

What bilateral damage of the superior parietal lobes tells us about visual attention disorders in developmental dyslexia

S. Valdois, D. Lassus-Sangosse, M. Lallier, O. Moreaud, L. Pisella

▶ To cite this version:

S. Valdois, D. Lassus-Sangosse, M. Lallier, O. Moreaud, L. Pisella. What bilateral damage of the superior parietal lobes tells us about visual attention disorders in developmental dyslexia. Neuropsychologia, 2018, 10.1016/j.neuropsychologia.2018.08.001. hal-02150810

HAL Id: hal-02150810 https://hal.science/hal-02150810

Submitted on 25 Oct 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

Version of Record: https://www.sciencedirect.com/science/article/pii/S0028393218304287 Manuscript 34e384aaa2b84ec1dc2e7ded6577e492

1	What bilateral damage of the superior parietal lobes tells us about visual attention			
2	disorders in developmental dyslexia			
3 4	S. Valdois ^{1,2} , D. Lassus-Sangosse ³ , M. Lallier ⁴ , O. Moreaud ^{1,3} et L. Pisella ⁵			
5				
6	1. CNRS, LPNC UMR 5105, 38040 Grenoble, France			
7	2. Université Grenoble Alpes, LPNC, 38040, Grenoble, France			
8	3. CMRR, Pôle Psychiatrie, Neurologie, Rééducation neurologique, CHU Grenoble-			
9	Alpes, France			
10	4. Basque Center on Cognition, Brain and Language, 20009, San Sebastián, Spain			
11	5. 1ImpAct – Lyon Neuroscience Research Center, Inserm U1028, CNRS UMR 5292,			
12	Bron, France,			
13				
14 15	Abstract ·			
16	Neuroimaging studies have identified the superior parietal lobules bilaterally as the neural			
17	substrates of reduced visual attention (VA) span in developmental dyslexia. It remains			
18	however unclear whether the VA span deficit and the deficits in temporal and spatial			
19	attention shifting also reported in dyslexic children reflect a unitary spatio-temporal deficit of			
20	attention - probably linked to general posterior parietal dysfunction- or the dysfunction of			
21	distinct attentional systems that relate to different neural substrates. We explored this issue			
22	by testing an adult patient, IG, with a specific damage of the bilateral superior parietal			
23	lobules after stroke, on tasks asessing the VA span as well as temporal and spatial attention			
24	shifting. IG demonstrated a very severe VA span deficit, but preserved temporal attention			
25	shifting. Exogenous spatial orientation shifting was spared but her performance was			
20 27	visual attention can be dissociated within the parietal lobe suggesting that different			
28	attentional systems associated with speficic neural networks can be selectively impaired in			
29	developmental dyslexia.			
30	Highlights: 3 to 5 on a separate file (85 characters max with blanks)			
31				
32	Investigation of a patient with bilateral superior parietal lobe (SPL) damage			
33 24	Evidence for poor visual attention span but normal temporal attention shifting			
35	Involvement of the SPLs in specific visual attention subskills			
36	Different visual attention systems are involved in developmental dyslexia			
37				
38 39				
40	Keywords: 6 Visual attention, brain damage, superior parietal lobule, developmental			
41	dyslexia, visual attention span, endogenous and exogenous spatial attention.			
42 42	Fundings: This research did not receive any specific grant from funding agoncies in the			
45 11	nublic commercial or not-for-profit sectors			

- public, commercial or not-for-profit sectors. 44
- 45

1. Introduction 46

A variety of visual attention deficits has been reported in dyslexic individuals, supporting the 2 3 visual attention account of developmental dyslexia (DD) (Vidyasagar & Pammer, 2010). More specifically, three main types of deficits have been reported: a visual attention (VA) 4 span deficit (Bosse et al., 2007), a temporal attention shifting deficit (Hari & Renvall, 5 2001) and a spatial attention orienting deficit (Facoetti et al., 2010). The Visual attention 6 7 (VA) span deficit results in poor multi-element parallel processing due to a reduction of the visual attention resources available for processing. Deficits in exogenous and endogenous 8 9 orienting of spatial attention are also reported, together with temporal attention shifting deficits that prevent the normal processing of rapidely presented stimulus sequences. 10 These deficits follow from difficulties to engage or disengage attention during spatial 11 processing or fast temporal processing. Although the existence of visual attention deficits in 12 13 DD is now well established, some authors propose that they reflect the dysfunction of one single attentional system (Facoetti et al., 2006; Vidyasagar & Pammer, 2010) while others 14 argue for two independent and functionally distinct, VA span and sluggish attentional 15 shifting, systems that could be selectively impaired in DD (Lallier et al., 2010, 2012). Here, 16 we will shed light on this debate from a neurobiological perspective. We reasoned that if 17 attentional shifting and VA span deficits reflect the dysfunction of a single attention system, 18 both skills should be implemented in the same neural network. Alternatively, two 19 functionnally distinct attentional systems should be implemented in two anatomically distinct 20 neural networks. 21

22 There is no doubt that the parietal cortex plays a key role in attentional processing (Friedrich 23 et al. 1998; Corbetta and Shulman 2002, 2011) and the sluggish attentional shifting skills of dyslexic individuals have been proposed to be underpinned by a dysfunction of this brain 24 structure (Hari and Renvall, 2001). However, attentional shifting involves several different 25 26 fronto-parietal sub-networks (Chica et al., 2013; Fan & Posner, 2002) that might be selectively affected in developmental dyslexia. From another line of research, bilateral 27 superior parietal lobules have been identified as the cerebral correlates of the VA span, thus 28 pointing to a dysfunction of a specific parietal area in relation to VA span deficits in DD 29 (Lobier et al., 2012, 2014; Peyrin et al., 2011, 2012; Reilhac, 2013). Since the role of the 30 SPLs in VA span performance is now well established, the aim of the current study was to 31 determine whether the SPLs are further involved in the other visual attention disorders 32 reported in developmental dyslexia. In order to do so, we will study the performance of a 33 34 brain-damaged patient to determine whether (i) a bilateral SPL damage impacts both VA span and temporal/spatial attention shifting performance or (ii) a SPLs damage selectively 35 impairs performance on VA span tasks without impacting temporal/spatial attention shifting. 36 Evidence for (i) would support the unitary account of the visual attention deficit in 37 developmental dyslexia, while evidence for (ii) would suggest that two distinct attentional 38 39 systems independently contribute to reading acquisition and developmental dyslexia.

40

1

41 1.1. A VA span deficit in developmental dyslexia

42

Dyslexic individuals who show a VA span deficit (Bosse et al., 2007; Dubois et al., 2010;
Germano et al., 2014; Zoubrinetzky et al., 2014, 2016) can only process a reduced number
of distinct visual elements simultaneously due to limited visual attention capacity (Bogon et al., 2014; Lobier et al., 2013). Behaviourally, the deficit is highlighted using tasks of letter

report in which a multi-consonant string is briefly and centrally presented on a computer 1 screen and subjects are required to orally report either all the consonant names (global 2 report) or the name of a single cued consonant (partial report). Some dyslexic children 3 exhibit impaired performance on these tasks despite preserved fast single letter 4 identification. However, the deficit is not specific to letter-strings since similar poor 5 performance is observed when using digits instead of letters within strings (Valdois et al., 6 7 2012). Importantly, children with poor performance in global and partial letter reports are similarly impaired in categorization tasks that do not require oral report and do not involve 8 9 verbal material (Lobier et al., 2012a). Thus, these individuals show difficulties at multielement parallel processing, regardless of the type of stimuli to be processed. In addition, 10 Lassus-Sangosse et al. (2008) showed that the VA span deficit is specific to parallel 11 processing: the dyslexic participants who were impaired at processing the 5-consonant 12 strings in parallel could report as many letters as the controls when the five consonants 13 were displayed sequentially, one at a time, at the center of the computer screen (see also 14 Valdois et al., 2011). There is also evidence that the deficit is not specific to horizontal 15 displays but extends to circular presentations (Dubois et al., 2010), which could explain why 16 poor VA span also affects performance in non-linear visual search tasks (Lallier et al., 2013) 17 and why VA span is higher in individuals who play action video games than in non-players 18 (Antzaka et al., 2017). Overall, the available behavioural data points towards a visually-19 driven deficit that reflects a limitation in the amount of attentional resources available for 20 multiple visual-element parallel processing. 21

22

23 Neuroimaging studies report the hypoactivation of the bilateral SPLs in dyslexic individuals with a VA span deficit (Peyrin et al. 2011, 2012; Reilhac et al. 2013). Indeed, the SPLs were 24 found activated under conditions of multi-element processing in normal readers (Lobier et 25 26 al., 2012) and selectively underactivated in VA span-impaired dyslexic individuals (Lobier et al., 2014). Importantly, similar hypoactivation of the SPLs was found regardless of the 27 alphanumeric or nonalphanumeric nature of the stimuli. These findings suggest that the 28 SPLs are specifically involved in the simultaneous processing of multiple visual elements. 29 Importantly, the link between VA span and the SPLs is specific. In the study of two 30 contrasted cases, Peyrin et al. (2012) showed that underactivation of the SPLs during multi-31 element processing was only found in the dyslexic participant with a VA span 32 deficit whereas the same brain regions were normally activated in the participant with a 33 34 selective phonological deficit. This link is further supported by training studies, showing that 35 the intensive use of a training program targetting VA span skills resulted in increased activation of the SPLs (Valdois et al., 2014). In sum, it is now well established that the SPLs 36 37 are the cerebral correlates of the VA span.

38

1.2. A temporal attention shifting deficit in developmental dyslexia

Temporal attention shifting deficits have been reported in developmental dyslexia in tasks that require the rapid (10 stimuli per second) sequential processing of stimuli, such as stream segregation and attentional blink tasks (Hari & Renvall, 2001). Deficits on these tasks are interpreted as reflecting sluggish attentional shifting, a difficulty to automatically disengage the focus of attention from one stimulus to reengage it onto another stimulus appearing briefly after (Hari & Renvall, 2001).

In visual stream segregation tasks, two visual stimuli alternate between two different 1 locations. When the two stimuli are temporaly far apart, they are perceived as one stream, 2 3 suggesting that attention covertly shifted from one location to the other. However, when the time interval between the two visual stimuli is shortened, they are perceived as two 4 alternating flashing stimuli at distinct positions --two distinct streams-because the 5 attentional focus can no longer shift fast enough. The stream segregation threshold 6 7 corresponds to the time interval at which perception switches from one to two streams and indicates the fastest visual attentional shifting speed. Dyslexic children were reported to 8 9 have higher visual stream segregation thresholds than normal readers, suggesting a reduced speed of automatic attentional shifting (Lallier et al., 2010a; see also Lallier et al. 10 2009 for evidence on adult participants). 11

Results from studies using the attentional blink paradigm yielded similar conclusions. In the 12 attentional blink paradigm, the visual stimuli presented in rapid succession are all displayed 13 at the same spatial location. The participants are engaged in a dual task procedure which 14 requires identifying a first target and then detecting a second target that is displayed at 15 different time lags from the first target's presentation time. The probability to detect the 16 second target decreases for the shortest lags -- participants are transiently blind to the 17 target-- and progressively improves for longer lags. While normal readers fail to detect a 18 second target that is presented within the first 500ms after the first target, the attentional 19 blink is abnormally prolonged in dyslexic individuals (Buchholz & Davies, 2007; Hari et al., 20 1999: Lallier et al., 2010b: Visser et al., 2004). 21

22

23 The parietal cortex has been proposed to be the neural substrate of the attentional blink deficit (Hari & Renvall, 2001; Hommel, Kessler, Schmitz, Gross, Akyürek, Shapiro & 24 Schnitzler, 2006). In healthy individuals, Marois et al. (2000) identified the right parietal 25 26 sulcus as the neural correlates of the attention system involved in the attentional blink paradigm, which is consistent with data from transcranial magnetic stimulation (Cooper, 27 Humphreys, Hulleman, Praamstra & Georgeson, 2004). Shapiro et al. (2002) showed that 28 the attentional blink more frequently followed from lesions of the temporo-parietal junction 29 than of the SPLs in brain-damaged patients. The overall findings suggest that a right 30 parietal sulcus dysfunction might account for visual sluggish attentional shifting skills in 31 dyslexic individuals. 32

33

1.3. A spatial attention shifting deficit in developmental dyslexia

35

Developmental dyslexia has also been associated with deficits in visuo-spatial attention. A 36 37 deficit in orienting spatial attention was hightlighted using the spatial cueing paradigm, a task originally designed by Posner (1980). In this paradigm, a target to be detected is 38 39 preceded by a briefly presented cue that orient attention towards the target location (valid condition) or the opposite side (invalid condition). Valid cues trigger faster reaction times 40 due to enhanced processing at the attended location. Invalid cues require to reallocate 41 attention towards the target location, thus yielding slower reaction times. In exogenous 42 cueing conditions, the cue is displayed in the periphery at the target location while it is 43 centrally displayed in the endogneous condition and is predictive of the target location which 44 45 appears in the periphery. Exogenous cueing involves automatic stimulus-driven attention

shifting towards the target location while endogenous cueing requires top-down voluntary
 attention.

3 Both adults and children with developmental dyslexia show orienting difficulties on spatial cueing tasks (Brannan & Williams, 1987; Facoetti et al., 2006). Reduced sensitivity to 4 exogenous cues was repeatedly reported in dyslexic individuals (Brannan & Williams, 1987; 5 Facoetti et al., 2000, 2005, 2010a; Roach & Hogben, 2004) and spatial attention inefficient 6 7 automatic orienting characterizes preliterate children with familial risk for developmental dyslexia (Facoetti et al., 2010b). Endogenous cueing was less systematically explored in 8 9 developmental dyslexia and yielded more discrepant results. In some studies, a reduced cueing effect was reported in both the exogenous and the endogenous spatial cueing 10 paradigms (Facoetti et al., 2006) but other studies reported normal cueing effect in the 11 endogenous condition (Facoetti et al., 2000). In the studies where stimulus onset 12 13 asynchrony (SOA) was manipulated, a reduced cueing effect was reported at short 100ms SOA but an amplified cueing effect at long 350ms SOA in the exogenous condition (Ruffino 14 et al., 2014). In many studies, the spatial cueing deficit was found to only characterize a 15 subset of dyslexic children who were severely impaired in pseudo-word reading accuracy 16 (Facoetti et al., 2006, 2010a; Ruffino et al., 2010). 17

18

Physiological studies of spatial attention orienting in healthy individuals have demonstrated 19 the involvement of a bilateral network with core regions in parietal and frontal areas (Nobre 20 et al., 1997; Woiciulik & Kanwisher, 1999; Yantis et al., 2002). Endogenous attention has 21 22 been postulated to be implemented in a dorsal fronto-parietal network which includes the SPL bilaterally (Corbetta & Schulman, 2002; Shomstein & Behrmann, 2010; Chica et al., 23 2013). Exogenous attention would rely on a ventral fronto-parietal network, including the 24 right temporo-parietal junction as a core region (Corbetta at al., 2000; Corbetta & Schulman, 25 26 2002; Shomstein & Behrmann, 2010). However, the anatomical segregation of the two exogenous and endogenous attention systems is still debated (Chica et al., 2013). One 27 cause of this ongoing debate is that the lesion of the ventral fronto-parietal network affects 28 the dorsal one (Corbetta & Shulman, 2011); Conversely, a lesion of the SPL does not seem 29 to affect the ventral fronto-parietal network (Gillebert et al. 2011 ; Shomstein & Behrmann, 30 2010). Thus, the exploration of the exogenous and endogenous attention systems in a 31 patient with bilateral SPL lesion is particularly relevant to distinguish the specific attentional 32 processes in which the SPL is critically involved. 33

34

35 1.4. Purpose of the current study

The purpose of the current study is to contribute to the ongoing debate on attentional 36 37 deficits in developmental dyslexia. Decades of behavioural research have provided evidence for the presence of VA span, temporal and spatial attention shifting deficits but 38 39 their co-occurrence or independence in the dyslexic population remains an unresolved issue. Only a few studies have explored temporal and spatial attention in the same dyslexic 40 participants. Ruffino et al. (2014) administered an exogenous spatial attention task and an 41 original temporal attention task to groups of dyslexic children. Only the subset of dyslexic 42 participants with the most severe pseudo-word reading accuracy deficit demonstrated 43 impaired performance in visual attention. Temporal and spatial attention deficits were found 44 45 to co-occur, suggesting that spatial exogenous and temporal attention may depend on the same functional system implemented by the same neural network. With respect to spatial 46

attention, current studies did not provide strong evidence for the independence or 1 association of exogenous and endogenous attention deficits in developmental dyslexia. 2 Some suggest deficits of the two spatial attention systems in developmental dyslexia 3 (Facoetti et al., 2006) while others suggest a selective deficit of exogenous (stimulus-diven) 4 attention (Facoetti et al., 2010b). Besides, evidence for a dissociation between VA span 5 deficit and attention shifting deficits in developmental dyslexia is also scarce. Lallier et al. 6 7 (2010c) administered tasks of attentional blink to a phonologically-impaired dyslexic participant and reported evidence for impaired temporal attention shifting despite preserved 8 9 VA span abilities. Strong behavioural evidence is thus lacking in support or against the independence of the various visual attention disorders reported in developmental dyslexia. 10

At the neurobiological level, the spatial and temporal attention deficit in developmental dyslexia, as well as the VA span deficit are attributed to a parietal dysfunction (Hari & Renvall, 2001; Gori & Facoetti, 2014; Peyrin et al., 2011, 2012; Vidyasagar & Pammer, 2010). In addition, there is strong evidence that the SPLs bilaterally are the neural substrates of the VA span deficit but the question remains whether these structures are also part of the neural substrates of temporal and spatial attention deficits in developmental dyslexia.

To address this issue, we will explore VA span, temporal attention and 18 endogenous/exogenous spatial attention skills in a brain-damaged patient who suffered a 19 bilateral SPL stroke. Based on previous evidence, this patient is expected to demonstrate a 20 severe VA span deficit. Additional deficits on temporal and spatial attention tasks would be 21 22 found if all these attentional components are implemented in the same parietal regions. 23 Alternatively, a dissociation between poor VA span but preserved temporal and spatial attention in our patient would demonstrate that several attentional systems may 24 independently contribute to developmental dyslexia. In addition, the performance of our 25 26 patient should be impaired in condition of endogenous cueing if primarily dependent on the dorsal fronto-parietal network including the SPLs but preserved in exogenous cueing if 27 implemented by the temporo-parietal junction of the ventral network (Chica et al., 2013; 28 Shomstein et al. 2010). 29

30

31 2. Case report

32

IG is a right-handed 44 year-old woman who suffered from an ischemic stroke when she 33 34 was 29. The lesion involved the whole area 7 and the intraparietal sulcus in both 35 hemispheres, as well as the upper part of Brodmann's areas 19, 18 and 39 (corresponding to area PEG as defined by Eidelberg and Galaburda 1984) but spared area 40 and the 36 37 tempo-parietal junction (see Figure 1a). Visual fields showed a partial right inferior homonymous quadrantanopia with temporal crescent sparing (Figure 1b), due to the 38 39 subcortical damage of the optical radiations below the parietal cortex in the left hemisphere. 40 Her visual acuity was normal (7/10 and 8/10 for the right and left eye respectively). She initially demonstrated a simultanagnosia and an impairment of online automatic visuo-motor 41 guidance (i.e., bilateral optic ataxia) but never showed any hemineglect syndrome. 42

The present experiment was conducted 15 years after IG's ischemic stroke long after clinical simultanagnosia had regressed. IG was administered a general neuropsychological screening battery to explore the cognitive long-term effects of her brain damage. Table 1 summarizes her performance on tests of intellectual efficiency, executive and attentional
 functions, language and visual abilities.

3

As shown on Table 1, IG's verbal IQ was well within the normal range (IQ=110). She shows 4 preserved executive functions (Wisconsin Card Sorting Test (WCST); Godefroy & Grefex, 5 2008), good verbal short-term memory (WAIS IV forward and backward digit span) and 6 good oral language abilities (Verbal IQ: Similitudes, Vocabulary, Informations; picture 7 naming from the Becs Greco, Merck et al. (2011); Pseudo-word Repetition, alphabetic and 8 9 semantic fluency). Her reading was functional and she did not complain for reading difficulties. However, she exhibited a reading age of 9 years 1 month on the Alouette 10 Reading Test (Lefavrais, 1967), suggesting limited reading skills. Note that IG reported no 11 history of reading or learning problems in childhood but she left school at the age of 16 12 (after having completed a professional certificate of dressmaker). Her limited reading 13 abilities may thus primarily reflect her socio-educational level but we cannot exclude some 14 additional effect due to her ischemic accident. 15

- 16
- 17

18

19 Figure 1: A-Horizontal section through IG's brain, visualised with structural MRI. B- Visual

- 20 field perimetry for patient IG showing quadrantanopia in the right lower quadrant of the
- visual field of both eyes.
- 22 23 A-









TASKS	IG	Mean (SD) controls	t modified, Z-score or standard score (SS)/percentile	
Intellectual efficiency				
Verbal IQ (WAIS IV)	110	100 (15)	z=0.66 σ	
Executive functions				
Wisconsin Category	6	5.95 (0.22)	t=0.22	
Wisconsin Errors	8	3.73 (2.66)	t= -1.59	
Verbal Short term memory				
Forward digit span	7	6.2 (1.3)	t= 0.61; ou SS=11/19	
Backward digit span	5	4.8 (1.4)	t= 0.14 ou SS =11/19	
Oral Language				
Picture naming	40/40	39.5 (0.7)	t=0.68	
Pseudo-word repetition	88/92	86.47 (2.62)	t=0.58	
Semantic fluency	53	39.83 (7.78)	t=1.68	
Alphabetic fluency	25	28.86 (5.55)	t=-0.69	
Written Language				
Reading Age	9yrs 1mth			
Auditory attention				
PASAT	58		75°ct	
Visual attention				
TMT A	105	42 (14)	<i>t</i> = -4.47**	
TMT B	208	66 (24)	t=-5.88 **	
Bells Test				
score	34/35		(30-50)° ct	
Time sec	222'		<5°Ct**	
Line bisection				
Long lines	+0.875cm		(50-60)°ct	
Short lines	-0.05cm		(40-50)°ct	
Visual processing (VOSP)				
Incomplete letters	17/20	Cut off : 17	=cut off	
Dot counting	4/10	Cut off : 8	z=-9.680**	
Position discrimination	15/20	Cut off : 18	<cut off**<="" td=""></cut>	
Single Letter identification				
threshold	33	33.3 (2 .25)	z=- 0.13 σ	

2 Table 1: IG general neuropsychological assessment (** = p < .05)

3

4 Assessment of her attention abilities revealed good auditory but poor visual attention skills. 5 She performed well within the normal range on the Paced Auditory Serial Addition Test (PASAT, Gronwall & Sampson, 1974; French adaptation: Mazza & Naegele, 2004), which 6 suggests normal auditory attention and good calculation skills. In contrast, her poor 7 performance on the two conditions of the Trail Making Test (TMT; Godefroy & Grefex, 2008) 8 and on the Bells Test (Gauthier, Dehaut & Joanette, 1986) suggests poor visual attention 9 processing abilities. Her performance on the latter task was characterized by good 10 accuracy, showing preserved abilities to identify a specific target among visually similar 11 distractors but at the cost of very long search time. In the same way, her very slow performance 12 in condition A of the Trail Making Test --that requires connecting numbers in sequential order -- may 13 primarily reflect a difficulty to locate the target numbers. Although performance on the TMT and 14 Bells Test suggests impaired visual attention, IG's normal performance on the line bisection 15 task (Schenkenberg et al. 1980) shows that her visual attention disorder is rather specific. 16 IG's shows no sign of visual neglect either investigated through the line bisection task or 17 through the very sensitive Bells Test. Actually, she never exhibited any hemineglect 18 19 syndrome, even when her visual attention abilities were first explored just after her stroke 20 when she was 29 (Pisella et al. 2000). The absence of hemineglect is well in line with her

very focal brain damage that spared the occipito-temporo-parietal junction. IG seems more 1 specifically impaired when the task requires the processing of multiple elements as in the 2 Bells test and TMT (but not in the line bisection task), a processing ability also involved in 3 the dot counting and position discrimination subtests of the VOSP (Visual Object and Space 4 Perception, Warrington & James, 1991) and in the incomplete letter subtest (Boucard & 5 Humphreys, 1992). In contrast, her performance in single letter visual processing is 6 7 remarkably preserved. In this task, single letters were randomly presented at the center of the screen for a short duration (varying from 33ms to 101 ms), immediately followed by a 8 9 mask. IG was able to identify 90% single letters presented for only 33ms (and all the letters displayed for longer durations) a performance well within the norm of adult skilled readers 10 (Valdois, Guinet & Embs, 2017). 11

12 **3. Experimental investigation**

IG was administered tasks of VA span, temporal and spatial attention shifting. The classical 13 paradigms of global and partial report that require parallel processing of briefly presented 5-14 consonant strings were used to assess her VA span abilities (*Experiment 1*) together with 15 an additional task of global report in which string-length and inter-letter spacing were varied 16 (Experiment 2) to determine whether the deficit extended to shorter strings and reflected 17 potential crowding effects. A sequential multi-letter processing task was further administered 18 to ensure that the deficit highlighted in the VA span tasks was specific to simultaneous 19 processing (Experiment 3). Temporal attention was assessed through visual tasks of stream 20 segregation (*Experiment 4*) and attentional blink (*Experiment 5*) while exogenous 21 (Experiment 6) and endogenous (Experiment 7) spatial cueing paradigms were used to 22 assess IG's spatial attention skills. 23

In each experiment, IG's performance was compared to that of a control participant, ERB, 24 and to control groups. ERB is a 46 year-old woman who reported no academic difficuly and 25 was matched in age and school level with IG. She was right-handed and reported no 26 medical or psychiatric illness. She exhibited a reading age of 14 years 3 months well within 27 the normal range of young adults, suggesting that IG reading age (RA= 9;1) may have been 28 affected by her brain damage. ERB was administered all the experimental tasks that were 29 30 proposed to IG. Groups of RA-matched control participants or healthy young adults were further used for comparison. In particular, IG performance was compared to the 31 performance of a group of reading age-matched (RA) controls in Experiment 1 and 2 to 32 ensure that her poor performance on the tasks was not just the consequence of her poor 33 reading level. The RA-matched control participants were administered the Alouette Reading 34 35 Test to ensure that their reading level, as a group, was comparable to that of IG (all ps>0.05). The control participants attended school regularly and had never repeated a 36 grade. None of them exhibited a reading disorder. Specific information on the control groups 37 is provided at the beginning of each experiment. 38

In each experiment, modified t-tests were used to compare IG performance with ERB performance by referencing the difference to the control sample (Crawford, Garthwaite & Wood, 2010). This method allowed testing whether the difference between ERB and IG performance was greater than the difference observed among pairs of controls, in which case they were said to differ significantly. In each task, IG performance was further compared to the control group performance using modified t-tests, as recommended when

the size of the normative sample is small (Crawford et al., 2002). Moreover, this method is

2 only minimally sensitive to departure from normal distribution (Crawford & Garthwaite,

- 3 2006), which was required here as a lot of measured abilities were within the competence of
- 4 most non-dyslexic control participants.
- 5 3.1. Visual attention span assessment
- 6 3.1.1. Experiment 1: Global and Partial Report
- 7 Participants: IG, ERB and a RA-matched control group of 108 5th Grade children (Mean
- 8 CA= 127.10; SD= 4.07) matched for reading age (Mean RA=120.89; SD= 20.06; p>.05) with
- 9 **IG participated in Experiment 1.**
- Stimuli: Random five letter-strings (e.g., RHSDM) were built up from 10 consonants (B, P, T, 10 F, L, M, D, S, R, H). The letters were presented in upper case (Geneva, 24) in black on a 11 white background. The 5-consonant strings never matched the skeleton of a real word (e.g., 12 FLMBR for FLAMBER "burn"). The strings included no repeated letter, no frequent bigram 13 (as BR or PL) and no meaningful substring (e.g., no frequent abbreviations as HLM in 14 15 French). To control for potential crowding effects, the space between adjacent letters was increased (0.6 cm). The whole line subtended an angle of 5.4°. Twenty five-letter strings 16 were successively presented in global report. Each letter was used ten times, twice in each 17 position. Fifty random five-letter strings were displayed in partial report. Each letter occurred 18 25 times and appeared five times in each position. 19

Procedure: At the beginning of each trial, a central fixation point was presented for 1000 ms 20 followed by a blank screen for 50 ms. Then, a letter-string was displayed at the center of the 21 screen for 200 ms, a duration long enough for an extended glimpse, yet too short for a 22 useful eye movement (Figure 2a). In global report, a white screen was presented at the 23 offset of the letter string and children had to report verbally as many letters as possible 24 25 immediately after the disappearance of the string. In partial report, a vertical bar indicating the position of the letter to be reported was presented for 50 ms (1.1° below the target letter) 26 at the offset of the letter-string (Figure 2b). Each letter was used as target once in each 27 position. Participants were asked to report the cued letter only. In the two global and partial 28 29 report tasks, the experimental trials were preceded of 10 training trials for which participants received feedback. No feedback was given during the experimental trials. The dependent 30 measure was the mean number of letters accurately reported (identity not location) across 31 the 20 trials (maximal score = 100) or the 50 trials for the global and partial report 32 respectively. The partial report task was adapted to account for IG's right guadrantanopia 33 that interfered with the presentation of the vertical cue when presented below the target in 34 the right visual field. The vertical bar was thus presented above instead of below the target 35 letter in both the left and right visual fields. 36

- 37
- 38
- 39
- 5.
- 40



Figure 2: (A) Procedure of global (a) and partial (b) report (Experiment 1); (B) Number or letters accurately identified in global and partial report as a function of letter position within strings for IG (black solid), ERB (grey solid) and the RA-matched control group (black hatch). For the controls, standard error bars are depicted for each position.

- 1
- 2

3 Results:

In global report (Figure 2B), IG reported 65 letters, which is significantly less (t_{modified(107)}=-4 2.24; p=0.014) than ERB (96 letters) and the RA-matched controls (mean RA score = 85.11 5 (9.8); t_{modified(108)} = -2.04, p<0.05). The ANOVA computed on the control data with letter 6 position as an intra-subject factor showed a significant letter position effect 7 (F(4;428)=156,33; p<.001), reflecting better performance on the leftmost than the rightmost 8 9 letters. Most of the variance (90.25%) in controls was explained by a linear function. A cubic function accounted for a small additional part of variance (6.77%). IG's response pattern did 10 not show the left-right gradient characteristic of the controls. She was poor at identifying the 11 letters in position 1, 2 and 3 ($t_{modified(108)}$: P1 =-5.2 ; P2 = -14.8 ; P3 = -2.66 ; all ps<0.05) but 12 showed preserved performance on the rightmost letters (P4 and P5). Her performance 13 pattern was deviant and well described by a cubic function that accounted for 97% of 14 variance. 15

In Partial Report (Figure 2B), IG only reported 16 target letters (out of 50), a performance 16 well below that of ERB (mean Partial Report score= 46; t_{modified(107)}=-4.93; p<10-6) and the 17 controls (mean=44,01 (4.3); $t_{modified(108)} = -6.48$, p<0.05). Letter report was significantly 18 impaired in IG whatever the position of the target in the string (all ts_{modified}<-1.66). In the 19 controls, the response pattern was characterized by a main position effect (F(4,404) = 5,66; 20 p<.0002). Regression analyses showed that 68% of the variance of RA controls' pattern 21 was accounted for by a linear function and an additional 31.33% by a quartic function. IG's 22 23 performance pattern was very atypical and could not be accounted for by either a linear or a 24 guartic function.

25

26 3.1.2. Experiment 2: Effect of String Length and Interletter Spacing in Global Report

Participants: The participants were IG, ERB and the same RA-matched control group as in
 Experiment 1.

29 Stimuli: Nineteen letter-strings made of 3, 4 or 5 letters were build up from 15 consonants (B, C, D, F, H, J, L, M, N, P, R, S, T, V). Each letter was around 7mm in height and 5.5 mm 30 in width. A given letter was never used twice in the same string and each letter was used 24 31 times: 10 times in the 30 5-letter strings, 8 times in the 30 4-letter strings and 6 times in the 32 33 30 5-letter strings. Each letter was used twice in each position for each of the three lengths. The 30 strings of each length were presented in two conditions of normal and large inter-34 letter spacing (Figure 3A). In the normally spaced condition, the inter-letter spacing was of 5 35 mms (close to 1 character space) whatever string length, so that strings of 3, 4 and 5 letters 36 subtended an angle of 3 degrees, 4.2 degrees and 5.4 degrees respectively (at a distance 37 of 50 cm from the computer screen). In the large spacing condition, the whole letter-string 38 subtended an angle of 10.23° (corresponding to 89.5 mm at a distance of 50 cm) whatever 39 the string length so that the distance between adjacent letters varied from 36.5 mm (around 40 6 character spaces) for the 3-letter strings, to 22.5 mm (around 4 character spaces) and 41

15.5 mm (between 2 and 3 character spaces) for the 4 and 5-letter strings respectively.
2 Letters were presented in upper case (Genova 24) in black on a white background.

Procedure: Each trial began with a 1000 ms fixation point, displayed at the center of the 3 computer screen (Figure 3). This was followed by a blank screen for 50 ms, then the letter-4 string was centrally displayed for 200 ms. The six conditions of length and spacing were 5 presented by blocks. After each trial, the participant was asked to orally report the name of 6 the letters he had identified. In each block, the participants carried out 10 training trials with 7 feedback. No feedback was provided during the experimental trials. The dependent 8 measure was the number of letters accurately identified (identity not location) out of a 9 maximal score of 75, 60 and 45 for the three length-by-spacing conditions. 10

11



12



14

15 Results

An ANOVA with Length (3, 4 and 5 letters) and Spacing (normal and large) as within-subject 16 factors was carried out on the accurracy rate of the RA-matched controls. Results (see 17 Figure 4) showed a significant main effect of length (F(2, 214) = 407.77; p<.001) and 18 spacing (F(1, 107) = 40.63, p<.001). The Length by Spacing interaction was significant (F(2, 100 Jm)). 19 214) = 13,04, p<.001). Planned comparisons showed that a spacing effect was only found 20 for 5-letter strings, the only condition in which more letters were accurately reported in the 21 normal than in the large spacing condition (81.05% vs. 77.48%). Performance of the 22 23 controls on the shorter strings (3 and 4 letters) was not affected by spacing. As shown on

- 1 Figure 4, IG performed lower than ERB in all conditions of length and spacing (all ts<-1.84,
- 2 all ps<.035), except for the 5 letter condition of large spacing (t_{modified(107)}=-0.78; p=.22).
- 3 Comparison with the RA-matched group of control participants showed that IG was severely
- 4 impaired. Her performance was poorer than for the controls in all conditions of length and
- ⁵ spacing (all ts<-2.1; p<.02), except again for the 5 letter condition of large spacing ($t_{(108)} = -$
- 6 **1.478; p=0.07**. IG did not show any disadvantage in the normal spacing condition as 7 compared to the large spacing condition (normal spacing=74.74 vs. large spacing=70.81),
- 8 thus showing the absence of crowding effect.



Figure 4 : Number of letters accurately identified by IG (black), ERB (grey) and the controls (hatch) in the two conditions of normal (NS) and large (LS) spacing for the 3-, 4- and 5-letter strings. The standard error bars are provided for the control group.

13 3.1.3. Experiment 3: Multi-letter Sequential Report

14 Participants: IG, ERB and a control group of 102 4th and 5th grade children were recruited as

- 15 controls. The control children had a mean chronological age of 11 years 4 months (mean
- 16 CA= 136.33 mths; SD=8.6) and a similar reading age as IG (mean RA=120.02, SD=14.57;
- 17 <mark>p>.05)</mark>

9

Stimuli: The strings of 5 consonants were constructed following the same constraints as in
 Experiment 1.

Procedure: A central fixation point was presented for 1000 ms followed by a blank screen 20 for 50 ms (see Figure 5A). Then 5 letters were successively displayed one at a time at the 21 center of the computer screen. Each letter was displayed for 200 ms and was immediately 22 23 followed by the next letter (ISI = 0). Participants were asked to report as many letters as possible, without any order or time constraints. They started naming letters at the end of the 24 sequential display. Ten training trials were proposed at the beginning of the task, for which 25 26 participants received feedback. No feedback was given during the 20 experimental trials. The dependent measure was the number of letters accurately reported (identity not location) 27 across the 20 trials (maximal score = 100). 28

Results: The response patterns of IG, ERB and the controls in the sequential report task are illustrated on Figure 5B. In the control group, performance was characterized by a main position effect (F(4, 404) = 128,05, p<.001), showing better report of the first than the last letters. IG reported as many letters as the controls in this sequential report task (84 vs. 72.95, SD=10.32, $t_{modified}=1.07$; p=0.21) and as many letters as ERB ($t_{modified}=0.22$; all p>0.05) p=0.21). IG scores were within the normal range for all positions ($t_{smodified}>0.22$; all p>0.05) and her response pattern was characterized by a cubic function as for the controls.

8



9

10 Figure 5: (A) The multi-letter sequential report procedure in Experiment 3; (B) Number of

11 letters accurately reported by IG (black line), ERB (grey line) and the controls (dotted line)

12 as a function of letter position within the sequential string. For the controls, standard error

- 13 bars are depicted for each position.
- 14 3.2. Temporal attention shifting
- 15 3.2.1. Experiment 4: Stream Segregation Threshold

Participants: IG, ERB and a control group of 19 healthy young adults (mean CA= 19.4 years; SD=0.6) who participated in Lallier et al. (2010; Experiment 2).

18 Stimuli: Two black dots subtending 0.1°x 0.1° of visual angle were displayed in alternance

2° above and below a fixation cross along the vertical median line of the screen (see Figure
6A). The participants were asked to fixate the central cross so that the dots were foveally

6A). The participants were asked to fixate the central cross so that the dots were for
 presented, and could be perceived accurately without eve mouvements.

Procedure: The procedure was the same as in Lallier et al. (2010, Experiment 2). Within 22 each trial, a fixation cross, subtending 0.5°x 0.5° of visual angle appeared at the center of 23 the screen for 500 ms, followed by the two dots that alternated at different time intervals 24 25 (SOAs). After each trial, the participants reported whether they had perceived one stream or two streams in a forced-choice paradigm. Each trial began with a 300 ms SOA, yielding a 26 systematic one-stream answer in all participants. The SOA was then decreased by steps of 27 40 ms, until the stimuli were perceived as two streams. The SOA was then increased or 28 decreased by steps of 20 ms until the next perception change. Steps of 10 ms and 5 ms 29 30 were then used to better estimate the segregation threshold that was defined as the mean

16

SOA over the last ten trials. Before the testing phase, a short training session was proposed to the participants. During this practice period, an unambiguous one-stream stimulus (SOA=400ms) and an unambiguous two-stream stimulus (SOA = 50 ms) were presented to be associated with the appropriate schematic drawings (see figure 6A). After each trial, participants answered by pointing at the drawing corresponding to the pattern they had perceived. When unsure, they were instructed to guess.



7

8 Figure 6: (A) Schematic representation of the visual stream segregation procedure 9 (Experiment 3). The dotted arrows symbolise the one-stream (longer SOAs) or two stream 10 (shorter SOAs) conditions. (B) Segregation threshold for IG (black), ERB (grey) and the 11 controls (hatch) in milliseconds. The standard error bars are provided for the control group.

12

Results: The segregation thresholds (Figure 6B) were estimated at 141.1 ms (SD=40.4) in the young adult healthy participants. The segregation thresholds of IG (threshold= 163.9 ms; $t_{modified}$ = -0,59) and ERB (threshold= 189.79 ms; $t_{modified}$ = -0.51) did not differ significantly from those of the controls and they did not differ between ERB and IG ($t_{modified(18)}$ =-0.45; p=0.33).

18

- 19 3.2.2. Experiment 5: Attentional Blink
- 20

Participants: IG, ERB and a RA-matched control group of 18 children who were 10 year 3
 month old on average (mean=123.72, SD= 6.14) and had a normal reading age (mean =
 123.83, SD= 11.99).

Stimuli: The stimuli were black or red digits (from 0 to 9; Arial font) that subtended a visual angle of 0.7°x0.7° at a viewing distance of 60 cm. They appeared on a grey background (red: 192; green: 192; Blue: 192). The digits were displayed at the center of the computer screen in rapid serial presentation. Each digit was presented 40 ms and was followed by a grey screen for 60 ms before the onset of the next digit, thus yielding a stimulus rate of 10 items/second. Each trial consisted of a sequence of 15, 19 or 23 digits which included a single red digit that was either a 1 or a 5 (50% probability).

Procedure: The procedure from Lallier et al. (2010) is illustrated on Figure 7A. Two 1 conditions were proposed, a dual and a single conditions. In the dual condition of attentional 2 blink, the participants had to identify a first target (T1), then detect a second target (T2). The 3 red digit ("1" or "5") was the T1 target to be identified. T1 occurred in all the sequences but 4 could appear with the same probability in the 7th, 11th or 15th position. The black digit "0" was 5 the T2 target to be detected. T2 was presented at varying time intervals following T1, from 6 7 lag 1 (SOA =100ms, T2 immediately follows T1) to lag 7 (SOA = 700ms, 6 intervening digits). T2 was present in half of the trials. In the single condition, there was no T1 targets 8 9 and the participants only had to detect the black digit "0" when present. The single condition was designed to ensure that participants could accurately process a single target (T2) when 10 presented in a stream of rapidly presented stimuli. Variations in the drop of performance 11 according to T2 temporal position was taken as an index of the attentional blink. Each 12 participant completed two successive blocks of 84 trials, the single task block first, then the 13 dual task block. A practice of 15 trials was proposed before the single task condition. The 14 experiment was administered in a dimly lit room, using E-Prime software on a PC computer 15 (computer screen=17-in.; refresh rate=85Hz). Each trial was initiated by a fixation cross, 16 presented for 500ms at the centre of the screen. The sequence of stimuli began 100 ms 17 after the fixation cross offset. After each trial, the participant had to report orally whether T2 18 was present or not for the single condition and to name the red digit ("1" or "5") and report 19 whether T2 was present or not in the dual condition. The experimenter initiated the next trial 20 by pressing the space bar on the keyboard. No feedback was given during the experimental 21 trials. 22



23

Figure 7: (A) illustration of the Attentional Blink protocole (Experiment 4); (B) Performance of IG (black line), the control group (dotted line) and ERB (grey line) in the dual condition. For the controls, standard error bars are depicted for each lag.

Results: Target detection accuracy was high in the single condition for IG ($M_{IG=}100\%$), ERB (M=86,9%) and the control participants (M =84%, SD=14,33) with no significant difference between IG and ERB ($t_{modified(17)}=0.65$; p=0.26). IG performed at ceiling and her performance did not differ from that of the controls at any lag (all $t_{modified} > .05$), showing very good ability to identify a target within rapid serial presentation. For the AB assessment in the control group, an analysis of Variance (ANOVA) with condition (single, dual) and lag (1, 2, 3, 4, 5, 6, 7) as within-subjects factors was carried out on T2 detection rate when T1 was correctly

identified. The results are provided on Figure 7B. The controls demonstrated a trend for a 1 Condition by Lag interaction (F(6; 102) = 2.036, p = .067), suggesting an attentional blink. 2 Planned comparisons (with Bonferroni correction) showed that they were less prone to 3 identify the target in the dual than in the single condition at lag 1 (F(1, 17) = 10.65; 4 p=0,005), lag 3 (F(1,17)= 8,83; p=0,0085) and lag 4 (F(1,17) = 11,13; p=0,0039). The 5 controls showed a lag-2 sparing. Overall, the performance of the controls was characterized 6 7 by an attentional blink during a time window of 100-400 ms after T1 presentation. The performance of IG (Figure 7B) did not differ significantly from that of the controls in any T2 8 9 temporal position (all ts_{modified} non-significant). In the same way, ERB performed as the controls (all modified t-tests non-significant). 10

To investigate the duration of the AB exhibited by IG, her dual condition performance was compared to the single condition performance of ERB and the controls at each lag. IG showed a performance similar as the controls and ERB performance (all ps>.05), which was also found for ERB. Lastly, in order to determine any AB depth deficit in IG, we computed the difference between the single and dual condition on T2 detection at each lag for each control participant, for ERB and for IG. The attentional blink depth was similar in IG, ERB and the controls at any lag (all modified t-tests non-significant).

- 18
- 19 3.3. Spatial Attention Shifting
- 20 3.3.1. Experiment 6 : Exogenous Orientation of Spatial Attention

Participants: Twenty-six children participated as controls. They were 10 year 10 month old
 on average (Mean= 130.11, SD = 3.65) for a reading age of 12 years on average (mean=
 144.46; SD = 16.61).

Procedure: As shown on Figure 8, each trial started with a fixation cross displayed at the 24 centre of the computer screen for 500 ms. Two circles (2.5° diameter) were then 25 simultaneously displayed for 500 ms at 8° of eccentricity from the fixation point in the left 26 and right visual fields. The rapid offset/onset (40 ms) of one of the two circles was used as a 27 peripheral cue to attrack attention randomly to the left or right circle. The target to be 28 detected was presented after two possible inter-stimulus intervals: ISI=60ms (SOA=100ms) 29 30 or ISI=160ms (SOA=200ms). The target was a dot (0.5°) displayed for 40 ms at the center of one of the two circles. The peripheral cue was either valid (corresponding to the location 31 of the following target; 50% of the trials) or invalid. Stimuli were white on a black 32 background and had a luminance of 24cd/m². Catch trials in which no target was presented 33 were intermingled with the response trials. 34

The participants were seated 50cm from the monitor with their head in a chin rest. They were asked to respond as fast as possible to the occurrence of the target by pressing the spacebar on the keyboard with their right hand. The maximum time allowed for response was 1500ms. The task began with 6 training trials. The experimental session consisted of 80 trials: 64 experimental trials (50%Valid, 50%Invalid), 16 catch trials (8 left and 8 right, 50% 100ms, 50% 200ms SOA).



1

2 Figure 8: Procedure of the exogenous spatial attention task (Experiment 5).

3 Results: RTs for ERB, IG and the controls depending on the SOA and Cue Condition are provided on Figure 9. A multifactorial ANOVA was performed on the control group RTs with 4 SOA (100ms vs. 200ms) and Cue Condition (valid vs. invalid) as within-subjects factors. 5 6 There was a significant main effect of Cue Condition (F(1, 25) = 65,033; p<.05), showing 7 that targets were detected faster when presented after a valid cue (mean $RT_{valid} = 300.41$ ms, SD_{valid} = 28,77 vs. mean RT_{invalid} = 335.34 ms, SD_{invalid} = 36,05). The SOA by Cue 8 Condition interaction was significant (F(1, 25) = 19,40; p<.002). The validity effect was 9 larger for a 100ms SOA than for a 200ms SOA (validity effect₁₀₀ = -46.65 ms, SD = 22.85 10 vs. validity effect₂₀₀ = -23.19 ms, SD = 27.75), suggesting an automatic engagement of 11 exogenous attention followed by a progessive disengagement. There was no significant 12 main effect of SOA (F<1). 13

IG and ERB exhibited similar validity effects at both the 100ms SOA (controls' mean = -14 46.65 ms, SD = 22.85 ms; validity effect in IG= -61.625 ms vs. ERB= -35.125 ms, p>0.0515 with $t_{modified}$ (25) =0.21) and the 200ms SOA (control mean = -23,19, SD = 27.75 ms; validity 16 effect in IG= -15.5 ms vs. ERB = -13.25ms, p>0.05 with $t_{modified(25)}$ =-0.06). As for the 17 controls, the validity effect of patient IG decreased with increasing SOA, indicating normal 18 engagement of exogenous attention (better at 100 ms than 200 ms) after bilateral SPL 19 damage. The index of temporal decrease (validity effect at 100 ms SOA minus validity effect 20 at 200 ms SOA) was computed for each participant. This index did not differ between IG 21 and ERB ((IG =-46.13 ms; Controls=-23.46 ms, SD_{control} = 26.63 ms; ERB=-21.88 ms, 22 23 p > 0.05 with $t_{modified(25)} = -0.64$).



1

2 Figure 9: Mean reaction times in target detection for the two exogenous conditions of cueing

3 (valid or invalid) and SOA (100 ms or 200 ms) for the control group (hatch), IG (black) and

4 ERB (grey). Standard error bars are depicted for the control group.

5

6 3.3.2. Experiment 6 : Endogenous orientation of spatial attention

7 Participants: IG, ERB and a control group of 71 healthy young adults (mean age = 19.8

8 years) whose performance on the voluntary orienting task was taken from a previous study

9 (Striemer et al. 2007) participated in Experiment 6.

Stimuli and procedure: A 80% predictive central arrow cue was used for voluntary orienting. 10 Some targets were presented without cues to examine response times for simple target 11 detection. Target location was indicated by green circles subtending 2° of visual angle, 12 presented 12° left and right of fixation (see Figure 10). The target was a red circle presented 13 within one of the green circles. The coloured stimuli were presented on a white background. 14 15 After a stimulus onset asynchrony (i.e. time between cue and target onset; SOA) of 300 or 500 ms, targets appeared at the cued (valid) or uncued (invalid) location. The participants 16 responded by pressing a button with the right hand on the keyboard. They were seated 17 50cm from the monitor with their head in a chin rest. For patient IG and her control ERB, a 18 19 longer SOA (800 ms) was additionally tested.

- 20
- 21
- 22
- 23
- 24
- 25



- 2 Figure 10: Procedure of the endogenous condition of visual attention orientation.
- 3
- 4 Results : Reaction times for target detection in the endogenous condition are provided on



5 Figure 11 for IG, ERB and the control group.

6

Figure 11: Mean reaction times for target detection in the endogenous condition of spatial
 orientation (Experiment 6) depending on cue condition (valid or invalid) and SOA (300 ms or

orientation (Experiment 6) depending on cue condition (valid or invalid
500 ms) for IG (black), the control group (hatch), and ERB (grey).

The ANOVA performed on the control RTs showed main effects of SOA and cue condition (F>12, p<0.05). RTs were faster in the long compared to the slow SOA and in the valid compared to the invalid cueing condition. There was no significant SOA by Validity interaction (F(1,70)=0.4; p>0.05), showing that, contrary to the exogenous condition, the validity effect was constant in time in the context of endogenous attention.

Contrary to the control group and ERB, IG was slower at detecting the target in the valid 6 than the invalid condition at the 500ms SOA, which suggests an inhibition of return. 7 Actually, IG and ERB exhibited different validity effects at both 300ms SOA (controls' mean 8 = -32.88 ms, SD=18.00; validity effect in IG=-40.64 ms vs. ERB =-86.0 ms, p<0.05 with 9 t_{modified(70)=}1.78) and 500ms SOA (control mean =-34.57 ms, SD= 17.67; validity effect in 10 IG=17.30 ms vs.ERB =-98 ms; p<0.05 with $\frac{1}{1000}$ =4.6, p<0.05). The index of temporal 11 decrease (validity effect at 300 ms SOA minus validity effect at 500 ms SOA) was computed 12 for each participant. IG exhibited a pathological decrease of the validity effect between 300 13 ms --where invalidly cued targets were detected slower than validly cued targets-- and 500 14 15 ms --where she showed an inhibition of return suggesting that she could not maintain her attention voluntarily as long as the controls and ERB (M_{controls}=1.54; SD_{controls}=20.66; IG_{index} 16 of decrease=-58.3; ERB_{index of decrease}=12.0, p<0.05 with $t_{modified(70)}$ =-2.41). 17

18 This temporal aspect of endogenous attention deficit was confirmed by an additional testing

of patient IG -- and comparison with ERB-- at a longer (800ms) SOA. Results are presented

20 on Figure 11. Contrary to ERB, IG did not benefit from the 80% valid cues. She did not

show faster reaction times for the valid than the invalid condition (IG validity effect=8.5 ms)

while ERB exhibited a large validity effect (ERB validity effect=-108 ms).



23

Figure 12: Reaction times of IG (black) and ERB (grey) in the valid and invalid cueing conditions of

- endogenous attention (Experiment 6) for an SOA of 800ms.
- 26

27 4. Discussion

IG, a bilateral SPL damaged patient, performed a series of visual attention tasks that 1 2 assessed the different facets of visual attention for which dyslexic participants have been shown to exhibit a deficit. Our main goal was to provide new insights on the facets of visual 3 attention that specifically relate to the superior parietal lobules. We reasonned that evidence 4 for deficits on the whole set of tasks in IG would support the existence of a single attentional 5 system related to a single neural network that includes the SPLs. As a direct consequence, 6 such a unitary account would predict that children with developmental dyslexia would show 7 simultaneous deficits on all types of tasks and that the visual attention deficits reported in 8 the dyslexic population do reflect a single visual attention dysfunction, as sometimes 9 suggested (Facoetti et al., 2006; Vidyasagar & Pammer, 2010). On the contrary. evidence 10 for a dissociation between two subsets of preserved vs. impaired VA tasks in IG would 11 12 clarify the type of visual attention skills that specifically relate to the SPLs. Such dissociation would support the existence of at least two attentional systems that relate to distinct neural 13 substrates. As a consequence, dyslexic children might show selective impairment of one or 14 the other attentional network. We could further expect these selective impairments to be 15 16 associated with distinct cognitive deficits in developmental dyslexia.

IG was first administered tasks of VA span (Experiment 1 and 2). In *Experiment 1*, she was 17 presented with briefly displayed 5-consonant strings in conditions of global and partial 18 report. In both conditions, her performance was far poorer than that of the controls, showing 19 that she could only identify a few consonants when simultaneously presented within 5-20 consonant strings. To better grasp the severity of her disorder, sequences of 3, 4 and 5 21 consonants were administered in *Experiment* 2 but in the global report condition only. 22 Results showed that her very poor performance extended to shorter strings of 3 and 4 23 consonants. Her poor performance on these two tasks of simultaneous multi-letter 24 processing (Experiment 1 and 2) contrasted with her very good report performance on 25 similar strings of five letters when presented sequentially in Experiment 3. IG performed at 26 the level of ERB and the controls on this later task and showed a similar response pattern. 27 She was guite good at processing letters when briefly presented one at a time, thus 28 showing that her poor performance on the VA span tasks (Experiment 1) can not be 29 attributed to poor oral report skills, poor single letter processing or poor verbal short-term 30 memory. A similar dissociation --poor multi-letter simultaneous processing but good multi-31 letter sequential processing—was previsouly reported in VA-span-impaired dyslexic children 32 (Lassus-Sangosse et al., 2008; Valdois et al., 2011). 33

IG's poor performance on the VA span tasks is well in line with previous evidence for a 1 2 SPLs dysfunction in children with VA span dyslexia (Lobier et al., 2014; Peyrin et al., 2011, 2012; Reilhac et al., 2013). Like IG, dyslexic children with SPL bilateral dysfunction show 3 poor performance in global and partial report tasks despite fast and normal single letter 4 processing. In line with previous neuroimaging data in healthy (Lobier et al., 2012) and 5 dyslexic (Lobier et al., 2014) individuals, the current findings confirm that the SPLs are 6 involved in multi-element (here letters, but true for non-alphanumaric characters as well, 7 Lobier et al., 2012, 2014) visual processing but not in the processing of the same elements 8 when presented in isolation. 9

Although our previous studies on developmental dyslexia mainly used sequences of 5 (or 6) items, other studies reported poor processing for shorter 4-digit strings in dyslexic children (Hawelka & Wimmer, 2005). The present findings show that a damage of the SPLs yields poor performance for even shorter letter strings. They suggest that future studies should explore VA span skills in dyslexic children more in depth, through systematic variations of string length.

Experiment 2 further assessed whether performance was sensitive to lateral interference 16 between adjacent letters, or crowding (Gori & Facoetti, 2015; Martelli et al., 2009; Whitney & 17 Levy, 2010). The classic global and partial report tasks have been initially designed to 18 19 minimize potential crowding effects by systematically increasing interletter spacing. To ensure that poor performance on VA span tasks in Experiment 1 was not just the 20 consequence of very severe crowding, spacing between letters was drastically increased in 21 the large spacing condition of Experiment 2. IG showed poor performance irrespective of 22 23 spacing for the 3 and 4 letter strings. A spacing effect was observed for 5-letter strings but showed an advantage for the small spacing condition, against any crowding interpretation. 24 25 The overall findings suggest that IG's poor performance on VA span tasks is free from lateral interference and primarily reflects a difficulty to process multiple letters within strings. 26 27 Another important consequence of the current findings is that bilateral SPL damage did not yield larger crowding effect, thus suggesting that the SPLs are not involved in crowding. 28

In Experiment 4 and 5, IG was administered tasks of stream segregation and attention blink to assess her temporal attention shifting skills. In *Experiment 4*, stream segregation threshold – corresponding to the inter-stimulus interval for which the participants could not decide whether they perceived one or two visual streams – was estimated. The segregation threshold was found similar in IG and the controls, showing that she had no difficulty to quickly engage her visual attention on a stimulus and disengage it automatically to process

1 the following one. Her preserved temporal attentional shifting skills were confirmed in 2 Experiment 5, in which IG showed similar attentional blink duration and similar attention blink depth as the controls. Overall, these findings demonstrate preserved temporal 3 attention shifting in IG despite a severe VA span deficit. A dissociation between VA span 4 and temporal attention shifting was previously reported in a case study of developmental 5 dyslexia (Lallier et al., 2010c). Evidence for the absence of correlation between visual 6 stream segregation threshold and VA span in children with developmental dyslexia further 7 supports the independence of the two underlying attentional systems (Lallier et al., 2009; 8 see also Lallier & Valdois, 2012). 9

Spatial attention shifting skills were explored in the two last experiments. In the exogenous 10 cueing condition of spatial attention (*Experiment 6*), IG showed a validity effect of the same 11 amplitude as the controls. She detected the target faster when the peripheral cue attracted 12 her attention toward the target location (valid condition), showing that she was guite efficient 13 at engaging attention on the right or left visual field following exogenous cueing. In contrast, 14 her performance was rather atypical in the endogenous condition of spatial attention 15 (Experiment 7). In this experiment where a central arrow pointing left or right indicated the 16 location of the upcoming target with 80% predictive power, the controls showed faster 17 responses and a stronger validity effect at longer SOAs. A validity effect was only found at 18 the shorter (300 ms) SOA in IG but not for longer SOAs (500 and 800 ms), suggesting that 19 20 she was unable to maintain her attention voluntarily as long as the controls. Many studies have reported a deficit in exogenous cueing in developmental dyslexia (Facoetti et al., 2005, 21 2010b; Roach & Hogben, 2004; Ruffino et al., 2014) or at risk pre-readers (Facoetti et al., 22 2010a; Franceschini et al., 2012) but the results were less consistent with respect to 23 endogenous cueing in the few studies that assessed the two cueing conditions in the same 24 participants (Facoetti et al., 2000, 2006). Reversely, a deficit specific to the endogenous 25 cueing condition was emphasized in Chinese dyslexic children in the absence of exogenous 26 spatial attention deficit (Liu, Liu, Pan & Xu, 2018). An additional key point here is strong 27 evidence in support of distinct anamo-functional attentional systems supporting endogenous 28 (goal-driven) vs. exogenous (stimulus-driven) spatial attention (Chica et al., 2013). 29

Overall, the present study provides strong evidence for the existence of distinct attentional systems that rely on distinct neural substrates. The exploration of IG visual attention skills clearly shows that bilateral superior parietal lobule damage does not result in a severe and general visual attention deficit but rather affects some specific dimensions of visual attention. IG shows impaired performance in VA span tasks of multi-element simultaneous

processing and in the endogenous condition of spatial attention shifting but preserved skills 1 2 in tasks of temporal attention shifting and exogenous spatial cueing. These findings strongly support the existence of distinct attentional systems, one of which involves the SPLs, 3 themselves part of the dorsal attentional network-DAN (Chica et al., 2013; Corbetta & 4 Shulman, 2002). They help clarifying the pattern of results reported in the scientific literature 5 on developmental dyslexia. Deficits of VA span and temporal and spatial attention shifting 6 have consistently been reported in individuals with DD but no study explored all three facets 7 of visual attention in the same participants, so that the question remains whether this 8 constellation of attentional deficits is systematically associated (or not) in the dyslexic 9 population. The current findings clearly suggest that selective deficits of VA span and 10 endogenous spatial cueing should characterize a first subset of dyslexic children who shows 11 12 a bilateral SPL dysfunction while at least a second subset might show selective deficit of temporal attention shifting and exogenous spatial attention. 13

Such a dichotomy is consistent with the hypothesis put forward by Lallier & Valdois (2012) 14 regarding the independence of the VA span theory (Bosse & Valdois, 2007) and the 15 sluggish attentional shifting (SAS, Hari & Renvall, 2001) theory of developmental dyslexia, 16 two theoretical accounts that further dissociate with respect to the contribution of visual 17 attention difficulties to phonological deficits. On one hand, a large body of research shows 18 that VA span and phoneme awareness deficits typically dissociate in developmental 19 dyslexia (Bosse et al., 2007; Germano et al., 2014; Lallier et al., 2010c; Zoubrinetzky et 20 al., 2014; See Saksida et al., 2016 for contradictory results and Reilhac et al., submitted, 21 for a response) and that VA span and phoneme awareness are independent unique 22 predictors of reading performance in typical readers (Bosse & Valdois, 2009; Lobier et al., 23 2013 ; Valdois et al., submitted ; van den Boer et al., 2013). On the other hand, sluggish 24 temporal attentional shifting typically cooccurs with phonological deficits in individuals with 25 developmental dyslexia (Lallier et al., 2009, 2010a, 2010b, 2010c) and exogenous spatial 26 attention deficits were reported in only a subgroup of dyslexic children with very poor 27 pseudo-word reading and poor phonological skills (Facoetti et al., 2010; Ruffino et al., 2014; 28 see however, Banfi, Kemény, Gangl, Schulte-Körne, Moll & Manderl, 2017). We would thus 29 expect the first subset of dyslexic children with SPLs dysfunction to show poor VA span, 30 atypical spatial endogenous attention but preserved phonological skills while dyslexic 31 children with poor temporal attention shifting and poor exogenous spatial attention would be 32 further impaired in phonological processing. Some recent findings further suggest that the 33 34 latter but not the former would show a categorical perception deficit (Zoubrinetzky et al., 2016). 35

The current findings also provide new insights on potential association/dissociation with an 1 2 asymmetric distribution of attention between the left and right hemifields --or minineglect-and atypical performance in visual search tasks depending on dyslexia subtypes. 3 Interhemispheric asymmetries have been searched for in developmental dyslexia and 4 sometimes reported (Facoetti & Molteni, 2001; Facoetti & Turatto, 2000; Hari, Renvall & 5 Tanskanen, 2001; Sireteanu et al., 2005). However, some case studies of dyslexic children 6 with a VA span deficit failed to report any asymmetry between the left and right visual 7 hemifields (Dubois et al., 2010; Valdois et al., 2011). IG's results are guite in line with these 8 later reports, showing that a bilateral SPL damage does not yield visuo-spatial neglect. 9 These findings are in line with the current litterature on the neural substrates of spatial 10 attention and unilateral spatial neglect. Lesions of the ventral attentional network-VAN -in 11 12 particular the right temporo-parietal junction-- and not of the dorsal attentional network-DAN, are traditionnaly associated with unilateral spatial neglect (Chica et al., 2013; Mort et al. 13 2003; Vallar & Perani, 1987). Moreover, although the VAN and DAN are known to interact, 14 this interaction is not symetrical. Recent neuroimaging results suggest that a structural 15 lesion of the VAN affects the functioning of the DAN (Corbetta et al. 2005) while conversely 16 a lesion of the DAN produces restricted deficits of spatial attention (Gillebert et al. 2011, 17 18 Shomstein et al. 2010) with features of Balint syndrome (e.g. visual disorientation, deficits of global perception, shape identification in a cluttered field, see Pisella et al., 2015, 2013), 19 20 ressembling more to VA span dyslexia than the clinical picture of spatial deficits in neglect. The current findings are further in line with evidence that the right-hemispheric regions of 21 the VAN are specifically involved in exogenous covert shifting of spatial attention, towards 22 both the left and right visual hemifields whereas the DAN is involved in the spatial selection 23 of objects for voluntary shifting of overt and covert attentional exploration (Chica et al., 24 2011; Corbetta et al., 2000; Corbetta & Shulman, 2011; Hopfinger et al., 2000; Kastner et 25 al., 1999). 26

IG's lesion matches the parietal regions of the DAN. She has been previously studied in visual search and shows a reduced visuo-attentional window specifically when she faces stimuli made by a combination of lines (Khan et al., 2016). This deficit in visual search after bilateral SPL damage is in line with the impaired search performance specific to « multifeatures shapes » combining separable features previously reported in poor readers (Casco & Prunetti, 1986) and with evidence for a visual search deficit in VA span impaired dyslexic children (Lallier et al., 2013).

34 **5. Conclusion**

Exploration of distinct facets of visual attention in patient IG, who suffers a bilateral SPL 1 2 lesion, supports the existence of distinct attentional systems that relate to distinct neural substrates. IG shows a lesion of the DAN yielding to poor multielement simultaneous 3 processing (i.e. poor VA span) and poor endogenous spatial attention while temporal 4 attention shifting and exogenous spatial attention are intact, showing that they relate to a 5 distinct attentional network, quite probably the VAN. Exploration of patient IG further 6 provides new insights on developmental dyslexia. A bilateral SPL dysfunction has been 7 reported in a subset of dyslexic children who show poor VA span but preserved 8 phonological skills. In line with the current findings, these dyslexic individuals are free from 9 sluggish temporal attention shifting but impaired in visual search. They should further show 10 impaired endogenous but preserved exogenous spatial attention, which remains to be 11 12 systematically investigated. In contrast, the current findings suggest that the exogenous spatial attention and temporal attention shifting deficits reported in developmental dyslexia, 13 actually defines another subset of dyslexic individuals who show associated phonological 14 deficits but preserved VA span and might be more prone to show a left-right hemifield 15 16 asymetry in tasks of spatial processing. While the temporal and spatial attentional functions of the VAN and the DAN are both involved in the development of reading, we argue that 17 they should nevertheless be distinguished. Evidence from IG allows making new predictions 18 about developmental dyslexia, which will help clarifying the role of each attentional 19 20 subsystem on reading acquisition and the consequences of a selective deficit of one or the 21 other attentional system in developmental dyslexia.

22

23 6. References

Banfi, C, Kenény, F., Gangl, M., Schulte-Körne, G., Moll, K. & Landerl, K. (2017). Visuao spatial cueing in children with differential reading and spelling profiles. Plos One, 12
 (7):e0180358.

Bogon, J., Finke, K. & Stenneken, P. (2014). TVA-based assessment of visual attentional
 functions in developmental dyslexia. *Frontiers in Psychology*, 5, 1172. doi:
 10.3389/fpsyg.2014.01172

Bosse, M.L., Tainturier, M.J. & Valdois, S. (2007). Developmental dyslexia: the visual attention span deficit hypothesis. *Cognition*, 104, 198-230.

Bosse, M.L. & Valdois, S. (2009). Influence of the visual attention span on child reading performance: a cross-sectional study. *Journal of Research in Reading*, 32, 2, 230-253.

Boucard, M. & Humphreys, G.W. (1992). The computation of conceptual structures from collinearity and closure: normality and pathology. *Neuropsychologia*, 30, 6, 527-546.

- Brannan, J. & Williams, M. (1987). Allocation of visual attention in good and poor readers.
 Perception & Psychophysics, 41, 23–28.
- Buchholz, J. & Davies, A. (2007). Attentional blink deficits observed in dyslexia depend on
 task demands. *Vision Research*, 47, 1292–1302.
- Chica, A.B., Bartolomeo, P. & Lupiáñez, J. (2013). Two cognitive and neural systems for
 endogenous and exogenous spatial attention. *Behavioral Brain Research*, 237: 107 123.
- 8 Chica, A.B., Bartolomeo, P. & Valero-Cabré., A. (2011). Dorsal and ventral parietal
 9 contributions to spatial orienting in the human brain. *Journal of Neuroscience*, 31, 22,
 10 8143-8149.
- Cooper, A.C.G., Humphreys, G.W., Hulleman, J., Praamstra, P. & Georgeson, M. (2004).
 Transcranial magnetic stimulation to right parietal cortex modifies the attentional blink.
 Experimental Brain Research, 155, 24-29.
- Corbetta, M., Kincade, M.J., Lewis, C., Snyder, A.Z. & Sapir, A. (2005). Neural basis and
 recovery of spatial attention deficits in spatial neglect. *Nature Neuroscience*, 8, 1603–
 1610.
- Corbetta, M., Kincade, J.M., Ollinger, J.M., McAvoy, M.P. & Shulman, G.L. (2000).
 Voluntary orienting is dissociated from target detection in human posterior parietal
 cortex. *Nature Neuroscience*, 3, 3, 292–297.
- Corbetta, M. & Shulman, G.L. (2002). Control of goal-directed and stimulus-driven attention
 in the brain. *Nature Reviews Neuroscience*, 3, (3): 201-215.
- Corbetta, M. & Schulman, G.L. (2011). Spatial neglect and attention network. *Annual Review Neuroscience*, 34, 569-599.
- Crawford, J.R., & Garthwaite, P.H. (2002). Investigation of the single case in
 neuropsychology: Confidence limits on the abnormality of test scores and test score
 differences. Neuropsychologia, 40, 1196-1208.
- Crawford, J. R. & Garthwaite, P. H. (2006). Methods of testing for a deficit in single case
 studies: evaluation of statistical power by Monte Carlo simulation. *Cognitive Neuropsychology*, 23, 6, 877-904.
- Crawford, J. R., Garthwaite, P. H. & Wood, L.T. (2010). Infrential methods for comparing
 two single cases, Cognitive *Neuropsychology*, 27, 377-400.
- Dubois, M., Kyllingsbaek, S., Prado, C., Peiffer, E., Lassus-Sangosse, D. & Valdois, S.
 (2010). Fractionating the multi-element processing deficit in developmental dyslexia:
 Evidence from two case studies. *Cortex*, 46, 6, 717-738.
- Eidelberg, D. & Galaburda, A.M. (1984). Inferior parietal lobule: Divergent architectonic
 asymmetries in the human brain. *Archives of Neurology*, 41, 843–852
- 39

- Facoetti, A., Corradi, N., Ruffino, M., Gori, S. & Zorzi, M. (2010a). Visual spatial attention
 and speech segmentation are both impaired in preschoolers at familial risk for
- developmental dyslexia. *Dyslexia*, 16, 226-239.
- Facoetti, A., Lorusso, M. L., Cattaneo, C., Galli, R. & Molteni, M. (2005). Visual and auditory
 attentional capture are both sluggish in children with developmental dyslexia. *Acta Neurobiol. Exp.* 65, 61–72.
- Facoetti, A. & Molteni, M. (2001). The gradient of visual attention in developmental dyslexia.
 Neuropsychologia, 39, 352–357.
- Facoetti, A., Paganoni, P., Turatto, M., Marzola, V., and Mascetti, G. G. (2000). Visual spatial attention in developmental dyslexia. *Cortex*, 36,109–123.
- Facoetti, A. & Turatto, M. (2000). Asymmetrical visual fields distribution of attention in
 dyslexic: A neuropsychological study. *Neuroscience Letters*, 290, 216 218.
- Facoetti, A., Turatto, M., Lorusso, M. L. & Mascetti, G. G. (2001). Orienting of visual
 attention in dyslexia: evidence for asymmetric hemispheric control of attention.
 Experimental Brain Research, 138, 46–53.
- Facoetti, A., Trussardi, A. N., Ruffino, M., Lorusso, M. L., Cattaneo, C., Galli, R. et al.
 (2010b). Multisensory spatial attention deficits are predictive of phonological decoding
 skills in developmental dyslexia. *Journal of Cognitive Neuroscience*, 22,1011–1025.
- Facoetti, A., Zorzi, M., Cestnick, L., Lorusso, M. L., Molteni, M., Paganoni, P. et al. (2006).
 The relationship between visuo-spatial attention and nonword reading in developmental
 dyslexia. *Cognitive Neuropsychology*, 23, 841–855.
- Fan, J., McCandliss, B.D., Sommer, T., Raz, A. & Posner, M.I. (2002). Testing the efficiency
 and independence of attentional networks. *Journal of Cognitive Neuroscience*, 14, 3,
 340-347.
- Franceschini, S., Gori, S., Ruffino, M., Pedrolli, K. & Facoetti, A. (2012). A causal link
 between visual spatial attention and reading acquisition. *Current Biology*, 22, 814–819.
- Friedrich, F.J., Egly, R., Rafal, R.D. & Beck, D. (1998). Spatial attention deficits in humans :
 a comparison of the superior parietal and temporal-parietal junction lesions.
 Neuropsychology. 1998, 12, 2, 193-207.
- Gauthier, L., Dehaut, F. & Joanette, Y. (1989). The Bells Test: A□ quantitative and
 qualitative test for visual neglect. *International Journal of Clinical Neuropsychology*, 11,
 49-54.
- Germano, D.G., Reilhac, C., Capellini, A.S. & Valdois, S. (2014). The phonological and
 visual bases of developmental dyslexia in Brazilian Portuguese. *Frontiers in Educational Psychology 5, 1169.*
- Gillebert, C.R., Mantini, D., Thijs, V., Sunaert, S., Dupont, P. & Vandenberghe, R. (2011).
 Lesion evidence for a critical role of the intraparietal sulcus in spatial attention. *Brain*,
 134, 1694-1709.

- Godefroy O. & le GREFEX (2008). Fonctions exécutives et pathologies neurologiques et psychiatriques: évaluation en pratique clinique. Marseille: Solal.
- Gori, S. & Facoetti, A. (2016). How the visual aspects can be crucial in reading acquisition?
 The intriguing case of crowding and developmental dyslexia. *Journal of vision*, 15, 1-20.
- Gronwall, D. & Sampson, H. (1974). *The Psychological effects of concussion*. Auckland
 University Press, Auckland, New Zeland.
- Hari, R., & Renvall, H. (2001). Impaired processing of rapid stimulus sequences in dyslexia.
 Trends in Cognitive Science, 5, 525–532.
- Hari, R., Renvall, H. & Tanskanen, T. (2001). Left minineglect in dyslexic adults. Brain, 124,
 1373-1380.
- Hari, R., Valta, M., & Uutela, K. (1999). Prolonged attentional dwell time in dyslexic adults.
 Neuroscience Letters, 271, 3, 202-204.□
- Hawelka, S. & Wimmer, H. (2005). Impaired visual processing of multiple-element arrays is
 associated with increased number of eye movements in dyslexic readers, *Vision Research*, 45, 855-863.
- Hopfinger, J.B., Buonocore, M.H. & Mangun, G.R. (2000). The neural mechanisms of top down attentional control. *Nature Neuroscience*, 3, 284–291.
- Khan, A.Z., Prost-Lefebvre, M., Salemme, R., Blohm, G., Rossetti, Y., Tilikete, C. & Pisella,
 L. (2016) The attentional fields of visual search in simultanagnosia and healthy
 individuals: How object and space attention interact. *Cerebral Cortex*, 26, 3, 1242-1254.
- Kastner, S., Pinsk, M.A., De Weerd, P., Desimone, R. & Ungerleider, L.G. (1999). Increased
 activity in human visual cortex during directed attention in the absence of visual
 stimulation. *Neuron*, 22, 751–761.
- Lallier, M., Donnadieu, S., Berger, C. & Valdois, S. (2010c). A case study of developmental
 phonological dyslexia: Is the attentional deficit in rapid stimuli sequences processing
 amodal? *Cortex*, 46, 231-241
- Lallier, M., Donnadieu, S., Thierry, G., Tainturier, M. J., Billard, C., Peyrin, C. et al. (2009).
 Auditory and visual stream segregation in children and adults: An assessment of the
 amodality assumption of the 'sluggish attentional shifting' theory of dyslexia. *Brain Research*, 1302, 132–147.
- Lallier, M., Donnadieu, S. & Valdois, S. (2010b). Visual attentional blink in dyslexic children: Parameterizing the deficit. *Vision Research*, 50, 1855-1861.
- Lallier, M., Donnadieu, S. & Valdois, S. (2013). Investigating the role of visual and auditory
 search in reading and developmental dyslexia. *Frontiers in Human Neuroscience*, 7,
 597. doi: 10.3389/fnhum.2013.00597
- Lallier, M., Tainturier, M. J., Dering, B., Donnadieu, S., Valdois, S. & Thierry, G. (2010a).
 Behavioral and ERP evidence for amodal sluggish attentional shifting in developmental dyslexia. *Neuropsychologia*, 48, 4125–4135.

- Lallier, M. & Valdois, S. (2012). Sequential versus Simultaneous processing deficits in
 developmental dyslexia. In T. Wydell & L. Fern-Pollak (Eds.), *Dyslexia: A comprehensive and international approach*. In Tech, DOI:10.5772/39042.
- Lassus-Sangosse, D., N'Guyen-Morel, M.A. & Valdois, S. (2008). Sequential or
 simultaneous visual processing deficit in developmental dyslexia. *Vision Research*, 48, 979-988.
- Lefavrais, P. (1967). *Test de l'Alouette*. [The "Alouette" Reading Test.] Paris: Editions du
 Centre de Psychologie Appliquée.
- Liu, S., Liu, D., Pan, Z. & Xu, Z. (2018). The association between reading abilities and
 visual-spatial attention in Hong-Kong Chinese children. Dyslexia, 10.1002/dys.1584.
- 11
- Lobier, M., Dubois, M. & Valdois, S. (2013). The role of visual processing speed in reading
 speed development. *Plos One*, 8, 4, e58097.
- Lobier, M., Peyrin, C., Pichat, C., LeBas, J.F. & Valdois, S. (2012). Neural correlates of preorthographic character string processing. *Neuropsychologia*, 50, 9, 2195-2204.

Lobier, M., Peyrin, C., Pichat, C., Le Bas, J.F. & Valdois, S. (2014). Visual processing of
 multiple elements in the dyslexic brain: Evidence for a superior parietal dysfunction.
 Frontiers in Human Neurosciences, 8, 479.

- Lobier, M., Zoubrinetzky, R. & Valdois, S. (2012). The visual attention span deficit in
 developmental dyslexia is visual and not verbal. *Cortex*, 48, 6, 768-773.
- Marois, R., Chun, M.M. & Gore, J.C. (2000). Neural correlates of the attentional blink,
 Neuron, 28, 299-308.
- Martelli, M. Di Filippo, G., Spinelli, D. & Zoccolotti, P. (2009). Crowding, reading and
 developmental dyslexia. *Journal of Vision*, 9, 4, 1-18.
- Mazza, S. & Naegele, B. (2004). *Test d'attention soutenue : PASAT modifié, adaptation française*, Collection : Tests et matériel en orthophonie, éd. Solal.
- Merck, C., Charnallet, A. & Auriacombe, S. (2011). La batterie d'évaluation des
 connaissances sémantiques du GRECO (BECS-GRECO): validation et données
 normatives. *Revue de neuropsychologie*, 3, 235-255.
- Mort, D.J., Malhotra, P., Mannan, S.K., Rorden, C., Pambakian, A. et al. (2003). The anatomy of visual neglect. *Brain*, 126, 1986–1997.
- Nobre, A.C., Sebestyen, G.N., Gitelman, D.R., Mesulam, M.M., Frackowiak, R.S. & Frith,
 C.D. (1997). Functional localization of the system for visuospatial attention using
 positron emission tomography. *Brain*, 120, 515–533.

Peyrin, C., Démonet, J.F., Baciu, M., LeBas, J.F. & Valdois, S. (2011). Superior parietal
 lobe dysfunction in a homogeneous group of dyslexic children with a single visual
 attention span disorder. *Brain & Language*, 118, 128-138.

Peyrin, C., Lallier, M., Démonet, J.F., Pernet, C., Baciu, M., LeBas, J.F. & Valdois, S.
 (2012). Neural dissociation of phonological and visual attention span disorders in

- developmental dyslexia: fMRI evidence from two case studies. *Brain & Language*, 120
 (3), 381-394.
- Pisella, L., Biotti, D. & Vighetto, A. (2015). Combination of attentional and spatial working
 memory deficits in Balint syndrome. Special issue on "Competitive visual processing
 across space and time: Interactions with memory", *Annals of the New York Academy of Sciences*, 1339, 165-175.

Pisella L., Blangero, A., Tilikete, C., Biotti, D., Rode, G., Vighetto, A., Mattingley, J.B. &
Rossetti, Y. (2013). Attentional disorders. In: K. Ochsner & S. Kosslyn (Eds): *The Oxford Handbook of Cognitive Neuroscience*, Volume 1 (pp. 319-350). Oxford
University Press, Oxford, UK.

- Pisella, L., Gréa, H., Tilikete, C., Vighetto, A., Desmurget, M., Rode, G., Boisson, D. &
 Rossetti, Y. (2000). An 'automatic pilot' for the hand in human posterior parietal cortex:
 toward reinterpreting optic ataxia. *Nature Neuroscience*, 3, 7, 729–736.
- Posner, M.I. (1980). Orienting of attention. *Quarterly Journal of Experimental Psychology*,
 32, 3–25.
- Reilhac, C., Peyrin, C., Démonet, J.F. & Valdois. S. (2013). Role of the superior parietal
 lobules in letter-identity processing within strings: fMRI evidence from skilled and
 dyslexic readers. *Neuropsychologia*, 51, 4, 601-612.
- Reilhac, C., Bosse, M.L. & Valdois, S. (submitted). Phoneme awareness, visual attention
 span and rapid automatized naming disorders in poor readers: the prevalence issue.
- Roach, N.W. & Hogben, J.H. (2004). Attentional modulation of visual processing in adult
 dyslexia. *Psychological Science*, 15, 10, 650-654.
- Roussel, M. & Godefroy O. (2008). *La batterie GREFEX : données normatives*. GREFEX
 (éd.), Fonctions exécutives et pathologies neurologiques et psychiatriques , 231-266.
- Ruffino, M., Gori, S., Boccardi, D., Molteni, M. & Facoetti, A. (2014). Spatial and Temporal
 attention in developmental dyslexia. *Frontiers in Human Neuroscience*, 8, 331.
- Ruffino, M., Trussardi, A. N., Gori, S., Finzi, A., Giovagnoli, S., Menghini, D. et al. (2010).
 Attentional engagement deficits in dyslexic children. *Neuropsychologia*, 48, 3793–3801.
- Saksida, A., Iannuzzi, S., Bogliotti, C., Chaix, Y., Démonet, J.F., Bricout, L. et al. (2016).
 Phonological skills, visual attention span and visual stress in developmental dyslexia.
 Developmental Psychology, 52, 10, 1503-1516.
- Schenkenberg, T., Bradford, D.C. & Ajax, E.T. (1980). Line bisection and unilateral visual
 neglect in patients with neurologic impairment. *Neurology*, 30 (5), 509-509.

Shomstein, S., Lee, J. & Behrmann, M. (2010). Top-down and bottom-up attentional
 guidance: investigating the role of the dorsal and ventral parietal cortices. *Experimental Brain Research*, 206, 197-208.

- Sireteanu, R., Goertz, R., Bachert, I. & Wandert, T. (2005). Children with developmental
 dyslexia show a left visual « minineglect ». Vision Research, 45, 3075-3082.
- Shomstein, S., Lee, J., & Behrmann, M. (2010). Top-down and bottom-up attentional
 guidance : investigating the role of the dorsal and ventral parietal cortices. *Experimental Brain Research*, 2006, 2, 197-208.
- Striemer C., Blangero A., Rossetti Y., Boisson D., Rode G., Vighetto A., Pisella L. &
 Danckert J. (2007) Deficits in peripheral visual attention in patients with optic ataxia.
 Neuroreport, 18, 11, 1171-1175.
- Vallar, G. & Perani, D. (1987). The anatomy of spatial neglect in humans. In: Jeannerod, M.
 (Ed.), *Neurophysiological and Neuropsychological Aspects of Spatial Neglect* (pp. 235 258). North-Holland: Elsevier Science Publishers.
- Valdois, S., Bidet-Ildei, C., Lassus-Sangosse, D., Reilhac, C., N'Guyen, M.A., Guinet E. &
 Orliaguet, J.P. (2011). A visual processing but no phonological disorder in a child with
 mixed dyslexia. *Cortex*, 47, 1197-1218.
- Valdois, S., Guinet, E. & Embs, J.L. (2017). *Evadys : Outil diagnostic des troubles de l'empan visuo-attentionnel.* Lyon : Happy Neuron (www.happyneuronpro.com).

Valdois, S., Lassus-Sangosse, D. & Lobier, M. (2012). Impaired letter string processing in
 developmental dyslexia : what visual-to-phonological code mapping disorder ? *Dyslexia*,
 18 (2), 77-93

- Valdois, S., Peyrin, C., Lassus-Sangosse, D., Lallier, M., Démonet, J.F. & Kandel, S.
 (2014). Dyslexia in a French Spanish bilingual child: Behavioural and neural modulation
 following a specific VA span intervention program. *Cortex, 53, 120-145*.
- Valdois, S., Roulin, J.L. & Bosse, M.L. (Submitted). Reading fluency depends on pre readers' ability to process multiple visual elements in parallel.
- Van den Boer, M., de Jong, P.F. & Haentjens-van Meeteren, M.M. (2013). Modeling the
 length effect : Specifying the relation with visual and phonological correlates of reading.
 Scientific Studies of Reading, 17, 4, 243-256.
- Vidyasagar, T. R. & Pammer, K. (2010). Dyslexia: a deficit in visuo-spatial attention, not in
 phonological processing. *Trends in Cognitive Science*, 14, 57–63.
- Visser, T. A., Boden, C. & Giaschi, D. E. (2004). Children with dyslexia: evidence for visual
 attention deficits in perception of rapid sequences of objects. *Vision Research*, 44, 21,
 2521-2535,
- Warrington, E.K. & James, M. (1991). *The Visual Object and Space Perception Battery.* Bury St Edmunds Thanes Valley Test Company.
- Whitney, D. & Levy, D.M. (2010). Visual crowding : a fundamental limit on conscious perception and object recognition. *Trends in Cognitive Sciences*. 15, 4, 160-168.
- Wojciulik, E. & Kanwisher, N. (1999). The generality of parietal involvement in visual attention. *Neuron*, 23, 4, 747–764.

- Yantis, S., Schwarzbach, J., Serences, J.T., Carlson, R.L., Steinmetz, M.A., Pekar, J.J. et
 al. (2002). Transient neural activity in human parietal cortex during spatial attention
 shifts. *Nature Neuroscience*, 5, 10, 995–1002.
- Zoubrinetzky, R., Bielle, F. & Valdois, S. (2014). New insights on developmental dyslexia
 subtypes: Heterogeneity of mixed reading profiles. *Plos One*, 9(6): e99337.
- Zoubrinetzky, R., Collet, G., Serniclaes, W., N'Guyen-Morel, M.A. & Valdois, S. (2016).
 Relationships between categorical perception of phonemes, phonological awareness, and visual attention span in developmental dyslexia. *Plos One, 11(3): e0151015*
- 10
- 11

.

.

- 12
- 13
- 14