Retrospective adherence data was collected from patients prior to remote monitoring.

Results Post establishment, initial data (3 patients) showed variable overnight adherence with mean use 38 minutes, 47 minutes and 7 hours 26 minutes respectively. Following the first phone contact with the parents, a personalized plan was agreed and arranged. The aim of each plan addressed issues such as adjusting/changing the mask interface, humidification and parental encouragement. Further scheduled contacts will occur on a personalized basis. Parents will receive a satisfaction questionnaire at the end of the monitoring period.

Discussion Remote monitoring technology has the potential to guide adjustments in NIPAPT therapy, monitor and improve adherence and reduce financial burden of hospital based review. Our preliminary work shows high uptake. We await results of the patient satisfaction questionnaire and cost breakdown following pilot study completion.

P034 A NOVEL APPROACH TO QUANTIFY SLEEP-RELATED RHYTHMIC MOVEMENT DISORDER USING AUTOMATIC 3D ANALYSIS

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Background Unlike other episodic sleep disorders, there are no agreed severity indices for sleep-related rhythmic movement disorder (RMD). Rhythmic movements (RMs) can be characterized by polysomnography, but sensor placement may inhibit movements. Actigraphy and 2D video can be used in the home, but have limitations. Actigraphy does not differentiate RMs from other movements, while manual scoring of 2D video is laborious.

Abstract P034 Figure 1 RM time of night distribution plot. Each data point shows how many 1.5 s segments are classified as RMs per 30 minutes, combining data across subjects. Automatic 3D analysis and manual 2D annotations show high agreement

We developed a sensitive method to detect and quantify RMs using automatic 3D video analysis.

Method Children with RMD (n=6, 4 male) aged 5–14 years were studied for two nights in a sleep laboratory. A ceiling-mounted camera captured 3D depth images, while another recorded 2D video, from lights off until lights on. We developed algorithms to analyze the characteristics of RMs and built a classifier to distinguish rhythmic from non-rhythmic movements based on 3D video data alone. Data from 3D automated analysis were compared to manual 2D video annotations in 1.5s segments to assess algorithm performance [figure 1]. Novel indices were developed: the RM index, frequency index and duration index to better characterize RMD severity.

Result Automatic 3D analysis demonstrated high levels of agreement with the manual approach (Cohen's Kappa >0.9; F1-score >0.9). We also demonstrated how RM assessment can be improved using plots of our novel indices for ease of visualization.

Conclusion 3D video technology is widely available and can be integrated into sleep laboratories. Our automatic 3D video analysis algorithm yields reliable quantitative measurement of RMs, reducing the burden of manual scoring. Furthermore, our novel RMD severity indices offer standardized measures of utility to clinical and research practice.

P035 THE UTILISATION OF A REMOTE MONITORING SYSTEM FOR THE IMPROVEMENT OF PATIENT CPAP COMPLIANCE

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Introduction First-line treatment for obstructive sleep apnoea hypopnoea syndrome (OSAHS) is continuous positive airway pressure (CPAP). Compliance with CPAP is essential for treatment to be effective. Excess air leakage from the CPAP mask can be uncomfortable for patients and is an important independent predictor of CPAP compliance. As the incidence of OSAHS is rising, novel treatment pathways must be utilised to maintain service standards whilst also maximising CPAP usage. The aim of this study was to determine whether a novel remote monitoring pathway could decrease excess CPAP mask leak and increase usage of the CPAP machine.

Methods Patients (n=48) started on CPAP were allocated to receive either Standard Care (SC) (n=22) or Remote Monitoring (RM) (n=26) follow-up according to their appointment time. Nurses consulted software (AirviewTM; Resmed Ltd) three times per week to review RM patients and contacted them if required. Additional contact could be initiated by patients from either group at any time. Patients were followed-up 4–6 weeks after CPAP issue; SC patients were reviewed in hospital, whilst RM patients were reviewed by telephone. Data was collected on median mask leak (L/min) and mean compliance (hrs/night).

Results There was no significant difference in compliance between SC (4.76±2.39 hrs) and RM (5.20±1.78 hrs) groups (p=0.486) (figure 1). There was a significant difference in mask leak between SC (13(8–25) L/min) and RM (3(1–7) L/min) groups (p<0.001) (figure 2).