Spalax ehrenbergi, under different light spectra PR One-way ANOVA Mesor Amplitude Acrophase $(pg ml^{-1} g^{-1})$ $(pg ml^{-1} g^{-1})$ $F_{25};P$ $(F_{6.36}; P)$ (h) (hh:mm) (%)M. socialis

Table S4. Factorial and cosinor analyses of urinary metabolites of cortisol (UMCort) in two rodent species, *Microtus socialis* and

Short wavelength	0.1; 0.44	12.9	33.4 (27.3-39.5) ^a	6.31 (1.95-14.6)	08:02	49	9.37; 0.04
					(05:55-10:08)		
Long wavelength	5.34; 0.0001	24	59.4 (51.4-67.4) ^b	20.4 (9.37-31.3)	10:04	43	11.68; 0.01
İ					(07:48-12:24)		

					(07:4
6. ehrenbergi					
Short wavelength	0.18: 0.98	12	49.0	1.56	

P>0.05.

					,		
S. ehrenbergi							
Short wavelength	0.18; 0.98	12	49.0	1.56	08:58	37	0.17; 0.84
Long wavelength	6.39; 0.0001	24	11.5 (9.24-13.8)	4.89 (1.46-8.32)	02:39	43	10.42; 0.02
					(00:06-05:24)		

τ, period length of the cosine curve approximated by spectral analysis; PR, percentage of the rhythm (represents the proportion of the total variance of the data accounted by the cosine approximation of a trial period).

The zero amplitude hypothesis was rejected at P<0.05. Different letters represent significant differences between treatments for each species

(P < 0.05).

Values in brackets for mesor, amplitude and acrophase are 95% confidence intervals (CI) of the group mean. CI values are not listed when