

# A 5-year follow-up study of Alfredson's heel-drop exercise programme in chronic midportion Achilles tendinopathy

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## ABSTRACT

**Background** Eccentric exercises have the most evidence in conservative treatment of midportion Achilles tendinopathy. Although short-term studies show significant improvement, little is known of the long-term (>3 years) results.

**Aim** To evaluate the 5-year outcome of patients with chronic midportion Achilles tendinopathy treated with the classical Alfredson's heel-drop exercise programme.

**Study design** Part of a 5-year follow-up of a previously conducted randomised controlled trial.

**Methods** 58 patients (70 tendons) were approached 5 years after the start of the heel-drop exercise programme according to Alfredson. At baseline and at 5-year follow-up, the validated Victorian Institute of Sports Assessment–Achilles (VISA-A) questionnaire score, pain status, alternative treatments received and ultrasonographic neovascularisation score were recorded.

**Results** In 46 patients (58 tendons), the VISA-A score significantly increased from 49.2 at baseline to 83.6 after 5 years ( $p < 0.001$ ) and from the 1-year to 5-year follow-up from 75.0 to 83.4 ( $p < 0.01$ ). 39.7% of the patients were completely pain-free at follow-up and 48.3% had received one or more alternative treatments. The sagittal tendon thickness decreased from 8.05 mm (SD 2.1) at baseline to 7.50 mm (SD 1.6) at the 5-year follow-up ( $p = 0.051$ ).

**Conclusion** At 5-year follow-up, a significant increase of VISA-A score can be expected. After the 3-month Alfredson's heel-drop exercise programme, almost half of the patients had received other therapies. Although improvement of symptoms can be expected at long term, mild pain may remain.

## INTRODUCTION

There are many conservative treatment options for chronic Achilles tendinopathy. Systematic reviews showed that the heavy load eccentric exercise programme has the most evidence of effectiveness in the treatment of chronic midportion Achilles tendinopathy.<sup>1–4</sup> The reviews demonstrated promising results on pain. However, there is limited evidence to show that eccentric exercises are more effective than other treatments in the management of tendinopathies. The exact mechanism involved in the eccentric exercises is unclear and there is an ongoing debate on the optimal load, frequency and duration.<sup>5,6</sup> It is suggested that eccentric training might stimulate

remodelling and tissue repair in tendons, the so-called therapeutic mechano transduction.<sup>7</sup> The classical heel-drop exercise programme first described by Alfredson *et al*<sup>8</sup> is time-consuming (180 repetitions per day for 12 weeks). The majority of the published studies have a maximum follow-up of 1 year, and there is a lack of evidence about the long-term (>3 years) outcome.<sup>9–13</sup> Three studies evaluated the long-term (>3 years) results of the eccentric exercises in chronic midportion Achilles tendinopathy.<sup>14–16</sup> Only one study used the validated Victorian Institute of Sports Assessment–Achilles (VISA-A) questionnaire, but in that study the exercise regimen differed from that described by Alfredson *et al*.<sup>6</sup> The VISA-A questionnaire is a validated score that measures pain and physical activity of the Achilles tendon.<sup>17</sup> The test is developed to create an index of severity of Achilles tendinopathy.

So far, no long-term VISA-A score results (>3 years) are known of the widely used heel-drop programme according to Alfredson. We evaluated the long-term clinical outcome of the heel-drop programme as described by Alfredson with the VISA-A score as the primary outcome measure. We also evaluated the ultrasonographic improvement 5 years after conservative therapy, alternative treatments received, pain status and patient satisfaction.

## METHODS

This study is the 5-year follow-up of an earlier published randomised controlled trial (RCT).<sup>13</sup> The RCT evaluated the effect of the addition of a night splint to eccentric exercises in patients with chronic Achilles tendinopathy. At baseline, 58 patients (70 tendons) with chronic midportion Achilles tendinopathy were included. Inclusion criteria at baseline were the presence of symptoms for more than 2 months and participation in sporting activities. All patients in the trial performed the heel-drop programme as described by Alfredson *et al*.<sup>8</sup> Half of the patients additionally received a night splint. The night splint had no significant influence on the VISA-A score at the 3-month and 1-year follow-up; all patients were therefore merged into one group for the 5-year follow-up.<sup>13,18</sup> After completion of the heel-drop programme, patients were free to choose additional treatment(s); no instructions or recommendations were given to the patients.



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The study protocol was approved by the Medical Ethics Committee of the Hospital (MEC05-21) and informed consent was given.

### Follow-up

All patients were approached by telephone for the 5-year follow-up. At the outpatient clinic, each patient completed the VISA-A score questionnaire and received an ultrasound examination by a single researcher. The VISA-A score was completed with minimal researcher assistance. Secondary outcome scores were obtained using a standardised questionnaire: (1) pain status (pain-free, not pain-free), (2) condition of the uninvolved tendon (affected, not affected), (3) subjective patient satisfaction towards the current status (categorised in poor, moderate, good, excellent), (4) received alternative treatments after the heel-drop programme and, (5) continuation of the eccentric exercises (continued, stopped after the heel-drop programme).

### Ultrasound examination

For ultrasound examination, patients were asked to lie prone with their feet hanging over the edge of the table. During the examination, both Achilles tendons were evaluated in the sagittal plane for thickness and neovascularisation according to the Öhberg scoring system. This score was evaluated as 0 (no vessels visible), 1+ (one vessel mostly in the anterior part), 2+ (one or two vessels throughout the tendon), 3+ (three vessels throughout the tendon) and 4+ (more than three large vessels throughout the tendon). Two different power Doppler machines were used to examine the tendons: the MyLab25Gold (Esaote Piemedical, Maastricht, The Netherlands) and the BK Pro Focus (BK Medical, Herlev, Denmark). Both power Doppler

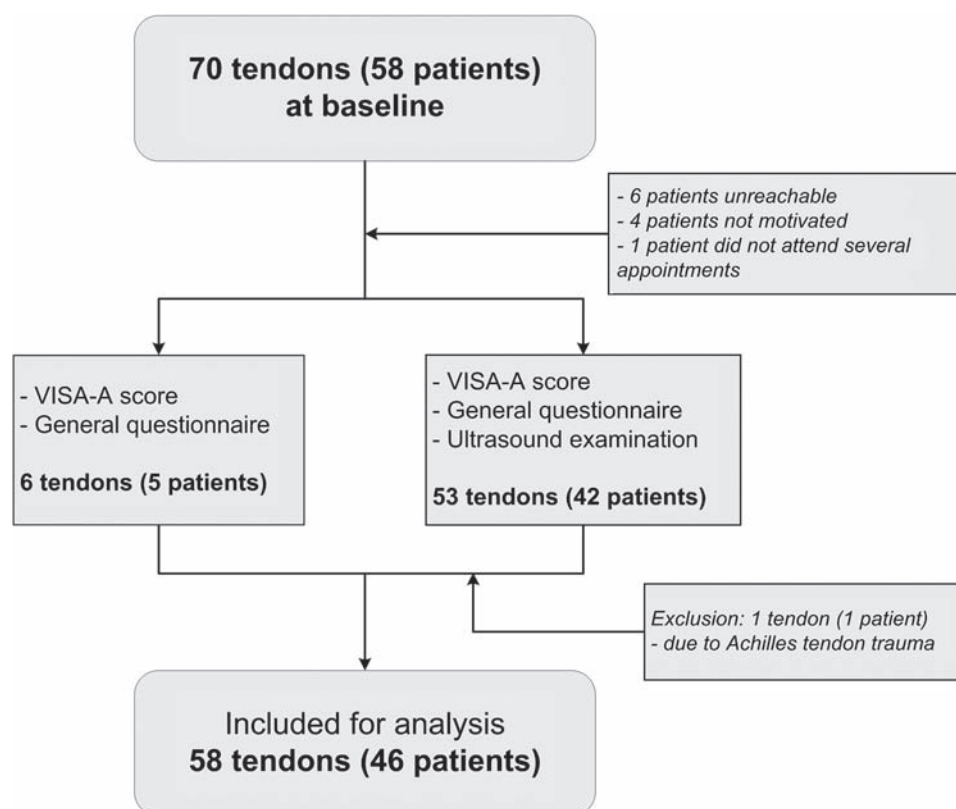
machines used a frequency of 12 MHz, 0.5 PRF (pulse repetition frequency). The colour gain setting was optimised for low flow; putting the setting just below that level produces random noise, varying from 66% to 79%.<sup>19</sup>

### Statistical analysis

Differences in VISA-A score and tendon thickness between baseline and follow-up were evaluated using a sample t test. Differences between two categorical variables were evaluated using a  $\chi^2$  test. Dependency analysis of the VISA-A score was performed using a non-repetitive analysis of covariance test. Statistical Package for the Social Sciences software (SPSS 17.0) was used. A P value of less than 0.05 was considered to be statistically significant.

### RESULTS

Attempts were made to contact all 58 patients (70 tendons). Six patients (six tendons) were not reachable for follow-up and five patients (five tendons) refused to participate (see Figure 1). Three out of these five patients reported having no or minimal complaints of the Achilles tendon; unfortunately, further details are unknown. The VISA-A score and questionnaire results were obtained from 47 patients (59 tendons): 12 bilateral and 35 unilateral. One patient (one tendon) was excluded from analysis due to a direct trauma to the Achilles tendon 2 days before the follow-up appointment, making the outcome non-representative. For general statistical analysis, 46 patients (58 tendons) were used. Five patients (six tendons) were not able to visit the hospital (no time or living abroad) but filled out the general questionnaire and the VISA-A score (see figure 1 for details). The ultrasonographic data consisted of information from 41 patients (52 tendons).



**Figure 1** Flowchart of included patients.

**Table 1** Patient characteristics at 5-year follow-up

	46 Patients (58 tendons)
Mean age (years (range))	50.9 (36–64)
Body mass index (kg/m <sup>2</sup> (range))	25.4 (19.9–33.5)
Unilateral/bilateral	34/12
No sports activity	4
Recreational sports	35
Competitive sports	7

At follow-up, the mean age was 50.9 years, with a mean body mass index of 25.4 kg/m<sup>2</sup> (see table 1 for details). Baseline specifics are described by de Vos *et al.*<sup>13</sup>

### Primary outcome: VISA-A score

The VISA-A score (n=58) improved significantly from 49.2 (SD 20.1) at baseline to 83.6 (SD 14.9) at the 5-year follow-up (p<0.001). The VISA-A score (n=55) improved significantly between the 1-year follow-up and the 5-year follow-up from 75.0 (SD 25.0) to 83.4 (SD 15.1) (p=0.006). See figure 2 for the trend line of the VISA-A score.

### Pain status

Of the patients, 39.7% reported being completely pain-free at the 5-year follow-up (mean VISA-A score 92.8, SD 8.8). The remaining 60.3% experienced some degree of pain, varying from only during extensive exercise to permanent pain (mean VISA-A score 77.5, SD 15.1). From the 34 patients with unilateral Achilles tendinopathy, 43.3% developed some degree of pain in the contralateral Achilles tendon.

### Patient satisfaction

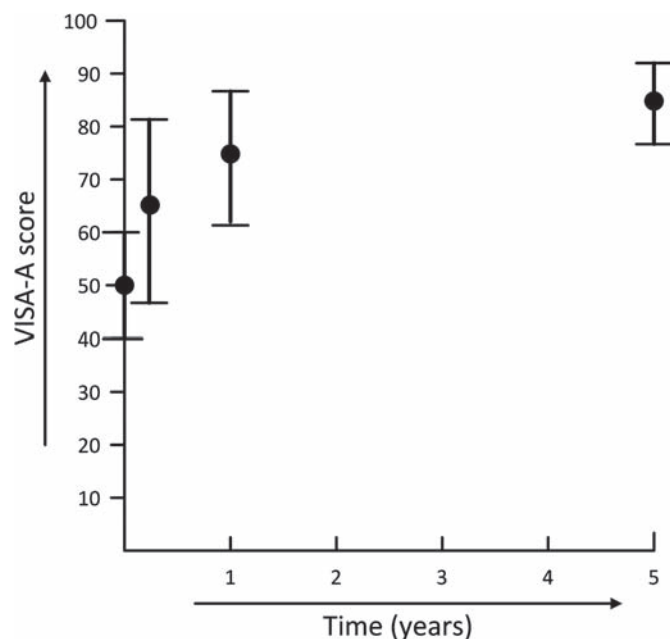
Patient subjective satisfaction towards the current status was excellent in 8.6% (mean VISA-A score 89.0, SD 8.0), good in 41.4% (mean VISA-A score 84.6, SD 16.3), moderate in 31.0% (mean VISA-A score 84.4, SD 15.2) and poor in 19.0% (mean VISA-A score 77.6, SD 13.5).

### Alternative treatments

In total, 48.3% (22 patients) had received one or more alternative treatments, non-custom-made inlays (11 patients), surgery (6 patients), physiotherapy/massage/friction (6 patients), shock-wave therapy (2 patients), immobilisation cast (2 patients), Polidocanol injection (1 patient), platelet-rich plasma injection (1 patient), glyceryl trinitrate patches (1 patient), acupuncture (1 patient), non-steroidal anti-inflammatory drugs (1 patient) and rest for 2 years (1 patient). Of the group of patients treated with an alternative treatment, 21.4% was completely pain-free. In the group of patients who did not receive an alternative treatment, 56.7% was completely pain-free. After completing the 3 months heel-drop programme in the original study, 67.2% (31 patients) never performed the eccentric exercises again. No correlation in pain status was found between patients who continued the eccentric exercises and those who did not (p=0.15).

### Ultrasonographic examination

The sagittal thickness decreased from 8.1 mm (SD 2.1) at baseline to 7.5 mm (SD 1.6) at the 5-year follow-up (n=50, p=0.051). At baseline, 58.8% had some degree of neovascularisation (Öhberg score 1 to 4+). After 5 years, 47% showed



**Figure 2** The VISA-A score in time.

neovascularisation during the ultrasonographic examination (n=51). There was no difference in increase in VISA-A score between the patients with neovessels at baseline (increase of 32.1 points, SD 22.1) and patients without neovessels at baseline (increase of 32.1 points, SD 23.7).

Fifty-four per cent of the patients without pain at follow-up had some degree of neovascularisation. No significant difference was found in tendon thickness at 5-year follow-up between the patients with and without symptoms (p=0.59).

### Factors influencing the VISA-A score

The VISA-A score at 5-year follow-up and the  $\Delta$ VISA-A (= difference between baseline and the 5-year follow-up) score were not influenced by age (p=0.61 resp. p=0.18), sex (p=0.50 resp. p=0.39), body mass index (p=0.83 resp. p=0.50), duration of symptoms at baseline (p=0.11 resp. p=0.27), degree of neovascularisation at baseline (p=0.53 resp. p=0.81) and sagittal tendon thickness at baseline (p=0.43 resp. p=0.39).

### DISCUSSION

This is the first study evaluating the 5-years results of the Alfredson heel-drop programme with the validated VISA-A score. The VISA-A score increased significantly from 49.2 at baseline to 83.6 at the 5-year follow-up and from the 1-year follow-up to 5-year follow-up from 75.0 to 83.4 (p<0.01).

Until now the VISA-A score has only been used in one long-term follow-up study on eccentric exercises in Achilles tendinopathy.<sup>16</sup> Silbernagel *et al* reported a VISA-A score of 90.5 at 5-year follow-up, 65% of symptom-free patients and approximately 6% receiving alternative treatments. In their study, the training programme lasted from 12 weeks to 6 months and was performed under the supervision of a physiotherapist. Patients in the current study received very accurate instruction for the heel-drop programme and were instructed to do the exercises with pain. The compliance for the exercises was good/excellent in the majority of the patients (>70%).<sup>13</sup> The basic population of Silbernagel *et al*

showed many similarities with our population, however, no competitive athletes were included in the study of Silbernagel *et al.* Maffulli *et al.*<sup>20</sup> suggested there could be a difference in response to eccentric training in the Scandinavian population. Several Scandinavian studies have demonstrated good/excellent short-term results from the eccentric exercises in the treatment of chronic Achilles tendinopathy.<sup>9–12</sup> In recent years, differing results have been found by non-Scandinavian studies showing only moderate short-term results.<sup>13 21 22</sup> The long-term Scandinavian studies evaluating eccentric exercises in chronic Achilles tendinopathy suggest a good long-term prognosis.<sup>14–16</sup> Further studies will be needed to reveal if there is a genetic explanation for this difference.<sup>23</sup>

Two other studies have reported long-term effects of the heel-drop programme used in this study in patients with Achilles tendinopathy.<sup>14 15</sup> However, neither of them used the validated VISA-A score. Ohberg *et al.* showed that 22 of 25 (88%) patients were satisfied with the treatment at a mean follow-up of 3.8 years (range 1.6–7.75). The second long-term follow-up study reported that 65% of the patients had no or mild pain after 4.2 years (range 29–58 months).<sup>14</sup> In our study, 39.7% of the study population was completely pain-free at the 5-year follow-up. Almost half of our patients had received one or more alternative treatments. These secondary subjective outcomes suggest a discrepancy with the significant increase of the primary outcome score. For advice to patients it implies that a quantified improvement can be expected, but mild remaining pain symptoms may be present at long term.

Ultrasonographic evaluation demonstrated a comparable decrease in tendon thickness from 8.8 mm before the heel-drop programme to 7.6 mm at follow-up. However, the clinical relevance of this minor decrease is questionable. In recent years, a debate has arisen about the meaning and the significance of neovascularisation in Achilles tendons. Some studies have found a correlation between the patient's symptoms and the degree of neovascularisation,<sup>24–26</sup> and others found that the presence of neovascularisation did not per se indicate a disease state.<sup>27–31</sup> Also the circumstances during the examination, for example, activity just before the measurement, was not controlled and is unknown. This study found that the VISA-A score was not related to the degree of neovascularisation. Further research is needed to understand the relevance of neovascularisation in Achilles tendons.

This study has some limitations. The 11 dropouts might have influenced the outcome. In a best case scenario analysis where the dropouts have the best outcome score, 50% of the patients would be pain-free. However, the value of this non-validated subjective outcome measurement is questionable. Patients had to answer questions concerning the last 5 years, which may have given recall bias. The long-term outcome could be influenced by the alternative treatments that the patients received after completion of the heel-drop programme. However, of the patients who did not receive alternative treatments, 56.7% were completely pain-free after 5 years. Robinson *et al.*<sup>17</sup> recommended to use the VISA-A score in a homogenous group of athletes. Especially at long-term follow-up, the VISA-A score questionnaire can be subject to other influences than solely the symptoms of the Achilles tendinopathy. Factors like general ageing and new comorbidity may have influenced the outcome in this study. Finally, the ultrasonographic examinations at baseline and at follow-up were performed by different observers. However, in an earlier study, an excellent interobserver reliability of grading the neovascularisation was found.<sup>32</sup>

## What this study adds

Patients with chronic Achilles tendinopathy treated with Alfredson's heel-drop exercise programme continue to improve, as measured by the VISA-A score, up to 5 years after completion of the programme.

## CONCLUSION

This is the first study evaluating the 5-year results of the widely used heel-drop exercise programme as described by Alfredson *et al.* with the validated VISA-A score. At 5-year follow-up, the VISA-A score increased significantly.

After the 3-month eccentric training programme, almost half of the patients had received other therapies. Although improvement of symptoms can be expected at long term, mild pain may remain.

**Contributors** Study concept and design: RJdV, JLT, AW, JANV. Acquisition of data: AvdP. Analysis and interpretation of data: AvdP, SdJ, RJdV, JLT, HJLvdH, JANV. Drafting of the manuscript: AvdP, SdJ, JLT, RJdV. Critical revision of the manuscript for important intellectual content: AvdP, SdJ, JLT, RJdV, AW, HJLvdH, JANV. Statistical analysis: AvdP, HJLvdH. Administrative, technical or material support: AvdP, SdJ, RJdV, JLT, AW. Study supervision: RJdV, JLT, HJLvdH, JANV.

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**Competing interests** None.

**Ethics approval** The study was approved by the Medical Ethics Committee of the Hospital.

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