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# What is the longitudinal profile of impairments and can we predict difficulty caring for the profoundly-affected arm in the first year post-stroke?

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1 **What is the longitudinal profile of impairments and can we predict difficulty caring for**  
2 **the profoundly-affected arm in the first year post-stroke?**

3 **Abstract**

4 Objective: To establish the longitudinal profile of impairments of body functions and activity  
5 limitations of the arm, and evaluate potential predictors of difficulty caring for the  
6 profoundly-affected arm post-stroke.

7 Design: Prospective cohort study.

8 Setting: Three UK stroke services.

9 Participants: People unlikely to regain functional use of the arm (N=155) were recruited at 2-  
10 4 weeks post-stroke, and followed up at 3, 6 and 12 months. Potential predictors at baseline  
11 were hypertonicity, pain, motor control, mood, sensation/perception, age and stroke severity.

12 Interventions: NA

13 Main Outcome Measures: Difficulty caring for the arm (LASIS), pain, hypertonicity, range of  
14 movement, arm function and skin integrity. Multi-variable linear regression identified the  
15 best fitting model for predicting LASIS at 12 months.

16 Results: One hundred and ten participants (71%) were reviewed at one year. There was a  
17 large variation in the profile of arm functions and activity limitations. Inability or severe  
18 difficulty caring for the arm affected 29% of participants. Hypertonicity developed in 77%,  
19 with severe hypertonicity present in 25%. Pain was reported by 65%, 94% developed  
20 shoulder contracture and 6% had macerated skin. Difficulty caring for the arm increased with  
21 age, greater level of hypertonicity and stroke classification; collectively these factors  
22 accounted for 33% of the variance in LASIS.

23 Conclusions: At one year post-stroke, there was a high incidence of impairments of body  
24 functions and activity limitations in people with a profoundly-affected arm. Individual

25 profiles were very variable and although some pre-disposing factors have been identified, it  
26 remains difficult to predict who is at greatest risk.

27 **Key words:** stroke, upper limb, spasticity, pain, contracture

28 **List of abbreviations:**

29 HAEM – Haemorrhage stroke

30 LACS lacunar stroke on Oxford classification

31 LASIS Leeds Arm Spasticity Impact Scale

32 MAL-14 Motor Activity log

33 MMAS Modified modified Ashworth Scale

34 PACS – Partial anterior circulation stroke on Oxford classification

35 POCS – Posterior circulation ischaemic stroke on Oxford classification

36 Q1, Q3 First quartile and third quartile of the inter quartile range

37 TACS total anterior circulation stroke on Oxford classification

38 Three quarters of people with stroke will experience arm weakness, and 62% of these will not  
39 recover dexterity at six months<sup>1</sup>. For the purposes of this research, the term 'profoundly  
40 affected arm' is used to describe the situation where a stroke survivor has no movement in the  
41 affected arm or when movement is not functionally useful<sup>2</sup>. While current physical therapies  
42 in stroke rehabilitation are based predominantly on exercise and task-specific training<sup>3,4</sup>,  
43 additional therapy and practice of tasks does not improve active function in those with the  
44 most significant weakness<sup>5</sup>. Hence for those most unlikely to regain active function, a focus  
45 on managing activity limitations and avoiding secondary complications may be more  
46 appropriate. This approach involves maintaining the ability to care for the arm including tasks  
47 such as hand-washing and nail-cutting (i.e. passive function activities<sup>6</sup> which may be  
48 conducted by the person themselves or their carer).

49 Previous research shows that hypertonicity is present as early as one week post-stroke<sup>7</sup> and  
50 affects up to 47% of survivors<sup>8</sup>. Pain can also occur within one week<sup>9</sup> with an incidence up to  
51 49%<sup>10</sup>. Contracture is apparent by two weeks and affects 50%<sup>11</sup>. Previously reported  
52 predictors of hypertonicity include reduced motor control<sup>7,8</sup>, and increased stroke severity<sup>8,12</sup>.  
53 The most common predictors of pain are reduced sensation<sup>13,14</sup>, and weakness<sup>13</sup>. The  
54 significance of depression is not clear, with some studies identifying a positive link with  
55 pain<sup>15</sup>, and others discounting this<sup>8</sup>. Contracture is most frequently predicted by  
56 weakness<sup>16,17</sup>. However, this previous research is limited as all of these studies have been  
57 conducted on general populations of stroke survivors and not targeted at those with the most  
58 significant weakness. Furthermore, none of these previous studies has evaluated the profile or  
59 potential predictors of difficulty caring for the arm after stroke in a systematic way<sup>2</sup>.

60 Despite the high proportion of people with a profoundly-affected arm post-stroke (62%<sup>1</sup>),  
61 there is currently no targeted research on (1) the longitudinal profile of activity limitation in  
62 caring for the arm, (2) the proportion of people who develop associated impairments of body

63 functions and (3) the relationship between initial clinical findings and subsequent difficulty  
64 caring for the arm. The aim of this study was to establish the longitudinal profile of  
65 impairment of body functions and activity limitation in people with a profoundly-affected  
66 arm, and evaluate potential predictors of difficulty caring for the arm, in the first year post-  
67 stroke.

## 68 **Methods**

### 69 **Participants**

70 In a prospective, longitudinal study, all adult patients with first or subsequent stroke admitted  
71 to three stroke units in the UK over 30 months from September 2011, and still under the care  
72 of the stroke team at 2 weeks post-stroke, were screened for inclusion. Criteria included  
73 stroke within the past 2-4 weeks and a Fugl-Meyer upper extremity score of equal to or less  
74 than 11 points at 2 weeks, or 15 points at 3 weeks, or 19 points at 4 weeks post-stroke. These  
75 scores are strongly associated with high probability of not regaining function in the arm<sup>1</sup>.  
76 Patients who were unable to use their arm before the stroke were excluded. Potential  
77 participants were assessed for their ability to consent using the Mental Capacity Act<sup>18</sup>. Those  
78 with capacity were asked for their consent to participate. If the potential participant was  
79 judged by the researcher not to have capacity to make this decision, a consultee was  
80 approached if available. A consultee is someone who knows the person well but is not acting  
81 in a professional capacity, who can consider the persons beliefs and provide assent on their  
82 behalf if this is in line with their interests<sup>19</sup>.

83 Participants' baseline data were collected at the point of consent and at 3, 6, and 12 months  
84 later in the setting of their choice. These time scales allow comparison with previous  
85 studies<sup>7,8,9,10</sup>. Throughout the study, all participants received usual care under the UK NHS.

### 86 **Baseline predictor variables and demographic variables**

87 Five potential predictors of difficulty caring for the arm and related impairments were  
88 identified<sup>2</sup>: motor control, mood, sensation/perception, hypertonicity and pain. As the  
89 primary outcome related to passive care activities, hand dominance was not considered as a  
90 predictor. To maximise inclusivity wherever possible the measures used were suitable for  
91 people with aphasia or cognitive impairment. This included using pictographic resources,  
92 observational tools and measures with evidence of validity when completed by proxy. The  
93 predictor measures are summarised in Table 1<sup>20,21,22,23,24,25,26,27</sup>. Scores for hypertonicity with  
94 the Modified Modified Ashworth Scale (MMAS)<sup>22</sup> were applied to the five arm muscles  
95 identified as commonly affected (i.e. shoulder adductors and internal rotators, and elbow,  
96 wrist and finger flexors)<sup>8</sup>. The single worst score of any muscle group (“*worst hypertonicity*”)  
97 and the summed score of hypertonicity in all five groups (“*total hypertonicity*”) were  
98 considered (independently) as predictors. Summary scores of this type have been developed  
99 and validated<sup>28</sup>.

100 In addition to these pre-specified predictors, demographic data were also collected including  
101 age, sex and type of stroke using the Oxfordshire Community Stroke Project Classification<sup>29</sup>.

### 102 **Outcome measures (3, 6 and 12 months post-stroke)**

103 The primary outcome measure was a scale of difficulty caring for the arm: the Leeds Arm  
104 Spasticity Impact Scale<sup>25</sup> (LASIS). This is an item bank of 12 tasks of caring for the arm  
105 including aspects of washing, nail-cutting and dressing. The participant rates each relevant  
106 task with degree of difficulty using a scale from 0 to 4, and scores are then averaged. Test-  
107 retest reliability has been established with a minimally detected change of 0.5<sup>30</sup>.

108 Secondary outcomes included passive range of movement, pain, hypertonicity, active  
109 function, and skin integrity. The measures are summarised in Table 1. A protocol for  
110 conducting the predictor and outcome measure assessments was developed and demonstrated  
111 a good degree of inter-rater reliability, with Kappa scores of 0.82 for MMAS scores, 1.0 for

112 pain, 0.8 for LASIS and 93% agreement for measuring range of movement to within 15  
113 degrees.

#### 114 **Statistical Analysis**

115 All statistical analyses were performed using the statistical programming language R<sup>31</sup>.

116 Summary statistics were produced. Where the data was normally distributed, means and  
117 standard deviations were used. Otherwise median and inter-quartile ranges were given.

118 Individual profile plots were constructed to visualise each participant's LASIS average across  
119 follow-up points. The linear association between each continuous predictor and LASIS  
120 average at 12 months was summarised using Pearson's correlation coefficient, whilst  
121 descriptive statistics for LASIS average at 12 months are presented for each level of each  
122 categorical predictor.

123 Multi-variable linear regression was used to identify models of predictors for LASIS average  
124 at 12 months post-stroke. For brevity, this paper reports only the overall best fitting model.

#### 125 **Sample size**

126 Sample sizes for multi-variable linear regression are based on the minimum  $R^2$  value of  
127 interest and the number of independent predictors. Whilst there were five potential predictors  
128 of interest, three are categorical: pain (three categories), sensation/perception (three  
129 categories) and hypertonicity (five categories), with two continuous predictors (Fugl-Meyer  
130 and mood scores). After recoding categorical predictors as indicator variables, as required  
131 for modelling, statistically, it may be considered that there are 10 possible explanatory  
132 variables/predictors. Assuming a significance level of 10%, a sample size of 120 participants  
133 was required to detect a medium effect size of 0.15 (which corresponds to  $R^2$  value of around  
134 13%) with 90% power. Based on previous studies<sup>32,33</sup>, it was estimated that there would be a  
135 potential drop off of 10% per measurement session. Therefore, the recruitment target was set

136 at 165 participants, with the aim of following-up at least 120 participants at the 12 months  
137 post-stroke time point.

### 138 **Ethics**

139 The study was approved by the NRES South West Ethics Committee (Reference:  
140 11/SW/0149).

### 141 **Results**

142 Figure 1 illustrates the process of recruitment and follow-up, including reasons for  
143 participants lost to follow-up: 833 people were screened for inclusion of which 216 (26%)  
144 fulfilled the inclusion criteria, and 155 gave consent or consultee assent to participate (72%  
145 of those eligible). At one year 110 participants (71%) were reviewed. Of the remaining 45  
146 participants, 6 declined reassessment, 33 had died and 6 were unavailable.

147 Participant demographic data at baseline and the predictor measures are summarised in Table  
148 2. The average age of participants was 74.7 years, with a higher proportion of women than  
149 men, and almost half of the participants had a total anterior circulation stroke (TACS). At  
150 baseline, 82.6% had already developed some hypertonicity, with 17.4% exhibiting pain and  
151 31.8% demonstrating impairment of sensation/perception. Outcome measures at each follow-  
152 up are summarised in Table 3 and briefly summarised below.

### 153 **Longitudinal profiles of difficulty caring for the arm**

154 LASIS outcomes were collected from 104 participants at all time points. The mean LASIS at  
155 3 and 6 months were similar (1.7 and 1.6 respectively) and by 12 months had increased to  
156 2.0. However, there was a large variation in the profiles of each participant's scores, as  
157 shown in the individual profile plots in Figure 2, with some showing increasing difficulty  
158 over time, some decreasing difficulty and some broader variation. At the 12 month time-point  
159 over half (59%) of participants reported no or little difficulty with care tasks but 12%

160 reported moderate difficulty and 29% indicated they either had a great deal of difficulty or  
161 were unable to perform tasks such as washing or dressing.

## 162 **Longitudinal profiles of related impairments of body functions and activity limitation**

### 163 *Active function*

164 As anticipated, the majority of participants had not recovered active use of the arm at 12  
165 months, with 73% scoring between 0 and 1 (inclusive) on the Motor activity log (MAL-14)  
166 and median values remaining at 0 across time points. However, fifteen participants (14%)  
167 regained some use of the arm (scoring two or more on MAL-14). The baseline characteristics  
168 of those who regained some use are shown in Table 4.

### 169 *Hypertonicity*

170 Individual profiles of hypertonicity were very variable over the three time-points, although  
171 median hypertonicity total score was 4.0 at all time points (see Table 3). Some participants  
172 showed trends for increasing hypertonicity over time, some decreasing and some with no  
173 discernible pattern. At one year 77% of survivors had developed some hypertonicity in at  
174 least one muscle group (MMAS score at least 1), with severe hypertonicity in at least one  
175 muscle group present in 25% of participants (MMAS score at least 3). The muscle groups  
176 most commonly affected by severe hypertonicity were elbow and wrist flexors (affecting  
177 14% of participants each), shoulder internal rotators (13%), finger flexors (10%) and shoulder  
178 adductors (6%).

### 179 *Pain*

180 Profiles of pain were also very variable within the group of participants, although a larger  
181 proportion reported pain at follow-up compared to baseline, when the vast majority (83%)  
182 were pain free. At 12 months, pain in some part of the arm was reported by 65% of  
183 participants.

### 184 *Range of movement*

185 Individual profiles of range of movement were variable over time at all the joints assessed,  
186 with some participants having increasing and some decreasing range between 3 and 12  
187 months. However, over the three time points, the mean range of movement, particularly at the  
188 shoulder and wrist, was less than would be expected in healthy older adults<sup>34</sup>. Range of  
189 movement in the fingers was less reduced. Table 3 includes range for the index finger  
190 proximal interphalangeal joint as an example. Other studies have defined contracture as the  
191 loss of at least 30% of the available range of movement<sup>17</sup>. Using these criteria, 94% of  
192 participants had developed shoulder contracture, 9% elbow contracture, 54% wrist  
193 contracture and 7% finger contracture at 1 year.

#### 194 *Skin integrity*

195 Seven participants (6%) developed macerated skin in the hand or elbow-crease at 12 months.  
196 None of the participants had broken skin at any point.

#### 197 **Predicting difficulty caring for the arm**

198 Table 5 summarises the bivariate relationships between LASIS average at 12 months and  
199 each of the predictors. There was evidence of a positive relationship between the LASIS  
200 average and age, hypertonicity total score and mood, although only the linear association  
201 between LASIS average and age was statistically significant. We used hypertonicity total  
202 scores in the best fitting model because they explained a greater percentage of variance for  
203 the LASIS average than worst hypertonicity.

204 The overall best fitting linear model was derived from the five pre-specified predictors and  
205 the four additional baseline variables. After the removal of three outliers, the final model was  
206 fitted to data from 106 participants and included age ( $p < 0.001$ ), hypertonicity total ( $p = 0.002$ )  
207 and stroke classification (participants who have suffered from lacunar stroke (LACS)<sup>29</sup>  
208 compared to (a) participants who have suffered from total anterior circulation stroke  
209 (TACS)<sup>29</sup> ( $p = 0.004$ ) and (b) participants who have suffered from a haemorrhage

210 (p=0.010))(see Table 6). Collectively, these three variables explained approximately one third  
211 (adjusted  $R^2=33\%$ ) of the variance in the LASIS average at 12 months. From the linear  
212 regression coefficients from this final best fitting model:

- 213 • A one year increase in age at baseline increases the LASIS average at 12 months by an  
214 average of 0.050 units (standard error (SE) 0.008);
- 215 • A one unit increase in hypertonicity total at baseline increases the LASIS average at 12  
216 months by an average of 0.109 units (SE 0.035);
- 217 • The mean LASIS average for the group of participants who had suffered from LACS was  
218 0.935 units (SE 0.314) lower than participants who had suffered from TACS and 0.962 units  
219 (SE 0.367) lower than the group of participants who had suffered from a haemorrhage.

## 220 **Discussion**

221 This is the first longitudinal study, to our knowledge, of people with a profoundly-affected  
222 arm after stroke. Whilst the sample included a high proportion of people with more severe  
223 classifications of stroke this was not surprising given the target population. Many studies  
224 restrict recruitment and do not involve people with severe communication or cognitive  
225 limitations but we have demonstrated it was possible to include them, by supporting them  
226 with enhanced communication resources or using proxies.

227 Given that participants were those with severe arm weakness at 2-4 weeks post-stroke,  
228 observable patterns between impairments and activity limitation were thought to be a  
229 possibility. However, this was not the case and longitudinal profiles of these factors were  
230 highly individual.

231 The incidence of impairments in the arm was high when compared to studies that have  
232 included general populations of stroke survivors. For example, 77% of our participants who  
233 had severe weakness at baseline presented with hypertonicity at one year compared to 49% of  
234 those who initially presented with milder weakness at baseline<sup>8</sup>. In addition, 65% of our

235 participants reported pain in the arm compared to 49% of a general population of stroke  
236 survivors reporting pain in any part of the body<sup>9</sup>. Incidence of contracture of the shoulder and  
237 wrist were also higher than that recorded in general populations of stroke survivors<sup>17</sup>,  
238 although this was not the case for the elbow. It is unclear why so many of our participants  
239 developed loss of range of the shoulder and wrist while the elbow and fingers remained less  
240 severely affected. The shoulder is typically held in adduction and internal rotation at rest so  
241 may be more vulnerable, while gravity may assist with extension of the elbow. The wrist is a  
242 complex joint and contracture of the finger flexor muscle-tendon units may impact on range  
243 of movement at the wrist in addition to the fingers. Differences in muscle architecture  
244 surrounding connective tissue may also contribute to the variation in contracture between  
245 muscles. Recent work in animal studies suggests that there is a direct relationship between  
246 muscle atrophy and fibrosis. Cytokine myostatin, for example, is not only central to the  
247 pathways that mediate muscle atrophy but can also activate fibroblasts and stimulate  
248 fibrosis<sup>35,36</sup>. Thus, differences in the weakness of individual muscles may impact on their  
249 relative degree of contracture development and increase in passive stiffness<sup>35,37</sup>.

250 The incidence of difficulty caring for the arm was high, as 29% of participants were either  
251 unable to care for their arm or described significant difficulty. A number of predictors of  
252 difficulty caring for the arm that can be assessed early after stroke were evaluated. The best  
253 linear predictive model based on these included age, hypertonicity and stroke classification,  
254 although these factors explained only 33% of the total variability in the LASIS average at one  
255 year post-stroke. Our previous review did not identify other impairments that are likely to  
256 influence longer-term outcome in caring for the arm<sup>2</sup>. Previous studies have not considered  
257 the use of biomarkers as predictors of outcome in this targeted group and it is possible they  
258 may add to the predictive value.

259 There are a number of clinical implications of this work. Whilst recognising that people with  
260 profoundly-affected arm gain little from active exercise to improve function<sup>4</sup>, given the high  
261 incidence of pain, hypertonicity and contracture they may benefit from an educational  
262 intervention to reduce the impact of these impairments, and from longer term monitoring.  
263 With regard to the important risk factors identified, age and stroke classification cannot be  
264 influenced in treatment after stroke but it is possible that early manifestations of hypertonicity  
265 can be altered and research could explore if targeting hypertonicity early after stroke can  
266 reduce the risk of difficulty caring for the arm longer term, particularly in those with other  
267 risk factors.

### 268 **Study limitations**

269 This study has a number of limitations. Whilst every attempt was made to include measures  
270 that had been validated in people with aphasia and cognitive impairment, this was not always  
271 possible. The Fugl-Meyer score, in particular, has not been validated in this group. Equally  
272 the anticipated sample size of 120 participants at 1 year was not achieved so adequate  
273 statistical power may be lacking. Hand dominance was not considered as a predictor variable  
274 and this may have an influence in self-care. Finally, no attempt was made to assess the  
275 amount or content of rehabilitation that participants received so it is possible that any such  
276 interventions may have impacted on outcomes. Therefore conclusions should be drawn with  
277 caution.

### 278 **Conclusions**

279 At one year post-stroke, there was a high incidence of difficulty caring for the arm (measured  
280 with LASIS) and of pain, hypertonicity and contracture. Notably, individual profiles were  
281 very variable and although some pre-disposing factors have been identified, it remains  
282 difficult to predict who is at greatest risk.

283 **References**

- 284 1. Kwakkel G, Kollen BJ, van der Grond J, Prevo AJH. Probability of regaining dexterity in  
285 the flaccid upper limb: Impact of severity of paresis and time since onset in acute stroke.  
286 *Stroke*. 2003;34:2181-2186.
- 287 2. Allison R, Shenton L, Bamforth K, Kilbride C, Richards D. Incidence, time course and  
288 predictors of impairments relating to caring for the profoundly affected arm after stroke: A  
289 systematic review. *Physiotherapy Research International*. 2015;21:210-227.
- 290 3. Royal Dutch Society for Physical Therapy. KNGF Guideline: Stroke. [http://www.fysionet-](http://www.fysionet-evidencebased.nl/images/pdfs/guidelines_in_english/stroke_practice_guidelines_2014.pdf)  
291 [evidencebased.nl/images/pdfs/guidelines\\_in\\_english/stroke\\_practice\\_guidelines\\_2014.pdf](http://www.fysionet-evidencebased.nl/images/pdfs/guidelines_in_english/stroke_practice_guidelines_2014.pdf).  
292 2014. Accessed January 11 2016.
- 293 4. Intercollegiate Stroke Working Party. National Clinical Guideline for Stroke. 5th ed.  
294 London, UK: Royal College of Physicians; 2016.
- 295 5. Parry RH, Lincoln NB, Vass CD. Effect of severity of arm impairment on response to  
296 additional physiotherapy early after stroke. *Clinical Rehabilitation*. 1999;13:187-198.
- 297 6. Sheean G. Botulinum treatment of spasticity: Why is it difficult to show a functional  
298 benefit? *Current Opinion in Neurology*. 2001;4:771-6.
- 299 7. De Jong LD, Hoonhorst MH, Stuive I, Dijkstra PU. Arm motor control as predictor for  
300 hypertonia after stroke: a prospective cohort study. *Archives of Physical Medicine and*  
301 *Rehabilitation* 2011; 92: 1411-1417.
- 302 8. Kong KH, Lee J, Chua KS. Occurrence and temporal evolution of upper limb spasticity in  
303 stroke patients admitted to a rehabilitation unit. *Archives of Physical Medicine and*  
304 *Rehabilitation*. 2012;93:143-148.
- 305 9. Ratnasabapathy Y, Broad J, Baskett J, Pledger M, Marshall J, Bonita R. Shoulder pain in  
306 people with a stroke: a population based study. *Clinical Rehabilitation* 2003; 17: 304-311.

- 307 10. Lundstrom E, Smits A, Terent A, Borg J. Risk factors for stroke-related pain 1 year after  
308 first-ever stroke. *European Journal of Neurology*. 2009;6:188–193.
- 309 11. Ada L, O'Dwyer N, O'Neill E. Relation between spasticity, weakness and contracture of  
310 the elbow flexors and upper limb activity after stroke: an observational study. *Disability and*  
311 *Rehabilitation* 2006; 28: 891–889.
- 312 12. Lundström E, Smits A, Terent A, Borg J. Time-course and determinants of spasticity  
313 during the first six months following first-ever stroke. *Journal of Rehabilitation*  
314 *Medicine*. 2010;42:296–301.
- 315 13. Gamble GE, Barberan E, Bowsher D, Tyrrell PJ, Jones AK. Post stroke shoulder pain:  
316 more common than previously realized. *European Journal of Pain*. 2000;4:313–315.
- 317 14. Sommerfeld DK, Welmer AK. Pain following stroke, initially and at 3 and 18 months  
318 after stroke, and it's association with other disabilities. *European Journal of Neurology*.  
319 2012;19:1325–1330.
- 320 15. O'Donnell MJ, Diener HC, Sacco RL, Panju AA, Vinisko R, Yusuf S, on behalf of the  
321 PRoFESS Investigators. Chronic pain syndromes after ischaemic stroke: PRo-FESS trial.  
322 *Stroke*. 2013;44:1238–1243.
- 323 16. Pandyan AD, Cameron M, Powell J, Stott DJ, Granat MH. Contractures in the post-stroke  
324 wrist: a pilot study of its time course of development and its association with upper limb  
325 recovery. *Clinical Rehabilitation*. 2003;17:88–95.
- 326 17. Kwah LK, Harvey LA, Diong JHL, Herbert RD. Half of the adults who present to  
327 hospital with stroke develop at least one contracture within six months: an observational  
328 study. *Journal of Physiotherapy*. 2012; 58:41–47.
- 329 18. Department of Health. Mental capacity act. London, UK: Department of Health; 2005.  
330 Retrieved from <http://www.legislation.gov.uk/ukpga/2005/9/section/3!>
- 331 19. Department of Health. Guidance on nominating a consultee for

332 research involving adults who lack capacity to consent. London, UK: Department of Health;  
333 2008.

334 20. Hsieh YW, Wu CY, Lin KC, Chang YF, Chen CL, Liu JS. Responsiveness and validity  
335 of three outcome measures of motor function after stroke rehabilitation. *Stroke*.  
336 2009;40:1386-1391.

337 21. Paci M, Nannetti L, Taiti P, Baccini M, Pasquini J, Rinaldi L. Shoulder subluxation after  
338 stroke: relationships with pain and motor recovery. *Physiotherapy Research International*.  
339 2007;12:95-104.

340 22. Ansari NN, Naghdi S, Hasson S, Mousakhani A, Nouriyani A, Omidvar Z. Inter-rater  
341 reliability of the Modified Modified Ashworth Scale as a clinical tool in measurements of  
342 post-stroke elbow flexor spasticity. *NeuroRehabilitation*. 2009;24:225-229.

343 23. Prescott RJ, Garraway WM, Akhtar AJ. Predicting functional outcome following acute  
344 stroke using a standard clinical examination. *Stroke*. 1982;13:641-647.

345 24. Leeds L, Meera RJ, Hobson JF. The utility of the Stroke Aphasic Depression  
346 Questionnaire in a stroke rehabilitation unit. *Clinical Rehabilitation*. 2004;18:228-31

347 25. Royal College Physicians. *Spasticity in adults: Management using botulinum toxin:*  
348 *National Guidelines*. London, UK:Royal College of Physicians; 2009.

349 26. Andrews AW, Bohannon RW. Decreased shoulder range of motion on paretic side after  
350 stroke. *Physical Therapy*. 1989;69:768-772.

351 27. Uswatte G, Taub E, Morris D, Vignolo M, McCulloch K. Reliability and validity of the  
352 upper-extremity motor activity log-14 for measuring real-world arm use. *Stroke*.  
353 2005;36:2496-2499.

354 28. Platz T, Vuadens P, Eickhof C, Arnold P, Van Kaick S, Heise K. REPAS, a summary  
355 rating scale for resistance to passive movement: item selection, reliability and validity.  
356 *Disability and Rehabilitation*. 2008;30:44-53.

- 357 29. Bamford J, Sandercock P, Dennis M, Burn J, Warlow C. Classification and natural  
358 history of clinically identifiable subtypes of cerebral infarction. *The Lancet*. 1991;337:1521-  
359 6.
- 360 30. O'Reilly F, Meldrum D. Test re-test reliability of the Modified Leeds Arm Spasticity  
361 Impact Scale (MLASIS) and the minimum detectable change (MDC) scores. *Irish Journal of*  
362 *Medical Science*. 2013;182:273.
- 363 31. R Core Team. *R: A language and environment for statistical computing*. R Foundation  
364 *for Statistical Computing*. <https://www.R-project.org/>. 2015. Accessed October 10 2015.
- 365 32. Sackley C, Brittle N, Patel S, Ellins J, Scott M, Wright C et al. The prevalence of joint  
366 contractures, pressure sores, painful shoulder, other pain, falls, and depression in the year  
367 after a severely disabling stroke. *Stroke*. 2008;39:3329-3334.
- 368 33. Leathley MJ, Gregson JM, Moore AP, Smith TL, Sharma AK, Watkins CL. Predicting  
369 spasticity after stroke in those surviving to 12 months. *Clinical Rehabilitation*. 2004;18:438-  
370 43.
- 371 34. Kalscheur JA, Emery LJ, Costello PS. Range of motion in older women. *Physical &*  
372 *Occupational Therapy In Geriatrics*. 1999;16:77-96.
- 373 35. Li ZB, Kollias HD, Wagner KR. Myostatin directly regulates skeletal muscle fibrosis. *J*  
374 *Biol Chem*. 2008;283:19371-8.
- 375 36. Desgeorges, M.M., et al., Molecular mechanisms of skeletal muscle atrophy in a mouse  
376 model of cerebral ischemia. *Stroke*, 2015. 46(6): p. 1673-80.
- 377 37. Bo Li Z, Zhang J, Wagner KR. Inhibition of myostatin reverses muscle fibrosis through  
378 apoptosis. *J Cell Sci*. 2012;125:3957-65.

379 Figure legends:

380 Figure 1: Flow diagram detailing recruitment and progression of participants

381 Figure 2 Individual participants LASIS scores at each time point. Each box contains 5  
382 participants in their order of recruitment- where data is missing the participant was lost to  
383 follow up.. (N=127 at 3 months, N= 117 at 6 months, N= 111 at 12 months).

384

385 **Table 1: Battery of predictor and outcome measures**

<b>Predictors</b>	<b>Name</b>	<b>Scoring</b>
Motor control	Fugl-Meyer Upper limb score <sup>14</sup>	0-66, higher score indicates better control
Pain	Yes/no response to pain at rest and on passive movement <sup>15</sup>	0,1 or 2, higher score indicates more pain
Hypertonicity	Modified Modified Ashworth scale <sup>16</sup>	0-4, higher score indicates higher tone
Perception/ sensation	Find the thumb test <sup>17</sup>	0,1 or 2, higher score indicates worse perception
Mood	Stroke Aphasic Depression Questionnaire-10 <sup>18</sup>	0-30, higher score indicates lower mood
<b>Outcomes</b>		
Difficulty caring for arm	Leeds Arm Spasticity Impact Scale <sup>19</sup>	0-4, higher score indicates more difficulty
Pain	As above	
Hypertonicity	As above	
Passive range of movement	Goniometry of shoulder flexion, abduction, external rotation; elbow flexion, extension; wrist extension, index, little finger and thumb extension at each joint <sup>20</sup>	Range measured in degrees of movement
Skin integrity	Axilla, elbow and hand, classified as dry/ intact; macerated or broken.	0,1 or 2, higher score indicates worse skin condition
Active arm	Motor activity log-14 <sup>21</sup>	0-70, higher score indicates better

function		use
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388 **Table 2: Descriptive statistics of participant characteristics and potential predictors at**  
 389 **baseline (n=155)**

<b>Age</b>	
Mean (SD)	74.7 (12.8)
Range	[38.0, 96.0]
<b>Sex</b>	
Female, n (%)	89 (57%)
Male, n (%)	66 (43%)
<b>Stroke classification</b>	
Not reported, n (%)	1 (0.6%)
Haemorrhage, n (%)	25 (16.2%)
Total anterior circulation stroke, n (%)	73 (47.4%)
Partial anterior circulation stroke, n (%)	30 (19.5%)
Lacunar stroke, n (%)	23 (14.9%)
Posterior circulation stroke, n (%)	3 (1.9%)
<b>Fugl-Meyer upper limb score</b>	
Median [Q1, Q3]	2.0 [2.0, 6.0]
Range	[0.0, 16.0]
<b>Hypertonicity (worse MMAS score)</b>	
0, n (%)	27 (17.4%)
1, n (%)	52 (33.5%)
2, n (%)	53 (34.2%)
3, n (%)	23 (14.8%)
4, n (%)	0 (0%)
<b>Hypertonicity (total score)</b>	

Median [Q1, Q3]	3.0 [1.0, 5.0]
Range	[0.0, 12.0]
<b>Pain</b>	
No pain at rest or movement, n (%)	128 (82.6%)
Pain on movement only, n (%)	21 (13.5%)
Pain at rest & on movement, n (%)	6 (3.9%)
<b>Mood (SADQ-10)</b>	
Mean (SD)	8.2 (4.5)
Range	[0.0, 23.0]
<b>Sensation/Perception (Find the Thumb test)</b>	
Not reported, n (%)	1 (0.6%)
Able to find affected thumb, n (%)	105 (68.2%)
Able to find affected arm only, n (%)	19 (12.3%)
Unable to find affected arm, n (%)	30 (19.5%)

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392 **Table 3: Descriptive statistics of outcome measures at each follow-up time point**

	3 months n=120	6 months n=113	12 months n=110
<b>LASIS average</b>			
Mean (SD)	1.7 (1.0)	1.6 (1.1)	2.0 (1.3)
Range	[0.08, 4.00]	[0.00, 4.00]	[0.00, 4.00]
<b>Hypertonicity (worse score)</b>			
Not reported, n (%)	2 (1.7%)	4 (3.5%)	1 (0.9%)
0, n (%)	16 (13.6%)	19 (17.4%)	24 (22%)
1, n (%)	31 (26.3%)	33 (30.3%)	29 (26.4%)
2, n (%)	52 (44.1%)	32 (29.4%)	29 (26.4%)
3, n (%)	19 (16.1%)	22 (20.2%)	24 (22%)
4, n (%)	0 (0%)	3 (2.8%)	3 (2.7%)
<b>Hypertonicity (total score)</b>			
Median [Q1, Q3]	4.0 [2.0, 8.0]	4.0 [1.0, 9.0]	4.0 [1.0, 7.8]
Range	[0.0, 15.0]	[0.0, 16.0]	[0.0, 15.0]
<b>Pain</b>			
No pain at rest or movement, n (%)	32 (26.7%)	31 (27.4%)	38 (34.5%)
Pain on movement only, n (%)	60 (50%)	56 (49.6%)	56 (50.9%)
Pain at rest & on movement, n (%)	28 (23.3%)	26 (23.0%)	16 (14.5%)
<b>Passive range shoulder abduction</b>			
Mean (SD)	76.8° (24.4°)	74.3° (22.6°)	79.9° (28.8°)
Range	[25.0°, 170.0°]	[10.0°, 160.0°]	[20.0°, 180.0°]
<b>Passive range shoulder external rotation</b>			
	22.8° (24.8°)	24.8° (22.7°)	25.0° (27.4°)

Mean (SD)	[-80°, 65.0°]	[-70.0°, 75.0°]	[-60.0°, 90.0°]
Range			
<b>Passive range elbow extension</b>			
Mean (SD)	165.0° (20.0°)	165.0° (20.8°)	164.8° (21.0°)
Range	[100.0°, 180.0°]	[90.0°, 180.0°]	[100.0°, 180.0°]
<b>Passive range wrist extension</b>			
Mean (SD)	38.6° (21.6°)	37.7 (26.7°)	43.7° (26.6°)
Range	[-60.0°, 90.0°]	[-50.0°, 80.0°]	[-60.0°, 80.0°]
<b>Passive range index PIP extension</b>			
Mean (SD)	175.3° (11.5°)	172.1° (17.3°)	174.2° (16.3°)
Range	[100.0°, 180.0°]	[90.0°, 180.0°]	[100.0°, 180.0°]
<b>Skin integrity</b>			
Not reported, n (%)	0	2 (1.8%)	0
Dry intact, n (%)	118 (98.4%)	105 (92.9%)	103 (93.6%)
Macerated, n (%)	2 (1.6%)	6 (5.3%)	7 (6.4%)
Broken, n (%)	0	0	0
<b>Active function (MAL-14)</b>			
Median [Q1, Q3]	0.0 [0.0, 0.8]	0.1 [0.0, 1.0]	0 [0.0, 1.2]
Range	[0, 4.21]	[0.0, 5.00]	[0.0, 4.38]

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**Table 4: Descriptive statistics at baseline by functionality at 12 months (n=109)**

	Participants with MAL less than two at 12 months (n=94)	Participants with MAL two or greater at 12 months (n=15)
<b>Age</b>		
Mean (SD)	72.7 (12.9)	68.1 (13.7)
Range	[38.0, 95.0]	[40.0, 96.0]
<b>Gender</b>		
Female, n (%)	49 (52%)	9 (60%)
Male, n (%)	45 (48%)	6 (40%)
<b>Stroke classification</b>		
Not reported, n (%)	0 (0%)	0 (0%)
Haemorrhage, n (%)	19 (20.2%)	1 (6.7%)
Total anterior circulation stroke, n (%)	45 (47.9%)	6 (40.0%)
Partial anterior circulation stroke, n (%)	16 (17.0%)	4 (26.7%)
Lacunar stroke, n (%)	12 (12.8%)	4 (26.7%)
Posterior circulation stroke, n (%)	2 (2.1%)	0 (0%)
<b>Fugl-Meyer upper limb scores</b>		
Median [Q1, Q3]	2.0 [2.0, 5.0]	5.0 [4.0, 10.5]
Range	[0.0, 15.0]	[2.0, 15.0]
<b>Hypertonicity (worse MMAS score)</b>		
0, n (%)	16 (17.0%)	3 (20.0%)
1, n (%)	26 (27.7%)	6 (40.0%)
2, n (%)	38 (40.4%)	5 (33.3%)
3, n (%)	14 (14.9%)	1 (6.7%)
4, n (%)	0 (0%)	0 (0%)
<b>Hypertonicity (total score)</b>		
Median [Q1, Q3]	3.0 [1.0, 5.8]	2.0 [1.0, 3.0]
Range	[0.0, 12.0]	[0.0, 7.0]

<b>Pain</b>		
No pain at rest or movement, n (%)	77 (81.9%)	13 (86.7%)
Pain on movement only, n (%)	12 (12.8%)	1 (6.7%)
Pain at rest & on movement, n (%)	5 (5.3%)	1 (6.7%)
<b>Mood (SADQ-10)</b>		
Mean (SD)	8.3 (4.8)	6.1 (4.2)
Range	[0.0, 23.0]	[1.0, 16.0]
<b>Sensation/Perception (Find the Thumb test)</b>		
Not reported, n (%)	1 (1.1%)	0 (0%)
Able to find affected thumb, n (%)	68 (72.3%)	11 (73.3%)
Able to find affected arm only, n (%)	10 (10.6%)	2 (13.3%)
Unable to find affected arm, n (%)	15 (16.0%)	2 (13.3%)

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399 **Table 5: Summary statistics of bivariate relationships between predictors and LASIS**  
 400 **average at 12 months**

	Mean (SD) [range]	Correlation Coefficient (95% CI)
Age	-	0.39 [0.22, 0.54]
Sex		-
Female	2.2 (1.4) [0.00, 4.00]	
Male	1.7 (1.3) [0.00, 4.00]	
Stroke classification		-
Lacunar stroke (LACS)	1.4 (0.9) [0.00, 3.82]	
Partial anterior circulation stroke (PACS)	1.5 (1.3) [0.09, 4.00]	
Posterior circulation stroke (POCS)	0.9 (0.7) [0.42, 1.44]	
Total anterior circulation stroke (TACS)	2.3 (1.4) [0.00, 4.00]	
Haemorrhage	2.2 (1.3) [0.45, 4.00]	
Fugl-Meyer upper limb score	-	-0.18 [-0.36, 0.01]
Hypertonicity (worse MMAS score)		-
0	1.7 (1.3) [0.27, 4.00]	
1	1.9 (1.4) [0.00, 4.00]	
2	1.9 (1.3) [0.00, 4.00]	
3	2.8 (1.3) [0.55, 4.00]	
4	NA	
Hypertonicity (total score)	-	0.19 [0.00, 0.37]
Pain		-
No pain at rest or movement	1.9 (1.4) [0.00, 4.00]	
Pain on movement only	2.6 (1.3) [0.10, 4.00]	

Pain at rest & on movement	1.7 (0.6) [0.80, 2.27]	
Mood (SADQ-10)	-	0.19 [0.00, 0.36]
Sensation/Perception (Find the Thumb)		-
Able to find affected thumb	1.8 (1.3) [0.00, 4.00]	
Able to find affected arm only	2.3 (1.1) [0.25, 4.00]	
Unable to find affected arm	2.3 (1.5) [0.36, 4.00]	

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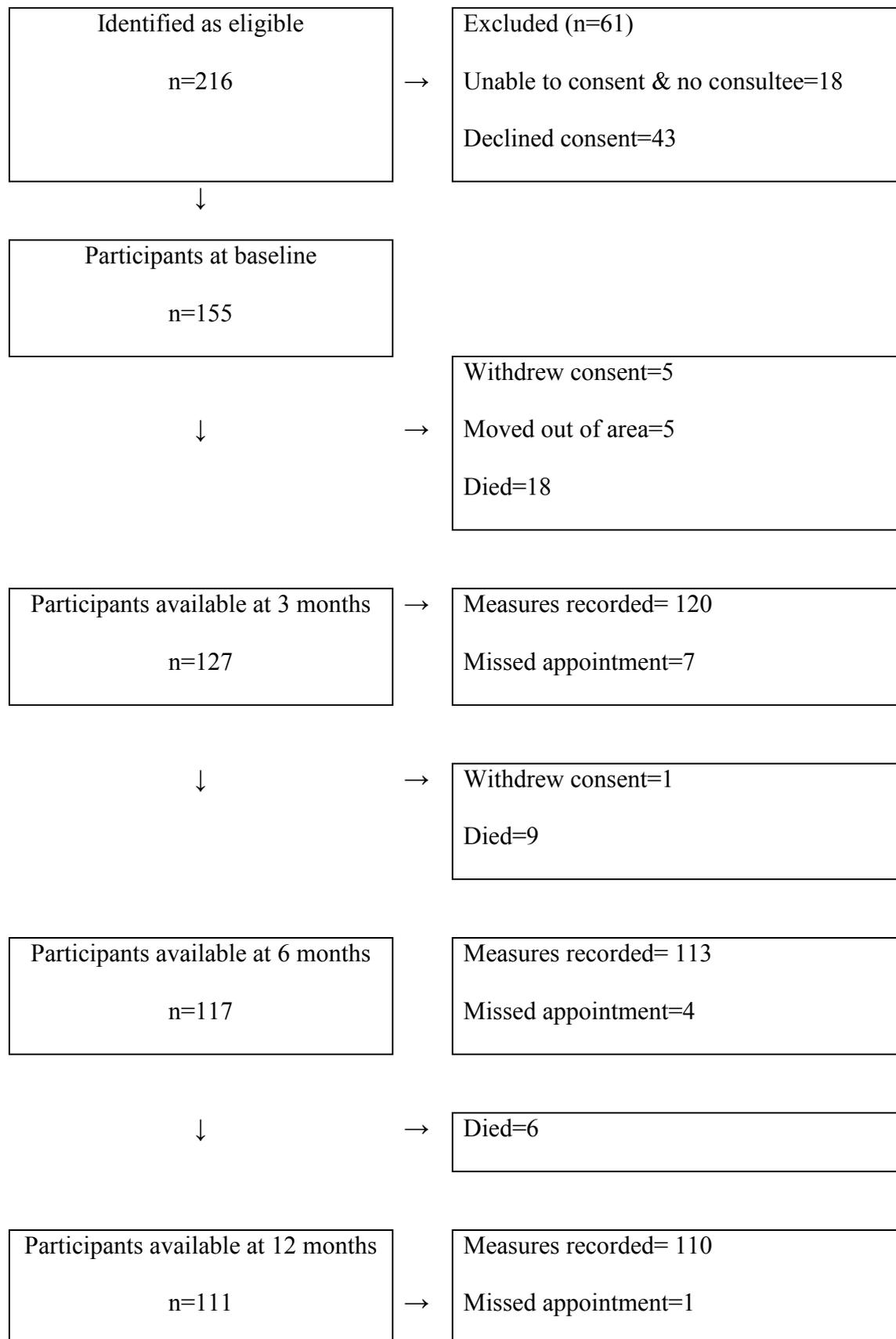
403 **Table 6: Regression statistics for the overall best fitting model for LASIS average at 12**  
404 **months**

	<b>Coefficient</b>	<b>95% confidence interval</b>	<b>p-value</b>
Intercept	-2.658	[-4.028, -1.288]	<0.001
Age	0.050	[0.034, 0.066]	<0.001
Hypertonicity total	0.109	[0.040, 0.178]	0.002
stroke class POCS	-0.200	[-1.809, 1.409]	0.808
stroke class PACS	0.121	[-0.608, 0.850]	0.744
stroke class TACS	0.935	[0.320, 1.550]	0.004
stroke class HAEM	0.962	[0.243, 1.681]	0.010
Residual standard error: 1.091 on 99 DF Multiple R-squared: 0.37 Adjusted R-squared: 0.33 F-statistic: 9.6 on 6 and 99 DF, p-value: <0.001			

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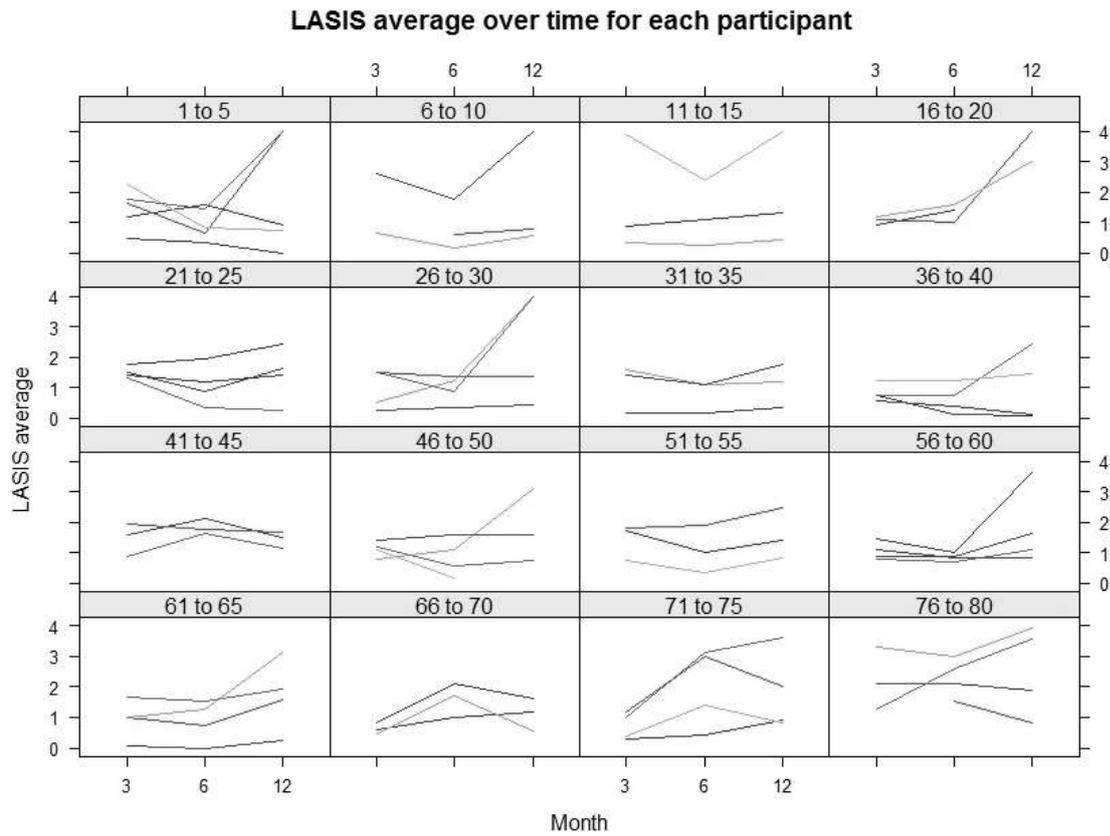
406 LACS (Lacunar stroke) is the baseline level for stroke classification.  
407

408 Figure 1  
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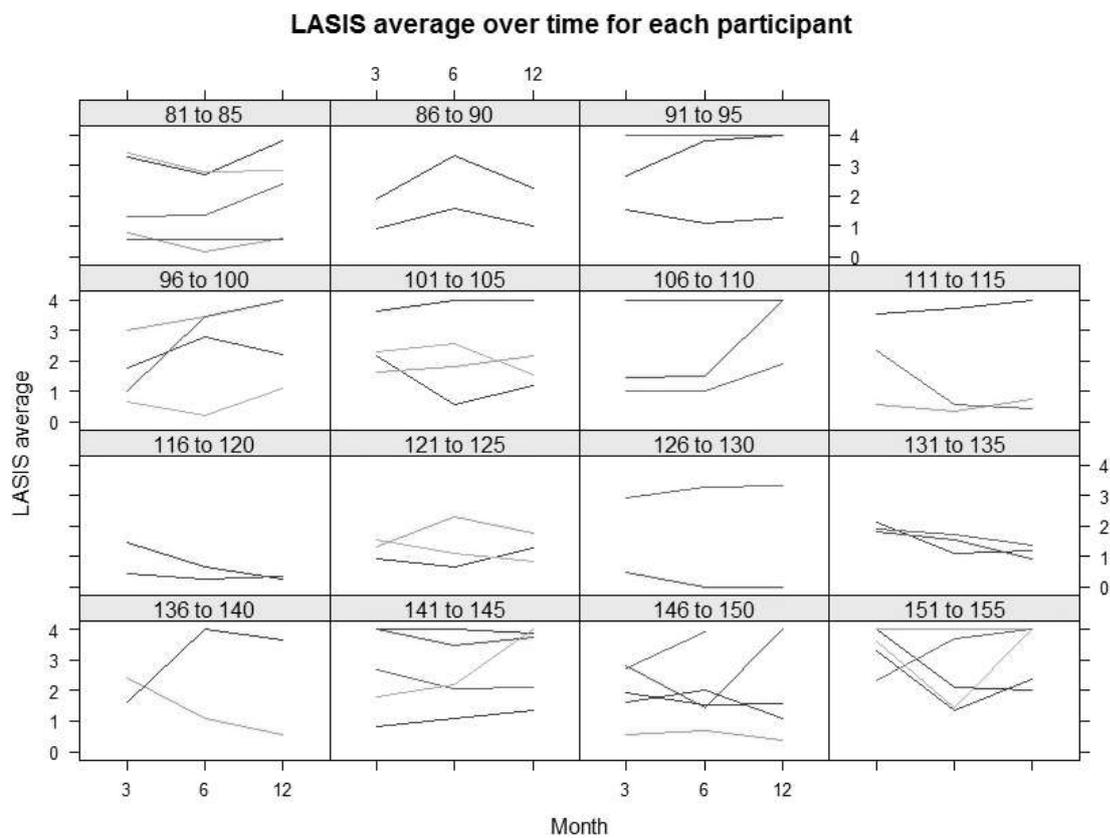


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412 Figure 2



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