sport and Exercise Scientist 2018.
618	54.	ACSM. ACSM's Guidelines for Exercise Testing and Prescription. 9th ed.
619		Baltimore: Lippincott, Williams, & Wilkins.; 2014.
620	55.	Clemes SA, Parker RA. Increasing our understanding of reactivity to pedometers
621		in adults. Medicine & Science in Sports & Exercise 2009;41(3):674-680.
622	56.	Lortz J SJ, Kuether T, Kreitschmann-Andermahr I, Ullrich G, Steinmetz M,
623		Rammos C, Jánosi RA, Moebus S, Rassaf T, Paldán K. Needs and requirements of
624		patients with peripheral arterial disease: Questionnaire study to design patient-
625		centred mobile interventions. JMIR Formative Research (pre-print) 2020.
626	57.	Rawstorn JC, Gant N, Direito A, Beckmann C, Maddison R. Telehealth exercise-
627		based cardiac rehabilitation: a systematic review and meta-analysis. Heart
628		2016;102(15):1183-1192.
629	58.	Maddison R, Rawstorn JC, Stewart RA, Benatar J, Whittaker R, Rolleston A, et al.
630		Effects and costs of real-time cardiac telerehabilitation: randomised controlled
631		non-inferiority trial. Heart 2019;105(2):122-129.
632	59.	Tew GA, Brabyn S, Cook L, Peckham E. The completeness of intervention
633		descriptions in randomised trials of supervised exercise training in peripheral
634		arterial disease. PloS one 2016;11(3):e0150869.
635		