Supplementary Materials

Pre-screening

One-day prior to the scanning session, participants were sent the Mood and Anxiety Symptom Questionnaire (Watson et al., 1995). The MASQ is a 77-item selfreport questionnaire that assesses General Distress: depressive (12 items), anxious (11 items) and mixed symptomatology (15 items). The questionnaire also included an anxiety-specific (Anxious Arousal, 17 items) and a depression-specific scale (Anhedonic Depression, 22 items). Higher scores reflect greater levels of symptomatology. The reported internal consistency for each scale is excellent with coefficient alphas ranging from 0.78 to 0.92. Factorial validity for the MASQ has been established in non-clinical samples, with three factors consistently found to best represent the data (Geisser et al., 2006; Reidy et al., 1997).

We used cut-off points based on Watson et al (1995, Table 1)

	Students		Ad			
MASQ scale	Male $(n = 438)$	Female $(n = 635)$	Male (n = 142)	Female (<i>n</i> = 186)	Patients (male; $n = 453$)	
GD: Mixed						
М	34.5 _a	35.2 _a	31.3_{b}	33.0 _{a.b}	34.9 _a	
SD	9.0	9.2	10.0	10.2	12.3	
GD: Anxiety						
М	22.3 _{a.b}	22.6_{a}	20.5_{b}	$20.8_{a,b}$	21.6 _{a,b}	
SD	6.4	6.3	7.5	6.7	7.5	
Anxious Arousal						
М	27.8_{a}	27.1_{g}	24.4 _b	24.2_{b}	28.3 _a	
SD	9.4	8.2	8.0	7.8	10.4	
GD: Depression						
Μ	$24.5_{b,c}$	25.8 _b	· 22.1 _c	$25.0_{b.c}$	28.0 _a	
SD	8.7	8.8	8.1	9.4	10.0	
Anhedonic Depression						
М	55.6 _b	54.2 _b	52.0 _b	55.2_{b}	65.5 _a	
SD	13.4	13.9	12.5	15.2	14.8	

 Table 1

 Descriptive Statistics for the MASQ Scales in Five Subject Groups

Note. Within a row, means not sharing a subscript are significantly different from one another (p < .05, two-tailed). MASQ = Mood and Anxiety Symptom Questionnaire; GD = General Distress.

The MSAQ questionnaire was emailed to each participant. For each item, they indicated to what extent they had experienced each symptom (1 = not at al, 5 =

extremely) during the past week including today. Participants were asked to return their completed answer list in the same day. Scores for each participants on the present study are presented in Table 1S.

Table 1S. MASQ scores for participants in the present sample. Two participants

	,		U		
ID	GD: Mixed	GD: Anxiety	Anxious	GD:	Anhedonic
			Arousal	Depression	Depressior
1	32	22	26	24	46
2	30	18	23	22	44
3	32	24	24	23	49
4	28	16	19	20	47
5	36	25	28	27	52
6	30	20	23	21	48
7	39	27	28	22	54
8	42	28	25	27	55
9	35	22	24	25	54
10	32	21	22	21	53
11	32	22	22	25	49
13	34	23	27	27	50
14	32	22	21	22	48
15	33	25	22	20	47
16	32	24	20	22	51
17	37	27	25	45	58
18	35	29	23	25	53
19	38	28	28	37	51
21	32	23	21	25	48
22	33	24	19	26	54
23	29	19	23	20	48
12	46	32	29	48	58
20	52	42	29	48	59
Mean	33.48	23.29	23.48	25.05	50.43
(SD)*	(3.4)	(3.4)	(2.8)	(5.9)	(3.5)

(marked in red) were not admitted to	he scanning session	. GD = General Distress
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*Mean and SD are based on participants data who were invited to a scanning session

Behavioural performance

Matched and mismatched pairs were analysed separately due to different responses being made in these cases.

Accuracy

The overall accuracy in responding to stimuli in the personal (Fig.S1, A) and emotion (Fig.S1, B) tasks was above 85%.





A one-way repeated measures ANOVA was carried out on accuracy data for matched and mismatched trials in each task. The results of these analyses are summarised in Table S1.

		Personal task
Matched	A main effect	*Post Hoc
	F(2,40)=3.62,	Self vs Friend
	p=0.036,	<i>t</i> (20)= 2.58, p _h =.04, MD= 0.03, 95% CI [0.009; 0.041]
	$^{\odot^2} = 0.07$	Self vs Stranger
		<i>t</i> (20)= 1.96, p _h = .12, MD= 0.02, 95% CI [-0.003; 0.042]
		Friend vs Stranger
		<i>t</i> (20)= -0.62, p _h = .54, MD= -0.01, 95% CI [-0.028; 0.0165]
Mismatche	F(2,40)=3.22,	Self vs Friend
d	p=0.05,	<i>t</i> (20)= 2.10, p _h = .08, MD= 0.03, 95% CI [-0.005; 0.061]

Table S2. The effects of Stimulus on accuracy performance

		Self vs Stranger
		t(20) = -0.18, ph = 0.86, MD = -0.002, 95% CI [-0.026; 0.021]
		Friend vs Stranger
		t(20) = -2.28, p _h = 0.84, MD=0.03, 95% CI[-0.055; -0.00]
		Emotion task
Matched	F(2,40)=0.015,	Post Hoc
	p=0.98,	Happy vs Sad
	$^{\odot}2 = 0.00$	t(20) = 0.16, p _h =1.0, MD= 0.0
		Happy vs Neutral
		t(20) = 0.13, p _h =1.0, MD= 0.0
		Sad vs Neutral
		<i>t</i> (20)= -0.03, p _h =1.0, MD= 0.0
Mismatche	F(2,40)=0.66,	Post Hoc
d	p=0.52,	Happy vs Sad
	1 $^{2}=0.02$	t(20) = -0.11, ph = .91, MD = -0.0
		Happy vs Neutral
		t(20)= -1.05, ph =.91, MD= -0.03
		Sad vs Neutral
		<i>t</i> (20)= -0.94, p _h =.91, MD= -0.02

*Post Hoc: paired sample *t*-test. We report the adjusted p-value for multiple comparisons using Holm Method (Holm, 1979). This method, in a stepwise way, computes the significance levels depending on the p-value based rank of hypotheses. MD- difference in means (the alternative hypothesis for these tests was that a true difference in means is not equal to 0).

A paired sample t-test showed that the differences in accuracy between

matched and mismatched trials in personal and emotion tasks were non-significant

(*t*(20)=1.42, p=.69; MD=0.031; 95%CI[-0.05; 0.08] and (*t*(20)= -1.68, p= 0.11; MD= -

0.03; 95%CI [-0.06; 0.01] for personal and emotion tasks respectively).

Reaction time

Means reaction time are displayed in Fig.S2



Fig. S2. Means reaction time in the personal (A) and emotion (B) tasks for matched (correct pairings) and mismatched (incorrect pairings) associations between shapes and labels. The error bars represent +/-SEM.

A one-way repeated measures ANOVA was carried out on RT data for

matched and mismatched trials in each task. The results of these analyses are

summarised in Table S2.

		Personal task
Matched	A main effect	Post Hoc
	F(2,40)= 27.62,	Self vs Friend
	p<0.001,	<i>t</i> (20)= -2.54, p _h = .02, MD= -30.58, 95% CI [-60.37, - 2.84]
		Self vs Stranger
		<i>t</i> (20)= -7.32, p _h < .001, MD= -88.24, 95% CI [-100.1; -
		72.71]
		Friend vs Stranger
		<i>t</i> (20)= -4.78, p _h < .001, MD= -57.66, 95% CI [-82.98; -
		32.33]
Mismatche	F(2,40)=27.96,	Self vs Friend
d	p<0.001,	<i>t</i> (20)= 7.22, p _h < .001, MD= 42.63, 95% CI [31.32; 53.93]
		Self vs Stranger
		<i>t</i> (20)= 1.91, p _h = .06, MD= 11.28, 95% CI [-0.79; 23.35]
		Friend vs Stranger
		<i>t</i> (20)= -5.31, p _h < .001, MD= -31.35, 95% CI [-44.84; -
		17.85]
		Emotion task

Table S3. The effects of Stimulus on Response Times

Matched	F(2,40)=29.70, p<0.001, ∞ ² = 0.10	Happy vs Sad t(20)= -0.32, ph = .75, MD= -3.24, 95% CI [-11.88; 10.36] Happy vs Neutral t(20)= -6.83, ph < .001, MD= -69.38, 95% CI [-84.93; - 47.35] Sad vs Neutral t(20)= -6.51, ph < .001, MD= -66.14, 95% CI [-82.97; - 47.78]
Mismatche d	F(2,40)=13.64, p<0.001, $^{\circ}^2 = 0.05$	Happy vs Sad t(20)= -4.73, ph < .001, MD= -25.22, 95% CI [-35.36; - 15.06] Happy vs Neutral t(20)= -4.29, ph < .001, MD= -22.85, 95% CI [-36.92; -8.77] Sad vs Neutral t(20)= 0.44, ph = .07, MD= 2.37, 95% CI [-6.00; 10.73]

A paired sample t-test showed that participants were faster in responding to matched trials compared to mismatched in the personal task (t(20)= -9.16, p <.001; MD= -188.71; 95%CI [-231.67; -145.75]. The difference between matched and mismatched trials in the emotion task did not reach significance (t(20)= -2.01, p= 0.06; MD= -54.84; 95%CI [-111.81; 2.11].

Reaction Time Advantages

To quantify the effects of personal relevance and emotions, we calculated the advantage in RT for self and friend compared to stranger ([RTstranger – RTself], [RTstranger – RTfriend]) and happy and sad emotional expressions compared to neutral ([RTneutral – RThappy], [RTneutral – RTsad]) (Table S3).

Table S4. Comparison RT advantages

Contrast	Results (Student's t-test)
[RTstranger – RTself] - [RTstranger – RTfriend]	<i>t</i> (20)=2.70, p=.01, MD = 39.39, SE
	difference = 14.54, 95%CI [8.95,

	69.62], Cohen's d =0.59, 95%CI
	[0.12, 1.05]
[RTneutral – RThappy] - [RTneutral – RTsad]	<i>t</i> (20)=0.29, p=.78, MD = 3.24, SE
	difference = 11.34, 95%CI [-20.43,
	26.90], Cohen's d =0.06, 95%CI [-
	0.37, 0.49]

Multiple regression

A multiple regression analysis was carried out to test whether the magnitude of the prioritization effects for positive and negative emotions could predict the magnitude of self-prioritization. Using the enter method it was found that prioritization of positive and negative emotions explained a significant amount of the variance of self-prioritization (F(2,20) = 9.04, p=0.002, R² = 0.50). The model summary and test statistics are presented below (all analyses were performed in JASP, 2020).

Model Summary	
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Mode	R	R ²	Adjusted				
1	IX.	IX.	R ² RI	MSE			
H_0	0.000	0.000	0.000	31.437			
H_1	0.708	0.501	0.446	23.403			
ANO	VA						
Mod	el		Sum of Squares	df	Mean Square	F	р
H_1	Regre	ssion	9906.980	2	4953.490	9.044	0.002
	Resid	ual	9858.829	18	547.713		
	Total		19765.810	20			

Note. The intercept model is omitted, as no meaningful information can be shown.

					95% ł	oca* CI
Model		Unstandardized	Bias	Standard Error	Lower	Upper
H ₀	(Intercept)	88.429	0.187	6.660	74.667	100.963
H_1	(Intercept)	46.141	-0.759	12.408	27.080	77.519
	Happy-bias	0.421	0.033	0.148	0.170	0.699
	Sad-bias	0.179	-0.015	0.141	-0.112	0.430

* Bias corrected accelerated

Note. Bootstrapping based on 5000 replicates.

Note. Coefficient estimate is based on the median of the bootstrap distribution.

In order to check for outliers, we assessed residual statistics (see table below).

An analysis of standard residuals was carried out, which showed that the data

contained no outliers (Std. Residual Min = -1.93, Std. Residual Max = 1.55).

Residuals Statistics

	Minimu		Moon	۶D	N	
	m	Maximum	Mean	SD	1 N	
Predicted Value	50.589	138.296	88.238	22.256	21	
Residual	-41.888	32.983	-7.615e - 16	22.202	21	
Std. Predicted Value	-1.692	2.249	3.192e -16	1.000	21	
Std. Residual	-1.930	1.553	-0.012	1.039	21	

Tests to see if the data met the assumption of collinearity indicated that

multicollinearity was not a concern (Happy-bias, Tolerance = .92,VIF = 1.09; Sad-

bias, Tolerance = .92, VIF= 1.09) (see table below).

							95% CI		Collinearity Statistics	
Model		Unstandardized	Standard Error	Standardized	t	р	Lower	Upper	Tolerance	VIF
H _o	(Intercept)	88.238	6.860		12.862	< .001	73.928	102.548		
H1	(Intercept)	47.502	11.371		4.178	< .001	23.613	71.391		
	Happy-bias	0.410	0.119	0.598	3.434	0.003	0.159	0.661	0.915	1.092
	Sad-bias	0.186	0.132	0.244	1.402	0.178	-0.093	0.464	0.915	1.092

Model Summary									
					Durbin-Watson				
Mode 1	R	R ²	Adjusted R ²	RMSE	Autocorrelation	Statistic	р		
H ₀	0.000	0.000	0.000	31.437	-0.435	2.719	0.083		
H_1	0.708	0.501	0.446	23.403	-0.413	2.628	0.161		

The data met the assumption of independent errors (Durbin-Watson value = 2.63) (see table below).

The balanced distribution of the residuals around the baseline (Fig. S3) suggests that the assumption of homoscedasticity has not been violated. The Q-Q plot (Fig. S3) shows that the standardized residuals fit along the diagonal suggesting that both assumptions or normality and linearity have also not been violated.



Fig. S3. Residuals vs. Predicted (on the right) and Q-Q Plot Standardized Residuals *Univariate fMRI*

The contrast [happy > neutral] (height threshold p<0.001, extended threshold = 30 contagious voxels, cluster FDR corrected <0.05) showed activation in the left precentral lobule (Table S4, Fig.S4). No voxels survived the threshold for the reverse contrast [neutral > happy].

Table S5. Clusters for contrast happy > neutral above the threshold.

Label	x	y	Z	k	Z
L-	-26	-22	64	289	3.67
Precentral					



Fig.S4. Activation results for the whole-brain univariate analyses for contrasts [happy>neutral]. The mask of clusters with significant univariate effects (a cluster corrected FDR-threshold of p<0.05, voxel-threshold p<0.001 uncorrected, extended threshold of 30 contiguous voxels) was created and overlaid on a MNI152 standard template using MRIcroGL (radiological convention)

The contrast [sad > neutral] (height threshold p<0.001, extended threshold = 30 contagious voxels, cluster FDR corrected <0.05) showed activation in the left precentral lobule (Table S5, Fig.S5). No voxels survived the threshold for the reverse contrast [neutral > sad].

Table S6. Clusters for contrast sad > neutral above the threshold.

Label	х	у	Z	k	Ζ
Parietal_Inf_l	-40	-38	40	543	4.65

Frontal_Inf_Tri_	-54	14	28	421	4.49
Supp Motor	-2	2	62	141	4.31
Area_L Frontal Sup_2_L	-24	0	64	306	4.09
Frontal_Inf_Tri_	46	0 16	22	207	3.93
R					



Fig.S5. Activation results for the whole-brain univariate analyses for contrasts [sad>neutral]. The mask of clusters with significant univariate effects (a cluster corrected FDR-threshold of p<0.05, voxel-threshold p<0.001 uncorrected, extended threshold of 30 contiguous voxels) was created and overlaid on a MNI152 standard template using MRIcroGL (radiological convention).

In the contrast [self > friend] (height threshold p<0.001, extended threshold = 30 contagious voxels, cluster FDR corrected <0.05) no voxels survived the threshold. Lowering the hight threshold to p<0.005 indicated activation in the MPFC (Rectus) and Precuneus. (Table S6). However, the activations were not significant at the FDR corrections.

Ζ k Label х у Z FDR_corr -2 Rectus -52 18 116 3.53 0.86 Precuneus 4 52 -18 82 3.22 0.86

Table S7. Clusters for contrast self > friend (height threshold p<0.005, extended

The reverse contrast [friend > self] did not reveal voxels above the threshold (height threshold p<0.001, extended threshold = 30 contagious voxels, cluster FDR corrected <0.05). Lowering the height threshold to p<0.005 indicate clusters in the left dorsolateral prefrontal cortex (DLPFC) (x/y/z = -28/36/10, k=104, Z =4.03) and small clusters in the left parietal cortex. However, the activations were not significant at the FDR corrections.

Defining a ROI in the dmPFC

threshold =30)

The dmPFC ROI was defined as a 7 mm sphere centred at x/y/z = 6/44/18 [41, 53] and containing 207 voxels (Fig S6).



Figure S6. The ROIs defined in the present study. The vmPFC (in green) is the ROI defined based on the contrast [self > stranger] (k= 213, centred at x/y/z = -6/52/-4).

The dmPFC (in red) is the ROI defined based on previous studies (Mitchel at al.,

2006) and Yankouskaya et al., 2018) (7 mm sphere centred at x/y/z = 6/44/18,

k=207).

References for Supplementary Materials

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