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7	Navigating the Factors Affecting Functional impairment in Spondyloarthritis
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15	Abstract
16	Objectives: To assess the predictive factors of functional impairment in SpA patients assessed
17	with BASFI (Bath Ankylosing Spondylitis Functional Index) and LI (Lequesne Index).
18	Methods: We conducted a retrospective study that included SpA patients. Socio-
19	demographics and disease-related data were reported (data collection spread over four months
20	from August 2019 to November 2019). Disease activity was assessed using the Bath
21	Ankylosing Spondylitis-Global score (BASG-s) and the Bath Ankylosing Spondylitis disease
22	activity Index (BASDAI). The spinal mobility was evaluated by The Bath Ankylosing
23	Spondylitis Metrology Index (BASMI). Structural progression was evaluated with The Bath
24	Ankylosing Spondylitis Radiologic Index (BASRI) and modified Stoke ankylosing
25	spondylitis spine score (mSASSS). We conducted a multivariate analysis to search for
26	predictive factors associated with BASFI and LI. Significance was accepted for a $p < 0.05$ for
27	all statistical tests. Results: Two hundred and sixty-three patients were included. Mean age of
28	the patients was 38.9 $\pm$ 12.7 years, sex-ratio=2.7. Mean age of onset of SpA was 27.6 $\pm$ 10.8.
29	Disease duration was 11.3 $\pm$ 9.5 years. Occupation was significantly associated with BASFI
30	score and LI. A significant functional impact was significantly correlated with a long duration
31	of the disease. The two scores were correlated with a limitation of spinal mobility (BASMI), a
32	greater disease activity (BASDAI and ESR) and a greater impact of the disease on health
33	status (BASG-s). Significant functional impairment was also correlated with structural

- impairment: mSASSS, BASRI and sacroiliitis grade. The variables independently related to
- 35 BASFI were the mSASSS score and the BASDAI. The variables independently related to LI
- were the profession (unemployed subjects had higher scores), mSASSS score, and the
- 37 BASMI. *Conclusion:* Occupation, disease activity, mobility and structural progression
- 38 predicted functional impairment in Tunisian SpA patients.
- *Keywords:* Spondyloarthritis; North Africa; Function; Impairment.

## Advances in Knowledge:

- *Comprehensive Assessment:* This article provides a thorough evaluation of functional impairment in spondyloarthritis (SpA) patients, utilizing both the BASFI and Lequesne Index (LI), shedding light on the underexplored LI in the context of SpA.
  - *Comprehensive Factors:* It identifies a wide range of factors influencing functional impairment, including disease duration, occupation, disease activity, spinal mobility, structural progression, and health status, providing a comprehensive view of the multifaceted nature of SpA.
  - *Unique Population:* This research focuses on North-African SpA patients, contributing valuable insights into the specific factors affecting this population, which may have distinct characteristics compared to other populations.

## **Application to Patient Care:**

- *Early Intervention:* The identification of disease duration as a significant factor highlights the importance of early diagnosis and intervention to minimize functional impairment and enhance patients' quality of life.
- *Holistic Approach:* Healthcare providers can use the insights from this study to adopt a holistic approach to SpA management, considering not only disease activity but also mobility, structural progression, and patient-reported outcomes.
- Tailored Care for North-African Patients: The study's focus on North-African SpA patients facilitates culturally sensitive and region-specific care, ensuring that interventions align with the unique needs of this population.

## Introduction

- 65 Spondyloarthritis (SpA) is a chronic inflammatory rheumatic disease that occurs particularly
- in young adult males. The prevalence of this disease varies across regions and data regarding

the North Africa and Middle East region is limited. North African patients with SpA exhibit 67 68 distinct clinical and genetic features. For one thing, North African patients tend to have more severe forms of SpA, with a higher prevalence of the axial form and of coxitis. 1,2 Another 69 important feature is the genetic trait of north African SpA patient with very low prevalence of 70 HLA-B27.<sup>3</sup> 71 72 73 Occurrence of SpA in a young and active subject is what makes this disease an extremely 74 challenging one. Thankfully, the perception of spondyloarthritis has shifted from a 75 debilitating and disabling disease to one that can be managed and controlled with appropriate 76 advanced therapy. However, although fascinating progress has been made the past decades in 77 the fields of pathophysiology -but still far to be understood- and treatment, patients are still 78 plaintive and still report symptoms of pain fatigue and functional impairment which are key patient-related outcomes (PROs).4 79 80 81 Patient age, sex, ethnicity, education level, smoking status, disease duration, medication use, family history of arthritis and inflammation emerged as potential risk factors for functional 82 83 limitation in the literature.<sup>5</sup> Functional impairment was also associated with increased levels of work disability and a decrease in work productivity. 6 This, in turn, results in considerable 84 85 economic and societal costs. Understanding how important these effects are, it becomes crucial for us to investigate the factors contributing to functional impairment among SpA 86 87 patients. 88 89 In light of this, our study aims to leverage these insights to develop targeted interventions that 90 can improve the quality of life and function of SpA patients. By doing so, we hope to 91 contribute not only to individual well-being but also to the overall economy and society. 92 Through this work, we propose to evaluate the functional impact in Tunisian SpA patients and 93 identify the factors that are most correlated with the alteration of their functional prognosis. 94 95 Methods 96 Study design and patient selection criteria 97 This is a retrospective study that included SpA patients, all fulfilling the assessment of 98 Spondyloarthritis International Society (ASAS) criteria, who were followed at the 99 rheumatology department of the Mohamed Kassab Institute of Orthopedics between 2008 and 100 2019 with data collection spread over four months (August 2019 to November 2019). A data

101 collection form was drawn up and we extracted the relevant information from medical 102 records. Patients with an associated chronic inflammatory rheumatism and those with pre-103 existing disabling conditions (Pre-existing disability independent of the disease and its 104 complications) were not included to limit confounding factors. A data collection sheet was 105 prepared. Significance was accepted for a p < 0.05 for all statistical tests. 106 107 Data collection We reported Socio-demographics. The reported clinical data included the age of onset of the 108 109 disease, the duration of SpA, the presence of spinal pain and coxitis, the search for extra-110 articular manifestations were recorded. 111 112 Disease activity, functional impact and spinal mobility were assessed using specific tools. 113 Disease activity was assessed by inflammatory biomarkers (Erythrocyte sedimentation rate 114 (ESR) and C-reactive protein (CRP), the Bath Ankylosing Spondylitis-Global score (BASG-115 s) and the Bath Ankylosing Spondylitis disease activity Index (BASDAI). The assessment of 116 functional discomfort in SpA patients was scored using the Bath Ankylosing Spondylitis 117 Functional Index (BASFI) and the hip Lequesne Index (LI). The presence of pain at enthesis 118 sites was evaluated by Maastricht Ankylosing Spondylitis Enthesitis Score (MASES). The 119 spinal mobility was (evaluated by The Bath Ankylosing Spondylitis Metrology Inde 120 (BASMI)). 121 122 Assessment of structural involvement was determined by: The Bath Ankylosing Spondylitis 123 Radiologic Index (BASRI) and modified Stoke ankylosing spondylitis spine score (mSASSS). The grading of sacroiliitis was done using the modified NY criteria.<sup>8,9</sup> The radiological form 124 125 of coxitis was specified (early coxitis, enveloping coxitis, pseudoarthritic coxitis, destructive 126 coxitis and synostotic coxitis). Therapeutic modalities were mentioned. 127 128 Statistical analysis 129 All calculations were made using the Statistical Package for Social Sciences (SPSS) version 130 25.0 for Windows. 131 132 For the analytical study, the comparison of the qualitative variables was carried out by the 133 analysis of variance (ANOVA with 1 factor). The study of correlation was performed by 134 Pearson's correlation coefficient (r). Significance was accepted for a p < 0.05 for all statistical

135 tests. In order to find the variables independently linked to the bath ankylosing spondylitis 136 functional index (BASFI) and the Lequesne index (LI), we conducted a multivariate analysis 137 in multiple linear regression, step-by-step descending method (in the first step, we introduce 138 all the variables and from step to step we remove the variable with the least significant "p"). 139 140 Ethics and conflict of interest 141 All patients gave consent to participate. The Ethics committee name and number are Kassab 142 institute ethical Committee, number IMKO018. This study was carried out in accordance with the guidelines in the Declaration of Helsinki. Standards of reporting: STROBE guidelines 143 144 were followed. We declare that we have no conflicts of interest in this work. All authors 145 contributed to this work. 146 147 **Results** Two hundred and sixty-three patients were included. The mean age of the patients was  $38.9 \pm$ 148 149 12.7 years [16-79], 192 men (73%) and 71 women (27%). The mean age of onset of SpA was 150  $27.6 \pm 10.8$  years [5-61]. Disease duration was  $11.3 \pm 9.5$  years, [1-62]. According to ASAS 151 criteria: SpA was axial in 43.1% of cases, peripheral in 44% of cases with enthesitic 152 involvement in 19.1% of cases. SpA was axial and peripheral in 44.7% of cases. Socio-153 demographic and disease-related baseline characteristics are summarized in table1. 154 The main clinical and paraclinical data are reported in Table 2. 155 156 Association of BASFI and LI with patient characteristics, disease characteristics, 157 paraclinical data, and therapeutic modalities Occupation was significantly associated with BASFI score and LI. The post hoc test showed a 158 159 higher BASFI score (p= 0.010) and LI (p<0.001) in the unemployed subjects. Regarding 160 baseline disease-related features, a significant functional impact was significantly correlated 161 with a long duration of the disease with p=0.007 for BASFI and p<0.001 for LI. 162 163 Peripheral involvement was also significantly associated with BASFI score and LI with 164 p=0.006 and 0.021 respectively. Extra articular involvement was not associated with higher 165 functional impairment (p=0.907 for BASFI and p=0.152 for LI). Neck and back pain were significantly associated with BASFI and LI scores. The same was true for coxitis, its bilateral 166 167 character and the limitation of the hip mobility (table 3). 168

169	The two scores were correlated with a limitation of spinal mobility (BASMI), a greater
170	disease activity (BASDAI and ESR) and a greater impact of the disease on health status
171	(BASG-s) (Table 4).
172	
173	Regarding the radiological data, radiological forms of coxitis (p=0.037 for BASFI score and
174	p<0.001 for LI) were significantly associated with functional discomfort. The post hoc test
175	showed that a destructive form of coxitis was associated with a higher BASFI score and LI.
176	Significant functional impairment was also correlated with structural impairment: mSASSS
177	p=0.001 for BASFI and p=0.003 for LI), BASRI (p<0.001 for both BASFI and LI) and
178	sacroiliitis grade (p<0.001 for both BASFI and LI).
179	
180	Table 3 and 4 show association of BASFI and LI with patient characteristics, disease
181	characteristics, paraclinical data, and therapeutic modalities
182	
183	Multiple linear regression
184	The variables independently related to BASFI were the mSASSS score and the BASDAI.
185	The variables independently related to LI were the profession (unemployed subjects had
186	higher scores), mSASSS score, and the BASMI.
187	Table 5 shows the results of the multiple linear regression of factors associated with the
188	BASFI score and the LI.
189	
190	Discussion
191	Our study aimed to determine the main factors associated with significant functional impact
192	in north-african SpA patients. BASFI and LI were higher in patients who were unemployed.
193	Moreover, greater functional impact was associated with peripheral involvement, neck pain,
194	back pain and coxitis. The elevation of both scores was also significantly correlated with a
195	longer duration of disease progression. The BASFI and LI were correlated with a limitation of
196	spinal mobility, greater disease activity. BASFI correlated with a higher MASES score and
197	the presence of low back pain.
198	
199	Regarding the association between radiological data and scores assessing functional
200	discomfort, a high BASFI and LI were significantly associated with the presence of a
201	destructive form of coxitis. They were also correlated with structural damage (mSASSS,
202	BASRI, sacroiliitis grade).

203 204 On multivariate linear regression, the variables independently related to BASFI were the 205 mSASSS score (p=0.016) and the BASDAI score (p<0.001). Variables independently related 206 to higher LI values were work occupation (unemployed subjects) (p=0.004), mSASSS score 207 (p=0.046) and BASMI (p<0.001). 208 209 To our knowledge, there have been very few studies in the literature evaluating the factors associated with a high LI, which is a French score developed in the 80s, <sup>10</sup> in SpA patients. 210 We were able to identify two published articles which used the Lequesne index to assess 211 functional impairment in SpA patients. 11,12 Interestingly, both articles were from Tunisia, a 212 francophone country. This highlights one particularity of our study: the search for all potential 213 214 factors associated with the LI in a population where it has been little used. However, this 215 index is frequently used in practice to assess the functional impact of hip and knee damage in 216 osteoarthritis patients. The search for all potential demographical, clinical and para clinical 217 factors associated with functional impairment in a population of SpA patients known to have 218 a higher prevalence of coxitis represents the other highlight of our study. However, the 219 retrospective nature may be a limitation. Furthermore, the disease was active (BASDAI was 220 >40) in 65.6% of cases. This could be explained by a recruitment bias because our study was 221 carried out in a university hospital that manages severe forms of SpA. 222 223 Regarding the socio-demographic features, profession was significantly associated with the LI 224 as well as the BASFI score. It was, indeed, independently associated to LI, with higher values 225 in the unemployed subjects. The absence of professional activity would be the consequence of 226 an important functional impact of the disease and not its cause. In the study by Ward MM et 227 al<sup>5</sup>, on the other hand, each increase in the occupational physical activity score (a score that 228 calculates the average of the level of activity of each job the patient has held in his or her 229 lifetime (1 mild, 2 moderate, 3 intense), adjusted by the number of years spent in each job 230 (score between 1 and 3)), the BASFI increased by 8.9 points. "Occupation" has emerged as an 231 important health-related outcome in SpA. In fact, bulk of studies showed that the prevalence 232 of work disability is high in these patients and is associated with both clinical and psychosocial factors. <sup>13,14</sup> Specifically, labor-intensive jobs and manual professions were 233 associated with poorer work outcomes. 13,15 Yasemin Ul et al compared AS patients with 234 235 healthy controls and found that functional impairment assessed by BASFI was a significant

predictor of work instability scores. 16 In our study, we found that unemployed patients had

237 higher scores on the BASFI and LI measures. This observation may be attributed to their 238 inability to maintain employment due to functional limitations. 239 240 Regarding clinical examination, peripheral involvement was also significantly associated with 241 LI and BASFI score. Indeed, the more enthesitic involvement, painful and swollen joints, the 242 higher the BASFI score. 17 Coxitis was particularly associated with a higher LI and BASFI. This is in line with the literature. 18,19 It was, in fact, increased by 1.6 compared to patients 243 244 without coxitis and higher values were found not only for all BASFI questions that seemed to 245 assess functional impact on the hips (e.g. difficulty in getting up from the floor or a chair, tying shoes, climbing stairs), but also for those that did not assess gestures involving the hips 246 such as looking over the shoulder without turning around. 18 According to a recent study of 247 248 patients with axSpA, it has been suggested that hip involvement has a greater impact on 249 functional disability than axial structural damage. In addition, coxitis may affect the ability to accommodate a rigid lumbar spine, which may exacerbate functional limitations. <sup>20</sup> Coxitis 250 251 remains a major and dreaded prognostic location in SpA patients, especially in north African 252 population where it exists in higher prevalence.<sup>2</sup> Indeed, in our series, more than half the 253 patients had coxitis. Fortunately, the treatment response seems similar to that of Western countries.<sup>2</sup> These data underline the importance of early management of coxitis. As the hip is 254 255 a weight-bearing joint, its damage affects not only the patients' daily movements but also 256 walking and thus the entire spinal statics, and would, hence, contribute to the aggravation of 257 the deformities already present. 258 259 In our study, disease activity, as assessed by the BAS-G and BASDAI and ESR, was 260 significantly correlated with LI. The same was true for the BASFI score. The association with 261 BASDAI was found after multivariate analysis by linear regression, which underlines the 262 pivotal role of this factor in the functional impact of the disease. Studies have supported the results of this analysis stating that BASDAI was independently associated with BASFI. 21,22 263 264 The observed increase in functional impairment among patients with higher disease activity 265 may be in part explained by a concomitant decrease in spinal mobility. In fact, BASDAI was reported to be associated with BASMI.<sup>23</sup> 266 267 268 In our study, the BASMI was significantly associated with LI and BASFI score which has also been reported by other studies in the literature.<sup>24</sup> These data show that axial mobility 269 limitation is closely related to function in these patients. More importantly, The BASMI was 270

271 independently associated with the LI after multivariate linear regression in our study. Spine 272 mobility is a key clinical feature that should be assessed at baseline. It is strongly dependent 273 on both inflammatory activity and structural damage, and the BASMI was proved to be more 274 contributory in assessing this parameter compared to other scores. <sup>23</sup> Moreover, results from the DESIR cohort show that spinal inflammation is independently and positively associated 275 276 with BASMI.<sup>25</sup> 277 278 Inflammation, assessed with ESR values, has a significant anabolic effect on bone in SpA and 279 based on longitudinal data, it appears that effectively managing inflammation may decelerate the radiographic advancement of axial spondyloarthritis.<sup>26</sup> 280 281 282 This appears to be even more relevant as the association of both BASFI and LI with mSASSS 283 was found after multivariate analysis by linear regression in our study. In fact, structural damage leads to decreased spinal mobility and difficulty to carry out daily tasks. Studies have 284 285 concurred with the results of this analysis stating that mSASSS was independently associated with BASFI.<sup>21</sup> Ankylosis in spondyloarthritis is the latest stage of structural damage. 286 287 Although the natural course of the disease has changed positively over the past years, joint ankylosis is not rare and affects 20 to 50% of patients. <sup>26</sup> In our study, radiographic structural 288 289 damage was assessed using mSASSS. However, this score measures anterior damage of the 290 vertebrae and underestimates postero-lateral vertebral rim ankylosis such as ankylosis of the 291 facet joint. Interestingly, J-Y jung et al showed that facet joint ankylosis may be more associated with functional impairment than syndesmophytes. <sup>26</sup> Consequently, both anterior 292 293 and posterior structural damage should be investigated to gain a better understanding of the patient's condition.<sup>27</sup> 294 295 296 Grade of sacro-iliitis was positively correlated with both BASFI and LI in our study. The impact of sacro-iliitis on clinical outcomes and functional status has been supported in some 297 studies. 28,29 In the German spondyloarthritis inception cohort, structural damage of the sacro-298 299 iliac joints influenced functional status and spinal mobility. The results suggested that an 300 increase of one grade of radiographic sacroiliitis in a single joint is associated with a 301 deterioration of 0.10/0.12 points in BASFI/BASMI, respectively, regardless of disease activity and structural damage in the spine. <sup>28</sup> As for the therapeutic modalities, neither 302

NSAIDS nor DMARDS appeared in the multiple regression analysis.

304

305 Several studies in the literature have shown the involvement of NSAIDs in functional impairment. This was found in the study by Kroon FP et al<sup>30</sup> where the mean BASFI in the 306 NSAID group decreased by 9.1 points (5.1 to 13) compared with the no treatment group. 307 308 However, few studies have evaluated the effect of cDMARDs on the functional impact of the 309 disease. In a Turkish case control study which included 51 patients with Ankylosing 310 spondylitis, <sup>31</sup> no improvement in BASFI score was noted when comparing patients receiving 311 NSAIDs alone to those receiving them in combination with methotrexate. Similarly, no 312 improvement in BASFI score was found after 16 weeks of methotrexate.<sup>32</sup> Whereas for Gonzalez-Lopez L et al<sup>33</sup> the group receiving methotrexate had a better BASFI than the group 313 314 receiving the placebo. 315 316 TNF alpha inhibitors did not influence the functional outcome of SpA in our study, whereas 317 several authors have reported the beneficial effect of anti-TNF alpha on function during SpA.<sup>34</sup> This could be explained by the fact that only 7.76% of the patients in our study 318 319 received TNF alpha inhibitors. 320 321 Our study has some strengths including the sample size which provides a good basis for 322 reliable statistical analysis and the use of functional indexes that have been infrequently 323 utilized in similar studies. Furthermore, the disease duration of patients in our series ranged 324 from 1 to 62 years, enabling a credible assessment of functional impact. However, it is 325 important to state the weaknesses of our study: First, the retrospective nature of the study does 326 not allow us to establish a causal link and draw solid conclusions. However, there was very 327 little missing data in the files. Second, the disease was active (BASDAI >40) in 65.6% of 328 cases. This could be explained by a recruitment bias, as our study was carried out in a 329 university hospital treating severe forms of SpA. 330 331 Conclusion 332 Our findings are consistent with the existing epidemiological data, which report a high 333 prevalence of coxitis in north-african patients with spondyloarthritis. Employment status, 334 spinal mobility, disease activity and structural damage emerged as predictive factors 335 independently associated with functional impairment. Functional impairment was assessed 336 using two robust and valid scores: BASFI and LI. The latter, more frequently used in 337 European and French-speaking countries, has been used in particular in osteoarthritis patients 338 and has proven its relevance and usefulness in spondyloarthritis patients. Early and effective

339	treatment is crucial for achieving rapid and sustained remission in spondyloarthritis. This
340	should be defined, ideally, as remission across all clinical, biological and radiological
341	domains. Thankfully, emerging therapies are targeting all three domains and have, therefore,
342	revolutionized management of spondyloarthritis, providing new opportunities to optimize
343	patient outcomes. Prospective studies with a long-term follow-up of the patients are needed
344	for a better evaluation of the effect of these new therapeutic modalities on the functional
345	impact of the disease.
346	
347	<b>Author Contributions</b>
348	KM contributed to the conceptualization and correcting the original draft, final approval and
349	agrees to be accountable for all aspects of the work. IC was involved in writing the original
350	paper and also agrees to be accountable for all aspects of the work. HF and DBN were
351	responsible for the analysis. RB acquired the data. DK and WH were involved in supervison.
352	All authors approved the final version of the manuscript.
353	
354	Conflicts of Interest
355	The authors declare no conflict of interests.
356	
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359	
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 Table 1: Socio demographic and disease-related features

Socio-demograp	phics and disease-related	
features		
Age (years) mean± SD [min-max]		$38.9 \pm 12.7 [16-76]$
Sex ratio		2.7
Status	Single	119,(45.2)
n,(%)	Married	136,(51.7)
	Divorced	6,(2.3)
	Widow(er)	2,(0.8)

Level of education	Illiterate	33,(12.6)
n,(%) Primary level		95,(36.1)
	Secondary level	95,(36.1)
	Higher level	40,(15.2)
	(university)	
Professional	Salaried	116,(44.1)
occupation	employment	
n,(%)	Self-employed	27,(10.3)
	Unemployed	92,(35)
	Student	18,(6.8)
	Retired	10,(3.8)
Social class, n,(%)	Disadvantaged	62, (23.6)
	Middle class	174, (66.1)
	Privileged	27, (10.3)
Age at onset (years) me	ean± SD [min-max]	$27.6 \pm 10.8$ [5-61]
Disease duration (years	) mean± SD [min-	11.3 ± 9.5 [1-62]
max]		
Classification of SpA	axial	113,(43.1)
n,(%)	peripheral	115,(44)
	Axial and peripheral	117,(44.7)
Extra-articular manifest	tation, n,(%)	62,(23.6)

498 SD:standard deviation, n=number

## **Table 2:** Main Clinical and paraclinical data

Clinical a	nd para-clin	ical data	
Number of	f tender joints	s, mean± SD [min-	$1.3 \pm 2.5$ [0-16]
max]			
Number of	f swollen join	ts, mean± SD [min-	$0.2 \pm 0.8$ [0-7]
max]			
Coxitis n,(	%)		281,(53.5)
BASMI m	ean± SD [min	n-max]	$3.9 \pm 2.3  [0-10]$
MASES m	nean± SD [mi	n-max]	$1.9 \pm 2.8  [0-13]$
HLA B27	allele (%)		56.3
Disease ac		BASG-s	38.1 ± 30.8 [0-100]
		mean± SD [min-	
	7	max]	
	,	BASDAI, mean±	49.4 ± 30 [0-100]
		SD [min-max]	
Ť		ESR mm/h mean±	36.6 ± 26.3 [2-125]
		SD [min-max]	
	mSASSS m	ean± SD [min-max]	$15 \pm 18.2  [0-72]$
Imaging	Sacroiliitis	Grade 1/grade 2	1.9/ 12.6
findings	(%)	Grade 3/ grade 4	38.8 / 46.7
	BASRI	Hips	$3.3 \pm 2.3  [0-4]$
	mean± SD	Sacro-iliac	$3.14 \pm 0.9  [0-4]$
	[min-max]	Spine	$3.3 \pm 2.3  [0-8]$
		Total	$8.2 \pm 3.9$ [0-16]

Functional impact	BASFI, mean± SD	$45.8 \pm 25.7$ [0-100]
	[min-max]	
	LI, mean± SD [min-	$7.5 \pm 6.4  [0-22]$
	max]	

BASFI: Bath Ankylosing Spondylitis functional index, LI: hip Lequesne Index, BASMI: Bath Ankylosing Spondylitis metrology Index, BASDAI: Bath Ankylosing Spondylitis disease activity Index, BASG-s: Bath Ankylosing Spondylitis global score, ESR: erythrocyte sedimentation rate, BASRI: Bath Ankylosing Spondylitis Radiologic Index, MASES: Maastrich Ankylosing Spondylitis Enthesitis Score, mSASSS: the Modified stoke ankylosing spondylitis spinal score.

Table 3: Association of BASFI score and LI with the qualitative parameters

			DACEL (m)	III(a)
			BASFI (p)	LI (p)
Gender			0.799	0.104
Level of educa	tion		0.076	0.003
Professional oc	ccupa	ation	0.010	<0.001
Classification	of	Axial	0.157	0.485
SpA		Peripheral	0.006	0.021
HLA B27 allel	le		0.153	0.651
Site of the	Cer	vical spine	<0.001	<0.001
spinal pain	Dor	sal spine	0.003	0.011
	Lun	nbar spine	0.025	0.371
	But	tock pain	0.274	0.3
Coxo-	cox	itis	<0.001	<0.001
femoral joint bilateral		<0.001	<0.001	
Joint limitation		<0.001	<0.001	
BASMI			<0.001	<0.001
MASES			<0.001	0.108
BASG-s			<0.001	0.025
BASDAI	BASDAI		<0.001	<0.001
CsDMARDS Yes/no		0.587	0.051	
Type		0.280	0.541	
bDMARDS Yes/no		0.812	0.175	
Type		0.236	0.282	
Surgical treatm	nent (	(THR)	0.019	<0.001

BASFI: Bath Ankylosing Spondylitis functional index, LI: hip Lequesne Index, BASMI: Bath Ankylosing Spondylitis metrology Index, BASDAI: Bath Ankylosing Spondylitis disease activity Index, BASG-s: Bath Ankylosing Spondylitis global score, BASRI: Bath Ankylosing Spondylitis Radiologic Index, MASES: Maastrich Ankylosing Spondylitis Enthesitis Score, CsDMARDS: Conventional synthetic disease-modifying antirheumatic drugs, bDMARDS: biologic disease-modifying antirheumatic drug, THR: total hip replacement, p: Significance one-factor ANOVA test

**Table 4:** Correlation of BASFI score and LI with the quantitative parameters

		BASFI		LI	
			P	r	P
Disease dura	ation	0.175	0.007	0.248	<0.001
BASMI		0.501	<0.001	0.573	<0.001
BASG-s		0.299	<0.001	0.141	0.025
BASDAI		0.46	<0.001	0.278	<0.001
ESR		0.253	0.001	0.215	0.003
mSASSS		0.238	0.001	0.212	0.003
BASRI	BASRI hips		< 0.001	0.610	<0.001
	Sacro-iliac	0.277	< 0.001	0.307	<0.001
	spine	0.286	<0.001	0.209	0.001
total		0.385	<0.001	0.468	<0.001
Grade of Sa	Grade of Sacroiliitis		<0.001	0.353	<0.001

BASFI: Bath Ankylosing Spondylitis functional index, LI: hip Lequesne Index, BASMI: Bath Ankylosing Spondylitis metrology Index, mSASSS: the Modified stoke ankylosing spondylitis spinal score, BASDAI: Bath Ankylosing Spondylitis disease activity Index, BASG-s: Bath Ankylosing Spondylitis global score, BASRI: Bath Ankylosing Spondylitis Radiologic Index, ESR: Erythrocyte sedimentation rate, r: Pearson correlation coefficient, p: Significance one-factor ANOVA test

**Table 5:** Multiple linear regression of factors associated with BASFI score and LI

	BASFI	BASFI			LI		
	β	95% confidence	р	β	95% confidence	р	
		interval of β[]			interval of β[]		
Profession	4.20	[-5.66,14.08]	0.396	2.72	[0.880, 4.564]	0.004	
BASMI	0.70	[-2.60, 4.01]	0.669	1.15	[0.626, 1.679]	< 0.001	
mSASSS	0.29	[0.05, 0.53]	0.016	-0.06	[-0.118, -0.001]	0.046	
BASDAI	0.62	[0.42, 0.83]	< 0.001	0.01	[-0.031, 0.061]	0.524	

BASFI: Bath Ankylosing Spondylitis functional index, LI: hip Lequesne Index, BASMI: Bath Ankylosing Spondylitis metrology Index, mSASSS: the Modified stoke ankylosing spondylitis spinal score, BASDAI: Bath Ankylosing Spondylitis disease activity Index.