External and circadian inputs modulate synaptic protein expression in the visual system of Drosophila melanogaster

Wojciech Krzeptowski†, Jolanta Górsko-Andrzejak†, Ewelina Kijak†, Alicja Górlich, Elżbieta Guzik, Gareth Moore and Elżbieta M. Pyza *

Department of Cell Biology and Imaging, Institute of Zoology, Jagiellonian University, Kraków, Poland

Edited by: Matti Weckstrom, University of Oulu, Finland
Reviewed by: Mikko Juusola, University of Sheffield, UK
Paivi Helena Torkkeli, Dalhousie University, Canada
*Correspondence: Elżbieta M. Pyza, Department of Cell Biology and Imaging, Institute of Zoology, Jagiellonian University, Gronostajowa 9, 30-387 Kraków, Poland e-mail: elzbieta.pyza@uj.edu.pl
† These authors have contributed equally to this work.

INTRODUCTION

In the first optic neuropil (lamina) of the fly’s visual system the majority of synaptic contacts is formed between the terminals of the compound eye photoreceptors and their postsynaptic partners. They are called tetrad synapses and use histamine (Hardie, 1987; Melzig et al., 1998; Gavin et al., 2007) to transmit photic and visual information to the brain. Using the specific antibodies against synaptic proteins: Bruchpilot (BRP), Synapsin (SYN), and Disc Large (DLG), the synapses in the distal lamina were specifically labeled. Then their abundance was measured as immunofluorescence intensity in flies held in light/dark (LD 12:12), constant darkness (DD), and after locomotor and light stimulation. Moreover, the levels of proteins (SYN and DLG), and mRNAs of the brp, syn, and dlg genes, were measured in the fly’s head and brain, respectively. In the head we did not detect SYN and DLG oscillations. We found, however, that in the lamina, DLG oscillates in LD 12:12 and DD but SYN cycles only in DD. The abundance of all synaptic proteins was also changed in the lamina after locomotor and light stimulation. One hour locomotor stimulations at different time points in LD 12:12 affected the pattern of the daily rhythm of synaptic proteins. In turn, light stimulations in DD increased the level of all proteins studied. In the case of SYN, however, this effect was observed only after a short light pulse (15 min). In contrast to proteins studied in the lamina, the mRNA of brp, syn, and dlg genes in the brain was not cycling in LD 12:12 and DD, except the mRNA of dlg in LD 12:12. Our earlier results and obtained in the present study showed that the abundance of BRP, SYN and DLG in the distal lamina, at the tetrad synapses, is regulated by light and a circadian clock while locomotor stimulation affects their daily pattern of expression. The observed changes in the level of synaptic markers reflect the circadian plasticity of tetrad synapses regulated by the circadian clock and external inputs, both specific and unspecific for the visual system.

Keywords: synaptic plasticity, circadian rhythms, light stimulation, locomotor activity, the first optic neuropil

In the visual system of Drosophila melanogaster the retina photoreceptors form tetrad synapses with the first order interneurons, amacrine cells and glial cells in the first optic neuropil (lamina), in order to transmit photic and visual information to the brain. Using the specific antibodies against synaptic proteins: Bruchpilot (BRP), Synapsin (SYN), and Disc Large (DLG), the synapses in the distal lamina were specifically labeled. Then their abundance was measured as immunofluorescence intensity in flies held in light/dark (LD 12:12), constant darkness (DD), and after locomotor and light stimulation. Moreover, the levels of proteins (SYN and DLG), and mRNAs of the brp, syn, and dlg genes, were measured in the fly’s head and brain, respectively. In the head we did not detect SYN and DLG oscillations. We found, however, that in the lamina, DLG oscillates in LD 12:12 and DD but SYN cycles only in DD. The abundance of all synaptic proteins was also changed in the lamina after locomotor and light stimulation. One hour locomotor stimulations at different time points in LD 12:12 affected the pattern of the daily rhythm of synaptic proteins. In turn, light stimulations in DD increased the level of all proteins studied. In the case of SYN, however, this effect was observed only after a short light pulse (15 min). In contrast to proteins studied in the lamina, the mRNA of brp, syn, and dlg genes in the brain was not cycling in LD 12:12 and DD, except the mRNA of dlg in LD 12:12. Our earlier results and obtained in the present study showed that the abundance of BRP, SYN and DLG in the distal lamina, at the tetrad synapses, is regulated by light and a circadian clock while locomotor stimulation affects their daily pattern of expression. The observed changes in the level of synaptic markers reflect the circadian plasticity of tetrad synapses regulated by the circadian clock and external inputs, both specific and unspecific for the visual system.
SYN has been shown to maintain vesicle release by recruiting synaptic vesicles from the reserve pool (Li et al., 1995; Pieribone et al., 1995; Sun et al., 2006; Akbergenova and Bykhovskaia, 2007).

The synaptic protein reported to be important for the synaptic assembly and plasticity is Discs large (DLG), a member of the MAGUK (Membrane-Associated Guanylate Kinase) family of scaffolding proteins. DLG has four mammalian members; SAP97/hDlg, PSD-93/Chapsyn-110, PSD-95/SAP90, and SAP102/NE-Dlg, which play important roles in the development and regulation of the nervous system and epithelia (Wang et al., 2011). In D. melanogaster DLG is involved in the development of epithelia (Woods et al., 1996), asymmetric cell division (Ohshiro et al., 2000), the development and plasticity of the larval NMJ (Lahey et al., 1994; Budnik et al., 1996; Guan et al., 1996; Tejedor et al., 1997; Thomas et al., 1997), as well as in other synaptic functions that are important for behavior of adult flies, including their circadian rhythms (Mendoza-Topaz et al., 2008). DLG is thought to play a key role in clustering GluRIIB receptors to external factors, such as direct light exposure and locomotor activity.

MATERIALS AND METHODS

ANIMALS

All experiments were conducted on a wild type Canton S strain of Drosophila melanogaster. The stock was maintained on a standard yeast-cornmeal-agar medium at 25 ± 1°C, under a day/night cycle (12 h of light and 12 h of darkness; LD 12:12). Following their eclosion, the experimental flies were kept in LD 12:12 for 1 week and the 7 days old males were decapitated at the specific time points of day and night. These time points were chosen to reflect the morning and evening locomotor activity peaks, which survive only to the pupal stage, the number of NMJ T-bars is increased and SSR is reduced (Lahey et al., 1994).

In the present study we examined circadian changes in the level of three crucial synaptic proteins; BRP, SYN, and DLG, and their gene expression in order to learn about their possible involvement in the circadian synaptic plasticity of tetrad synapses in the lamina and in the brain. In addition we analyzed their responses to external factors, such as direct light exposure and locomotor stimulation.

IMMUNOHISTOCHEMISTRY

The heads of the flies collected at the experimental time points (ZTs or CTs) were processed in parallel, under the same conditions. Flies were immobilized with CO2 and decapitated in a drop of fixative—4% formaldehyde (PA) in a 0.1 M phosphate buffer (PB). Following tissue fixation and overnight infiltration in a 25% solution of sucrose, the cryostat sections of heads were cut and incubated with one of the three mouse monoclonal antibodies: nc82, 3C11, or 4F3 (Developmental Studies Hybridoma Bank (DSHB)) detecting the synaptic proteins BRP, SYN, and Disc Large (DLG), respectively (Kittel et al., 2000; Wagh et al., 2006; Hofbauer et al., 2009; Hamanaka and Meinertzhagen, 2010). The 4F3 antibody was produced against the PSD-95/DLG/zona occludens-1 2 (PDZ-2) (Farnas et al., 2001). After several washes in a 0.1 M sodium phosphate buffer (PBS) containing 0.2% Triton-X (Sigma), the tissue was incubated with the goat anti-mouse secondary antibody conjugated with Cy3 (Jackson ImmunoResearch Laboratories Inc.). Sections were examined using a Zeiss LSM 510 Meta confocal microscope following washing and mounting in a Vectashield medium (Vector).

QUANTIFICATION OF IMMUNOLABELING

In each experiment, the images of the lamina of flies decapitated at different time points were collected at non-saturated settings, using identical image acquisition parameters. The changes in the level of fluorescence of BRP, SYN, and DLG-specific immunolabeling (brightness), corresponding to the changes in the amount of the given protein (BRP, SYN, or DLG), were quantified using Image] software (NIH). In each experiment the intensity of fluorescence was measured in the cartridges of the distal part of the cross-sectioned lamina (the second and the third row of cartridges from the lamina cortex) (Figure 1B), as represented by the Mean Gray Value, defined by the sum of the gray values of all pixels in the selected area, divided by the number of pixels within the selection. In the Image software, the range of gray values in 8-bit images is divided into 256 bins. For each time point (ZTs or CTs) we collected data from 7 to 12 individuals. The background signal was subtracted.
FIGURE 1  |  (A) The synaptic neuropils of the lamina (L), medulla (M) and the central brain of *Drosophila melanogaster* visualized by anti-BRP immunostaining using Mab nc82. There is a strong fluorescent signal in the mushroom body pedunculi in the central brain as well (arrow heads). R, retina. Scale bar: 50 μm. (B) The array of cartridges (arrows) in the cross sectioned neuropil of lamina as revealed by immunostaining using Mab 4F3 against DLG. Scale bar: 20 μm. Insert: the magnification of a single cartridge. Scale bar: 1 μm. (C–E) Cross sections of cartridges of GMR-Gal4 x UAS-S65T-GFP transgenic flies showing targeted expression of a green fluorescent protein (GFP) in photoreceptors (green) and immunolabeling (magenta) with Mab 3C11 against BRP (C), 3C11 against SYN (D) and 4F3 against DLG (E). R—the terminals of six photoreceptors, L1, L2—interneurons receiving photic and visual information from epithelial glial cells that surround lamina cartridges. Scale bar: 1 μm.

DETERMINATION OF mRNA

For brain dissection we used the method described by Fujita et al. (1987). To summarize, experimental flies, some 15–20 individuals per time point, were frozen in liquid nitrogen. Subsequently, flies were transferred into glass tubes filled with 2 ml of cold acetone (−80°C) for dehydration, and stored at −80°C for 1 week. During that time, the acetone was replaced twice. Then, flies brains were dissected and total RNA was isolated using a NucleoSpin RNA XS kit (Macherey-Nagel) according to the manufacturer’s protocol. To avoid contamination by genomic DNA, samples were treated with rDNase. RNA was eluted in 11 μl of RNase-free water. Reverse transcription was performed according to the manufacturer’s protocol using a SuperScript III First-Strand Synthesis System for RT-PCR kit (Invitrogen), for *dlg* analysis or a High Capacity cDNA Reverse Transcription Kit (Applied Biosystems) for analysis of *brp* and *syn*. cDNA was prepared in 20 μl volumes using an oligo(dT)20 primer or random primers. Gene quantification was performed using the Step One Plus Real-Time PCR System (Applied Biosystems) and TaqMan Gene Expression Assays (Applied Biosystems). *dlg* (Assay ID: Dm01799278_g1), *brp* (Assay ID: Dm0194336_g1), and *syn* (Assay ID: Dm02148572_m1) genes were examined. Quantitative PCR was performed with 1 μl of cDNA as a template in a final reaction volume of 20 μl. Thermal cycling conditions were as follows: 2 min at 50°C, 10 min at 95°C followed by 40 repeats of 15 s at 95°C, and 1 min at 60°C. Data collection was performed during each annealing phase. Raw C_T values were collected and analysis was performed according to the 2^−ΔΔC_T method, using *rpl32* (Assay ID: Dm02151827_g1) as a reference gene. Gene quantifications were completed using the software supplied by, and according to the instructions of the manufacturer. All experiments were performed independently at least three times, and each experiment comprised three parallel samples. In each run a negative control (distilled water instead of cDNA) was included.

WESTERN BLOT ANALYSIS

Flies collected at different time points (ZTs and CTs) were immediately frozen in liquid nitrogen. Their heads (20–30 heads per time point) were cut off, frozen and homogenized by sonication in a RIPA buffer (1.5 μl/1 head) containing a complete protease inhibitor cocktail (Boehringer). After being gently shaken for 1 h on ice, the obtained homogenates were stored at −20°C until centrifugation. Samples were centrifuged at 13,200 rpm for 1 h at 4°C and supplemented with a 2 × Laemmli buffer (1:1) before denaturation at 100°C for 5 min. The protein level was measured with a Quant-iT Protein Assay Kit using a Qubit fluorometer (Invitrogen). Protein extracts were subjected to electrophoresis using the NuPAGE SDS-PAGE Gel System. The 4–12% Bis-Tris gel was loaded with 20 or 10 μg (SYN and DLG, respectively) of the total protein per lane. The proteins were blotted by electrotransfer onto a PVDF membrane that was blocked with 5% nonfat dry milk in PBS with 0.1% Tween-20. Following blotting, the membrane was immunoprobed either with Mab 3C11 (1:10,000) or Mab 4F3 (1:20,000). The effectiveness of protein loading (loading control) was defined by probing the blots with the mouse monoclonal anti-α tubulin antibody (AA4.3), diluted 1:20,000 (DSHB). The ECL detection system (Perkin Elmer Inc.) was applied for immunodetection, whereas the AlphaEaseFC Stand Alone image analysis program (Alpha Innotech, Cell Bioscience,) was used for the densitometric analysis of the obtained Western blots. The collected data derived from three independent experiments providing three distinct protein preparations.

STATISTICS

All statistical tests were performed using the Statistica v.10 (StatSoft, Inc.). The significance of differences observed between two independent groups (e.g., day and night) were statistically analyzed using the *T*-test or Mann–Whitney Test, depending on the result of the Shapiro–Wilk *W*-Test. Statistical significance
across groups, ZT or CT time points, was analyzed using One-Way ANOVA followed by a Tukey post-hoc Test or the nonparametric counterpart of ANOVA—Kruskal–Wallis Test (one-way test), followed by the Multiple Comparison Test. To assess differences between LD 12:12 and DD we performed the Two-Way ANOVA regarding the light conditions and the time points (ZTs and CTs) as two key factors.

Immunolabeling quantification data are shown as a percentage of the highest value (100%) that was obtained in a given experiment (means ± SD).

For visualization of qPCR data, gene expression was normalized on an arbitrary scale, where the ZT1 time point was set to 1.00 (means ± SE).

RESULTS
The patterns of immunolabeling of BRP, SYN, and DLG in the whole brain of *D. melanogaster* was similar, if not identical. In each case, the immunofluorescence was confined to neuropils of the visual system and the central brain, as displayed by an exemplary image of BRP–specific immunolabeling in Figure 1A.

Analysis of these labelings in the cross sectioned lamina showed that each of the studied proteins is concentrated within the synaptic units or cartridges, revealing the array of cartridges that constitute this neuropil (Figure 1B). However, the pattern of expression within the cartridge (Figures 1C–E) is different for each of these proteins, forming well defined spots in case of BRP (which localizes to T-bar ribbons) and being more dispersed in the case of SYN or DLG (Figures 1C–E).

BRUCHPILOT
The *brp* mRNA did not cycle in the brain of flies raised either in LD 12:12 (Figure 2A) [Kruskal–Wallis Test: *H*(3, *N* = 12) = 2.63, *p* = 0.453] or in DD (Figure 2B) [Kruskal–Wallis Test: *H*(3, *N* = 12) = 3.04, *p* = 0.38].

After 1 h locomotor stimulation, the level of BRP fluctuated (ANOVA, *p* < 0.0001), being higher at ZT1 and ZT13 in comparison to ZT4 and ZT16 (Figure 3A) (Tukey’s HSD Test, differences between ZT1 and ZT4 at *p* < 0.05; between ZT1 and ZT16 at *p* < 0.01; between ZT4 and ZT13 at *p* < 0.001, and between ZT13 and ZT16 at *p* < 0.001). However, the amplitude of the observed changes was larger than in non-stimulated flies (Górski-Andrzejak et al., 2013).

Light stimulation profoundly increased the level of BRP in the lamina of *D. melanogaster* [Kruskal–Wallis Test: *H*(2, *N* = 57) = 11.08, *p* = 0.004] (Figure 3B). When compared to the non-stimulated controls, the intensity of BRP-specific immunofluorescence in the lamina of flies exposed to the light pulse at CT1 for either 15, or 60 min, increased by 58 and 63%, respectively (Multiple Comparison Test, *p* = 0.02 and *p* = 0.005, respectively).

SYNAPSIN
The level of *syn* mRNA did not cycle in the brain of flies held either in LD 12:12 [Kruskal–Wallis Test: *H*(3, *N* = 12) = 1.37, *p* = 0.7] or in DD [Kruskal–Wallis Test: *H*(3, *N* = 12) = 3.82, *p* = 0.5] (Figures 4A,B). Also the total amount of SYN in whole head homogenates studied using the Western blotting method, did not change significantly during the 24 h period, either in LD 12:12 (ANOVA, *F* = 0.18, *p* = 0.9) or in DD [Kruskal–Wallis Test *H*(3, *N* = 12) = 5.1, *p* = 0.16] (data not shown).

Similar results were obtained for SYN protein in the lamina of flies held in LD 12:12. The analysis of the fluorescence intensity of immunolabeled SYN showed that the abundance of this protein did not change significantly in the course of day and night in LD 12:12 (ANOVA, *F* = 0.54, *p* = 0.7) (Figure 5A). The average level of SYN-specific fluorescence during the day and the night was the same (*t*-test, *t* = 0.02, *df* = 26, *p* < 0.98). In DD, on the other hand, there were significant changes in the level of SYN-immunospecific fluorescence in the lamina over the 24 h period [Kruskal–Wallis Test (3, *N* = 31) = 14.7, *p* = 0.002] (Figure 5B). The fluorescence intensity was the highest in the middle of the subjective day, at CT4, and the lowest in the middle of the subjective night, at CT16 (Figure 5B). The difference between these two time points was statistically significant (Multiple Comparison Test, *p* = 0.002). The average level of fluorescence during the subjective day was the same as during the day in LD 12:12, but was 31% lower during the subjective night, than during the night in LD 12:12. The difference between the subjective day and the subjective night was statistically significant (Mann–Whitney Test, *U* = 32, *p* = 0.0005) and so was the difference between the daily patterns of SYN-specific immunofluorescence in DD and LD 12:12 (MANOVA, *F* = 3.2, *p* = 0.03). Moreover, the average level of fluorescence in DD was 22% lower than in LD 12:12 (Mann–Whitney Test, *U* = 254.5, *p* = 0.007).
After 1 h locomotor activity stimulation at different times of the day, the intensity of SYN-specific immunofluorescence in the lamina displayed significant changes in the course of the day (Kruskal–Wallis Test (4, N = 53) = 18, p = 0.001) (Figure 5C). It was the lowest in the lamina of flies stimulated at the beginning of the day (ZT1—47%). At that time the fluorescence intensity was considerably lower (Multiple Comparison Test, p = 0.02) than at the beginning of the night (ZT13), when it reached the highest value measured for this set of data (65%). Also the average level of fluorescence during the day (ZT1 and ZT4) was significantly lower (11%) than during the night (ZT13 and ZT16) (Mann–Whitney Test, U = 88, p = 0.001).

After light stimulation at CT1 the level of SYN in the lamina increased but only after the short 15 min light pulse (Kruskal–Wallis Test (2, N = 35) = 11.05, p = 0.004) (Figure 5D). The short pulse increased SYN-specific immunofluorescence intensity by 21% when compared with the control. However, after a longer, 60 min light pulse, the level of SYN was similar to the control. The differences between the 15 min light pulse and the control (67 vs. 47%), as well as between the 60 min and 15 min pulses (67 vs. 45%), were significant (Multiple Comparison Test, p = 0.02 and p = 0.006, respectively).

**DISC LARGE**

Real-time PCR quantification revealed significant changes in the expression of dlg in the brain of *D. melanogaster* collected at different time points of the LD 12:12 cycle (Kruskal–Wallis Test: $H(3, N = 23) = 17.7, p = 0.0005$) (Figure 6A). The highest level of *dlg* mRNA was detected in the middle of the day (ZT6) and at the end of the day (ZT12), whereas the lowest was typical for the end of the night (ZT18). The differences between ZT1 and ZT6 (63%) or ZT1 and ZT18 (103%) were statistically significant (Multiple Comparison Test, $p = 0.01$ and $p = 0.005$ respectively), as was the difference between ZT12 and ZT18 (86%, Multiple Comparison Test, $p = 0.05$). The expression of *dlg* in the brain of flies raised in constant conditions of darkness did not change during the subjective day and night (Kruskal–Wallis Test: $H(3, N = 28) = 4.83, p = 0.18$) (Figure 6B). In turn, the total amount of DLG in the whole head homogenates obtained from flies kept in LD 12:12 or DD, was the same during the 24 h cycle (Kruskal–Wallis Test (3, N = 12) = 1.17, p = 0.76; Kruskal–Wallis Test (3, N = 8) = 1.13, p = 0.72, respectively) (data not shown).

The abundance of DLG in the distal lamina of flies held in LD 12:12, showed significant changes during the light/dark cycle (Kruskal–Wallis Test: $H(3, N = 174) = 34.2, p < 0.0001$) (Figure 7A). The pattern of changes in the DLG level in the lamina was bimodal with two peaks, at ZT1 and ZT12, as in the case of BRP (Górski-Andrzejak et al., 2013). The average intensity of DLG-specific fluorescence at these two time points of the cycle was the same (Multiple Comparison Test: $p < 0.1$), whereas it was lower in the middle of the day (ZT6) and in
FIGURE 5 | Synapsin-specific immunofluorescence (brightness) measured from confocal images of the fly’s lamina. The level of this protein is the same during the day and night of LD 12:12 (A), oscillates in constant darkness (B), changes after 1 h of locomotor stimulation (C) and increases after 15 min of light stimulation (D) (*p ≤ 0.05, **p ≤ 0.01).

FIGURE 6 | The relative level of dlg mRNA in the brain of flies held in LD 12:12 or DD. In LD 12:12 the mRNA of dlg changes during 24 h cycle with a peak in the middle of the day (ZT6) and a trough in the middle of the night (A). No rhythm was observed in DD (B); *p < 0.05, **p < 0.01.

The above described pattern of DLG rhythm in the lamina of flies kept in LD 12:12 was changed by locomotor stimulation (Figure 7C). After a 1 h long locomotor stimulation of flies, administered prior to their decapitation, the intensity of DLG-specific fluorescence was still fluctuating during the middle of the night (ZT18). The differences between ZT1 and ZT6 (16%) or ZT1 and ZT18 (15%) were statistically significant (Multiple Comparison Test: p < 0.0001 and p < 0.0001, respectively), as well as the differences detected between ZT6 and ZT12 (13%, Multiple Comparison Test: p = 0.002) or ZT12 and ZT18 (12%, Multiple Comparison Test: p = 0.006). Similar oscillations in the level of DLG-specific immunofluorescence were also observed in the distal lamina of flies kept in constant darkness [Kruskal-Wallis Test: H(3, N = 198) = 27.3, p < 0.0001] (Figure 7B). Interestingly, the circadian pattern of changes was the same as the one observed in LD 12:12, since the fluorescence intensity was significantly higher at CT1 and CT12 than at CT4 and CT18. The differences between CT1 and CT6 (12%) or CT18 (8%), although slightly smaller than between ZT1 and ZT6 or ZT16, were statistically significant (Multiple Comparison Test: p = 0.0008 and p = 0.04, respectively). Also the differences between the second peak of DLG abundance at CT12 and CT6 (15%) or CT18 (12%) were significant (Multiple Comparison Test: p < 0.0001 and p = 0.004, respectively).
FIGURE 7 | DLG-specific immunofluorescence (brightness) measured from confocal images of the fly’s lamina. In both LD 12:12 (A) and DD (B) the level of DLG oscillates and its pattern is similar in both conditions with two peaks at ZT1 or CT1 and ZT12 or CT12. One hour of locomotor stimulation changes the rhythm of DLG (C). Light stimulation increases DLG abundance in the lamina of wild type flies (D) (*p \leq 0.05, **p \leq 0.01, ***p \leq 0.001).

24 h cycle [Kruskal–Wallis Test: \(H(3, N=149) = 24.8, p < 0.0001\)]. However, the daily pattern was altered with a peak at the end of the night (ZT18) when the fluorescence level was 20, 22, and 17% more intensive than at ZT1, ZT6, and ZT12, respectively (Multiple Comparison Tests: \(p = 0.002, p = 0.0001, p = 0.02\), respectively).

To examine the direct exposure of light on the DLG level in the lamina, we examined DLG-specific immunofluorescence in flies after 15 or 60 min light pulses in DD (Figure 7D). The results revealed the significant influence of such stimulation on the DLG level in the lamina [Kruskal–Wallis Test: \(H(2, N=254) = 51.4, p < 0.0001\)]. A significant increase in the intensity of fluorescence (in comparison to the non-stimulated control group), was observed not only after the 60 min light pulse (21%), but also after the 15 min pulse (11%) (Multiple Comparison Test: \(p < 0.0001\) for both comparisons). A significant difference (10%) in the level of fluorescence was also revealed between the two stimulated groups (Multiple Comparison Test: \(p = 0.004\)).

DISCUSSION

All three synaptic markers examined in our previous (Gór ska-Andrzejak et al., 2013) and present study showed both cyclic and stimuli-depended changes in their abundance in the lamina of D. melanogaster. This is the first report that the abundance of the synaptic structural proteins (BRP and DLG) and the proteins involved in synaptic vesicle release (SYN) is controlled by the circadian clock not only in neurons of the circadian clocks (Shapiro-Reznik et al., 2012) but also in other brain regions, such as the first optic neuropil (lamina). Moreover, the observed changes in the abundance of BRP, DLG, and SYN were specific for each protein. In contrast, no oscillations in these proteins, except BRP (Gór ska-Andrzejak et al., 2013), were detected in the whole head. Also their gene expression, except dlg, was not cycling when measured as the mRNA levels in the brain of flies sacrificed at different time points in the 24 h cycle. These results indicate that the expression of the synaptic proteins is specifically regulated in different regions of the brain.

Of the three synaptic proteins, BRP has already been studied in the brain and lamina of D. melanogaster and its abundance displayed both daily and circadian rhythms (Gór ska-Andrzejak et al., 2013). In the lamina the level of BRP increases twice during the 24 h cycle of LD 12:12; at the beginning of the day (ZT1) and at the onset of the night (ZT13). The same changes were observed in whole head homogenates analyzed using Western blotting. In DD the level of BRP increases only once during the 24 h cycle—at the beginning of the subjective night (CT13) (Gór ska-Andrzejak et al., 2013). The circadian rhythm of BRP is correlated with the circadian rhythm in the locomotor activity of D. melanogaster and the circadian plasticity of monopolar cells in the lamina (Pyza and Meinertzhagen, 1999; Weber et al., 2009).

In the present study we also found that the daily rhythm of BRP is not altered by locomotor stimulation. The stimulation, however, increases the level of BRP, especially at ZT1 and ZT13 when BRP normally peaks during the 24 h light/dark cycle. The level of BRP is also affected by light, since it increases after light pulses, and the circadian rhythm of BRP is entrained by light (Gór ska-Andrzejak et al., 2013). Moreover, among two peaks
observed in BRP abundance during the day and night of LD 12:12, the morning peak is controlled by light but the evening one by the circadian clock, although both vanish in mutants with the null mutation of the clock gene period (per) (Górskaj et al., 2013).

Our present results, as well as our previous findings (Górskaj et al., 2013) indicate that BRP abundance at tetrad synapses is controlled by the circadian clock and modulated by light and other stimuli. Therefore, BRP organizes the tetrad presynaptic elements according to the activity of a fly and flow of visual and photic information from the eye photoreceptors.

The BRP level is also affected by sleep deprivation (Gilestro et al., 2009) and by activity stimulation, however, a short, 1 h locomotor stimulation does not change the daily pattern of BRP but magnifies the amplitude of the rhythm.

In the case of SYN the results are different than those obtained for BRP in LD 12:12 (Górskaj et al., 2013). The level of SYN in the lamina, measured as SYN-specific immunofluorescence intensity, during the 24 h cycle of LD 12:12 did not change. There are also no significant changes in SYN level in the head analyzed using the Western blot technique. These results are surprising, since SYN, like BRP, constitutes presynaptic machinery. However, SYN is involved in the regulation of the reserve pool of vesicles (Akerbergenova and Bykhovskaya, 2007), which is known to be released exclusively upon prolonged stimulation and high synaptic activity (Denker et al., 2011). During the normal activity of flies, the majority of these vesicles do not undergo either exocytosis, or recycling. It has been shown that during such circumstances only a surprisingly small percentage of vesicles (approximately 1–5%) maintains synaptic activity and is used repeatedly. Taking into account that normally active synapses appear not to require numerous reserve vesicles to sustain neurotransmitter release (Denker et al., 2011), it seems plausible that the level of SYN stays the same during the day and night. In DD on the other hand, the circadian rhythm was detected, with a higher level of SYN during the subjective day than during the subjective night. This reveals the influence of both the circadian clock and light on the expression of SYN. It also indicates that the circadian input increases the level of SYN synthesis during the subjective day. In LD 12:12 the circadian regulation seems to be masked by light, suggesting that the visual system is ready to recruit synaptic vesicles at any time during the day and night. In DD the SYN level was increased only by the short (15 min) light pulse and not by the long (60 min) one, which indicates the fast adaptation of SYN to light exposure.

SYN has been used, as a synaptic marker, in many studies in vertebrates and in D. melanogaster. The level of SYN increases during wakefulness and after sleep deprivation in the brain of Drosophila (Gilestro et al., 2009) and after exercise in the hippocampus and dentate gyrus of rats (Vaynman et al., 2004; Bechara et al., 2013). After locomotor stimulation, the increase in the SYN level in the lamina was observed only during the night. At the beginning of the day, the forced activity decreased the expression of SYN. This decrease may protect the visual system against overstimulation by light and the visual stimuli during high locomotor activity at the beginning of the day. This is also consistent with known data on SYN responses to intense stimulation. The SYN-dependent maintenance of synaptic release occurs under high-frequency nerve stimulation. The Ca2+ triggered increase in the phosphorylation of SYN, promotes recruitment of vesicles from the reserve pool to the active zone of the presynaptic element (Pieribone et al., 1995; Bloom et al., 2003; Menegon et al., 2006; Sun et al., 2006; Gaffield and Betz, 2007). In synapsin-null mutant of D. melanogaster, the proportion of vesicle recycling in vivo is increased by 30% (Denker et al., 2011).

Therefore, the two proteins, BRP and SYN, although both presynaptic and engaged in synaptic vesicle release during neurotransmission, appear to function in distinct modes that are activated upon low—or high—frequency stimulations (Knapek et al., 2011).

The third studied protein, DLG, belongs to the MAGUK family of proteins, which play a central role in molecular mechanisms involved in the regulation of spine morphogenesis and contact actin regulating proteins that are crucial for controlling the size and shape of dendritic spines (Zheng et al., 2011).

In D. melanogaster several transcripts are coded by the dlg gene, whilst two proteins, DLGA (PSD-95-like protein) and DLGS97 (SAP97-like protein) are main isofoms (Mendoza et al., 2003). The DLGA is expressed in the epithelia, somatic muscles and nervous system and the dlg mutation is lethal in the development (Woods and Bryant, 1991).

The dlns97 mutants survive until the adult stage but their behavior, including circadian rhythms, are disrupted (Mendoza-Topaz et al., 2008). In the present study we used the antibody which labels both DLGA and DLGS97 proteins, however, the observed changes are related to DLGS97 since this isoform predominates at the synapses in the brain of the adult flies (Mendoza-Topaz et al., 2008). In the lamina, we detected the circadian rhythm of DLG. Moreover, the pattern of the rhythm in DD was the same as in LD 12:12, with two morning/subjective morning and evening/subjective evening peaks. This pattern is characteristic for the daily rhythms in locomotor activity, plasticity of the lamina monopolar cells (Pyza and Meinertzhagen, 1999), activity of the sodium pump in the lamina (Górskaj et al., 2009; Damulewicz et al., 2013) and for the rhythm of BRP (Górskaj et al., 2013).

We also found that dlg shows daily expression in the brain, which suggests that the cyclic expression of dlg might be crucial for rhythms in brain functions and in behavior. In dlns97 mutants, defects in phototaxis and courtship behavior, as well as disruption of the circadian rhythm in locomotor activity have been observed (Mendoza-Topaz et al., 2008). However, the rhythm in dlg expression was not maintained in constant darkness, so the circadian clock does not control its expression directly. Surprisingly the level of DLG, measured using Western blotting, does not oscillate in whole head homogenates. It may result from different rhythms of DLG isoforms or different patterns of DLG rhythm in various brain regions. In the lamina we observed DLG—immunofluorescence in pre- and postsynaptic cells but we were not able to examine pre- and postsynaptic changes in the abundance of DLG separately.

Having analyzed the daily rhythms of DLG and BRP, we can conclude that both scaffolding proteins organize proper synaptic contacts during high activity of the visual system and locomotor
activity of flies. The correlation between activity and expression of DLG has also been found after sleep deprivation and long periods of wakefulness (Gilestro et al., 2009). In the first neuropil of the visual system we observed that the DLG level increases after light pulses in DD, activating the visual system but not after locomotor stimulation. The 1 h forced motor activity at four time points of LD 12:12 increased the abundance of DLG in the middle of the night. These results indicate that DLG, like SYN, may protect the visual system against overstimulation during the day by two stimuli, light, and locomotor activity, so DLG expression is suppressed during the day when active flies are additionally forced to fly.

Each of the three synaptic proteins that were studied appears to be endogenously regulated at synapses and additionally modulated by external stimuli. In case of the visual system, light is a strong stimulus that can synchronize, modulate, and/or mask the circadian expression of synaptic proteins. However, these proteins can also be modulated by non-visual stimuli, such as motor stimulation, which nonetheless have a stronger impact on the visual system than the visual stimulation (Kula and Pyza, 2007). In the lamina of D. melanogaster, light and motor stimulations not only affect synaptic protein expression but also the morphology of postsynaptic cells, which are remodeled in their structure during the day and night (Pyza and Meinertzhagen, 1995, 1999; Weber et al., 2009). The mechanisms of BRP, SYN, and DLG circadian regulation, and the effects of light and motor stimulations on their abundance are unknown. In our previous study on BRP we found that the circadian rhythm of this protein is abolished in the clock mutant per0 and its pattern is changed in another clock mutant tim0, in the circadian photoreceptor mutant cry0, and in the phototransduction mutant norpA (Gorska-Andrzejak et al., 2013). These cyclic and stimulodependent changes of synaptic proteins may be responsible for several daily and circadian rhythms in the morphology of neurons (Pyza and Meinertzhagen, 1995, 1999; Weber et al., 2009), glial cells (Pyza and Gorska-Andrzejak, 2004), activity of ion pumps (Pyza et al., 2004; Gorska-Andrzejak et al., 2009b; Damulewicz et al., 2013) and in other rhythms detected in the fly’s lamina (Pyza and Gorska-Andrzejak, 2008; Pyza, 2010).

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