

# How Dialysis Patients Live: A Study on Their Depression and Associated Factors in Southern Italy

Maria Sinatra<sup>1</sup>, Antonietta Curci<sup>1</sup>, Valeria de Palo<sup>1</sup>, Lucia Monacis<sup>2</sup>, Giancarlo Tanucci<sup>1</sup>

<sup>1</sup>Department of Psychology, University of Bari, Bari, Italy; <sup>2</sup>Department of Human Sciences, University of Foggia, Foggia, Italy. Email: m.sinatra@psico.uniba.it Received August 12<sup>th</sup>, 2011; revised September 20<sup>th</sup>, 2011; accepted October 29<sup>th</sup>, 2011.

Depression is an independent risk factor of poor outcomes for Chronic Kidney Disease (CKD) patients. Perceived social support and alexithymia are psychosocial variables identified by previous studies as predictive of depression in normal controls and CKD patients. Repetitively thinking and socially sharing emotional experiences have been investigated in association with depression in normal populations. Our cross-sectional study aimed to assess the effects of perceived social support, alexithymia, mental rumination, and social sharing on depression in CKD patients and controls. 103 CKD patients (age =  $61.9 \pm 7.2$ , 54 men) and 101 controls (age = 64.51 ± 6.56; 47 men) completed a questionnaire of 5 sections: Pluridimensional Inventory for Haemodialysis Patients (IPPE), Multidimensional Scale of Perceived Social Support (MSPSS), Geriatric Depression Scale (GDS), Toronto Alexihymia Scale (TAS-20), Social Sharing and Mental Rumination. Multiple regression analysis models with dummy variables assessed the effects of IPPE, MSPSS, TAS-20, Social Sharing, and Mental Rumination on GDS across the subgroups of participants. SPSS software was used. Depression levels resulted higher for patients than controls, especially in patients dialyzed for less than 4 years. The effects of perceived social support and alexithymia differed with respect to the subsamples. Rumination was positively associated with depression in normal controls, but negatively related with depression in patients dialyzed for 4+ years. The study confirmed high levels of depression in CKD patients. Depression was influenced by perceived social support, alexithymia, and the cognitive elaboration of emotional troubles associated with the disease. Rumination appeared as a dysfunctional consequence of emotions for normal controls, but had an adaptive function for patients dialyzed for 4+ years.

Keywords: Depression, Perceived Social Support, Alexithymia, Mental Rumination, Social Sharing, CKD

## Introduction

Depression has been shown to be a common psychiatric disorder in patients affected by chronic kidney disease (Hedayati et al., 2009). It has been found to be prevalent in patients with end-stage renal disease (ESRD) on long-term haemodialysis or following kidney transplantation, with an estimated prevalence rate between 15% and 40% (Ricardo et al., 2010). Previous studies have demonstrated that depression is an independent risk factor for poor outcomes in CKD patients (Hedayati et al., 2008; Hedayati et al., 2010).

Social support has been defined as the availability of significant interpersonal relationships which make individuals feel cared for, esteemed, and valued (Sarason et al., 2009). The concept involves two broad aspects: 1) social support as actually received (functional and structural support); 2) the individual's subjective perception of availability of social support (perceived social support). It has been demonstrated that the availability of social support is an important protective factor against depression in chronic illnesses (Hoth et al., 1997). Further, the perception of social support has been found to be associated with reduction of depression, increased perception of quality of life, increased access to health care, compliance with prescribed therapies, and direct physiological effects on the immune system for patients with ESRD (Cohen et al., 2007).

Individuals with available social support have more opportunities to talk about their state of health, diagnosed disease and symptoms, share their worries and concerns, and manage the

emotional burden of their illness with significant others, such as spouses, family members, close friends. Among the behavioral consequences of the availability of social support there is a phenomenon called social sharing, which entails the description, in a socially shared language, of emotional events to an addressee by the person who experienced them (Zech & Rimé, 2005). Social sharing has been demonstrated to be a highly pervasive phenomenon, since it occurs after an emotional experience in 80% to 95% of cases in all countries and cultures (Rimé, 2009). In samples drawn from normal populations social sharing has been found to be associated with increased health benefits as assessed by physician visits, reported symptoms, immunological functions, and indices of subjective well-being (Frattaroli, 2006).

Social sharing is generally associated with another process called mental rumination, i.e. the intra-individual persistence of thoughts, images, and physiological symptoms related to an emotional experience. Rumination has been originally regarded as a strategy for coping with emotional experiences. According to this view, rumination arises when individuals experience interruptions while pursuing their goals, and emotions originate as a consequence of these interruptions. The more relevant the interrupted goal, the higher the need for the individual to restore it. Rumination is the strategy adopted by individuals to deal with the interruption of progress towards their goals, and it terminates when the objective is reached or is deemed impracticable and then abandoned (Martin & Tesser, 1989). On the other hand, numerous studies have suggested a close relation-

ship between rumination and depression since, through mental rumination, depressed individuals keep focusing on their symptoms and the possible consequences. This has in turn the effect of increasing the state of depression in a self-perpetuating cycle of self-focused thoughts (Roberts, Gilboa, & Gotlib, 1998).

The constructs of social sharing and mental rumination have been broadly investigated in the field of psychology of emotions. Another important line of research focuses on the association between rumination and a personality trait called alexithymia, which has been conceptualized as an individual deficit in identifying, communicating, and regulating emotions. Alexithymia is a multifaceted construct including three salient features: 1) difficulty identifying feelings and discriminating them from bodily sensations of emotional arousal; 2) difficulty expressing emotions and describing them to others: 3) limited capacity to engage in fantasy and other imaginative activities (Taylor, 1994). The construct of alexithymia has relevant implications for individuals' health, since it has been found to be associated with a broad range of psychiatric symptoms, psychosomatic problems, and chronic illnesses (Taylor, 1994; Kauhanen et al., 1996). Alexithymia has also been demonstrated to be a stable risk factor for the development of depressive disorders in cross-sectional and cohort studies on general and clinical samples (Grabe, Spitzer, & Freyberger, 2004).

Studies have examined the independent contribution of social support and alexithymia on depression (Kojima et al., 2007) and on long-term mortality (Kojima et al., 2010) of CKD patients. However, to our knowledge, no studies have compared the conditional effects of social support, alexithymia, sharing and rumination processes upon depression of ESRD patients and controls. The present study has two main goals: 1) to confirm the findings concerning the higher prevalence of depression in ESRD patients; 2) to evaluate the relationship between depression, social support, alexithymia, mental rumination, and social sharing of emotionally relevant experiences. Indeed, we expected that ESRD patients would suffer from depressive disorders to a higher extent than controls, and that social support and sharing would be valuable protective factors against patients' depressive disorders, while alexithymia and mental rumination would have the opposite effect.

### **Subjects and Methods**

## **Subjects**

The data for this cross-sectional study were collected by means of a computerized database containing demographical and clinical information regarding all consecutive ESRD patients, older than 18 years, entering maintenance dialysis treatment programs at three dialysis units in Southern Italy from 1 January 1995. This group was made up of 103 subjects who have been undergoing dialysis treatment programs for at least 1 year. Table 1 contains their demographical characteristics. Matched for age, gender and race with the patients' subsample, a group of 101 elderly participants were also involved in the study (age =  $64.51 \pm 6.56$ ; 47 men).

## **Data Collection**

On the basis of previous reports, the covariates identified *a priori* as possible risk factors for depression were age, gender and, for the clinical subsample, time elapsed since the beginning of haemodialysis programs. Indeed both the clinical his-

tory and examination of the patients were taken into consideration in order to assess the comorbidities according to the ICED score (Miskulin et al., 2003). The ICED score aggregates the presence and severity of 19 medical conditions and 11 physical impairments within two scales: the Index of Disease Severity (IDS) and the Index of Physical Impairment (IPI). The final ICED score (ranged from 0 to 3 reflecting increasing severity) is determined by an algorithm combining the peak scores for the IDS and IPI. All the patients underwent a 4 h dialysis session thrice weekly, on high and low-flux membranes, on standard bicarbonate dialysate, using 15-gauge fistula needles. The patients were anticoagulated by using systemic heparin. Table 1 shows the clinical characteristics of patients involved in the study.

### **Design and Procedure**

The present study was performed in accordance with the Helsinki declaration for studies on humans and it was approved by the local ethics committee. All participants were involved in the study on a voluntary basis. Before answering the questionnaire they were required to give informed consent.

The whole session lasted 30 minutes on average. Patients were recruited while waiting to undergo routine medical treatments in hospital; controls were recruited from among the researchers' acquaintances. Participants were asked to fill in a questionnaire made up of different sections.

The first section contained the IPPE (Pluridimensional Inventory for Haemodialysis Patients) (Biasioli & Ballaben,

Table 1.

Patients demographical and clinical characteristics.

N		103			
Age (years)		$61.9 \pm 7.2$			
Gender (M/F)		59/44			
Race		100% Caucasian			
Renal Diseases (% of t					
	Glomerular	20.9			
	Hypertensive	52.9			
	Vascular	10.0			
	Tubulo-Interstitial	16.2			
Comorbidities (n/% of patients)					
	Diabetes	24/22.9			
	Hypertension	44/41.9			
	Chronic Heart failure	29/27.6			
	Arrhythmia	7/6.7			
	Stroke	28/26.6			
	Peripheral Vasculopathy	20/19.3			
	Chronic lung diseases	8/7.6			
	Neoplasms	10/9.5			
Biasioli, S., & Ballaben, P.	Liver diseases	6/5.7			
ICED Score	(%)				
	0	16.2			
	1	17.1			
	2	33.3			
	3	33.4			

2003), an instrument specifically intended to assess the quality of the psychological life of haemodialysis patients through 24 sentences to be rated on 4-point scales. The IPPE comprised 6 dimensions (family relationships, body, need to drink, daily life, medical support, perception of CKD state). This section was omitted for control participants.

The second section contained the MSPSS (Multidimensional Scale of Perceived Social Support) (Zimet et al., 1988), a 12-item inventory of 7-point scales, assessing 3 dimensions of perceived social support concerning family, friends, and significant others.

The third section contained the GDS (Geriatric Depression Scale) (Yesavage et al., 1983), an inventory consisting of 30 dichotomies (yes/no) designed to evaluate depressive symptoms in elderly people.

The fourth section contained the TAS-20 (Toronto Alexithymia Scale) (Parker, Taylor, & Bagby, 2003), a 20-item inventory of 5-point scales, assessing 3 dimensions of alexithymia, concerning difficulty in identifying feelings, difficulty in describing feelings, and externally oriented thinking.

Finally, the fifth section enclosed measures of intensity of emotion (11-point scale, ranging from 0 = no emotion to 10 = the highest degree of emotion in life), and the consequent social sharing and mental rumination processes (5-point scales ranging from 1 = never to 5 = very often, concerning need for sharing, actual social sharing, frequency of thinking, frequency of spontaneous images) (Rimé, Mesquita, Philippot, & Boca, 1991). For the patients' groups, the measures in Section 5 related to the experience of first hearing of the CKD diagnosis; for the control group, the questions related to the highest degree of emotional event experienced in recent years.

## **Statistical Analyses**

Analyses were run on the aggregated indices obtained for each scale in the questionnaire. More specifically, for the IPPE, six indices were computed by averaging the items of the scales (Biasioli & Ballaben, 2003), i.e. Family relationships, Body, Need to drink, Daily Life, Medical Support, Perception of CKD state (Cronbach's alphas = .80 on average). For the MSPSS, the three dimensions of Family, Friends, and Significant Others were obtained by averaging four items for each dimension (average Cronbach's alpha = .96) (Zimet et al., 1988). GDS items were dichotomies summed to obtain a general score for Depression. This score ranges from 0 (=not depressed) to 30 (= max depression), with a cut-off of 11 corresponding to the presence of clinically relevant depressive symptoms, and a cut-off of 18 corresponding to the presence of severe depressive symptoms (Yesavage et al., 1983). For the TAS-20, items were selectively summed to obtain three aggregated scores corresponding to Difficulty in Identifying Feelings (DIF), Difficulty in Describing Feelings (DDF), and Externally-Oriented Thinking (EOT) (average Cronbach's alpha = .70) (Parker, Taylor, & Bagby, 2003). Finally, for the fifth section of the questionnaire, scores for need for sharing and actual social sharing were averaged into a composite index of Social Sharing; scores for frequency of thinking and frequency of spontaneous images were averaged into a composite index of Mental Rumination (average Cronbach's alpha = .87).

Preliminarily, patients were classified in two groups: Those who have been undergoing haemodialysis treatment programs for less than 4 years ( $2.05 \pm .87$  years; 56.3%), and those who have been treated with haemodialysis programs for more than 4 years ( $10.00 \pm 5.01$  years; 43.7%). This choice was made for

two reasons. The first was that after an average of four years undergoing haemodialysis treatment, CKD patients are eligible for kidney transplantation (Leichtman et al., 2008). It follows that 4 years represent an important cut-off point in the life of dialyzed patients. The second reason is statistical: in the subsequent analyses the cut-off of 4 years was found to maximize the effect size for all comparisons.

In a second step, GDS levels (absence of depressive symptoms vs. mild depressive symptoms vs. severe depressive symptoms) were compared across the three groups of participants (dialyzed patients for less than 4 years vs. dialyzed patients for 4+ years vs. controls). Further, responses to Section 5 of the questionnaire were analyzed to verify whether patients' experience of hearing of the CKD diagnosis was comparable with the occurrence of a high-level emotional experience for controls.

In a third step, average IPPE scores were compared across the two groups of patients through a Multivariate model of Analysis of Variance (MANOVA). A second MANOVA was then run to compare the average MSPSS, GDS, TAS-20, Social Sharing, and Mental Rumination scores across the two subgroups of patients and the control group. Post-hoc analyses were carried by applying Tukey's HSD test on the average scores for the three subgroups.

Finally, multiple regression analysis models with dummy variables were employed to assess the effects of IPPE, MSPSS, TAS-20, Social Sharing, and Mental Rumination on GDS scores across the subgroups of participants. First, a multiple regression model was run on the total sample data, with the GDS scores as dependent measures. Two dummy variables-Condition (control = 0, patients = 1), and Gender (female = 0, male = 1)—and a set of continuous predictor variables (Age, the three MSPSS scores, the three TAS-20 scores, Social Sharing, Mental Rumination) were entered in the model, as well as their 2-way, 3-way, and 4-way interaction terms. However, only the test of the simple slopes (i.e., the 4 conditions in Table 2) was analyzed, and the conditional effects for all predictors in the model (dummy and continuous variables) were retained. The main advantage of the procedure consists in the opportunity to examine the conditional effects of each predictor in the model, i.e. the effect of a given variable when other interacting variables are included in the model. To make easier the interpretation of the effects, all continuous predictor variables were centred before running the analysis. The reference categories for the dummy variables in the model were control = 0 and female = 0. Second, a multiple regression model was run on the subsample of patients to assess the effects of the Duration of their haemodialysis program (dummy variable with 0 = lessthan 4 years vs. 1 = 4+ years), IPPE, MSPSS, TAS-20, Social Sharing, and Mental Rumination scores on their GDS scores. In this model, Gender and Age were not considered as covariates since introducing their interaction terms with the other predictors would have dramatically affected the degrees of freedom of the model. It follows that only the 2-way interactions of the dummy variable with the other continuous predictors were analyzed, and the conditional estimates of the simple slopes for the two subgroups of patients were reported in Table 3. All data were processed using SPSS software.

## **Results**

As shown in Table 1, most patients were characterized by an ICED score higher than 2, reflecting the usual features found

Table 2.

Multiple regression analysis with dummy and continuous predictor variables on the GDS scores—total sample.

Predictors					
R-squared = .81	Beta (SE)				
F(69, 129) = 8.12****					
	Gender				
	Female		Male		
Condition					
0 = control	9.71 (3.05)***		1.86 (1.68)		
1 = dialysis					
	Condition				
	Control		Dialysis		
Gender					
0 = female	2.48 (2.13)		-5.37 (2.39)*		
1 = male					
	Female control	Male control	Female dialysis	Male dialysis	
Age	.57 (1.18)****	.02 (.24)	.74 (.29)*	.18 (.12)	
MSPSS – Family	79 (.46)	-1.02 (.42)**	.48 (.70)	.25 (.35)	
MSPSS – Friends	.38 (.38)	.49 (.29)	-1.05 (.82)	56 (.18)***	
MSPSS – Others	.57 (.37)	.19 (.30)	.06 (.96)	47 (.30)	
TAS-20 – DIF	.22 (.19)	.26 (.30)	.10 (.51)	1.02 (.19)****	
TAS-20 – DDF	.58 (.37)	1.08 (.39)**	.98 (.85)	-1.46 (.43)***	
TAS-20 – EOT	.19 (.28)	29 (.49)	42 (.45)	1.02 (.25)****	
Social sharing	.48 (.99)	-1.03 (1.36)	2.23 (3.95)	-1.76 (.93)	
Mental rumination	3.69 (1.15)***	3.91 (1.58)*	-1.55 (3.46)	68 (1.25)	

Notes: All regressions have as reference categories controls = 0 and female = 0; Beta coefficients are shown with their SE in parenthesis, and the significance level of the associated t-test with 1 df; The sign of the beta coefficient is positive for the cell corresponding to dialyzed and female participants, accounting for a positive association between these two conditions, and negative for the cell of dialyzed and male patients accounting for a negative association between these two conditions; \*p < .05, \*\*p < .01, \*\*\*p < .005, \*\*\*\*p < .001.

worldwide in the dialysis units during everyday clinical practice. Thus our cohort of patients is to be considered as representative of the typical uremic population, making the results found in this paper easily applicable to all real situations.

The preliminary analysis of the GDS levels across the three subgroups of participants (dialyzed patients for less than 4 years vs. dialyzed patients for 4+ years vs. controls) yielded significant results (chi-square (4, N=204)=21.21, p<.001). Figure 1 showed that the absence of depressive symptoms was more frequent for controls than for patients, while dialyzed patients for less than 4 years appeared generally more depressed than patients undergoing haemodialysis programs for 4+ years (respectively 79.3% and 64.4% mild to severe depressive symptoms).

With respect to the data from Section 5 of the questionnaire, for the patients, the experience of first hearing of the CKD diagnosis dated back  $9.73\pm7.87$  years, while for the control group, the recalled emotional event was significantly more recent, dating back  $2.52\pm2.32$  years on average (t(167)=7.62, p<.001). Among the control participants, 30.7% recalled a personal or family health problem due to illness or accident, 23.8% reported the death of a close relative or friend, 12.9% relational troubles (i.e. relationship breakdown, divorce, etc.), 11.9% problems at the workplace, 6.9% family problems, 4.0% victimization experiences (i.e. being robbed, assaulted, etc.),

while 9.9% of controls failed to recall a specific emotional experience. Only 8 of the CKD patients were not able to recall the moment they first knew of their diagnosis.

No significant difference was found between the emotional impact of the first hearing of the CKD diagnosis for the patients' subgroup and the emotional experience of the controls (respectively,  $9.05 \pm 1.85$ , and  $8.77 \pm 2.10$ , t(195) = 1.01, n.s.). Similarly, no significant differences were found between the patients and the control subgroups in the measures of Social Sharing (respectively,  $3.38 \pm 1.10$ , and  $3.48 \pm 1.08$ , t(198) = .70, n.s.), and Mental Rumination (respectively,  $4.42 \pm .63$ , and  $4.20 \pm .94$ , t(197) = 1.92, n.s.).

In a subsequent step, average measures of IPPE were compared across the two subgroups of patients (less than 4 years vs. 4+ years dialyzed). The multivariate Fisher's F for the MANOVA was found to be significant (F(6, 96) = 7.28, p < .001). As shown in Figure 2, this effect was due to the significant differences in three of the IPPE dimensions, corresponding to Body (F(1, 101 = 6.30, p < .05), Medical Treatments (F(1, 101) = 7.72, p < .01), and Perception of the CKD State (F(1, 101) = 22.58, p < .001) in those patients undergoing haemodialysis treatment programs for 4+ years reporting higher scores on these subdimensions than patients undergoing haemodialysis for less than 4 years. The overall result of the MANOVA on the MSPSS, GDS, TAS-20, Social Sharing, and

Table 3. Multiple regression analysis with dummy and continuous predictor variables on the GDS scores—patients' subsample.

Predictors	
R-squared = .89	Beta (SE)
F(29, 71) = 19.35****	
Dialysis duration	
0 = Up to 4 years	1.92 (1.49)
1 = 4 + years	

	Less than 4 years	4+ years
	Less than 4 years	4+ years
Ippe—familiar relationships	61 (1.43)	-4.84 (1.80)**
Ippe—body	-1.09 (.63)	1.05 (.95)
Ippe—need to drink	.98 (.77)	42 (1.31)
Ippe—daily life	-1.47 (.99)	-2.33 (1.73)
Ippe—medical support	3.71 (1.18)***	1.90 (1.57)
Ippe—Perception of CKD state	15 (.67)	1.01 (1.31)
MSPSS—Family	.66 (.27)*	-1.05 (.45)*
MSPSS—Friends	22 (.15)	39 (.16)*
MSPSS—Others	45 (.23)	.41 (.39)
TAS-20—DIF	.68 (.13)****	.34 (1.44)
TAS-20—DDF	.70 (.24)***	17 (.47)
TAS-20—EOT	36 (.19)	.75 (.29)*
Social sharing	-1.38 (.85)	1.49 (.75)
Mental rumination	.52 (.99)	-2.97 (1.25)*

Notes: All regressions have as a reference category less than 4 years haemodialysis = 0; Beta coefficients are shown with their SE in parenthesis, and the significance level of the associated t-test with 1 df; \*p < .05, \*\*p < .01, \*\*\*p < .005, \*\*\*p < .001

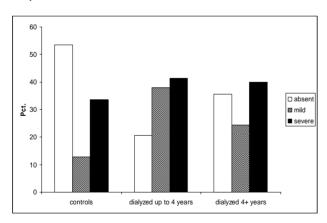


Figure 1.

Distribution of GDS levels across the three subgroups of participants.

Mental Rumination scores by condition (patients dialyzed for less than 4 years vs. patients dialyzed for 4+ years vs. controls) were also found to be significant (F(18, 376) = 7,26, p < .001). As shown in Figure 2, this was due to the significant differences across groups in the GDS (F(2, 196) = 4.33, p < .05) and TAS-20-DIF scores (F(2, 196) = 15.83, p < .001), with the highest scores reported by patients who had been undergoing haemodialysis programs for less than 4 years, and the lowest reported by controls (Tukey's HSD ps < .05).

The results of the first multiple regression analysis run on the

total sample GDS scores are reported in Table 2. The model generally has a good fit. Women dialyzed appeared as the most depressed subgroup, and age affected depression for both dialyzed and control females. Social support had a negative impact on depression for male participants: More specifically, social support from Family negatively affected GDS scores for male controls, while social support from Friends negatively affected GDS scores for male patients. All three dimensions of TAS-20 impacted upon GDS scores for male patients, with DFI and EOT being positively associated, and DDF negatively associated. By contrast, DDF was associated with a positive beta coefficient for male controls. Finally, Social Sharing was never associated with GDS scores, while Mental Rumination was found to have a significant positive impact upon GDS for control participants.

The results of the multiple regression analysis run on patients' GDS scores are reported in Table 3. This model has a good general fit. For patients undergoing haemodialysis programs for less than 4 years, significant predictors of GDS appeared to be the IPPE-Medical Support, DIF, and DDF dimensions. For patients undergoing haemodialysis treatment programs for 4+ years, significant predictors proved to be IPPE-Familiar Relationship, MSPSS-Family, MSPSS-Friends, EOT, and Mental Rumination. All these predictors exhibited negative associations with GDS scores except the EOT dimension, whose beta coefficient was found to be positive.

### **Discussion**

Results of the present study confirmed the higher prevalence of depression among CKD patients as compared with normal controls. In addition, patients undergoing haemodialysis treatments for less than 4 years appeared comparatively more depressed than patients dialyzed for 4+ years. This finding is consistent with the assumptions of the Kübler-Ross (2009) fivestage model of attitudes toward death. According to this model, terminally ill patients undergo five stages of elaboration of their grief: Denial, Anger, Bargaining, Depression, and Acceptance. Depression precedes the final stage of Acceptance. In the stage of Depression, terminally ill patients are aware of the losses they have suffered because of their state of health (reactive depression), and became progressively more aware of the fact that these losses are going to increase (preparatory depression). Subjective pain and practical implications of the disease and necessary medical treatments are amplified. Patients experience a sense of defeat. In the subsequent stage, called Acceptance, patients have had the chance to elaborate their experience, so that they become aware of its inevitable ending. Feelings of anger and depression may still be experienced, nonetheless they have moderate intensity. At this stage patients may experience a sense of profound intimacy with their close relatives. It should be noted that Acceptance does not necessarily overlap with the very final phase of the illness, when patients may still experience denial, anger or depression. Besides the drawbacks mainly due to the limited empirical support for the model (Maciejewski, Zhang, Block, & Prigerson, 2007), the Kübler-Ross conceptualization represents a useful outline of the psychological processes involved in the individual's coping with a terminal disease. As such, it could be applied to ERSD patients. In the present study we adopted a cut-off of 4 years to divide patients into two groups. We reasoned that after 4 years of haemodialysis treatments, patients are sufficiently aware of the serious implications of their chronic disease. After 4 years on average

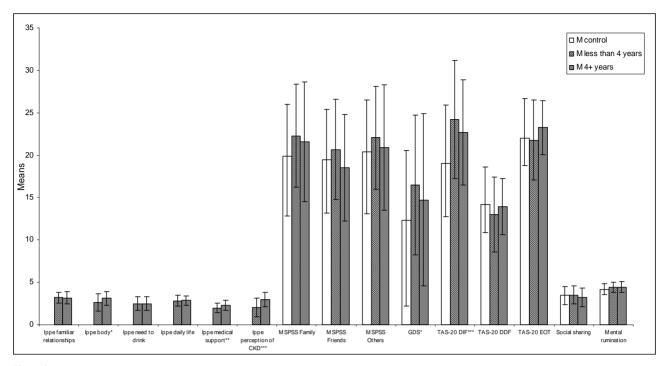


Figure 2. Ms and SDs IPPE, MSPSS, GDS, TAS-20, social sharing, and rumination. Note: \*p < .05, \*\*p < .01, \*\*p < .001.

from the beginning of their programs, patients generally have a chance of getting kidney transplantation (Leichtman et al., 2008). Whether or not they accept a transplant, in 4 years patients have had the opportunity to elaborate their experience and accept the disease and its consequences. It follows that the cut-off of 4 years roughly parallels the distinction of the 5-stage model between the Depression and Acceptance phases. This intuition was confirmed by the statistical analyses run in the present study that showed that the cut-off of 4 years maximized the effects for all comparisons.

Dialyzed women appeared significantly more depressed than dialyzed men. Age also played a significant effect for women in both the dialyzed and control groups, since their depression increased with age. These findings are consistent with the literature on gender differences in depression. It is well established that women are about twice as likely as men to develop depression across different cultures, and ethnicities (Nolen-Hoeksema, Larson, & Grayson, 1999; Weissman et al., 1996).

Even in presence of the same stressors, women are more likely to react with depressive disorders than men. Many explanations have been proposed for this fact, such as interpersonal stress (Conger et al. 1993), self esteem (Cambron, Acitelli, & Pettit, 2009), cognitive and affective processes (Mezulis & Fanasaki, 2009; Nolen-Hoeksema & Morrow, 1993), biological and hormonal factors (Grigoriadis & Robinson, 2007), coping style (Nolen-Hoeksema, Larson, & Grayson, 1999; Broderick & Korteland, 2002), social and cultural constraints (Leach et al., 2008), although evidence is rather inconclusive on this regard. Additionally, in previous studies the effect of gender was also found to be more pronounced as age increases (Umberson, Wortman, & Kessler, 1992).

Perceived family support had a significant effect upon depression of male controls, while friends' support and alexithymia had a significant impact on depression in dialyzed men. These findings are of interest for the understanding of the protective factors against depression and the treatment of depressive disorders. Elderly men suffer more from social isolation than women (Kendler, Myers, & Prescott, 2005). Further, alexithymia and related emotional difficulties have typically been found to be associated with male gender (Parker, Taylor, & Bagby, 2003; Lane, Sechrest, & Riedel, 1998). The perception of low availability of social support and alexithymia make men susceptible to developing depression. By contrast, women generally have larger networks of significant others, and are affected by alexithymia to a lesser extent than men. This implies that treatment for depression focusing on reducing the sense of isolation and difficulty in expressing emotions would be more successful for men than for women. Finally, no significant effects were found for social sharing, while mental rumination increased depression in both male and female controls. This probably happens because individuals have failed to conclude the psychological elaboration of their emotional experiences.

Repetitive thinking leads to the maintenance of a disturbing cycle of perseverating thoughts which in turn increase the negative emotional state following the original experiences and the concurrent depression (Nolen-Hoeksema, 1991; Nolen-Hoeksema & Morrow, 1993). With respect to ESRD patients, although their level of depression was generally higher than for controls, ruminating about the first hearing of CKD diagnosis was not found to have effects on increasing their level of depression.

Focusing on CKD patients, no significant differences were found in their level of perceived social support, social sharing, and mental rumination. Patients who had been undergoing haemodialysis treatment programs for 4+ years appeared less satisfied with their physical condition, required more medical treatments, were more concerned by their CKD state, and experienced more difficulties in identifying feelings than patients who had been undergoing such treatments for less than 4 years. With respect to the effects of the perceived quality of life and social support variables in the regression model, it should be

noted that the signs of the coefficients are different for the two subgroups of patients: while perceived quality of life and perceived social support diminished the level of depression in patients dialyzed for 4+ years, these variables had either the opposite or no effects at all on depression in patients dialyzed for less than 4 years. This accounts for the fact that, although the patients' perception of quality of life and the level of perceived support do not change during the course of the treatment program, these perceptions have the beneficial effect of reducing patients' depression only in the very last stages of the disease. This is consistent with what was reported above concerning the stages of psychological elaboration of the disease by terminally ill CKD patients (Kübler-Ross, 2009). In the stage called Acceptance, patients might rely upon their individual and social resources to cope with their disease. Depression could be alleviated by the perception of significant others being available to provide support, and by an ultimate adjustment with the ESRD difficulties.

Alexithymia also had a significant effect on CKD patients' depression, but this effect involved different dimensions with respect to the duration of patients' haemodialysis program: while DIF and DDF increased depression in patients dialyzed for less than 4 years, the EOT dimension intensified depression in patients who had been undergoing haemodialysis program for 4+ years. The first two dimensions of TAS-20 have frequently been found associated with depression in a number of studies run on both normal and psychiatric samples (Marchesi, Brusamonti, & Maggini, 2000; Hendryx, Haviland, & Shaw, 1991). With respect to the EOT dimension, it could be possible that 4+ years dialyzed patients prefer to employ introspective strategies of thinking to regulate their depressive symptoms. Focusing on their internal psychological states might be useful for long-term dialyzed patients to reduce the psychological burden of living with their disease. On the other hand, it has been variously shown that the subscale EOT has a lower internal consistency and more problematic psychometric properties than the other two TAS-20 dimensions (Gignac, Palmer, & Stough, 2007; Kooiman, Spinhoven, & Trijsburg, 2002). Also in this study, the EOT Cronbach's alpha (.52) is lower than the alphas of the DIF and DDF dimensions (respectively, .90 and .70). Therefore, these measurement issues may have interfered with our results (Gignac, Palmer, & Stough, 2007).

Finally, social sharing processes did not exhibit significant associations with CKD patients' depression, but mental rumination had the effect of reducing depression in patients dialyzed for 4+ years. Unlike controls, CKD patients dialyzed for 4+ years employed rumination strategies in an adaptive way. Far from being the effect of a dysfunctional persistence of unwanted thoughts, rumination appeared to be a useful way of reducing the symptoms of depressive disorder and appease the individual in the terminal phases of CKD. One possible theoretical explanation for these findings is that mental rumination serves as a strategy for coping with interruptions in pursuing significant goals. Through rumination, individuals attempt to reduce disruption, and reinstate the unaccomplished goals (Martin & Tesser, 1989). The unpredictability of CKD and the imminence of death certainly represent for ESRD patients the ultimate interruption of their life goals (Schipper & Abma, 2011). Ruminating may help patients to restore this interruption and create a sense of prolongation of their life beyond illness and death. A definitive empirical evaluation of this interpretation would be possible by analyzing the contents of CKD patients' ruminative thoughts, and comparing them with the selfevaluated coping strategies adopted by patients to deal with

health issues. This would be an interesting avenue for future research on the factors associated with depression in CKD patients in order to prevent poor outcomes in their treatment.

Some comment is required on the absence of significant relationships between social sharing processes and depression in both CKD patients and controls. These findings contradict the expectations of the present study, and leave unsolved the problem of defining the behavioural processes through which the perception of social support reduces depressive symptoms. It might be that individuals do not need to share the content of their emotional experiences to reduce depression, and that simply figuring out that significant others are available to provide support when needed is sufficient to alleviate the state of depression. Alternatively, it is possible that social sharing is not the only way people exchange and receive support, and that the mere presence of significant others, their availability to provide material goods or practical assistance are important instances of social support. Again, future research work could help to clarify the issue concerning social and psychological factors contributing to the alleviation of depression in CKD patients.

The present study has some limitations. First, the relationships between patients' depression, perceived social support, alexithymia, social sharing, and mental rumination of emotions need to be investigated through a cohort design in order to evaluate the temporal evolution of these relationships and the different impact of the psychosocial factors with respect to the progression of CKD. Employing a cohort design would also help to clarify the issue of the causal relationship between depression and adverse outcomes in CKD patients. Second, assessing CKD patients' depression has often been recommended as a clinical means of ensuring accurate diagnosis, effective treatment, and follow-up (Watnick, 2007). However, no specific instruments are available to evaluate depression in CKD patients. The most frequently used inventories for this purpose are the Beck Depression Inventory (BDI) (Craven, Rodin, & Littlefield, 1988) the Center for Epidemiologic Studies Depression Scale (CESD) (Andresen et al., 1994), and the Patient Health Questionnaire-9 (PHQ-9) (Watnick, Wang, Demadura, & Ganzini, 2005). These instruments are brief self-report scales for the screening of depression that have also been validated on samples of CKD patients. The sample for the present study consisted of individuals—dialyzed and controls—aged over 60. As a consequence, we employed the GDS as a specific instrument to assess depression in elderly individuals (Yesavage et al., 1983). Using GDS is not a common practice in research on CKD patients, thus the psychometric properties of the scale applied to CKD populations require careful consideration. Third, the sample of the present study needs to be enlarged in order to allow for an accurate analysis of the interactions between the variables associated with depression and the covariates of the design (gender and age in the present study). Fourth, the sample for the present study consisted of elderly people, therefore results cannot be generalized to the entire population of patients undergoing haemodialysis programs. Fifth, in the present study the indices of quality of life for haemodialysis patients were included in the set of potential predictors of depression in ESRD patients. However, the perception of quality of life could also be considered a mediator variable in the relationship between social support, alexithymia, social sharing, mental rumination, and patients' depression. To test this hypothesis, mediation analysis or structural models of direct and indirect relationships should be employed and this in turn requires a considerable enlargement of the sample size. Finally, a deeper analysis of the content and characteristics of mental

rumination is required. The pattern of results concerning the effects of rumination on depression found in the present study could be due to the fact that, when ruminating about past emotional events, individuals focus on different aspects of their experience and adopt different styles of thinking, which can in turn impact upon their level of depression (Watkins & Teasdale, 2001).

To sum up, the present study offers interesting indications for the understanding of the relationship between psychosocial factors and depression in patients with CKD. As long as the state of disease progresses and patients' need for medical treatments increases, psychological adjustment evolves towards an acceptance of the disease and a significant reduction in the psychological symptoms of distress. This process is influenced by the way people perceive others supporting them, individual personality traits, and the cognitive elaboration of emotional troubles associated with the disease. A purposeful analysis of the way these psychosocial factors impact during the progress-sion of CKD could bring important benefits for the therapy of ESRD patients.

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