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Research letter

Sleep apnoea management in Europe during the COVID-19 pandemic: data from the European Sleep Apnoea Database (ESADA)

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Research letter

Sleep apnoea management in Europe during the COVID-19 pandemic – data from the

European Sleep Apnoea Database (ESADA)

Short title: OSA management during COVID-19 pandemic

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Research letter

Sleep disordered breathing (SDB) is highly prevalent with a male to female predominance of two to one, and is more common in middle-aged and elderly subjects (1). Affected patients often present with comorbidities such as obesity, cardiovascular disease (systemic hypertension, heart failure, atrial fibrillation), and diabetes mellitus Type II (2). The strong overlap between the profile for SDB patients and the identified risk factors for adverse outcomes of COVID-19 infection that include age, male gender, and cardio-metabolic comorbidity (3) suggest that SDB patients may benefit from effective therapy if confronted with COVID-19 infection (4).

The COVID-19 pandemic has fundamentally changed the operation of health care systems around the globe. Resources have understandably been shifted from elective care to acute management of severely ill patients with life-threatening COVID-19 infection (5). According to current recommendations, sleep medicine services are advised to reduce in-house services, and to provide medical care by remote contact using phone, video-calls, and telemedicine solutions (6, 7, 8, 9).

In the current study, we assessed the impact of the COVID-19 pandemic on the management of SDB patients in Europe. We approached the centres of the European Sleep Apnoea Database (ESADA) cohort, a well-established network of sleep centres in 19 European countries (2), and a subsample of accredited sleep centres of the German Sleep Society (DGSM). The aim was to analyse how recommendations from expert organisations were applied across various European regions, and whether specific mitigation strategies were already in practice.

A purpose-built questionnaire addressed information about changes in clinical routines imposed during the COVID-19 pandemic compared with routine practices. Specifically, details on the diagnosis of SDB, the procedures for titration of positive airway pressure (PAP) treatment, and the follow up of PAP treatment were addressed. One question sought the estimated proportion of staff still active in the sleep service during the pandemic compared with beforehand (0-100%, each for "physicians" and "nurses/technicians"). Finally, access to regional or national COVID-19 related guidelines for patients with SDB and for caregivers in sleep medicine services was evaluated. Actual numbers of confirmed COVID-19 infected inhabitants and reported COVID-19 deaths were computed per million inhabitants for each country based on information provided by the Johns Hopkins University COVID-19 statistics website (https://coronavirus.jhu.edu/map.html, April 12, 2020).

Data were provided by 25 of 29 ESADA centres and 15 of 283 DGSM-accredited sleep centres. Most centres were linked to a pulmonary department (**table**) and all bar one were hospital based. In 31 of the total 40 participating centres, patients were unable to physically attend because of sleep centre or travel restrictions.

Diagnostic routines were changed substantially by the pandemic. Prior to the pandemic, laboratory-based polysomnography was performed "rarely or regularly" in 92.5% of centres but only in 20% during the pandemic (p<0.001, **table**). On the other hand, telemedicine-based sleep apnoea diagnosis was used in 30.0% of centres prior to the pandemic and this number was only marginally reduced to 27.5% during the pandemic. From the 10 centres regularly using telemedicine-based diagnosis, three stopped the service during the pandemic, four maintained the level of use, and three increased use from rarely to regularly. Only two centres started this routine during the pandemic.

Prior to the pandemic, laboratory-based CPAP or bi-level PAP titration and initiation were practiced in almost all centres but less than one fifth of centres continued this routine during the pandemic (p<0.001, **table**). In the 13 centres already practicing telemedicine-based CPAP titration, four stopped the service, one reduced the use, six maintained the

level of use, and two increased the use from rarely to regularly. Four centres started this routine during the pandemic.

Prior to the pandemic, 39 centres report regular follow-up procedures in patients with SDB. This service continued in 36 centres during the pandemic but the mode of follow up had changed (**table 1**). Specifically, three centres offered in-laboratory and seven centres performed ambulatory follow-up sleep recordings in selected patients. Thirty centres provided follow-up by phone or video-calls. Out of the nine centres without phone access, two practiced telemedicine follow-up and four centres provided the support via home care providers. The remaining three centres offered no access to follow-up during the pandemic. Telemedicine based follow up was practiced by a minority of centres.

Staffing levels in the sleep medicine service was reduced to 25% (Interquartile range (IQR) 40) for physicians and to 19% (IQR 28) for nurses/technicians compared to pre-pandemic levels. Staffing reductions varied across European regions, being least in Northern and most in the Southern and Central parts of Europe. However, there was no association identified between remaining staff and numbers of infected individuals or numbers of deaths relating to COVID-19 infection in each country. In addition, national recommendations or guidelines for patients with SDB were available in 45% of centres (0% in Eastern, 39% in Central, 60% in Southern and 100% in Northern and Western European centres). Only 28% of centres had guidelines for sleep service personnel.

This descriptive report highlights several important findings. First, sleep medicine services have been reduced by almost 80% during the first 1-2 months of the COVID-19 pandemic in Europe. Second, more comprehensive sleep studies using polysomnography or in-laboratory PAP titrations have been completely interrupted, or practiced only to a very limited extent in highly selected patient groups. Third, commencement of treatment for SDB by various

types of PAP therapy is equally reduced in the vast majority of centres and countries. Fourth, patient follow-up is mainly managed by phone based patient contacts. Fifth, the full potential of mitigation strategies available by telemedicine has not been explored .

The sharp reduction in sleep medicine services in this study was expected and corresponds to the practice in other areas of medicine (3). Prevention of virus spread at the sleep centres such as by PAP-induced aerosol spread (10) as well as quarantine restrictions are plausible explanations for this decline. Nonetheless, we identified sizeable variations within and between countries in the lockdown of sleep medicine services. Rather unexpectedly, the use of telemedicine and other innovative technical solutions including disposable diagnostic tools and non-contact sleep surveillance for sleep apnoea diagnosis, were not reported to any major extent. Uncertainty relating to data protections laws and inadequate practical experience are believed to limit dissemination of such new technology.

There is a medical need for the continued management of SDB patients. Most clinical case series regarding severely affected COVID-19 patients report clinical features including male predominance, obesity, and cardiometabolic disorders (3), all of which are strongly associated with SDB. Furthermore, there are as yet unexplored potential mechanistic links between an imbalance of the Angiotensin II receptor and Angiotensin II Converting Enzyme (ACE2) and severe COVID-19 infections, which may also apply in SDB (11, 12).

In conclusion, our findings suggest that the sleep medicine community needs to collaborate in developing strategies for care of patients with both suspected and established SDB during major events such as the COVID-19 pandemic. Activities need to focus on the recognition of severe cases of SDB and how to initiate treatment in already identified severe cases. The potential of new technologies that enable remote monitoring to optimize treatment may be more frequently applied during times of restricted health care resources in order to obtain better management of severely affected COVID-19 patients.

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Abbreviations

AASM: American Academy of Sleep Medicine

BTS: British Thoracic Society

CI: Confidence Interval

ESADA: European Sleep Apnoea Database

IQR: Interquartile range

OSA: Obstructive sleep apnoea

PG: Polygraphy

PSG: Polysomnography

SD: Standard Deviation

SPOR: Swedish Perioperative Registry

TABLES

Table 1: Characteristics of the participating sleep centres and change of activities duringthe COVID-19 pandemic

Variable	Result		Commentary	
Number of centres	40		European Sleep Apnoea Database (ESADA) (25), German Sleep Society (15)	
Distribution across Europe	5 regions, 15 countries	8	North (3), West (2), Central (22), South (10), East (3)	
Clinical affiliation of the sleep centre	Pulmonary Departmen any other department (Other departments (18): Neurology (1), ENT (3), Cardiology (1), Independent unit at hospital (12), Independent unit outside a hospital (1)	
Patient spectrum of the slee centre				
Sleep apnea diagnostic proce	dures used prior to Covid-19) pandem	ic (Answer: Rarely or regularly)	
Polysomnography in lab 92.5%	Polygraphy at home 87.5%		Telemedicine based 30.0%	
Sleep apnea diagnostic proce	dures during Covid-19 pando	emic		
Polysomnography in Lab 20.0%	Polygraphy at home 32.5%	Teleme 27.5%	dicine based	
CPAP treatment start proced	ures used prior to Covid-19 p	pandemic	: (Answer: Rarely or regularly)	
In lab titration – 90.0%	Ambulatory titration – 55.0%		dicine based APAP titration – 32.5% rly 6 centres)	
CPAP treatment start proced	ures used during the Covid-1	.9 pander	nic (Answer: Rarely or regularly)	
In lab titration – 17.5%	Ambulatory titration – 22.5% (all rarely)		dicine based APAP titration – 32.5% rly 4 centres)	
BiLevel-PAP treatment start procedures used prior to the Covid-19 pandemic (Answer: Rarely or regularly)				
In lab titration – 87.5%	Ambulatory titration –		Telemedicine based BiLevel-PAP titration – 20.0% (Regularly 4 centres)	
BiLevel-PAP treatment start procedures used during the Covid-19 pandemic (Answer: Rarely or regularly)				
In lab titration – 17.5%			Telemedicine based BiLevel-PAP titration 12.5% (Regularly 3 centres)	
Follow up routines for PAP treatment prior to the Covid-19 pandemic (Answer: Rarely or regularly)				
In lab follow up – 82.5%	Ambulatory titration -	Distanc	e follow up:	

	92.5%	Base on phone calls: 70% Based on telemonitoring – 50.0% ("Regularly use" at 7 centers)		
Follow up routines for PAP treatment during the Covid-19 pandemic (Answer: Rarely or regularly)				
In lab Follow up – 7.5%	Ambulatory titration – 17.5% (all rarely)	Distance follow up: Base on phone calls: 75.0% Based on telemonitoring – 57.5% ("Regularly use at 12 centers, 8 centers started, 4 centers stopped telemonitoring for PAP follow up)		

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*= center provided data for the study

Accredited sleep centres of the German Sleep Society (DGSM) participating in the study

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Aachen	J Schiefer	Sleep Centre, Neurological Department, University Hospital
Arolsen	W Knüppel	Sleep Centre, Krankenhaus Bad Arolsen
Berlin	H Danker-	Competence Centre for Sleep Medicine, Charité
	Hopfe	Universitätsmedizin Berlin, Campus Benjamin Franklin
Düsseldorf	M Minuth	Florence-Nightingale Krankenhaus, Kaiserswerther Diakonie
Essen	G Nilius	Sleep Centre, Pulmonary Medicine, Kliniken Essen Mitte
Frankfurt	H Schneider	Zentrum für Schlafmedizin Frankfurt
Garmisch	R Püschel	Klinikum Garmisch-Partenkirchen
Halle	S Schädlich	Krankenhaus Martha-Maria Halle
Hannover	K Meyer	Sleep Centre, Medizinischen Hochschule
Heidelberg	A Benz	Thoraxklinik Heidelberg, University Hospital
Landshut	B Schneider	Sleep Centre, Pediatric Hospital St. Marien
Nordhausen	J Büntzel	Sleep Centre, Suedharz Klinikum
Oldenburg	l Koper	Sleep Centre at Sana Klinik Oldenburg
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