

Title: Time-varying surface electromyography topography as a prognostic tool for chronic low back pain rehabilitation

Yong Hu^{*}, PhD, Jerry Weilun Kwok, MPhil, Jessica Yuk-Hang Tse, Keith Dip-Kei Luk, FRACS, FRCSEd, FRCS(Glas), FHKCOS, FHKAM(Ortho Surg),

Department of Orthopaedics and Traumatology, The University of Hong Kong, Pokfulam, Hong Kong

****** Correspondence Author

Dr Yong Hu

Dept. Of Orthopaedics and Traumatology,

The University of Hong Kong

Address: 12 Sandy Bay Road, Pokfulam, Hong Kong

Email address: yhud@hku.hk;

Tel: (852) 29740359; Fax: (852) 29740335

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1 **Abstract**

2

3 **BACKGROUND CONTEXT:** Non-surgical rehabilitation therapy is a commonly used
4 strategy to treat chronic low back pain (LBP). The selection of the most appropriate
5 therapeutic options is still a big challenge in clinical practices. Surface
6 electromyography(EMG) topography has been proposed to be an objective assessment of
7 LBP rehabilitation. The quantitative analysis of dynamic surface EMG would provide an
8 objective tool of prognosis for LBP rehabilitation.

9

10 **PURPOSE:** to evaluate the prognostic value of quantitative sEMG topographic analysis, and
11 to verify the accuracy of the performance of proposed time-varying topographic parameters
12 for identifying the patients who have better response towards the rehabilitation program.

13 **STUDY DESIGN:** A retrospective study of consecutive patients.

14 **PATIENT SAMPLE:** 38 patients with chronic non-specific LBP and 43 healthy subjects.

15

16 **OUTCOME MEASURES:** The accuracy of the time-varying quantitative sEMG topographic
17 analysis for monitoring LBP rehabilitation progress was determined by calculating the
18 corresponding ROC curves.

1

2 Physiologic Measure: sEMG during lumbar flexion and extension

3

4 METHODS: Patients who suffered from chronic non-specific LBP without the history of
5 previous back surgery and any medical conditions causing acute exacerbation of LBP during
6 the clinical test were enlisted to perform the clinical test during the 12-week physiotherapy
7 treatment. LBP patients were classified into two groups: "responding" and "non- responding"
8 based on the clinical assessment. The "responding" group referred to the LBP patients began
9 to recover after the physiotherapy treatment whereas the "non- responding" group referred to
10 some LBP patient who did not recover or got worse after the treatment. The results of the
11 time-varying analysis in the "responding" group were compared with those in the "non-
12 responding" group. In addition, the accuracy of the analysis was analyzed through ROC
13 curves.

14

15 RESULTS: The time-varying analysis showed discrepancies in the root-mean-square
16 difference (RMSD) parameters between the "responding" and "non- responding" group. The
17 RMSD RA and RMSD RW at flexion and extension in the "responding" group were
18 significantly lower than those in the "non- responding" group ($P < 0.05$). The areas under ROC
19 curves of RMSD RA and RMSD RW at flexion and extension were greater than 0.7 and were

1 statistically significant.

2

3 **CONCLUSIONS:**

4 The quantitative time-varying analysis of sEMG topography shows significant difference
5 between the healthy and LBP groups. The discrepancies in quantitative dynamic sEMG
6 topography of LBP group from normal group, in terms of RMSD RA and RMSD RW at
7 flexion and extension, are able to identify those LBP subjects who are responsive to
8 conservative rehabilitation program focused on functional restoration of lumbar muscle.

9

10 **Key words:** chronic low back pain (LBP), rehabilitation therapy, prognosis, Surface
11 electromyography, time-varying topography

12

13

14

15 **Introduction**

16

17 In the majority of persons with low back pain (LBP), a specific diagnosis cannot be made [1,
18 2]. Without knowledge of the underlying cause, finding an efficacious match between any
19 individual LBP patient and an almost infinite selection of therapeutic options is highly
20 problematic [2, 3]. Consequently, the resulting trial and error approach to matching patients
21 and treatment perpetuates the expense and prevalence of LBP [4-7].

22

23 While the various etiologies of LBP await discovery, investigators have attempted to
24 improve treatment efficacy by developing diagnosis-independent techniques to match LBP
25 patients to treatments that are likely to succeed [2, 8-10]. To date, several baseline variables
26 have been identified that predict which patients are likely to respond preferentially to a
27 specific therapeutic intervention. For example, Childs et al.[8] formulated a clinical
28 prediction rule based on a constellation of five variables (symptom duration, symptom
29 location, fear–avoidance beliefs, hip rotation range of motion and lumbar mobility). In
30 persons who were positive for 4 or more of the 5 prediction variables, the estimated
31 probability of treatment success using spinal manipulation was estimated at 92% of those
32 subjects [8].

33

34 Musculoskeletal dysfunction is one of the causes of LBP and surface EMG (sEMG) is
35 widely used in clinical experiments for biomechanical and musculoskeletal analysis. sEMG
36 has been renowned for being non-invasive and dynamic application, a gold standard for
37 measuring muscle function [4, 11-13]. With use of surface electrodes (sEMG), this painless
38 and easily applied technique has been used extensively to document muscle impairments [4,
39 12, 14]. The objective sEMG measurement of global muscle groups is potential to offer a
40 reliable reference for physiotherapy treatment of LBP and so to play a role as diagnostic and
41 monitoring tools. In the past few decades, many researchers have been working in
42 quantizing sEMG signal for LBP assessment, such as raw sEMG, median frequency, reflex
43 latency and positions of standing, trunk flexion/extension and sitting, etc[15-19]. Increasing
44 number of literature reports that there are significant differences in sEMG between the LBP
45 patients and the normal people which offer potential clinical application of sEMG for
46 diagnosis of LBP [13, 20-22].

47

48 Although sEMG is used commonly in the spine, interpretation of its results can be
49 problematic given the spine's multiple layers of overlapping muscles. As a result, several
50 investigators have developed spatial arrays of sEMG electrodes to describe regional muscle
51 activity rather than activity on a per muscle basis. From data derived in this way, the
52 localized sEMG root mean square [23] value of an array point can be estimated by a 2-D

53 topographic representation of muscle electrical activity using a linear cubic spline
54 interpolation [13]. The result is a visual representation of muscle activity over a two
55 dimensional region [13]. Our hypothesis was that topography sEMG testing may prove
56 more valuable to assess the lumbar muscle function during dynamic flexion-extension and
57 its potential use to predict the prognosis of functional restoration rehabilitation in a
58 population of chronic LBP subjects. It would be helpful to classify those patients who have
59 good respond to conservative care.

60

61

62 **Methods**

63

64 *Subjects:*

65 A total of 43 healthy subjects (mean age = 32 ± 6.5 years, 23 males and 20 females) and 38
66 patients with chronic non-specific LBP (mean age = 42 ± 9.7 years, 28 males and 10
67 females) were recruited based on inclusion and exclusion criteria (Table 1). Approval for the
68 study was received in advance of testing by the Institutional Review Board for clinical
69 research ethics review. A written consent was collected from each participant.

70

71 *Surface EMG Test:*

72 All subjects received lumbar muscle sEMG test after enrolment. The sEMG data were
73 collected from the lumbar region using a 7x3 array of electrodes applied evenly in the
74 lumbar region from the spinal level of L2 to L5 (Figure 1). Each sEMG electrode was 1.5
75 cm in diameter and applied to alcohol-cleaned skin having impedance of less than 10 k Ω as
76 measured by a multimeter (UT611, Uni-T LTD , Shenzhen, China). sEMG signals were
77 amplified by 2000 times and filtered between 15-950 Hz. The data were acquired at a
78 sampling rate of 2000 Hz by a data acquisition card (DAQ6063, National Instruments Inc.,
79 Austin, Texas, USA). Then, subjects were asked to perform a trunk-bending motion which has
80 been suggested as one of the useful dynamic tasks for evaluating lumbar muscle function[13].

81 The trunk-bending motion consisted of three phases: flexion, relaxation and extension.
82 Subjects were asked to bend their trunk forward for 1 second with the range of the flexion
83 angle between 20-30 degrees as estimated by utilizing a protractor. Subsequently, they held
84 their flexed posture for 2 seconds and then returned to the original straight standing posture
85 for 2 second. The whole sEMG measurement was carried out under a constant room
86 temperature so that the effect of temperature on the active potential conduction velocity and
87 contractility in the muscle fibers was eliminated.

88

89 *Rehabilitation Program:*

90 All enrolled LBP patients completed a 12-week in-patient rehabilitation program (5 days per
91 week) [24]. The standard exercise therapy and mobilization technique was performed in this
92 study. The dosage, intensity and other factors related to these activities were prescribed and
93 re-evaluated at each session by the hospital physical therapy staff. This individualized
94 program is based on a “functional restoration program” [25, 26] that is divided into three
95 phases: physical conditioning (5 weeks), working conditioning (4 weeks), and work readiness
96 (3 weeks). In the physical conditioning phase, patients received 4 hours of physiotherapy (PT)
97 and 2 hours of occupational therapy (OT) each day. These therapies focused on spinal
98 mobilization, back muscle strengthening, cardiovascular and work skill training. In the work
99 conditioning phase (PT: 3 hours/day, OT: 3 hours/day) and work readiness phases (PT: 2

100 hours/day, OT: 4 hours/day), patients continued with work simulated tasks as well as
101 strengthening exercises, treadmill activities and pelvic stabilization training.

102

103 *Clinical assessments:*

104 At enrollment, LBP patients were asked to complete a standard intake questionnaire to obtain
105 self-reports of age, gender, weight, height, medical history, the location and nature of their
106 symptoms. Before and after a 12 week rehabilitation program (see below), subjects completed
107 1) an 11-point visual analog pain-rating scale (VAS) ranging from 0 (no pain in the last 24
108 hours) to 10 (worst imaginable pain in the last 24 hours)) and the 2) Oswestry Disability
109 Questionnaire (ODQ)[27].

110

111 According to the results of the ODI and VAS evaluations, the LBP patients were categorized
112 to 2 sub-groups as either “responding” or “non-responding” based on the minimal clinically
113 important difference (MCID) reported for the VAS (2 points decrease)[28] or the ODQ (10
114 points improvement) [29]. In the present study, LBP patients exceeding the MCID of the
115 VAS, ODQ or both were considered to be responders. Otherwise, they are regarded as
116 “non-responding”.

117

118 *Dynamic sEMG topography analysis*

119 A total of 16 channels sEMG signals were recorded from array-electrode. A sliding analysis
 120 window of 0.2 s was employed to segment the sEMG signals along the lumbar flexion and
 121 extension. With a moving window interval of 0.1 s, a total of 50 blocks of sEMG signals
 122 was segmented from a whole circle of flexion-extension(flexion:10 blocks, relaxation:20
 123 blocks, extension:20 blocks). In each block, root-mean-square [23] values of sEMG signals
 124 were calculated for each channel by the following equation:

$$X_{\text{rms}} = \sqrt{\frac{x_1^2 + x_2^2 + \dots + x_i^2 + \dots + x_n^2}{n}}$$

125 (1)

126 where x_i is sEMG signal, and n is the sampling number within the analysis window ($n=400$
 127 in this study). The RMS values of each analysis window were normalized to the maximum
 128 RMS value among all of the analysis windows of a whole flexion-extension circle. To
 129 construct a 2-D SEMG topography, the RMS values of the 16 SEMG channels within a
 130 definite time interval were calculated as per a 160 x 120 matrix, using a linear cubic spline
 131 interpolation of each scan as described in a published report [13]. During the whole
 132 flexion-extension circle, each block of sEMG signals can generate a frame of topography
 133 colour map. Therefore, a sequence of 50 topography frames (10 frames in flexion, 20
 134 frames in relaxation, 20 frames in extension) can be created. Fig. 2 (a) demonstrated the 5
 135 continuing frames of sEMG topography in flexion action. The topography represents the
 136 intensity of sEMG distribution by the colour gradient, in which a blue colour means the

137 lowest value and a red colour is the highest value. In each frame, three topographic
 138 parameters, namely relative area (RA), relative width (RW), and relative height (RH), as
 139 proposed by a previous report[13], were used to measure the features of the highest 60%
 140 RMS value region in sEMG topography as shown in Fig. 2 (b).

141

142 After measuring 3 topographic parameters in all 50 frames of whole circle, can be plot a
 143 time-varying curve. Figure 3 demonstrated a plot of a time-varying RA curve from a normal
 144 subject. Mean and standard deviation of time-varying topography parameters from
 145 forty-three healthy subjects were calculated as the normal values. To quantify the
 146 discrepancies in the variation patterns of these parameters during the flexion and extension
 147 phases between the normal and LBP patient groups, the root-mean-square difference
 148 (RMSD) of each parameters variation pattern in LBP patient with respect to the normal
 149 group was evaluated according to the following equation.

$$\text{RMSD} = \sqrt{\frac{\sum_{i=0}^{i=N} (b_i - a_i)^2}{N}}$$

150

(2)

151 where a_i is a set of the mean value from normal data (reference data),

152 b_i is a set of the LBP patient group data (compared data), and N is the

153 sampling number.

154

155 In this study, all the topographic parameters of the RMSD during relaxation phase was not
156 taken into consideration since the sEMG signals in the relaxation phase was lack of lumbar
157 myoelectric activities. Therefore, the parameters of RMSD RA, RMSD RW, RMSD RH in
158 both flexion and extension were calculated.

159

160 *Statistical Analysis:*

161 All presented data were analyzed using SPSS 16.0 software. RMSD RA, RMSD RW,
162 RMSD RH in both flexion and extension from normal group and the “responding” and
163 “non-responding” groups were compared by one-way ANOVA. The sensitivity and
164 specificity of parameters were determined by the ROC curve. P-value < 0.05 was
165 considered as statistically significant.

166

167 **Results**

168

169 *Time-varying topography of healthy subjects*

170 Figure 3 presented a sample time-varying relative area (RA) curve of a healthy subject.

171 Time-varying curves of topography parameters, i.e. RA, RW and RH, were calculated in all

172 healthy subjects. Then, normal patterns of time-varying RA, RW and RH can be obtained

173 and presented in Figure 4.

174

175 *Comparisons between the “responding” and “non- responding” groups*

176 As shown in figure 4, a sample curve from a LBP patient was plotted on the normal pattern.

177 It showed an obvious bias between LBP and normal curves. An ANOVA group comparison

178 of time-varying RA, RW and RH showed significant difference ($p < 0.05$) between the

179 healthy and LBP groups. To each LBP patient, RMSD parameter can be calculated as a

180 quantitative measure of the discrepancies in comparison to healthy normal data.

181

182 In this study, 16 LBP was classified as “responding” group and 23 patients as

183 “non-responding” group. All RMSD parameters of “responding” group were consistently

184 lower than the “non-responding” group as shown in Fig. 5. The lower the RMSD parameter

185 value was, the closer to the normal condition, and hence the better respond to the

186 physiotherapy treatment. Significant differences were found in RMSD RA and RMSD RW
187 in both flexion and extension phase between the “responding” and “non- responding”
188 groups ($p < 0.05$). However, RMSD RH did not show significant difference between the
189 “responding” and “non- responding” groups.

190

191 *Accuracy test*

192 ROC curves of RMSD RA and RMSD RW at flexion and extension were plotted out for
193 testing the accuracy of discriminating the responding cases from non-responding cases by
194 the RMSD parameters as shown in Fig. 6. The area under the ROC curve (AUC) of RMSD
195 RW at extension (0.723) was the largest and followed by RMSD RA at extension (0.699).
196 They were also found significantly different ($p = 0.023$, $p = 0.043$). All four RMSD RA and
197 RMSD RW at flexion and extension had AUC larger than 0.5.

198

199

200 **Discussion**

201 There are various therapeutic interventions to be recommended for the treatment of LBP[9],
202 in which exercise therapy is a commonly used management of chronic LBP[2]. The goal of
203 this study is to evaluate the prognostic value of dynamic sEMG topography in chronic LBP
204 patients to take an intensive non-surgical rehabilitation programme. The results support our
205 hypothesis that topography sEMG testing is able to predict the prognosis of functional
206 restoration rehabilitation.

207

208 While exercise rehabilitation can be considered to have better efficacy than most other
209 interventions [10, 30], it would be unrealistic to expect a single therapeutic intervention to
210 be capable of resolving all LBP complaints in all subjects. In various manipulative and
211 exercise therapy[10], this study attempted to evaluate the prognostic value of dynamic
212 sEMG topography in an intervention focused on treating excessive muscle activity. In this
213 study, 16 of 38 LBP patients experienced pain relief or functional improvement after 12
214 weeks of intensive. Therefore, by understanding which subjects may respond to care in
215 advance of its provision, the potential exists to prescribe the intervention only to those most
216 likely to benefit[31] and thereby reduce the direct and indirect costs associated with
217 treatment [32].

218

219 A previous report proposed the quantitative analysis of sEMG topography as an objective
220 method for LBP rehabilitation assessment [13]. The present study further developed the
221 time-varying quantitative analysis of sEMG topography rather than a static topography in
222 each sub-action of the forward bending motion. Findings in time-varying parameters of
223 sEMG topography showed significant difference in LBP ($p < 0.05$), which support the
224 previous results that LBP showed different topography from healthy subjects [13]. In
225 addition, the aim of this study is to observe whether the time-varying sEMG topographic
226 parameters can differentiate the “responding” group and the “non- responding” group of
227 LBP, so as to evaluate its prognostic value.

228

229 To measure the dynamic surface EMG topography, time-varying RA, RW and RH patterns
230 could reflect the dynamic change of lumbar muscular contraction patterns during the
231 flexion-extension motion. In the present study, RMSD was proposed to evaluate the
232 discrepancy of time-varying sEMG topography between any individual LBP and the mean
233 value of healthy group. The lower of RMSD indicate the most similar pattern of the sEMG
234 topography. As shown in figure 5, RMSD RA and RMSD RW of responding group were
235 found to be significantly lower than that of non-responding group. Even without statistical
236 significance, RMSD RH also showed an obvious lower value in the responding group than
237 in the non-responding group. It suggests that the LBP patient, with the dynamic sEMG

238 topography pattern close to normal healthy, would most likely respond to rehabilitation
239 therapy.

240

241 Prognosis of LBP has been discussed in a lot of literatures [31, 33-39], but most of the
242 prognostic variables are not specific with individual patient. In this study, the discrepancy of
243 every individual patient from normal data can be quantitatively measured and plotted as
244 figure 4. The merit of this prognosis tool is that it can provide a valuable prediction to
245 clinician and the patient to select the most appropriate treatment on the early stage. ROC
246 analysis On the other hand, the results of ROC curves showed that the area under curves
247 (AUC) of four RMSD parameters, RMSD RA and RMSD RW at both flexion and extension
248 are greater than 0.5 , which prove the prognostic value of RMSD parameters.

249

250 There were two limitations in this study. The healthy subjects in the control group were
251 younger than patients because of the difficulty to recruit healthy subjects older than 45 years
252 old. The second limitation was that we did not collect body mass index (BMI), so as to not
253 analyze the effect of BMI on the surface EMG topography. Four prognostic parameters of
254 time-varying surface topography were proposed, but it is still to determine a clear threshold
255 as well as a optimal combination of parameters for predicting prognosis in a separate study
256 of large scale LBP population.

257

258 **In summary, the quantitative analysis of time-varying sEMG topography showed**
259 **significant difference between healthy and LBP groups. The discrepancies in quantitative**
260 **dynamic sEMG topography from normal healthy curves, in terms of RMSD RA and RMSD**
261 **RW at flexion and extension, was able to identify those LBP subjects who would respond**
262 **to a conservative care program focused on functional restoration of lumbar muscle. The**
263 **use of quantitative analysis of time-varying sEMG topography would help to select the**
264 **most appropriate conservative treatment for chronic LBP patients.**

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- 362

364 **Legends:**

365

366 Figure 1 Placement of the 3×7 electrode-array on lumbar surface

367

368 Fig. 2 A sample sequence (5 frames) of dynamic topography during flexion phases and a

369 measurement of relative area (RA), relative width (RW), and relative height (RH)

$$\text{Relative Area (RA)} = \frac{\text{High activity area}}{\text{Total topography area}},$$

$$\text{Relative Width (RW)} = \frac{\text{Width of high activity area}}{\text{Total topography width}},$$

$$\text{Relative Height (RH)} = \frac{\text{Height of high activity area}}{\text{Total topography height}}$$

370

371

372 Fig. 3 A sample time-varying relative area curve of sEMG topography from a normal

373 subject

374

375 Fig. 4 Patterns of Time varying sEMG topography parameters: RA (a), RW(b) and RH(c)

376 (Normal range of healthy subject in red and green regions)

377

378 Fig. 5 Comparison of RMSD parameters (RMSD RA, RW, RH at flexion and extension)

379 between “responding” and “non-responding” groups.

380 Values of the six tested RMSD parameters of “responding” groups are consistently lower
381 than those of “non-responding” groups. RMSD RA and RMSD RW flexion and extension
382 are found significantly different between groups ($p < 0.05^*$)

383

384 Fig. 6 ROC curves of RMSD RA and RMSD RW flexion and extension.

385 The area under curve (AUC) of each ROC curve is >0.7 and is statistically significant ($p < 0.05^*$).

386

Table 1. Inclusion and exclusion criteria

Healthy subjects
<p>Inclusion criteria:</p> <ul style="list-style-type: none"> • Age from 18 to 60 years • Normal physical and neurological examinations <p>Exclusion criteria:</p> <ul style="list-style-type: none"> • With occurrence of low back pain in the past 6 months. • Previous spinal surgery • Pregnancy
Low back pain patients
<p>Inclusion criteria:</p> <ul style="list-style-type: none"> • Age 18 to 60 years • A primary symptom of low back pain, with or without referral into the lower extremity • A minimum Oswestry Disability Questionnaire (ODQ) score of 30%. <p>Exclusion criteria:</p> <ul style="list-style-type: none"> • The presence of "red flags" as obtained from questionnaire and/or physical exam (see below), • Signs consistent with nerve root compression (e.g. positive straight-leg increase < 45 degrees or diminished reflexes, sensation, or lower-extremity strength) • Pregnancy • Previous spinal surgery • Cancer • Unexplained weight loss • Immunosuppression • Prolonged use of steroids, Intravenous drug use, Urinary tract infection, Pain that is increased or unrelieved by rest, Fever, Significant trauma related to age (e.g., fall from a height or motor vehicle accident in a young patient, minor fall or heavy lifting in a potentially osteoporotic or older patient or a person with possible osteoporosis), Bladder or bowel incontinence, Urinary retention (with overflow incontinence). Open sores, Saddle anesthesia, Loss of anal sphincter tone, Major motor weakness in lower extremities, Fever, Vertebral tenderness, Limited spinal range of motion, other neurologic findings persisting beyond one month.

Figure1
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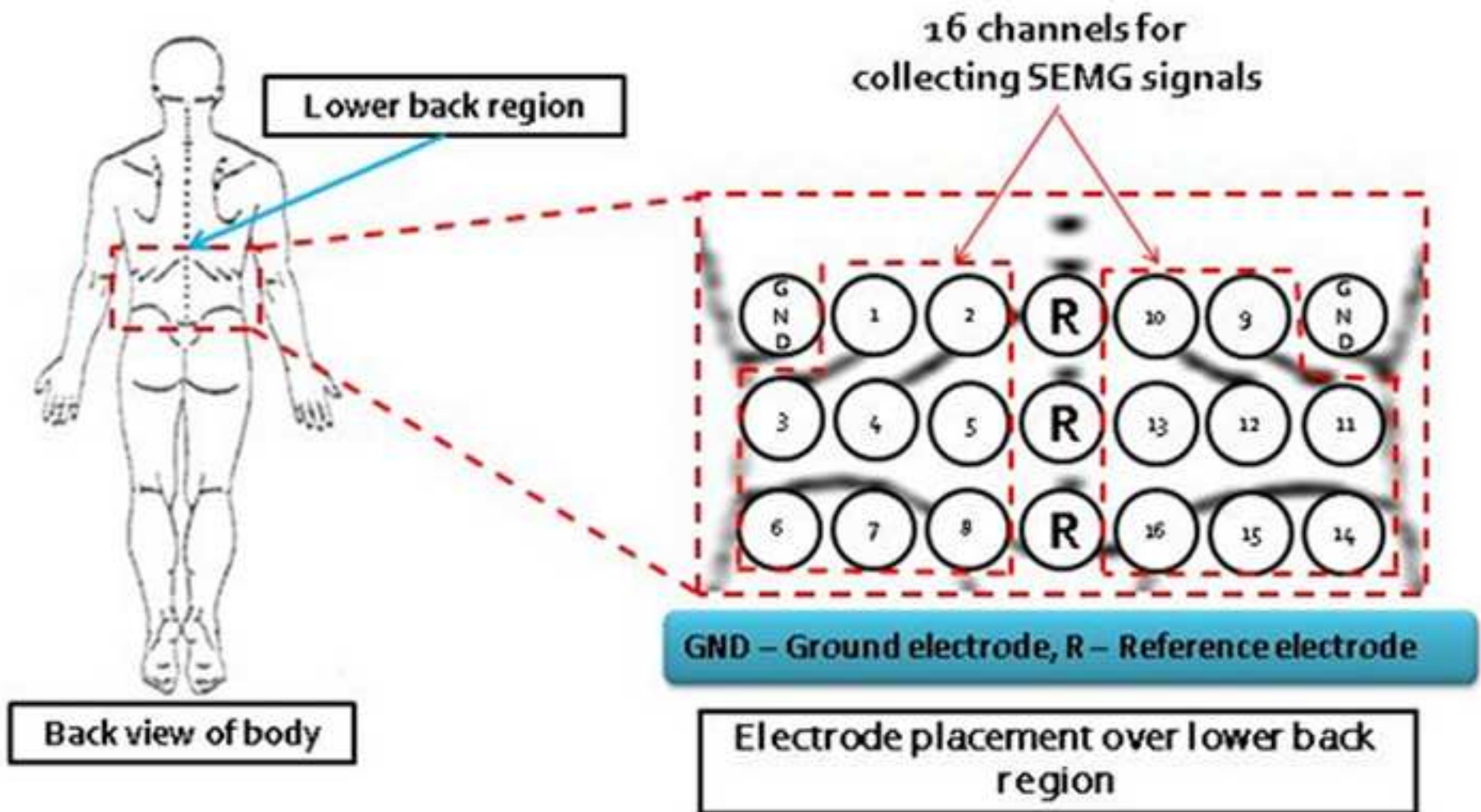


Figure2(a)

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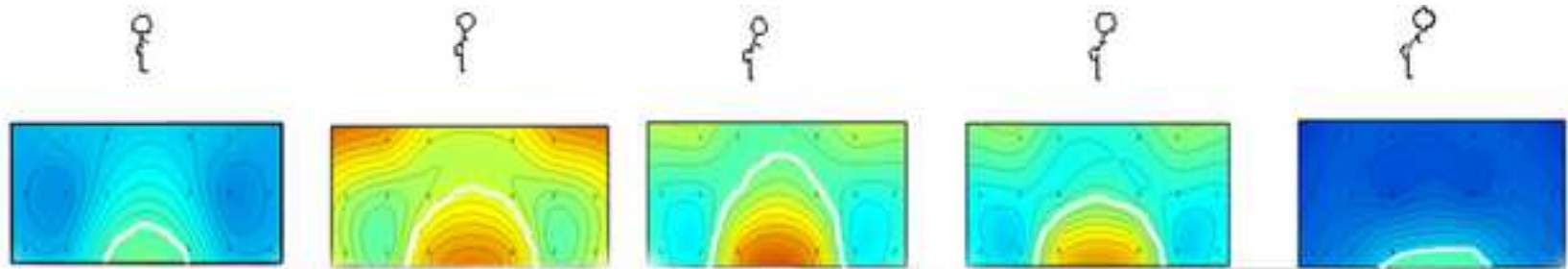


Figure2(b)

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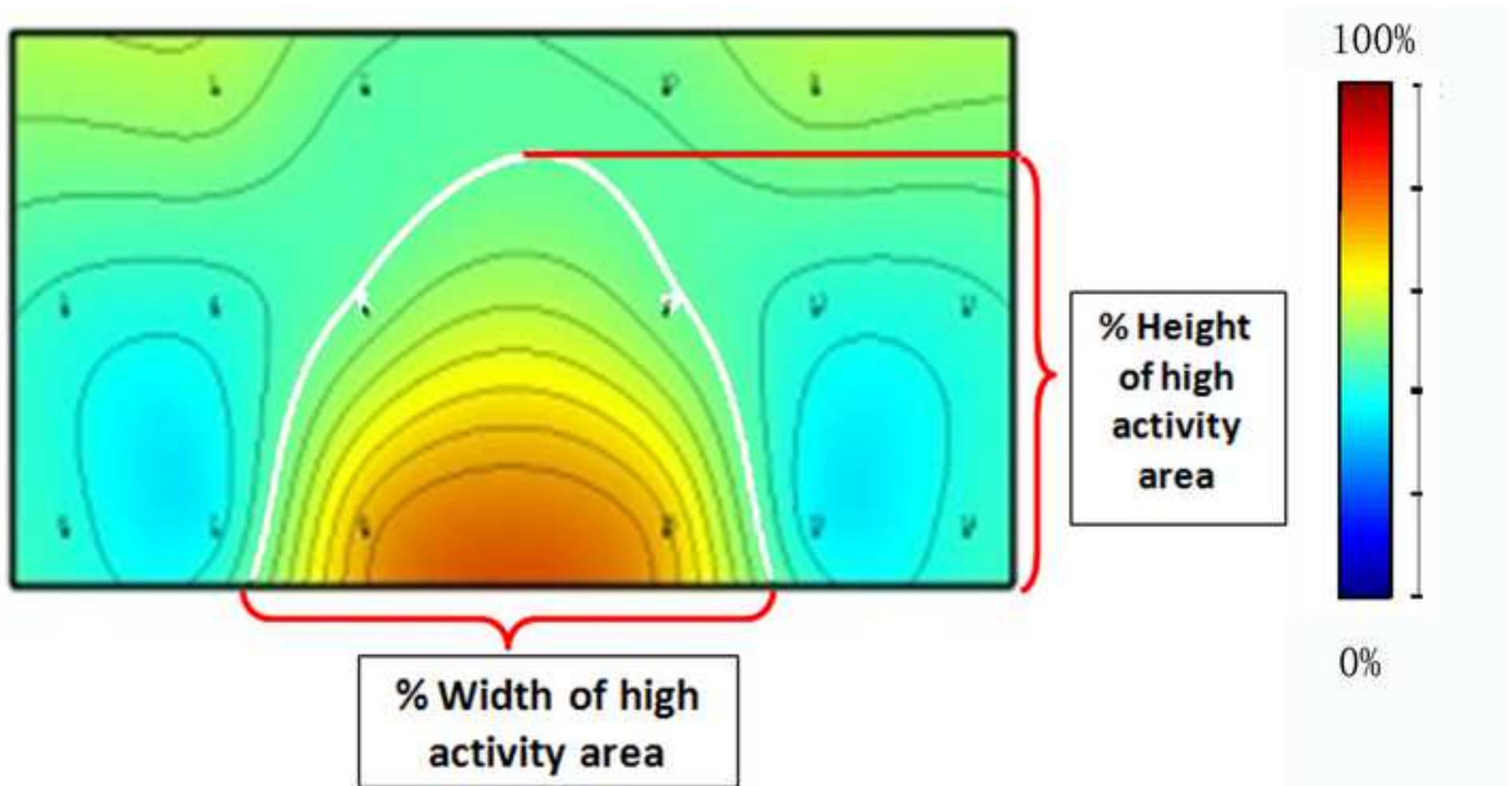


Figure4(a)
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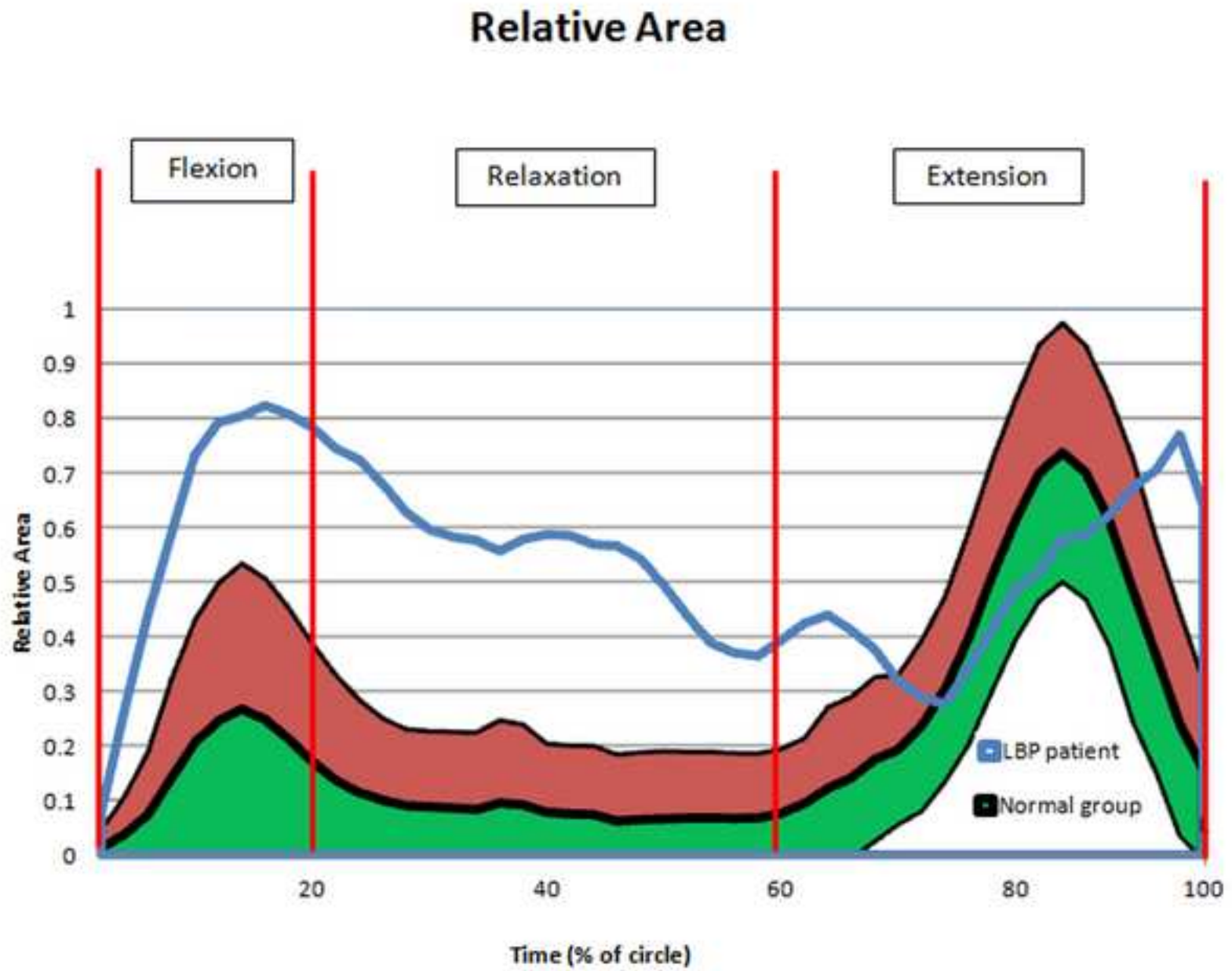


Figure4(b)
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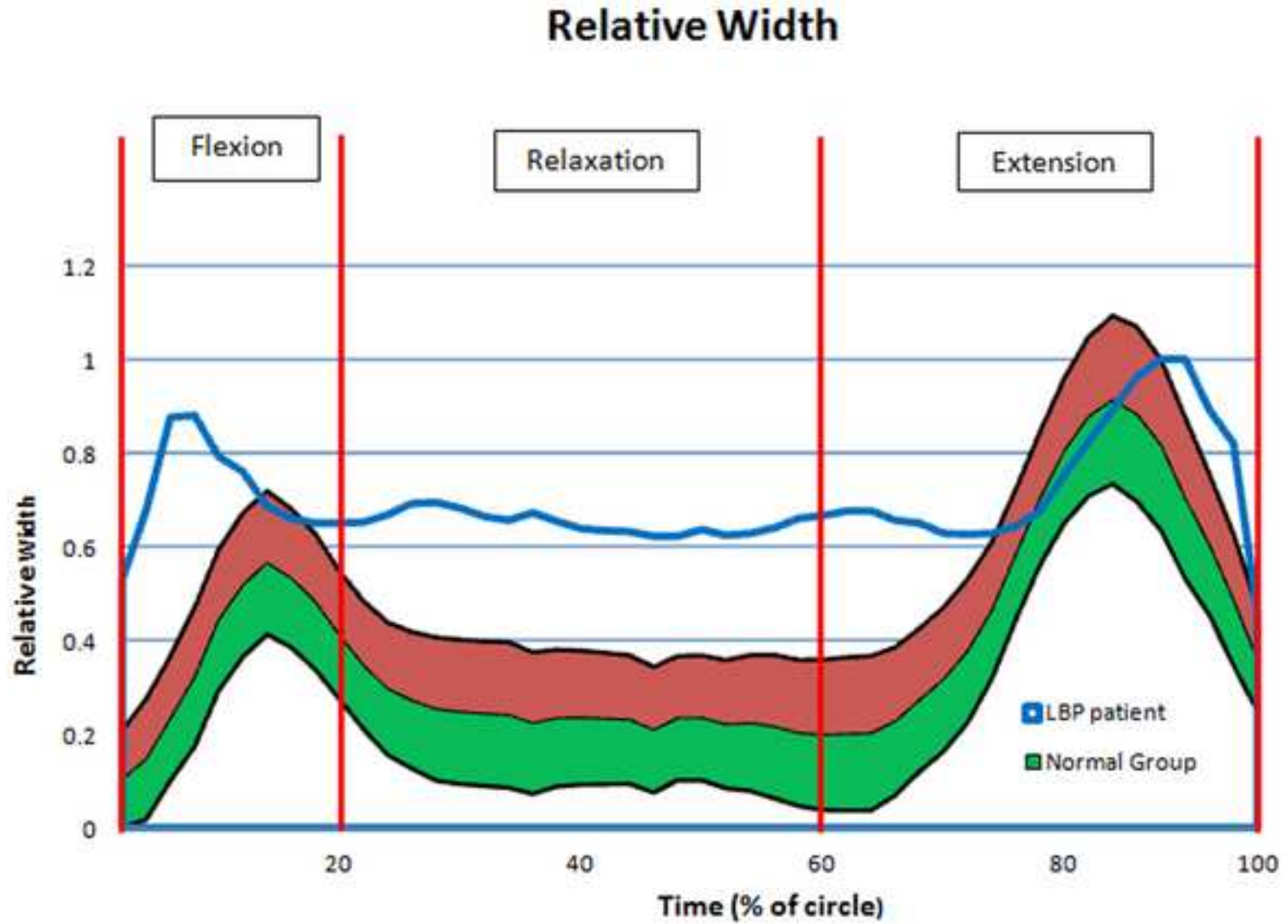
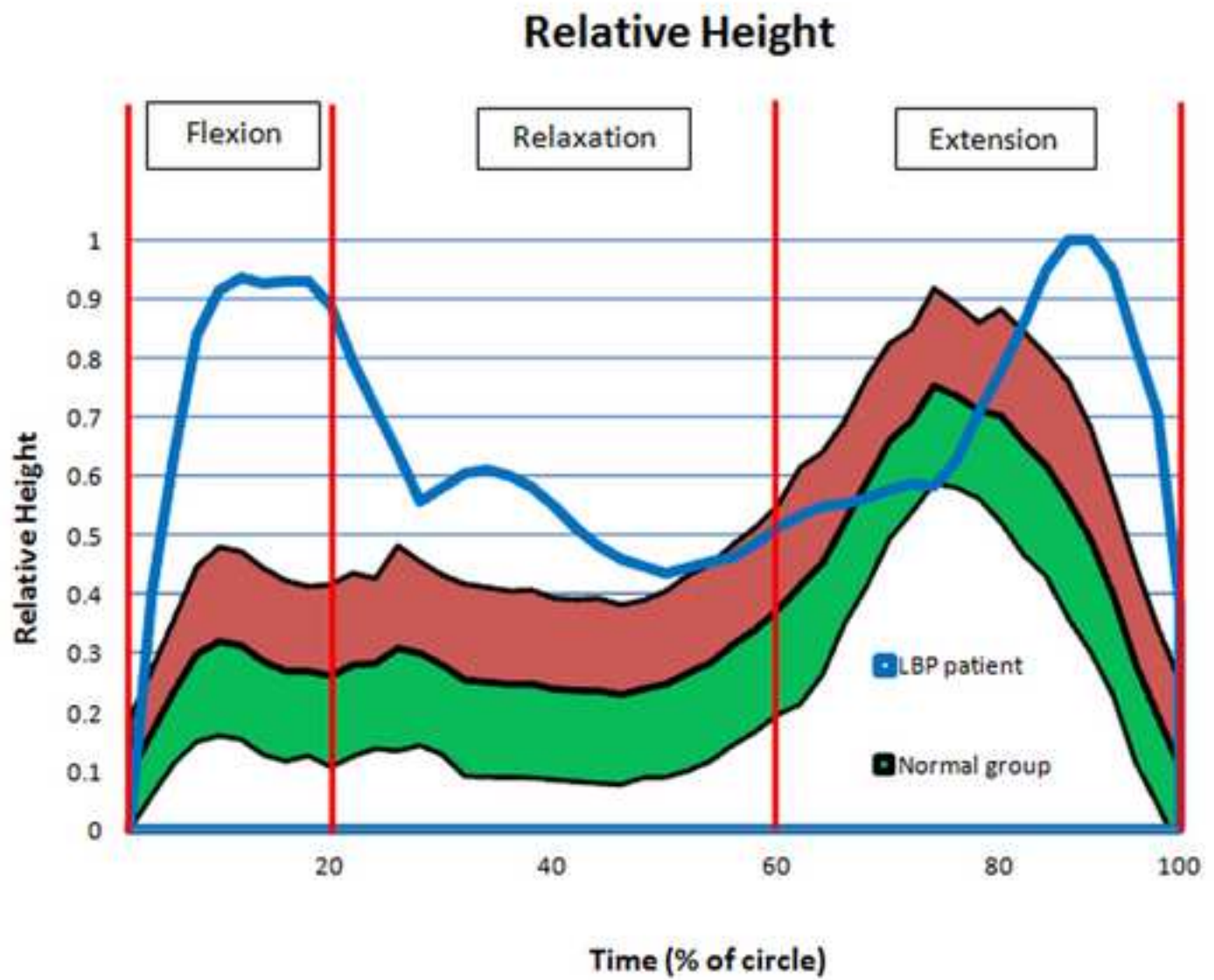


Figure4(c)
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Comparison of RMSD parameters between "responding" and "non-responding" groups

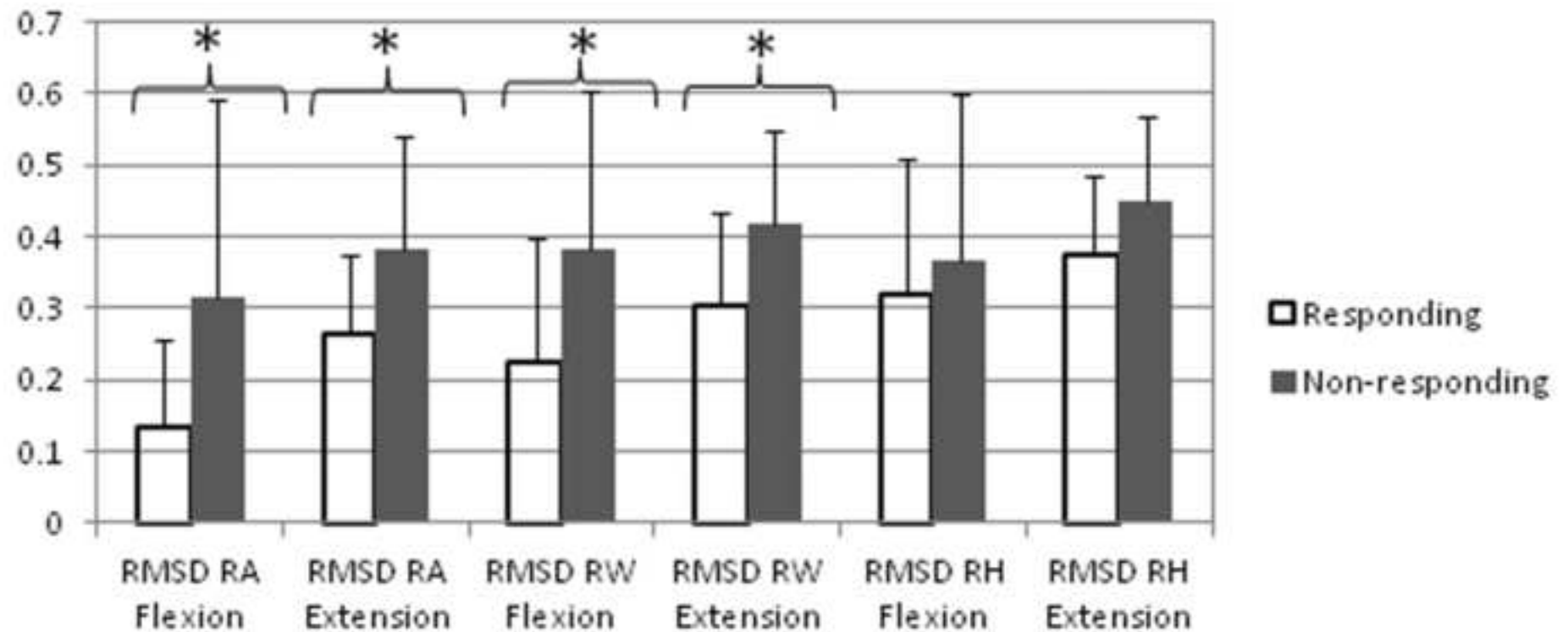


Figure6
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