Bilag 8a. Summary of Findings

Kortvarig landbaseret træning af aerob kapacitet til patienter med Reumatoid Artrit.

Patient or population: patients with Rheumatoid Arthritis

Settings: hospital, outpatient (rheumatology)

Intervention: Short-term land-based aerobic capacity training

	Illustrative comparative risks* (95% CI)					
	Assumed risk	Corresponding risk			Quality of	
Outcomes	Control	Short-term land-based aerobic capacity training	Relative effect (95% Cl)	No of Participants (studies)	the evidence (GRADE)	Comments
Functional ability AIMS . Scale from: 0 to 10. Follow-up: mean 12 weeks	The mean functional ability in the control groups was 0.9 points ¹	The mean Functional ability in the intervention groups was 0.06 standard deviations lower (1.33 lower to 1.2 higher)		56 (1 study)	$\oplus \oplus \ominus \ominus$ low ²	Absolute % change: AIMS -22%, relative % change: AIMS -43%, NNT: n.a., SMD: 0.03 (-0.46 to 0.51)
Muscle strength Isometric extension Follow-up: mean 12 weeks	The mean muscle strength in the control groups was 11 foot points	The mean Muscle strength in the intervention groups was 0.38 standard deviations lower (1.67 lower to 0.9 higher)		10 (1 study)	$\oplus \oplus \ominus \ominus$ low ²	Absolute % change: isometric extension 22%, relative % change: isometric extension 18%, NNT: n.a., SMD: -0.38 (-1.67 to 0.9)
Self-reported pain AIMS. Scale from: 0 to 10. Follow-up: mean 12 weeks	The mean self- reported pain in the control groups was -0.7 points	The mean Self-reported pain in the intervention groups was 0.27 standard deviations lower (0.79 lower to 0.26 higher)		56 (1 study)	⊕⊕⊕⊝ moderate ²	Absolute % change: AIMS -23%, relative % change: AIMS -10%, NNT: n.a., SMD: -0.27 (-0.79 to 0.26)
Disease activity	See comment	See comment	Not estimable	0 (3 studies)	⊕⊕⊝⊝ low ²	Absolute % change: ERS 5%/tender joints -36%, relative % change:ESR 19%/tender joints - 29%, NNT:n.a., SMD: statistical

					heterogeneity, pooling data not possible
Radiological damage - not measured	See comment	See comment	Not estimable -	See comment	Was not assessed in included studies

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI). **CI:** Confidence interval;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Mean change from baseline in control group

² Small patient numbers