VRAAG 5B: Pain

Primaire studies

Study ID	Method	Patient characteristics	Interventions	Results	Critical appraisal of study quality
Atalay 2013 Biyik 2013	Design: Randomized crossover trial Funding/Col: supported by Selcuk Scientific Research Project Coordinating Office Project Nr 08102027/ No competing interests Setting: Konya, Turkey Sample size: N=50 Duration: 14 weeks	Eligibility criteria: hemodialysis patients with neuropathic pain A priori patient characteristics: intervention vs. control Age mean: 58.2y Male 30% Hemodialysis duration: 55.1m	Vs. Pregabalin	Pain: CRITICAL OUTCOME SFMPQ Total: (p<0.001) Gabapentin: before 18.9 ± 4.3, after 9.3 ± 4.3 Pregabalin: before 18.5 ± 3.9, after 9.8 ± 3.6 Change in % (NS): Gabapentin: -8.9 +/- 4.1 Pregabalin: -9.3 +/- 4.0 SFMPQ VAS: (p<0.001) Gabapentin: before 68.8 ± 12.8, after 33.0 ± 15.6 Pregabalin: before 67.0 ± 11.8, after 32.9 ± 12.8 Change in % (NS): Gabapentin: -33.5 +/- 13.2 Pregabalin: -36.3 +/- 12.4 SFMPQ PPI: (p<0.001) Gabapentin: before 2.8 ± 0.8, after 1.4 ± 0.7 Pregabalin: before 2.8 ± 0.8, after 1.4 ± 0.7 Change in % (NS): Grabapentin:-1.3 +/- 0.8 Pregabalin: -1.4 +/- 0.6 Quality of life: CRITICAL OUTCOME PSQI: (p<0.001) Gabapentin: before 8.7 ± 4.2, after 5.9 ± 3.0 Pregabalin: before 8.8 ± 4.6, after 6.1 ± 4.2 BDI: (p<0.001) Gabapentin: before 13.61 ± 5.9, after 10.9 ± 5.9 Pregabalin: before 13.61 ± 5.9, after 10.9 ± 5.9 Pregabalin: before 42.6 +/- 18.2, after 57.1+/- 18.9 Pregabalin: before 42.7 +/- 17.9, after 57.3 +/- 17.1 Change in % (NS): Gabapentin: 13.0 +/- 9.2 Pregabalin: 16.1 +/- 11.2 SF-36 mental component scale score: (p<0.001) Gabapentin: before 51.6 +/- 19.5, after 63.2 +/- 18.3 Pregabalin: before 50.5 +/- 18.6, after 63.1 +/-	Level of evidence: high risk of bias • Unclear allocation concealment • Open label study • No ITT analysis: 10 dropouts excluded from analysis

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				15.8 Change in % (p=0.043): Gabapentin:9.6 +/- 11.2 Pregabalin: 14.6 +/- 11.6	