CASE SERIES



Impact of COVID-19 on the pain and disability of patients with adult spinal deformity

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Received: 16 October 2020 / Accepted: 16 February 2021 / Published online: 2 March 2021 © Scoliosis Research Society 2021

Abstract

Purpose To evaluate the pain and functional effect of the COVID-19 pandemic on patients with ASD reflected by their response to SRS-22, ODI, and SF-36 questionnaires.

Methods Patients who had stable pain and functional outcome scores over the preceding 2 years were enrolled in a local prospectively collected adult spinal deformity (ASD) database. A reanalysis of their SRS22, ODI and SF-36 data 14 days into confinement were compared to their last pre-confinement scores.

Results 89 patients were included in this study (average age 60.7 years, 91% female) with an average time from last FU until confinement of 9.6 months. The ODI total score worsened by 5 points post-confinement with no difference seen in personal care, walking and social life. In contrast, the SRS-22 score showed small improvements in function/activity and satisfaction, but no significant differences for the other domains. Similarly, the SF-36 showed small improvements in physical function, physical and emotional role, vitality and PCS.

Conclusion The global COVID-19 pandemic and ensuing confinement had variable overall effects on ASD patients, without the expected marked worsening. In addition, this study illustrates that the SRS-22 questionnaire is less influenced by environmental and psychological factors than the ODI supporting its objectivity and accuracy in the evaluation of the QoL of ASD patients.

Keywords Spine · COVID-19 · Coronavirus · Surgery · Scoliosis

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Introduction

Spinal conditions are a major cause of pain and disability worldwide. Chronic spinal pathology, such as adult spinal deformity (ASD) is common, affecting a third of patients aged over 50 years and two-thirds of patients aged over 70 years [1]. ASD is known to affect the quality of life (QoL), functional capacity and pain of patients [2, 3]. In fact, ASD has the worst patient reported QoL of all common chronic medical conditions, including arthritis, chronic lung disease, congestive heart failure, diabetes and ischaemic heart disease [4]. However, various factors may influence the degree of this disability and QoL including psychological distress [5–7].

2020 saw a pandemic infection from COVID-19. The effect of this pandemic is far reaching and includes marked psychological distress to populations worldwide [8-10]. In fact, COVID 19 has incited such fear that new scales, such as the fear of COVID-19 scale, have been necessitated

to understand a population's phobia [11]. This includes patients personally unaffected by the virus.

Clearly, the psychological effect of COVID-19 has also affected patients with ASD, even in those not personally infected. This pandemic has, therefore, provided an unprecedented opportunity to further understand the effect of a global event on the pain, function and QoL of patients with ASD. It may be predicted that the overwhelming psychological effect of COVID-19 may improve pain and function in these patients as their perspective shifts from their spinal pathology to a more imminent risk. Alternatively, the stress exerted by COVID-19 may exacerbate symptoms. Compounding this, populations were confined to their homes, permitting less physical exertion on their spinal deformity, but potentially more sedentary time to focus on their health.

Thus, the aim of this study was to determine the effect of the COVID-19 pandemic on the pain, function and QoL of patients with ASD not personally infected by COVID-19 in France during confinement, and in doing so, to explore whether variations in responses to questionnaires in the undercurrent condition are affected by non-spinerelated issues.

Methods

Consecutive patients included in this study were extracted from a prospective database of patients who had previously undergone deformity correction for ASD. The inclusion criteria were: patients with degenerative or idiopathic ASD, defined as a coronal Cobb angle > 20° , thoracic kyphosis > 60° , sagittal vertical axis (SVA) > 5 cm or pelvic tilt (PT) > 25° treated with deformity correction; a minimum of 2 years follow-up post-surgery, with "clinical stability" during the last two follow-ups prior to confinement and the last follow-up visit being within 1 year of the start of confinement. Patients were excluded if they had been previously or were currently diagnosed as COVID-19 positive.

"Clinical stability" over the last two follow-ups was defined using the Oswestry Disability Index (ODI), Scoliosis Research Society 22 (SRS-22) and Short Form 36 (SF-36) Physical Component Score (PCS). Only patients with a difference in ODI less than or equal to 15 points [ODI last follow-up (FU)-ODI previous FU] and a variation in the absolute value of SRS-22 total score and SF-36 PCS [(Last FU-Previous FU)/Previous FU] less than or equal to 20% were considered to have "clinical stability".

All patients were from the same geographical area, affected by the same governmental restrictions and answered all three questionnaires within 15 days of confinement.

Statistical analysis

Cross-tabulation was generated and either a Fisher exact or Pearson Chi² test was used to compare all the distributions. Paired *t* tests were used to assess differences in means for the same cohort between different follow-up time points. A *p* value < 0.05 was considered statistically significant.

Results

89 patients were included in this analysis (average age 60.7 years (SD 16.8), 91% female). The average time from last FU until confinement was 9.6 months (SD 4.9). The comparison between the last pre-confinement FU and post-confinement FU is shown in Table 1.

Overall the ODI, which is an index derived to quantify disability from low back pain, was seen to worsen by five points post-confinement with no difference seen in personal care, walking and social life. In contrast, the SRS-22 score, which is the most widely used patient reported outcome score for spinal deformity, showed small improvements in function/activity and satisfaction, but no significant differences for the other domains. Similarly, the SF-36, which is a health-related quality of life score, showed small improvements in physical function, physical and emotional role, vitality and PCS.

Discussion

To our knowledge, this is the first study to assess the effects of a global pandemic on patients with ASD by analysing its consequences on health-related QoL scores. We hypothesised that confinement would have detrimental psychological impacts and a lack of rehabilitation that would result in patients experiencing worsened pain, function and QoL. This hypothesis was confirmed when using the ODI, with the overall score, pain, lifting, sitting, standing, sleeping, sex life and travelling all being rated statistically (p < 0.05) and clinically (> 10% change) significantly poorer during confinement. Furthermore, an unchanged social life on ODI scores does not correlate with the societal effects of confinement during this time.

However, both the SRS-22 and SF-36 questionnaire had no significant change in the patient's reported pain and showed improvements in the SRS-22 function and satisfaction, as well as the SF-36 physical function, emotion and vitality. This contrasted our expectation and necessitates a reconsideration of the detrimental effect of confinement during a global crisis. In addition, it may raise the question

Table 1 Comparison of outcome scores pre- and post-confinement

M	NT

	Mean	N	SD	Р
ODI				
Pain intensity				0.003
Last follow-up	1.1	89	1.1	
Post confinement	1.4	89	1.3	
Personal care				0.661
Last follow-up	0.4	89	0.9	
Post confinement	0.5	89	1.0	
Lifting				0.000
Last follow-up	2.3	89	1.5	
Post confinement	3.2	89	1.7	
Walking				0.680
Last follow-up	0.9	89	1.3	
Post confinement	0.9	89	1.3	
Sitting				0.014
Last follow-up	0.8	88	1.0	
Post confinement	1.1	88	1.4	
Standing				0.002
Last follow-up	1.3	89	1.3	
Post confinement	1.9	89	1.7	
Sleeping				0.043
Last follow-up	0.5	89	0.8	
Post confinement	0.8	89	1.3	
Sex life				0.004
Last follow-up	0.3	45	0.7	
Post confinement	0.8	45	1.6	
Social life				0.171
Last follow-up	1.0	89	1.3	
Post confinement	0.8	89	1.2	
Travelling	0.0	07		0.025
Last follow-up	0.9	89	1.1	01020
Post confinement	1.2	89	1.6	
Total score	1.2	07	1.0	0.000
Last follow-up	20.1	89	15.2	0.000
Post confinement	25.8	89	20.5	
SRS-22	25.0	07	20.5	
Function/activity				0.016
Last follow-up	3.9	89	0.9	0.010
Post confinement	3.9 4.1	89	0.9	
Pain	4.1	09	0.7	0.870
Last follow-up	3.7	89	1.0	0.879
Post confinement	3.7	89	1.0	
	5.0	09	1.0	0.052
Self-image/appearance	2.0	80	0.0	0.052
Last follow-up Post confinement	3.8	89 80	0.8	
	3.7	89	0.9	0.220
Mental health	2.0	00	0.0	0.220
Last follow-up	3.8	89	0.9	
Post confinement	3.9	89	0.9	0.004
Satisfaction with management		00	0 -	0.004
Last follow-up	4.4	89	0.7	
Post confinement	4.6	89	0.7	

	Mean	Ν	SD	Р
Total score				0.314
Last follow-up	3.9	89	0.7	
Post confinement	3.9	89	0.7	
SF-36				
Physical function				0.012
Last follow-up	44.9	89	9.2	
Post confinement	46.5	89	8.8	
Physical role				0.00
Last follow-up	45.8	89	10.9	
Post confinement	49.7	89	10.9	
Pain				0.05
Last follow-up	46.5	89	9.3	
Post confinement	48.2	89	11.0	
Vitality				0.00
Last follow-up	49.4	89	10.1	
Post confinement	46.0	89	9.9	
Social functioning				0.76
Last follow-up	48.9	89	9.8	
Post confinement	48.6	89	11.4	
Emotional role				
Last follow-up	47.1	89	11.5	
Post confinement	52.5	89	8.6	
Mental health				
Last follow-up	48.5	89	10.5	
Post confinement	49.1	89	11.9	
General health				
Last follow-up	51.9	89	9.2	
Post confinement	53.9	89	1.0	
PCS				0.024
Last follow-up	46.6	89	9.2	
Post confinement	48.4	89	10.1	
MCS				0.55
Last follow-up	49.6	89	10.5	
Post confinement	50.3	89	9.6	

Table 1 (continued)

of the specificity of each score as an objective indicator of the QoL of spinal deformity patients [12-14]. Considering the "clinical stability" of these patients, the psychological influence of the pandemic and staying at home during the lockdown period can be considered the main factors that influenced any modification in the score results.

Clearly, these results cannot be extrapolated to times when normal daily activity is feasible and does not represent potential benefits for patients to remain at home permanently, but rather offers insight into the effect of the COVID-19 pandemic on the ASD community. In addition, this study cannot be extrapolated to patients affected by back pain without underlying deformity, where the interpretation

would wrongly suggest potential benefits of remaining at home.

In addition, this study has a multitude of limitations that need to be recognised when interpreting the results. First, only operative patients were included. Second, despite the previous 2 years FU showing stable scores and it being felt unlikely that the spinal related complications arose during the time between latest FU and confinement, particularly considering that complications would have impacted all three questionnaires simultaneously, radiological follow-up was not performed to assess for complications [14]. Third, there is no stratification of age, pulmonary status or radiographic severity of spinal deformity that could be influenced by the COVID-19 infections. Fourth, this is a single-centre study, therefore, differences among other countries and territories due to the differences of the COVID-19 pandemic need to be considered. Lastly, we did not correlate the outcome scores to any psychological well-being score.

Conclusion

The global COVID-19 pandemic and ensuing confinement had variable overall effects on patients, without the expected marked worsening. In addition, this study illustrates a higher specificity for the SRS-22 questionnaire when compared to the ODI in assessing ASD patients with less influence from environmental or psychological factors, emphasising its objectivity and accuracy in the evaluation of the QoL of ASD patients.

Acknowledgements Glynny Kieser for her editorial input.

Author contributions DCK: acquisition, conception, design, analysis, interpretation, draft, critical revision, approved version, agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved; AB, IO: acquisition of data, design, critical revision, approved version, agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved; DTC, DL, KH, SJ, JP, LB: acquisition of data, critical revision, approved version, agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the accuracy or integrity of any part of the work are appropriately investigated and resolved; DTC, DL, KH, SJ, JP, LB: acquisition of data, critical revision, approved version, agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Funding No funding was received for this work.

Data availability On request from the corresponding author.

Declaration

Ethical approval This study was approved by the IRB of Pellegrin Hospital (CE-GP-2020-31) and the study was performed in accordance with the ethical standards as laid down in the 1964 Declaration of Helsinki.

Consent to participate Informed consent was obtained from all individual participants included in the study.

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