Νεφρολογικό Τμήμα ΓΝ Πτολεμαΐδας ΕΓΚΕΦΑΛΟΣ ΚΑΙ ΝΕΦΡΟΣ 10-11/11/2023

Γνωσιακή λειτουργία στην περιτοναϊκή κάθαρση.

Ντέμκα Αλεξάνδρα Νεφρολόγος, ΓΝ Παπαγεωργίου

Μέθοδοι υποκατάστασης νεφρικής λειτουργίας

ΜΕΤΑΜΟΣΧΕΥΣΗ





ΠΕΡΙΤΟΝΑΙΚΗ ΚΑΘΑΡΣΗ



Γνωσιακή Λειτουργία - Cognition Function

<u>Γνωσιακή Λειτουργία</u> (Cognition Function)

Προσοχή, Αντίληψη, Μνήμη, Γλώσσα, Σχεδίαση, Επίλυση προβλήματος, Λήψη απόφασης Πολλαπλές εργασίες



<u>Γνωσιακή Δυσλειτουργία</u> (Cognition Impairment)

(1) Normal cognitive aging

(2) Mild cognitive impairment

(3) <u>Major cognitive impairment</u> (Dementia)

γνωσιακή δυσλειτουργία 🗲 καθημερινή δραστηριότητα

Diagnostic and Statistical Manual of Mental Disorders, 5th Edn. American Psychiatric Association (2013).

Cognition in Advanced Kidney Failure l A COMMON BUT POORLY RECOGNIZED PROBLEM

1839, Journal of Practice Medicine:

". . . reciprocal action of the brain on the kidney and the kidney on the brain, has long been known."

Dr. Thomas Addison



Karakizlis H. BMC Nephrol. (2021) 22:205.

*ΡΣΔ: ρυθμός σπειραματικής διήθησης *ΓΛ: γνωσιακή λειτουργία

- ✓ για κάθε 10 ml/min/1,73m2 μείωση του ΡΣΔ σε > 55 ετών
 → 11% αύξηση στον επιπολασμό έκπτωσης ΓΛ
- ✓ eGFR < 45 mL/min/1,732: ΓΛ μειώνεται σημαντικά</p>
- ✓ eGFR < 15mL/min: επιπολασμός κυμαίνεται από 27-77%

Γενικό πληθυσμό: επιπολασμός 12% (68,5–78,3 ετών)

Crowe K. Front. Neurol. 12:787370.

Cognition in Advanced Kidney Failure 2 A COMMON BUT POORLY RECOGNIZED PROBLEM

Επιπτώσεις



Self Efficacy Engagement with health care

Decision making Quality of Life







Dialysis withdrawal

Depression & Stress

Hospitalisation



Uremia

Inflammation

Structural Changes

Dialysis



Παράγοντες κινδύνου



Age Vascular Disease Diabetes Depression & Sleep

Disorders



Crowe K. Front. Neurol. 12:787370.

Cognition in Peritoneal Dialysis

Complex activities - Lifestyle changes - Adaptation to dialysis - Medication regimens

All above are partly dependent on normal cognitive function self-monitoring, self-care

Most patients who choose peritoneal dialysis ✓ independent and capable of self-administering dialysis and medications



 ✓ a need to assess patients' ability to choose and maintain PD therapy before and regularly after dialysis therapy initiation

Kalirao, P. (2011). AJ of Kidney Diseases, 57(4), 612–620.

<u>Prevalence and Risk factors of</u> <u>Cognitive Impairment</u>

Peritoneal Dialysis

Vs

Non-CKD, Pre-CKD, Hemodialysis, Transplantation

Cognitive Impairment in Peritoneal Dialysis Patients

51 PD compared with 338 HD and 101 controls without CKD 9 validated neuropsychological tests

Cognitive Impairment: 75% PD patients (50%: moderate, 50%: severe)



PD: > 2.5 times more likely than non-CKD and as was HD moderate to severe cognitive impairment

*dementia in people without ESRD > 65 years old: 10%, > 85 years old: 35%-40%

*PD cohort was on average 11 years younger than the HD cohort (57.5 vs 68.5 years)



Kalirao, P. (2011). AJ of Kidney Diseases, 57(4), 612–620.

Frequency and risk factors for cognitive dysfunction in peritoneal dialysis patients

149 PD patients at a single center between 2016 and 2020 Neurocognitive screening (ACE-R and MMSE)

Addenbrooke's Cognitive Examination – Revised (ACE-R) incorporated Mini-Mental State Examination (MMSE)



Risk factors: age, female gender, diabetes, depression, PD > 12m

ACE-R score by cohort

Residual urine output and ACE-R score

MMSE is ineffective in detecting subtle NCI in this population compared with ACE-R

Gamage I. Nephrology. 2022;27:945–952

Comparison of the extent and pattern of cognitive impairment among pre-dialysis, dialysis and transplant patients: a cross sectional study from Australia

Observational cross sectional study compare the extent of CI and the types of cognitive deficits 155 patients eGFR < 30 ml/min per 1.73m2



PRE PD HD KΤ Total n=24 n=25 n=54 n=52 P value n=155 Cognitively impaired, $4(16.7)^{a}$ 30 (55.6) ab 10 (19.2)^b 12 (48.0) 56 (36.1) < 0.001* Proportion, n, (%) Total MoCA score 27.07 24.8023.12 26.77 25.23 < 0.001* (22.11-24.13) ab Mean (95% CI) (25.55-28.58)^a (23.32-26.28) $(25.74-27.80)^{b}$ (24.58-25.88)

Memory → directly impact on patient's ability to learn and recall information provided

Dialysis, Age ≥65, Male gender, Diabetes, Cerebrovascular disease

Lambert K. Nephrology, 22 899-906.

High Prevalence of Leukoaraiosis in Cerebral Magnetic Resonance Images of Patients on Peritoneal Dialysis

a large vascular ischemic component combined with neurodegenerative pathologic states

57 PD patients relatively young (mean age, 48.4 years) without diabetes / cerebrovascular disease

✓ prevalence of leukoaraiosis was significantly greater in PD than controls (68.4% v 17.5%; P<0.001)

Old age, poorly controlled Hypertension, PD procedure itself and/or ESRD

Leukoaraiosis presence





Increased risk of stroke, Disability, Cognitive impairment decline

*ποσοστό απώλειας όγκου είναι συνήθως 0,5% ανά έτος μετά την ηλικία των 40 ετών

Kim CD. American Journal of Kidney Diseases, Vol 50, No 1 (July), 2007: pp 98-107

Prevalence of cognitive impairment among peritoneal dialysis patients: a systematic review and meta-analysis

- ✓ a search of the literature on CI in PD patients published between 1 Jan 1980 and 25. April 2019
- ✓ 8 studies were included and the relevant data from 1736 patients (PD controls)

Variables	Number of studies included	Coefficient (95% CI)	R^2 analogue	p value
PD first policy (reference group: yes)	8	-0.897 (-2.543-0.749)	0	0.29
Prevalence of DM	7	-0.212 (-7.801-7.377)	0	0.96
Sex ratio (male: female)	7	- 1.663 (- 3.128-0.795)	0	0.24
Mean age in years	8	-0.014 (-0.112-0.090)	0	0.79
Study design (reference group: prospective study)	8	-0.251 (-1.851-1.350)	0	0.76
Publication year (reference: on or before 2014)	8	0.251 (-1.350-1.851)	0	0.76
Mean PD duration in months	8	0.002 (-0.061-0.065)	0.08	0.95

- ✓ Risk factors for CI: older age, female sex and lower education
- ✓ Potential reversible factors for CI: *electrolytes disturbances, depression and vitamin D defciency*.
- ✓ CI was associated with a higher risk of hospitalization, mostly due to PD-related peritonitis.

Shea YF. Clinical and Experimental Nephrology https://doi.org/10.1007/s10157-019-01762-1

Prevalence of Cognitive Impairment in Peritoneal Dialysis Patients and Associated Factors

PD patients $(n = 6)$			controls $(n = 4)$		
<65 years (n = 3)	≥65 years (n = 3)	p value	<65 years (n = 1)	\geq 65 years (n = 3)	p value
23%	60%	0.14	10%	60%	0.02

15 Control group: CI more common in ≥65 years 18 PD group: may occur earlier (<65 years)

Variable	PD patients		p value*
	CI (n = 6)	NCI (n = 12)	
ACE III test			
Total score, mean±SD (min, max)	78.5±11.6 (61-88)	94.8±2.7 (89-99)	0.018
Attention, mean±SD (min, max)	91.5±10.4 (72-100)	94±8.7 (72-100)	0.626
Memory, mean±SD (min, max)	60.8±8.9 (46-73)	94.8±6.3 (81-100)	0.00
Fluency, mean±SD (min, max)	62±24.5 (28-86)	88.9±7.6 (79-100)	0.042
Language, mean±SD (min, max)	91±11 (77-100)	98.3±2.7 (92-100)	0.165
Visuospatial, mean±SD (min, max)	86.5±18 (56-100)	94.8±9.9 (75-100)	0.327

a significant trend for PD patients with CI to score lower in the domains of **memory** and **verbal fluency**

			ACE III total score (%)	Attention (%)	Memory (%)	Verbal fluency (%)	Language (%)	Visuospatial (%)
PD patients Years of Pearson (n = 18) education Correlation Significance	Pearson	0.78**	0.19	0.63**	0.84	0.73**	0.56*	
	Significance	0.00	0.45	0.005	0.00	0.00	0.02	
	Dialysis	Pearson	-0.19	-0.08	-0.33	-0.05	-0.08	0.2
	(months)	Significance	0.46	0.76	0.18	0.84	0.75	0.93
Controls	Years of education	Pearson	0.53	0.41	0.52*	0.44	0.52*	0.31
(1 - 13)	concertori	Significance	0.04	0.13	0.046	0.1	0.049	0.26

no correlation between the duration of dialysis and the results of the ACE III test,

Golenia A. Kidney Blood Press Res 2023;48:202–208

Prevalence of mild cognitive impairment in automated peritoneal dialysis patients

71 patients on APD Mean age: 42 years, Mean dialysis duration: 17 months



Mini Mental State Examination (MMSE) Montreal Cognitive Assessment (MoCA)

CI (mild deterioration) was present in 7%: MMSE and 68%:MoCA, and 4 and 37% in the healthy controls

Diabetes more frequently Higher serum glucose Lower serum phosphorus

Variable	В	95% CI	P-value
Education (years)	0.53	0.19-0.87	0.003
Serum sodium (mmol/L)	0.56	0.04 - 1.08	0.03
Serum creatinine (mg/dL)	0.48	0.06-0.91	0.03
Age (years)	-0.10	-0.20-0.00	0.05

Older age, Less education Lower serum sodium Lower serum creatinine

Coexisting Frailty and Cognitive Impairment in Patients on Continuous Ambulatory Peritoneal Dialysis

784 CAPD patients started to enroll from 2014 to 2016 and ended follow-up by 2017

Cognitive impairment: 55.5% Frailty: 27.6% Frailty + Cognitive Impairment: 23.9%



CFS score: associated with MoCA score $(\beta=-0.69, P<0.001)$

Frailty: Clinical Frailty Scale (CFS), and Cognitive function: Montreal Cognitive Assessment (MoCA).



Coexisting frailty and cognitive impairment was associated with 1. decreased patient survival rate

2. increased peritonitis rate

Yi C. Scientific Reports. (2018) 8:17305

Cognitive Changes in Peritoneal Dialysis Patients: A Multicenter Prospective Cohort Study

	Baseline	2-y Follow-up	Ρ
3MS score	84.8 ± 12.2	83.1 ± 14.8	0.006ª
Cognitive impairment	58 (19.8%)	70 (23.9%)	0.2
Trail-A, s	65 [49-90]	56 [41-80]	<0.001ª
Trail B, s	144 [103- 227]	144 [102- 193.5]	0.03ª
Immediate memory score	74.0 ± 18.1	76.6 ± 18.4	0.01ª
Delayed memory score	89.3 ± 17.7	87.8 ± 20.5	0.2
Visuospatial skill score	86.7 ± 23.7	91.2 ± 19.9	0.004ª
Language ability score	93.5 ± 13.7	94.6 ± 14.1	0.2

458 PD patients were enrolled and followed up for 2 years.

CI prevalence increased from 19.8% to 23.9% Executive function, immediate memory, visuospatial skill improved

	-		~	-			-		•		-			
	3MS Score		Trail-A (s)		Trail-B (s)		Immediate M Score	/lemory	Delayed Mer Score	nory	Visuospatial Score	skill	Languag Score	e Ability
Variable	β (SE)	Р	β (SE)	Ρ	β (SE)	Р	β (SE)	Р	β (SE)	Р	β (SE)	Ρ	β (SE)	Р
Age	-0.12 (0.05)	0.01ª	0.65 (0.19)	<0.001ª	1.59 (0.45)	<0.001*	-0.20 (0.07)	0.006	0.13 (0.07)	0.07	-0.10 (0.08)	0.2	-0.02 (0	.06) 0.7
Female sex	0.46 (1.20)	0.7	-2.98 (4.99)	0.6	-1.56 (11.34)	0.9	2.02 (1.85)	0.3	-2.05 (1.87)	0.3	-2.17 (2.21)	0.3	0.83 (1	.53) 0.6
BMI (kg/m ²)	-0.25 (0.18)	0.2	1.25 (0.70)	0.08	1.20 (1.60)	0.5	-0.01 (0.27)	0.9	-0.21 (0.27)	0.5	-0.55 (0.32)	0.09	-0.20 (0	.22) 0.4
DM	-0.36 (1.56)	0.8	19.38 (6.44)	0.003	46.94 (14.42)	0.001 ^ª	-4.56 (2.43)	0.06	-1.70 (2.47)	0.5	1.43 (2.94)	0.6	-0.37 (2	.01) 0.9
CVD	1.02 (1.53)	0.5	10.50 (6.20)	0.09	3.04 (14.20)	0.8	2.32 (2.40)	0.3	0.78 (2.44)	0.7	-1.27 (2.85)	0.7	0.12 (2	.00) 0.9
Depression score	-0.14 (0.07)	0.04ª	-0.07 (0.27)	0.8	0.03 (0.61)	0.9	-0.22 (0.10)	0.03 ^a	-0.30 (0.11)	0.005ª	-0.19 (0.12)	0.1	-0.20 (0	.09) 0.02
Educational level	2.86 (0.65)	< 0.001	-5.82 (2.55)	0.02ª	-17.70 (5.70)	0.002	2.64 (0.98)	0.007ª	3.54 (1.03)	0.001ª	5.21 (1.21)	< 0.001	1.45 (0	.79) 0.07
hs-CRP (mg/L)	-0.05 (0.05)	0.4	0.33 (0.25)	0.2	0.53 (0.55)	0.3	-0.11 (0.09)	0.2	-0.17 (0.10)	0.09	0.03 (0.11)	0.7	-0.06 (0	.07) 0.4
SNa (mmol/L)	-0.02 (0.07)	0.8	-0.08 (0.27)	0.8	-0.46 (0.61)	0.5	-0.05 (0.11)	0.7	-0.24 (0.11)	0.03ª	-0.30 (0.12)	0.02ª	-0.01 (0	.09) 0.9
SAlb (g/L)	0.64 (0.11)	< 0.001	-0.07 (0.46)	0.9	0.70 (1.04)	0.5	-0.13 (0.17)	0.5	0.90 (0.18)	<0.001ª	0.80 (0.21)	< 0.001	0.44 (0	.14) 0.003

Lower serum albumin level Advanced age, Lower education, Diabetes Depression Hyponatremia An association of cognitive impairment with diabetes and retinopathy in end stage renal disease patients under peritoneal dialysis

Liao J-L. (2017) PLoS ONE 12(8): e0183965

✓ Diabetes and retinopathy: risk factors of CI
 ✓ 424 clinically stable PD patients were enrolled

	-				
Variables	Total	non DM	DM	DM and Retinopathy	P
N (%)	424	307(72.4%)	36 (8.5%)	81(19.1%)	-
3MS score	83.4score	84.3score	85.5scor ^t	79.1scor1 ^d	0.004
Cognitive impairment	113(27.4%)	72(24.2%)	7(21.2%)9	34(42%) ^d	0.004
Trails A,s	88.1ls A,	82.0ls A,	92.6ls A,	109.9s A,s ^d	0.008
Trails B,s	207.4s B,s1	191.8s B,s3	204.9s B,s	268.0s B,s1°	0.001
Executive dysfunction	127(32%)	73(25.2%)	13(41.9%)	41(53.9%) ^e	<0.001
Immediate memory score	73.1diate	74.9diate	70.0diate	67.7diate ^d	0.003
Impaired immediate memory	275(67.2%)	185(62.7%)	26(76.5%)	64(80.0%) ^d	0.007
Delayed memory score	88.6yed m	88.9yed m	89.4yed m	87.0yed m	0.66
Impaired delayed memory	83(21.3%)	62(21.8%)	7(22.6%)	14(18.9%)	0.85
Language ability score	92.4uage	93.2uage	92.4uage	89.2uage	0.08
Impaired language ability	69(16.8%)	43(14.5%)	8(23.5%)	18(22.5%)	0.13
Visuospatial skill score	85.2ospat	87.8ospat	84.3ospat	75.6ospat ^e	<0.001
Impaired visuospatial skill	194(48.9%)	129(44.3%)	16(51.6%)	49(65.3%) ^d	0.005

PD Patients with DM and Retinopathy had **significantly higher** prevalence of CI compared with patients in non-DM group.

DM and retinopathy rather than DM only were significantly associated with increased risk for CI, ORs 2.09[1.11,3.92]

Retinopathy is associated with impaired cognition in patients undergoing peritoneal dialysis

Liao J.(2023) Renal Failure, 45:2, 2258989

- ✓ a relationship between retinopathy and cognition with and without CKD
- ✓ a cross-sectional design 107 PD participants



Retinopathy was associated with global CI.

Retinopathy: a valuable primary screening tool for assessing the risk of cognitive decline.

Self-Care Peritoneal Dialysis Patients with Cognitive Impairment Have a Higher Risk of Peritonitis in the Second Year

Comparison of Clinical Features and 2-Year Outcomes Between PD Patients With and Without Cognitive Impairment at First Year

	Who	e group		Self-care PD group		
	With CI	Without CI		With CI	Without CI	
	(<i>n</i> =23)	(<i>n</i> =120)	p value	(<i>n</i> =16)	(<i>n</i> =98)	p value
Age in years, mean	66.1±9.3	57.9±13.4	0.006*	64.5±9.1	55.5±12.9	0.008ª
Different age strata, n (%)						
<65	10 (43.5)	85 (70.8)		8 (50)	75 (76.5)	
65-74	6 (26.1)	19 (15.8)	0.03 ^b	5 (31.3)	15 (15.3)	0.09 ^b
74-85	7 (30.4)	16 (13.3)		3 (18.8)	8 (8.2)	
Female, n (%)	16 (70.0)	53 (44.2)	0.036	10 (62.5)	48 (49.0)	0.32 ^b
HK-MoCA at 1 year, mean	14±5.1	25±3.5	<0.001ª	14±5.4	25±3.3	<0.001
Education						
Illiterate	3 (13.0)	16 (13.3)		2 (12.5)	11 (11.2)	
Primary	7 (30.0)	23 (19.2)	0.62 ^b	3 (18.8)	15 (15.3)	0.97 ^b
Secondary	9 (39.1)	50 (41.7)		7 (43.8)	42 (42.9)	
Tertirary	4 (17.4)	31 (25.8)		4 (25.0)	30 (30.6)	
Baseline comorbidities						
DM, n (%)	13 (0.57)	51 (42.5)	0.22 ^b	9 (56.3)	39 (40.0)	0.22 ^b
HI, n (%)	22 (95.7)	109 (90.8)	0.69 ^b	15 (93.8)	90 (91.8)	1.0 ^b
Hyperlipidemia, n (%)	15 (65.0)	72 (60)	0.64 ^b	12 (75.0)	58 (59.2)	0.28 ^b
IHD, n (%)	2 (8.7)	22 (18.3)	0.37 ^b	2 (12.5)	18 (18.4)	0.73 ^b
Previous stroke, n (%)	2 (8.7)	15 (12.5)	1.0 ^b	2 (12.5)	10 (10.2)	0.68 ^b
PVD, n (%)	7 (30.4)	12 (10)	0.008 ^b	5 (31.3)	6 (6.1)	0.002 ^b
Use of sedative medications ^d , n (%)	6 (26.1)	39 (32.5)	0.53 ^b	4 (25.0)	32 (32.7)	0.77 ^b
ical outcomes in the second year						
PD-related peritonitis, n (%)	11 (47.8)	23 (19.2)	0.003 ^b	7 (43.8)	20 (20,4)	0.04 ^b
Suffered from both peritonitis and exit-site infection, n (%)	6 (26.1)	8 (6.7)	0.004 ^b	4 (25.0)	7 (7.2)	0.049 ^b
PD-related peritonitis rates (episodes per year)	0.64	0.25	0.004 ^c	0.50	0.27	0.048 ^c
With exit-site infection, n (%)	9 (39.1)	51 (42.5)	0.76 ^b	7 (43.8)	39 (30.6)	0.79 ^b
Exit-site infection rates (episodes per year)	0.39	0.42	0.800	0.44	0.42	0.90 ^c
Unplanned hospitalization rates (episodes per year) ^e	1.85	1.32	0.05°	1.66	1.19	0.08 ^c
With emergency admissions ^e , n (%)	17 (73.9)	59 (49.2)	0.03 ^b	12 (75.0)	42 (42.9)	0.03 ^b
Number of emergency admissions ^e , median (IQR)	1 (0-3)	0 (0-2)	0.05°	1(0-3)	0 (0-2)	0.07°
Duration of admissions ^e , median (IOR)	5 (0-18)	1 (0-9)	0.03°	5 (0-23)	0 (0-8)	0.04°

CI

Increasing age, Female sex, Anemia, PVD presence

Comparison of	f Self-Care PD	Patients With	and Without	Peritonitis in the	e Second Year
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	With peritonitis	Without peritonitis	
	(<i>n</i> =27)	(<i>n</i> =86)	p value
Age in years, mean	60.2±11.4	55.7±13.1	0.12 ^a
Different age strata, n (%)			
<65	18 (66.7)	64 (74.4)	
65-74	6 (22.2)	14 (16.3)	0.72 ^b
74-85	3 (11.1)	8 (9.3)	
HK-MoCA at 1 year, mean	22±6.8	24±4.9	0.07ª
Cognitive impairment at 1 year, n (%)	7 (25.9)	9 (10.5)	0.04 ^b
Smoking status, n (%)			
Non-smoker	19 (70.4)	68 (79.1)	
Ex-smoker	7 (25.9)	16 (18.6)	0.64 ^b
Chronic smoker	1 (3.7)	2 (2.3)	
Depression ^e , n (%)	2 (7.4)	2 (2.3)	0.24b
Usage of sedative medications, n (%) ^d	9 (33.3)	27 (31.4)	0.85 ^b
Albumin at first year (g/dL), mean	36.7±4.4	35.3±3.8	0.13ª
Exit-site infection in the second year, n (%)	10 (37.3)	36 (41.9)	0.66 ^b
Exit-site infection rate in the second year (episodes per year)	0.26	0.89	0.85°

CI at 1 year on self-care PD is associated with a higher risk for PD-related **peritonitis** in the 2nd year.

<u>Cognitive assessment should be part of</u> <u>the overall assessment in older PD patients</u>

Shea YF. Peritoneal Dialysis International, Vol. 39, pp. 51–58

Sleep Disorders and Cognitive Impairment in Peritoneal Dialysis: A Multicenter Prospective Cohort Study

- ✓ the relationship between sleep disorders and CI, and predictors for declining cognitive function
- ✓ a multicenter prospective cohort study
- ✓ 458 clinically stable PD patients followed up for 2 years.

	All participants (n = 458)	Participants who completed the 2 assessments (n = 293)		р
		baseline	2 years later	-
3MS score	84.7±12.3	84.8±12.2	83.1±14.8	0.006**
Cognitive impairment	90 (19.7)	58 (19.8)	70 (23.9)	0.161
Trail-Making Test A duration, s	65.5 (47.25-95)	65 (49-90)	56.0 (41.0-80.0)	< 0.001***
Trail-Making Test B duration, s	150 (106-233)	144 (103-227)	144.0 (102-193.5)	0.025*
Immediate memory score	74.0±17.9	74.0±18.1	76.6±18.4	0.010*
Delayed memory score	89.3±17.0	89.3±17.7	87.8±20.5	0.155
Visuospatial skill score	86.0±23.5	86.7±23.7	91.2±19.9	0.004**
Language ability score	93.3±13.1	93.5±13.7	94.6±14.1	0.207
Insomnia score	3.0±2.4	3.0±2.5	3.3±2.4	0.087
Restless legs syndrome score	1.6±6.0	1.6±5.8	1.9±6.7	0.416
Excessive daytime sleepiness score	8.5±4.7	8.3±4.5	9.6±5.5	< 0.001***
Possible narcolepsy	21 (4.7)	10 (3.4)	16 (5.5)	0.189
Sleepwalking and nightmares	85 (19.1)	56 (19.6)	49 (16.9)	0.610
Possible rapid eye movement behavior				
disorder	32(7.2)	20 (6.8)	26(8.9%)	0.337

Sleep disorders were common among PD patients (excessive daytime sleepiness, Insomnia)

Prevalence of CI increased from 19.8 to 23.9%.

Possible narcolepsy was associated with general CI

During follow-up, sleepwalking and nightmares were associated with higher risks of **declined delayed memory**

Sleep disorders: Insomnia, Restless legs syndrome, Sleep apnea syndrome, Excessive daytime sleepiness, Narcolepsy, Sleep walking and nightmares, Rapid eye movement behavior disorders

Zhao et al. DOI: 10.1159/000502355

The comparison of cognitive function and risk of dementia in CKD patients under peritoneal dialysis and hemodialysis

15 cohort or cross-sectional studies comparing cognitive functions using neuropsychological tests

By qualitative analysis, more studies are inclined to PD compared with HD with better cognitive functions.

MMSE														RAVLT											
	Haen	odialysis		Periton	al dialy	ysis		Mean Difference		Mea	Difference				Hae	modial	ysis	Perito	neal dialy	sis		Mean Difference	Mean	Difference	
Study or Subgroup	Mean	SD T	tal	Mean	SD	Total	Weight	IV. Fixed. 95% Cl.		IV. F	ixed. 95% Cl	_		Study or Subgroup	Mean	50	Total	Mean	SD	Total	Weight	IV. Fixed, 95% CI	IV. Eb	ed. 95% Cl	_
Cukor 2013	27.4	1.7	25	28.2	1.2	6	2.4%	-0.80 [-1.97, 0.37]	_	-	-			Griva 2003	2.64	2.09	77	3.03	1.61	68	84.5%	-0.39 [-0.99, 0.21]	-	+	
Tiki 2003	28	0.3	25	29	0.3	17	97.6%	-1.00 [-1.18, -0.82]						Wolcott 1988	5.1	2.6	17	6.5	1.5	17	15.2%	-1.40 [-2.83, 0.03]		1	
Total (95% CI)			50			23	100.0%	-1.00 [-1.18, -0.81]		٠				Total (95% CI)			94			85	100.0%	-0.54 [-1.10, 0.01]	-		
Heteropeneity: Chi ² = 0	0.11. df =	= 1 (P = 0.	(4); P	= 0%							+ +	_		Heterogeneity: Chi# = 1	1.63. df	1=1(P	= 0.20)	P = 395	2			+	-	+ +	-
Test for overall effect: 2	Z = 10.6	8 (P < 0.0	1001)						~2	-1	0 1		2	Test for overall effect:	Z = 1.9	2 (P=0	1.06)					-4	-2	0 2	
A										ravours	PU Pavours n			F									Pavours H	J Pavours PD	
														PV/PT											
MoCA														BVRI											
	Haen	odialval		Periton	tab les	vala		Mean Difference		Mea	n Difference				Haer	modialy	rais	Peritor	real dialy	sis		Mean Difference	Mean	Difference	
Study or Suboroup	Mean	SD T	tal.	Mean	SD	Tota	Weight	IV. Fixed, 95% CI		IV.F	fixed, 95% CI			Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV. Random, 95% CI	IV. Rat	idom, 95% Cl	_
Ivasare 2017	23	3.5	41	24	2	25	18.8%	1 00 1-2 33 0 331			-			Griva 2003	5.97	2.31	77	4.97	1.74	68	51.1%	1.00 [0.34, 1.66]	-		
Lambert 2017	23.12	1.01	54	24.8	1.48	25	81.2%	-1.68 (-2.32, -1.04)		-				Kalirao 2011	4.7	3	338	6.4	3.6	51	48.9%	-1.70 [-2.74, -0.66]	-		
										-				B-1-1 (1941) (194											
Total (95% CI)			95			50	100.0%	-1.55 [-2.13, -0.98]		٠				Total (90% CI)			415			119	100.0%	-0.32 [-2.96, 2.33]			
Heterogeneity: Chi ² = 0	0.82, df	= 1 (P = 0	37); P	= 0%					+	-	-	-	+	Heterogeneity: Tau* =	3.45; C	79" = 18	48, df =	=1(P<0	10001); P	= 95%			4 -2	0 2	
Test for overall effect:	Z = 5.28	(P < 0.00	(100						-4	-2 Enume	DD Emmo	2	4	Test for overall effect:	Z = 0.2	4 (P = 0	.81)						Favours H	D Favours PD	
B										ravours	PD Pavours P	-		F											
TMT-B														SIT											
	Manage	adiabasis		Indiana	al dish.	ala.		Mean Difference		BRos.	Difference						1.2.1	-							
Study or Substant	Mean	SD T	-	lean	SD.	Total	Walabi	IV Random 65% Cl		IV.P	andom 95% C	1			Haer	modialy	1515	Peritor	seal dialy	115 March 1		Mean Difference	Mean	Difference	
000 2002	00.02	61 75	**	13.36	40.74		10.15	36 66 130 63 53 681	-			1		Study or Bubgroup	Mean	- 30	10121	Mean	50	10121	meight	IV. Fixed, 92% CI	IV. FI	red. 95% CI	-
Neumann 2018	79.2	45.8	96	77.8	51.2	101	30.1%	1 40 6 12 15 14 951			-			Kalirao 2011	110.4	43.2	338	84.4	21.7	51	81.1%	26.00 [17.11, 34.89]			_
Wolcoff 1988	163.5	0.6	17	97.3	32.3	17	22.8%	71 20 123 05 118 453					_	Williams 2004	11/.4	14	20	90.3	28	10	18.9%	22.10 [3.64, 40.51]			
There are a series	100.0	-			01.0			i nao pano, monaj						Total (95% CD			358			61	100.0%	25 26 117 26 33 271		-	
Total (95% Cl)		1	90			188	100.0%	30.73 [-2.66, 64.11]			-	-		Hotermenaity Chil a (114 11	= 1 (P)	0.711	12 = 0%					+ +	+ + +	_
Heterogeneity: Tau ^a = 6	693.90; 0	2hif = 15.8	3, df =	2(P=0	0.0004);	12 = 87	\$6			-			-	Test for owerall effect	7 = 6.11	0/0 + 0	000011						-20 -10	0 10 20	
Test for overall effect: 2	2 = 1.80	(P = 0.07)							-100	Environme	ND Envours E	0	00	G									Favours H	D Pavours PD	
C										1.0001.0	The Farmers			G											
SUMT														Risk of de	men	tia									
SDIVIT															Haem	nodialy	sis P	eritonea	I dialysis			Odds Ratio	Ode	is Ratio	
	Haem	odialysis		Peritone	al dialy	sis		Mean Difference		Mea	n Difference			Study or Subgroup	Ever	nts 1	Total	Events	Tot	al Web	oht M-	H. Random, 95% CI	M-H. Rat	dom. 95% Cl	_
Study or Subgroup	Mean	SD To	aL_f	fean	SD	Total	Weight	IV. Random, 95% CI		IV. R	andom, 95% C			Lin 2015	37	75 5	2332	181	325	2 47.	7%	1.34 [1.15, 1.56]			
Griva 2003	47.1	15.2	77 4	4.73	14.56	68	39.1%	2.37 [-2.48, 7.22]			_			Sithinamsuwan 2005		5	60	1	. 3	0 2	4%	2.64 [0.29, 23.64]			-
Radic 2011	29.04	11.7	22 :	4.33	14.9	20	31.7%	-5.29 [-13.45, 2.87]	-	-				Wolfgram 2014	82	46 11:	2960	338	866	3 49.	9%	1.94 [1.74, 2.17]			
Wolcott 1988	29.6	13.8	17	41.6	13.9	17	29.2%	-12.00 [-21.31, -2.69]																	
Total (BEN CD							100.00			-				Total (95% CI)		165	1352		1198	5 100.	0%	1.64 [1.15, 2.32]		•	
roum (acre Ci)						105	100.0%	-4.20 [-12.79, 4.28]	-				-	Total events	120	26		520					249		
menerogeneity: Tau ^a = 4	- 0.00	m = 8,10,	01-2	fr = 0.0	with a l	1078			-20	-10	0 1	0	20	Heterogeneity: Tau ² = 0	0.06; CI	hP = 15	.08, df =	2 (P=0	0005); P	= 87%		0.0	2 01	1 10	-
rest for overall effect 2	c = 0.98	(r = 0.33)								Favours	HD Favours F	^o		Test for overall effect: 2	= 2.78	5 (P = 0.	006)					0.0	Emergen M	Envrore PD	

By quantitative analysis,

- PD showed better performance in the tests of
- 1. Mini-Mental State Examination (MMSE),
- 2. Montreal Cognitive Assessment (MoCA),
- 3. Stroop interference test

PD exhibited lower risk of dementia compared with HD.

Cognitive Dysfunction and Health-Related Quality of Life in Patients with End-Stage Renal Disease Undergoing Hemodialysis in Comparison with Patients Undergoing Peritoneal Dialysis: A Cross-Sectional Study

✓ cross-sectional study, 265 patients who received hemodialysis or peritoneal dialysis

✓ CF: MoCA – HRQOL: SF-36 and KDTA

Variable	Model 1 [β (95% Cl)]	Model 2 [β (95% CI)]	Model 3 [β (95% Cl)]
MOCA	-8.35 (-9.85 to -6.86)***	-7.16 (-8.61, -5.70)***	-7.02 (-8.87, -5.26)***
SF-36	-10.12 (-11.94 to -8.45)***	-9.00 (-10.73, -7.28)***	-8.67 (-10.76, -6.58)***
KDTA	-8.67 (-10.10 to -7.23)***	-7.64 (-9.05, -6.24)***	-7.55 (-9.25, -5.85)***

PD patients: worse cognitive dysfunction and worse HRQOL compared to HD patients

MoCA: Montreal Cognitive Assessment SF-36: Kidney Disease Quality of 36-Item Short Form Survey KDTA: Kidney disease-related quality of life assessment

Zeng B. Med Sci Monit 2022; 28:e934282

Peritoneal dialysis is associated with better cognitive function than hemodialysis over a one-year course

271 patients (96 hemodialysis VS. 101 peritoneal dialysis patients)

Peritoneal dialysis was associated with better outcomes than hemodialysis at baseline and follow-up



PD as the gentler, more continuous, and potentially more efficient dialysis modality might be more beneficial for restoring CF.

Executive functioning: Trail Making Test-B *Attention*: d2-Revision-Test *QoL*: Kidney Disease Quality of Life Short Form Cognitive Function (KDQOL-CF)

Neumann D. Kidney International (2018) 93, 430–438

Cognitive Impairment in Peritoneal Dialysis

Self-monitoring and Self-care are partly dependent on normal cognitive function

Risk factors

Depression, Hyponatremia, Vitamin D deficiency, Diabetes, Daily prolonged exposure to the high glucose load, Inflammation, Malnutrition, Diffuse vasculopathy, Midabdominal obesity, Metabolic syndrome



Cognitive impairment

Risk of peritonitis Hospitalization Mortality Medical costs Treatment compliance Need for dialysis assistance Transfer to HD

Crowe K. Front. Neurol. 12:787370.

Συμπέρασμα



 Η γνωσιακή δυσλειτουργία είναι καλά αναγνωρισμένη ως ανεξάρτητος προγνωστικός παράγοντας θνησιμότητας σε άτομα υπό ΠΚ και μπορεί να επηρεάσει αρνητικά την ικανότητα λήψης αποφάσεων και την κρίση τους

2. Η γνωσιακή λειτουργία θα πρέπει να αξιολογείται **τακτικά** στους ΠΚ ασθενείς **ανεξάρτητα από την ηλικία** τους.

3. Ιδιαίτερη προσοχή πρέπει να δοθεί στην ανίχνευση και τα **προληπτικά μέτρα ήπιας** γνωστικής εξασθένησης σε ασθενείς υπό ΠΚ.

ΕΠΙΣΤΗΜΟΝΙΚΗ ΕΚΔΗΛΩΣΗ Εγκέφαλος και Νεφρός

10-11 Νοεμβρίου 2023 Pantelidis Hotel, Πτολεμαίδα

ΟΡΓΑΝΩΣΗ ΝΕΦΡΟΛΟΓΙΚΟ ΤΜΗΜΑ "Χ. ΚΑΤΣΙΝΑΣ" ΓΕΝΙΚΟ ΝΟΣΟΚΟΜΕΙΟ ΠΤΟΛΕΜΑΙΔΑΣ "ΜΠΟΔΟΣΑΚΕΙΟ"

ΥΠΟ ΤΗΝ ΑΙΓΙΔΑ





AL PEPEBEREIAT ANTICHE MAKEADAMAT

Ευχαριστώ !



