## **VRAAG 5G: Depression**

## Systematic reviews

Study ID	Method	Patient characteristics	Intervention(s)	Results	Critical appraisal of review quality
Nagler 2012	SR Funding/Col: None declared Search date: December 2011 Databases:Cochrane Renal Group Specialised Register, CENTRAL, MEDLINE, EMBASE, PsychINFO, International Pharmaceutical Abstracts, Clinical trial registries Study designs:RCTs and observational studies N included studies: 28	Eligibility criteria: Adults or children with chronic kidney disease stages 3-5	Antidepressant drug treatment	Depression: CRITICAL OUTCOME no MA-results  Quality of life: IMPORTANT OUTCOME no MA-results	Moderate quality: only one reviewer, inclusions and exclusions not transparent     Included RCTs: Pervin (2006), Blumenfield (1997)
Rabindranath 2005a	SR Funding/Col: Funded by National Kidney Fund (UK) Search date: March 2006 Databases: Medline, Embase, Psychinfo, The Cochrane Library Study designs: RCTs N included studies: 1	Eligibility criteria:     Patients with ESRD on chronic dialysis and older than 18 years     Patient characteristics:     Age range: 18-70 years	Antidepressants vs. placebo or no treatment or a comparison of drugs	Depression: CRITICAL OUTCOME no MA-results  Quality of life: IMPORTANT OUTCOME no MA-results	High quality     Included RCTs: Blumenfield (1997)

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Rabindranath 2005b	SR Funding/Col: funded by the National Kidney Research Fund Search date: October 2003 Databases: Medline, Embase, PsycInfo, The Cochrane Library Study designs: RCTs N included studies: 0	Eligibility criteria: patients who are dialysed for ESRD older than 18 years diagnosed with depression	Psychosocial interventions vs. control or no intervention	Depression: CRITICAL OUTCOME no MA-results  Quality of life: IMPORTANT OUTCOME no MA-results	High quality     Included RCTs: -

## Primaire studies

Study ID	Method	Patient characteristics	Interventions	Results	Critical appraisal of study quality
Cukor 2014	Design: Randomized crossover trial     Funding/Col: Supported by National Institute of Health (K23DK076980) /none     Setting: 2 dialysis units in Brooklyn, USA     Sample size: N=65     Duration: 6 months	Eligibility criteria:     Haemodialysis patients with     ESRD and with elevated     depressive affect     A priori patient characteristics:     intervention vs. control     Male 27%     Mean dialysis treatment: 50     months	Cognitive behavioural therapy first (n=33) vs. Wait-list control first (n=26)	Depression: CRITICAL OUTCOME BDI-II: Treatment first: baseline 24.7 (9.8), after treatment 11.7 (9.8), after 2 <sup>nd</sup> phase 9.9 (8.5) Wait-list first: baseline 21.9 (8.9), after wait-list 14.5 (8.5), after treatment 9.1 (6.5) Model-estimated mean change score during treatment: treatment first -11.7 (SD 1.5; p<0.001), wait-list first -4.8 (SD 1.4; p<0.001) Model-estimated mean change score during wait- list: untreated group -6.7 (1.7; p<0.001)  HAM-D: Treatment first: baseline 15.7 (6.8), after treatment 6.5 (6.8), after 2 <sup>nd</sup> phase 6.7 (5.8) Wait-list first: baseline 12.9 (5.3), after wait-list 10.9 (5.4), after treatment 5.0 (4.3) Model-estimated mean change score during treatment: treatment first -9.1 (SD 1.1; p<0.001), wait-list first -5.9 (SD 1.1; p<0.001) Model-estimated mean change score during wait- list: untreated group -1.9 (1.2; p<0.17)  SCID: Treatment first: baseline 54, after treatment 5, after 2 <sup>nd</sup> phase 10 Wait-list first: baseline 33, after wait-list 31, after treatment 4  Quality of life: IMPORTANT OUTCOME	Level of evidence: high risk of bias  Randomization method and allocation concealment not described Patients not blinded, but blinded assessors  6 drop-outs, no ITT analysis

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Duarte 2009	Design: Randomized clinical trial     Funding/Col: project supported by Fundacao de Amparo a Pesquisa do Estado de Sao Paulo (04/08710-8)./ authors declare no competing interests     Setting: 2 dialysis units in Brasil     Sample size: N=85     Duration: 9 months	Eligibility criteria: Patients with ESRD receiving outpatient hemodialysis treatment     A priori patient characteristics: intervention vs. control     Age mean: 53 years     Male 41%     Diabetes 34%	Cognitive-behavioural group therapy (n=41) vs. Control (n=44)	KDQOL: Treatment first: Baseline: 99.5 (27.9) Treatment: 115.3 (25.5) Follow-up: 118.3 (27.7)  Wait-list: Baseline: 105.1 (23.7) Wait-list: 110.6 (25.1) Delay: 119.7 (24.7)  Pooled estimated treatment effect: 11.7 (2.0)  Depression: CRITICAL OUTCOME BDI Cognitive Subscale Intervention: Baseline: 13.7±7.1 After 3 mths: 7.1±5.9 After 9 mths: 6.3±7.1  Control: Baseline: 16.7±7.9 After 3 mths: 12.1±6.4 After 9 mths: 10.8±7.1 (intervention vs. control at 3 months: p<0.001)  BDI Somatic Subscale Intervention: Baseline: 10.6±4.0 After 3 mths: 7.0±3.8 After 9 mths: 6.1±3.2  Control: Baseline: 10.6±4.1 After 3 mths: 9.1±3.8 After 9 mths: 9.5±3.9 (intervention vs. control at 3 months: p=0.012)  BDI total Intervention: Baseline: 24.2±9.7 After 3 mths: 14.1±8.7 After 9 mths: 10.8±8.8  Control: Baseline: 27.3±10.7 After 3 mths: 10.8±8.8  Control: Baseline: 27.3±10.7 After 9 mths: 17.6±11.2 (intervention vs. control at 3 months: p=0.001)	Level of evidence: high risk of bias  Central randomization Patients not blinded, but blinded assessors No ITT analysis

Study ID	Method	Patient characteristics	Interventions	Results	Critical appraisal of study quality
				Major depression module MINI: Intervention: Baseline : 6.4±1.3 After 3 mths: 1.9±2.8 After 9 mths: 2.0±3.1	
				Control: Baseline : 6.4±1.2 After 3 mths: 4.3±2.9 After 9 mths: 3.5±2.9 (intervention vs. control at 3 months: p<0.001)	
				Suicide Risk module MINI: Intervention: Baseline : 2.2±5.1 After 3 mths: 1.2±4.2 After 9 mths: 0.6±1.2	
				Control: Baseline : 1.4±3.5 After 3 mths: 0.7±1.9 After 9 mths: 0.6±2.0 (intervention vs. control at 3 months: p=0.433)	
				Quality of life: IMPORTANT OUTCOME Burden of kidney disease: Intervention: Baseline : 28.7±22.4 After 3 mths: 43.6±27.1 After 9 mths: 43.2±28.8	
				Control: Baseline : 22.9±22.8 After 3 mths: 27.0±27.3 After 9 mths: 27.3±26.8 (intervention vs. control at 3 months: p=0.004)	
				Cognitive function: Intervention: Baseline : 64.4±23.0 After 3 mths: 77.2±25.1 After 9 mths: 81.1±20.5	
				Control: Baseline : 69.1±24.7 After 3 mths: 71.4±26.3 After 9 mths: 76.0±23.8 (intervention vs. control at 3 months: p=0.261)	

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				Quality of social interaction: Intervention: Baseline : 65.2±23.3 After 3 mths: 81.1±19.3 After 9 mths: 81.7±18.7	
				Control: Baseline: 70.0±22.2 After 3 mths: 66.5±22.3 After 9 mths: 71.2±24.4 (intervention vs. control at 3 months: p=0.002)	
				Sleep: Intervention: Baseline : 58.1±21.5 After 3 mths: 67.6±23.0 After 9 mths: 73.1±19.1	
				Control: Baseline : 58.4±18.7 After 3 mths: 58.4±17.8 After 9 mths: 62.8±19.3 (intervention vs. control at 3 months: p=0.034)	
				Mental component summary: Intervention: Baseline : 37.4±11.6 After 3 mths: 47.3±12.1 After 9 mths: 46.3±12.3	
				Control: Baseline: 41.1±11.2 After 3 mths: 39.3±11.9 After 9 mths: 38.6±11.7 (intervention vs. control at 3 months: p=0.002)	
Hosseini 2012	Design: Randomized controlled trial     Funding/Col: supported	Eligibility criteria: Hemodialysis patients with ESRD     A priori patient characteristics:	Citalopram (n=22) vs.	Depression: CRITICAL OUTCOME HADS Depression	Level of evidence: high risk of bias
	by grant from Mazandaran University of Medical Sciences / none declared	intervention vs. control  o Age mean: 50.5 years  o Male 42%	psychological training (n=22)	Psychol. Training: Pretest: 9.58 ± 3.47 Posttest: 7.33 ± 4.80	Randomization method and allocation concealment not described     No blinding
	Setting: Imam Khomeini Hospital, Iran     Sample size: N=44     Duration: 3 months			Citalopram: Pretest: 9.42 ± 3.11 Posttest: 6.26 ± 4.18	No ITT analysis
	Duration, 3 months			Quality of life: IMPORTANT OUTCOME Not reported	

Study ID	Method	Patient characteristics	Interventions	Results	Critical appraisal of study quality
Erdley 2014	Design: Randomized controlled trial     Funding/Col: without funding/ no Col     Setting: Geisinger medical center, USA     Sample size: N=36     Duration: 6 weeks	Eligibility criteria:     haemodialysis patients with     age 60 or older     A priori patient characteristics:     intervention vs. control	Problem-solving therapy (n=15) vs. Usual care (n=18)	Depression: CRITICAL OUTCOME BDI PS-therapy: Baseline: 15.7 (8.0) 6 weeks: 9.3 (3.1) Usual care: Baseline: 10.7 (6) 6 weeks: 11.3 (7.4) (PS-therapy vs. Usual care, p=0.6)  PHQ-9 PS-therapy: Baseline: 10.5 (4.9) 6 weeks: 3.3 (1.9) Usual care: Baseline: 6.1 (4.1) 6 weeks: 5.83 (4.2) (PS-therapy vs. Usual care, p=0.1)  Quality of life: IMPORTANT OUTCOME Not reported	Level of evidence: high risk of bias  • Allocation concealment not described • No blinding