Data Supplement for Bauer et al. (10.1176/appi.ps.201200121) Characteristic Caregivers Patients Detail (%) (%) n n 39 42 Female 65 25 (41.7) Sex 21 35 35 (58.3) 58 Male \leq 30 years 5 8 28 (46.7) 47 Age 31 to 40 years 13 22 11 (18.3) 18 19 32 41 to 50 years 9 (15.0) 15 51 to 60 years 13 11 18 8 (13.3) > 60 years 12 29 4 (6.7) 7 Mean age in years (SD) 48.8 37.3 (15.4)(15.0)Highest level of Secondary school to end of 24 40 21 (35.0) 35 education Grade 9 37 17 (28.3) 28 22 Secondary school to end of 7 12 10 (16.7) 17 Grade 10 7 20 12 12 (20.2) Secondary school to end of Grade 12 University/College Full time/Part time/In Occupational 34 57 31 (51.7) 52 situation education or training 42 Not working/Retired 26 43 25 (41.7) 7 No information 4 (6.7) 15 25 Single Marital status Living with partner 45 75 Degree of kinship: Spouse/Partner 21 35 Patient is ... 17 Parent 38 7 12 Child Sibling 104 17 7 Other relative/Friend Common household Yes 37 62 with patient No 23 38 4 45 Size of household 7 1 person 77 2-4 people 10 5 or more people 17 39 65 Support in caring Yes for the patient 21 35 No 9 15 Family's monthly ≤€1000 net income €1001 - €2000 28 47 €2001 - €3000 27 16 > €3000 10 6 2 Not specified 1 Contact with patient ≤ 2 hours 5 15 11 5 8 per week More than 2 hours to 5 25 hours 18 More than 5 hours to 15 8 hours 40 More than 15 hours to 35 hours

	> 35 hours				
Distance between	≤ 20 km	24	40	_	
place of residence	More than 20 km to 50 km	15 21	25		
and hospital	> 50 km	10 21	35		
Caregiver has legal	Yes	8 52	13	_	
responsibility for	No	0.52	87		
the patient	110		07		
Participation in	Yes	7 53	12	-	
caregiver group	No		88		
Duration of illness	≤ 15 years	-		37 (61.6)	62
	> 15 years			12 (20.0)	20
	No information			11 (18.3)	18
Previous	Yes	-		13 (21.7)	22
psychotherapeutic	No			37 (61.7)	62
treatment	No information			10 (16.7)	17
Ever attempted	Yes	_		14 (23.3)	23
suicide	No			41 (68.3)	68
	Unknown/Unclear			5 (8.3)	8
Clinical Global	"Moderately ill" or less			8 (13.3)	13
Impression (CGI)	More than "moderately ill"	-		49 (81.7)	82
Part 1 (Admission)	No information			3 (5.0)	5
Clabal Assessment					
Global Assessment of Functioning	$\leq 40 \text{ points}$	-		37 (61.7)	62
(GAF) (Admission)	41-100 points			22 (36.7)	37
, , , , , ,	No information			1 (1.7)	2
	Mean score (SD)	-		38.6 (12.2)	-
Care situation	No care			38 (63.3)	63
24 hours after		-			28
admission	Care arranged			17 (28.3)	
) (No information			5 (8.3)	8
Mental illness in first degree relative	Yes	=		13 (21.7)	22
inst degree relative	No			27 (45.0)	45
	No information			20 (33.3)	33
Duration of current hospital stay	\leq 50 days	-		25 (41.7)	42
nospitai stay	> 50 days			32 (53.3)	53
	No information			3 (5.0)	5
Informative	Yes	-		22 (36.7)	37
discussions with caregiver/	No			35 (58.3)	58
psychoeducation	No information			3 (5.0)	5
Clinical Global	"Moderately ill" or less	-		25 (41.7)	42
Impression (CGI)	More than "moderately ill"			30 (50.0)	50
Part 1 (Discharge)	No information			-5 (8.3)	8
	Mean score (SD)	-		3.2 (0.9)	-
Global Assessment	$\leq 40 \text{ points}$	_		7 (11.7)	12
of Functioning	41-100 points			50 (83.3)	83
(GAF) (Discharge)	No information			3 (5.0)	5
	Mean score (SD)	_		57.2	-
	Tyledii scole (SD)			((15.9)	_
Living situation of	Alone	-		16 (26.7)	27
patient after	With family member(s)			42 (70.0)	70
discharge	No information			2 (3.3)	3
	1.5 momenton	<u> </u>	1	12 (3.3)	٦

Table 2: Frequency of burdens and rewards named by caregivers in each diagnostic group

		ICD-10 diagnosis					
	Total	Schizophrenia (F20) (n=15)	Depression (F31) (n=15)	Bipolar Disorder (F32/33) (n=15)	Personality Disorder (F60/61) (n=15)		
Number of burdens	787	180	233	197	177		
Number of rewards	413	88	107	93	125		
Ratio of "Number of burdens" to "Number of rewards"	1.9	2.1	2.2	2.1	1.4		

Table 3: Coping strategies of caregivers (n=60)

Coping style (assessed	ICD-10 dia	agnosis			Total				
with the Freiburg Questionnaire on Coping with Illness by F. Muthny, 1989)	F2 (n=13)	F31 (n=14)	F32/33 (n=15)	F60/61 (n=14)	(n=56)				
	Mean	Mean	Mean	Mean	Mean	Mode	SD	Min	Max
Depressive coping	2.5	2.4	2.4	2.2	2.4	2.2	0.80	1.0	4.6
Active, problem- oriented coping	3.6	3.5	3.3	3.4	3.5	3.4	0.90	1.0	5.0
Distraction and building self-esteem	2.6	2.8	2.5	3.0	2.7	2.8	0.71	1.0	4.4
Religiousness and searching for meaning	2.8	2.7	2.6	2.8	2.7	3.0	0.88	1.4	4.8
Trivialization and wishful thinking	2.7	2.3	1.8	2.1	2.2	1.0	0.89	1.0	4.7

Abbreviations: SD=standard deviation; Min= Minimum; Max=Maximum

Table 4: Linear regression analysis with the dependent variable "Frequency of citing rewards"

Predictor variables	Beta	P value
Increased "religiousness and searching for meaning" as strategy to cope with illness	0.425	0.001
Younger age of caregiver	0.378	0.002
More namings of burdens	0.343	0.005

Multiple linear regression: N=52; F(3.0)=10.410; p<0.000; R²=0.394