Table 13: Telemonitoring and in person follow up versus in person follow up – Severe OSAHS

| Quality assessment | | | | | | | No of patients | | | Effect | Quality | Importance |
|--|----------------|------------------|-----------------------------|--------------------------------------|----------------------|----------------------|--------------------------------------|---------|-------------------------|--|---------------------|------------|
| No of studies | Design | Risk of bias | Inconsistency | Indirectness | Imprecision | Other considerations | Telemonitoring + in person follow up | Control | Relative (95% CI) | Absolute | | • |
| Systolic blood pressure - morning (follow-up 3-6 months; Better indicated by lower values) | | | | | | | | | | | | |
| 2 | | , | no serious inconsistency | no serious indirectness | none | none | 209 | 197 | - | MD 0.33 higher (3.1 lower to 3.75 higher) | ⊕⊕OO LOW | IMPORTANT |
| Adherenc | e- h per day (| follow-up | 3 - 12 months; ra | nge of scores: 0 | -8; Better indica | ated by higher val | ues) | | | | | |
| 6 | | very serious¹ | very serious ³ | serious indirectness ⁴ | serious ² | none | 200 | 205 | - | MD 0.6 higher (0.12 lower to 1.31 higher) | ⊕OOO VERY LOW | IMPORTANT |
| Adherenc | e-on nights P | AP used(| h per day) (follow | -up mean 2-3 mo | onths; range of | scores: 0-8; Bette | r indicated by higher va | alues) | | | | |
| 2 | | | no serious inconsistency | serious indirectness ⁴ | serious ² | none | 48 | 46 | - | MD 1.22 higher (0.03 lower to 2.48 higher) | ⊕OOO VERY LOW | IMPORTANT |
| Mean % n | ights CPAP u | ise >4 hoι | irs (follow-up mea | an 2 months; ran | ige of scores: 0 | -100; Better indica | ted by higher values) | | | | | |
| 1 | | | no serious inconsistency | serious indirectness ⁴ | serious² | none | 20 | 20 | - | MD 15 higher (4.03 lower to 34.03 higher) | ⊕OOO VERY LOW | IMPORTANT |

| | randomised | very | no serious | serious | | none | 28 | 26 | - | MD 10 higher (10.81 | \oplus OOO | IMPORTAN |
|-------|----------------------|------------------------------|--------------------------|--------------------------------------|---------------------------|--------------------|-------------------------|------------|---|---|--------------|----------|
| | trials | serious ⁴ | inconsistency | indirectness ⁴ | serious ² | | | | | lower to 30.81 higher) | VERY LOW | |
| ality | of life (Physica | l compos | ite)difference (fo | llow-up mean 4 r | months; range o | f scores: 0-100; E | etter indicated by high | er values) | | | | |
| | randomised trials | very serious ¹ | no serious inconsistency | serious indirectness ⁴ | | none | 40 | 42 | - | MD 0.3 higher (3.1 lower to 3.7 higher) | ⊕000 VERY | CRITICAL |
| | | | | | very serious ² | | | | | | LOW | |
| ality | of life (mental) | difference | e (follow-up mea | n 4 months; rang | ge of scores: 0-1 | 00; Better indicat | ed by higher values) | | | | | |
| | randomised trials | very serious ¹ | no serious inconsistency | serious indirectness ⁴ | very serious ² | none | 40 | 42 | - | MD 0 higher (4.15 lower to 4.15 higher) | ⊕000 VERY | CRITICAL |
| | | | | | | | | | | | LOW | |
| ality | of life EQ5D (fo | ollow-up n | nean 3 months; r | range of scores: | 0-1; Better indic | ated by higher va | lues) | | | | | |
| | randomised trials | very serious ¹ | no serious inconsistency | serious indirectness ⁴ | | none | 52 | 48 | - | MD 0 higher (0.07 lower to 0.07 higher) | ⊕000 VERY | CRITICAL |
| | lilais | serious | inconsistency | liidirectriess | very serious ² | | | | | lower to 0.07 fligher) | LOW | |
| ality | of Life-GHQ12 | (follow-up | mean 12 month | s; range of score | es: 0-12; Better | indicated by lowe | r values) | | | | | |
| | randomised | very | no serious | serious | very serious ² | none | 39 | 49 | - | MD 0.2 higher (2.31 | ⊕000 | CRITICAL |
| | trials | serious ¹ | inconsistency | indirectness ⁴ | | | | | | lower to 2.71 higher) | VERY LOW | |
| epin | ess Epworth (E | SS) (follo | w-up mean 2-12; | range of scores | : 0-24; Better inc | dicated by lower | values) | | | | | |
| | randomised | very | no serious | serious | | none | 127 | 137 | - | MD 0 higher (1 lower | | IMPORTAN |
| | trials | serious ¹ | inconsistency | indirectness ⁴ | no serious | | | | | to 1 higher) | VERY LOW | |

| _ | randomised trials | very serious ¹ | , | serious indirectness ⁴ | very serious ² | none | 87 | 95 | | MD 0.44 lower (3.21 lower to 2.33 higher) | | IMPORTANT |
|---|----------------------|------------------------------|---|--------------------------------------|---------------------------|------|----|----|--|---|--|-----------|
| Functional outcome of sleep A. questionnaire (follow-up mean 2 months; Better indicated by higher values) | | | | | | | | | | | | |
| 1 | randomised trials | | | serious indirectness ⁴ | very serious ² | none | 20 | 20 | | MD 0.8 higher (2.06 lower to 3.66 higher) | | CRITICAL |
| Mortality | | | | | | | | | | | | |
| Not reported | | | | | | | | | | | | CRITICAL |

¹ Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias 2 Downgraded by 1 increment if the confidence interval crossed one MID or by 2 increments if the confidence interval crossed both MIDs, MID for machine usage (adherence)-1 hour; MID for Systolic and Diastolic BP – 5 mm hg. For mean % of nights that the CPAP was used >4 hours outcome, clinically important difference was considered to be 10 % or 1 hour.. Established MIDs for SF-36 physical/mental- 2/3; ESS- 2.5; EQ5D- 0.03; FOSQ- 2GRADE default MIDs (0.5XSD) used for all other continuous outcomes.

³ Downgraded by 1 or 2 increments because: The point estimate varies widely across studies, unexplained by subgroup analysis. The confidence intervals across studies show minimal or no overlap, I2=50% unexplained by subgroup analysis. Subgroup analyses were tested for BMI < or >30 kg/m^2 , ESS < or >9, coexisting conditions, high risk occupation and type of treatment. Random effects analysis used.

⁴ Downgraded by 1 or 2 increments because the majority of the evidence included an indirect or very indirect population respectively. The study included a mixed OSHAS severity population based on mean baseline AHI.