



Contents lists available at ScienceDirect

Schizophrenia Research

journal homepage: www.elsevier.com/locate/schres

Auditory versus visual neuroscience-informed cognitive training in schizophrenia: Effects on cognition, symptoms and quality of life

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ARTICLE INFO

Article history:

Received 19 February 2020

Received in revised form 1 May 2020

Accepted 6 May 2020

Available online xxxx

Keywords:

Schizophrenia
Cognitive training
Visual training
Auditory training
Neuroplasticity
Computerized training

ABSTRACT

Background: Cognitive impairments are related to deficits in primary auditory and visual sensory processes in schizophrenia. These impairments can be remediated by neuroscience-informed computerized cognitive trainings that target auditory and visual processes. However, it is not clear which modality results in greater improvements in cognition, symptoms and quality of life. We aimed to investigate the impact of training auditory versus visual cognitive processes in global cognition in patients with schizophrenia.

Methods: Seventy-nine schizophrenia participants were randomly assigned to either 40 h of auditory or visual computerized training. Auditory and visual exercises were chosen to be dynamically equivalent and difficulties increased progressively during the training. We evaluated cognition, symptoms and quality of life before, after 20 h, and after 40 h of training. [ClinicalTrials.gov](https://clinicaltrials.gov/ct2/show/study/11032700) (1R03TW009002-01).

Results: Participants who received the visual training showed significant improvements in global cognition compared to the auditory training group. The visual training significantly improved attention and reasoning and problem-solving, while the auditory training improved reasoning and problem-solving only. Schizophrenia symptoms improved after training in both groups, whereas quality of life remained unchanged. Interestingly, there was a significant and positive correlation between improvements in attention and symptoms in the visual training group.

Conclusions: We conclude that the visual training and the auditory training are differentially efficient at remediating cognitive deficits and symptoms of clinically stable schizophrenia patients. Ongoing follow-up of participants will evaluate the durability of training effects on cognition and symptoms, as well as the potential impact on quality of life over time.

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1. Introduction

Schizophrenia is a complex and highly incapacitating mental disorder that has the highest disability weight, according to the Global Burden of Disease (Salomon et al., 2012). This has a significant impact on affected individuals and their families, resulting in important societal and economic costs (Knapp et al., 2005). Schizophrenia is often characterized by cognitive impairments that are important predictors of poor quality of life, occupational and social functioning (Green, 2006; McGurk and Meltzer, 2000; Ritsner, 2007). Cognitive impairments in

schizophrenia have been shown in the domains of attention, memory, executive functions, and social cognition (Bildler et al., 2002; Hutton et al., 1998; Szoke et al., 2008), and are associated with brain structural abnormalities (Crespo-Facorro et al., 2007).

Over the last decades it has been proposed that abnormalities in primary sensory brain regions in schizophrenia induce dysfunctions in basic visual and auditory processes, which may contribute to impairments in higher-order functions (Javitt, 2009). Deficits in the auditory system are apparent in basic sensory processing, such as tone encoding and detection of pitch deviation (Javitt et al., 1993, 1997; March et al., 1999). Deficits in the visual system are evident in smooth eye movements, motion detection, and contrast sensitivity (Butler et al., 2008; O'Donnell et al., 1996). Importantly, deficits in basic processing are associated with impairments in higher-order processes such as phonological

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processing, auditory emotion recognition, recognition of objects and facial expressions (Javitt, 2009), which may ultimately contribute to poor social outcomes (Arnott et al., 2011; Dale et al., 2010; Javitt, 2009; Kantrowitz et al., 2015; Koro et al., 2002; Martínez et al., 2015; Morrison et al., 1988; Revheim et al., 2006; Tek et al., 2002).

Over the past decade, Dr. Vinogradov's group and others have shown that neuroscience-informed cognitive training targeting auditory processes improves global cognition, speed of processing, verbal learning and memory, as well as reasoning and problem-solving in early onset and chronic schizophrenia subjects (Fisher et al., 2009). Importantly, participants who were able to make the most progress through basic auditory training exercises also showed the most improvement in higher-level cognitive outcome measures (Adcock et al., 2009; Biagianni et al., 2016). The brain of trained schizophrenia participants recovered impaired sensory gating and oscillations in neural activity related to auditory processing (Popov et al., 2011, 2015); and restored neural patterns of response in a source monitoring task (Subramaniam et al., 2012). A significant positive relationship was evident between training-induced enhancements of prefrontal cortical activity and better ratings on social and occupational domains of the Quality of Life Scale six months after completion of training (Subramaniam et al., 2014). Cognitive improvements in schizophrenia were also associated with increases in serum levels of pro-cognitive molecules such as BDNF (Vinogradov et al., 2009) and D-serine (Panizzutti et al., 2009), as well as variations in the cognition-related gene COMT (Panizzutti et al., 2013).

On the other hand, fewer studies evaluated cognitive training targeting visual processes in schizophrenia. In one study that evaluated schizophrenia subjects undertaking different trainings (visual, auditory and cognitive control) concomitantly, improvements in the visual perceptual exercises were associated with gains in visual but not verbal memory (Surti et al., 2011). Also, a specific training on motion perception improved impairments in the visual modality, but with no transfer of effect to other cognitive measurements (Norton et al., 2011).

In sum, the evidence indicates that neuroscience-informed cognitive training targeting auditory and visual processes can remediate brain dysfunctions present in schizophrenia. However, it is not known which sensory modality (visual or auditory) of cognitive training will generalize more efficiently to the global cognition in schizophrenia. Global cognition is defined as a composite measure encompassing the Measurement And Treatment Research to Improve Cognition in Schizophrenia (MATRICS)-defined domains affected in schizophrenia: speed of processing, attention, working memory, verbal learning, visual learning, reasoning and problem-solving, and social cognition. To investigate the comparative impact of cognitive training with auditory versus visual processes in schizophrenia, we posed two questions: 1) Which modality of intensive sensory cognitive training - auditory or visual - has a larger impact on global cognition in schizophrenia participants? 2) Are the cognitive gains in participants who underwent auditory or visual cognitive training associated with changes in symptoms and quality of life?

To answer these questions, we selected two groups of cognitive training exercises that were dynamically identical, with the exception of the sensory modality that was trained (i.e. auditory or visual). We compared the effect of the auditory versus visual neuroscience-informed cognitive training exercises in schizophrenia patients, using global cognition as the primary outcome. We evaluated the effect of training on the different cognitive domains affected in schizophrenia, clinical symptoms and quality of life as secondary outcomes.

2. Methods

2.1. Participants

Seventy-nine schizophrenia subjects were recruited from the day-hospital and outpatient clinic of the Institute of Psychiatry (IPUB) at the Federal University of Rio de Janeiro from September 2013 until December 2016.

Participants were self-referred or referred by a psychiatrist and signed a written consent form after being informed about the study procedures. We included participants between 18 and 60 years old with a schizophrenia or schizoaffective disorder diagnosis. Participants were clinically stable and had an outpatient status for at least one month before starting the training. Participants continued to take their medication as usual during the study and had no change in dosage >10%. At study entry, participants received an evaluation to determine eligibility that included the WAIS-IV Information and Matrix Reasoning IQ test (Wechsler, 2008), the Mini International Neuropsychiatry Interview Plus diagnostic interview (Sheehan et al., 1998) and the Simpson-Angus Scale (Simpson and Angus, 1970) for extrapyramidal symptoms. The Positive and Negative Syndrome Scale (PANSS) (Kay and Fiszbein, 1987), the Hamilton Depression and Anxiety rating scales and the World Health Organization Quality Of Life (WHO, 1998) questionnaires were administered at baseline and after 40 h of training. Participants were excluded if they were illiterate or had any history of another psychiatric diagnosis, intellectual disability or brain damage. We also excluded participants with extrapyramidal symptom scores above 19, IQ below 80 or any serious visual or auditory impairment that precluded their participation in the study.

The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2008. All procedures involving patients were approved by the Brazilian National Committee of Ethics in Research (12990013.0.0000.5263) and pre-registered at ClinicalTrials.gov (1R03TW009002-01).

2.2. Study design

This study used a randomized, double-blind, parallel design, where participants performed 40 h of auditory or visual cognitive training. Participants were randomized to either the auditory or the visual cognitive training using a hierarchical stratification by gender, age, IQ, years of education, diagnosis, PANSS scores, ethnicity, BMI, and recruitment site. Participants were blind to group assignment, and the research team performing the assessments had no access to participants' group assignment. Participants completed cognitive assessments at baseline and after 20 and 40 h of training. All the assessments took place in our research facilities. However, participants were given the choice to perform the online trainings in our research facilities or at home. They were asked to practice for 1 h per day, at least 3 days a week, which enabled the training to last between 2 and 4 months. The research team monitored training compliance weekly using the electronic records available online at the training platform.

2.3. Cognitive assessments

We assessed the seven cognitive domains defined by the Measurement and Treatment Research to Improve Cognition in Schizophrenia (MATRICS), using the MATRICS Consensus Cognitive Battery recommended tests (MCCB) available in Portuguese and the Cambridge Neuropsychological Test Automated Battery (CANTAB). We administered the MCCB Category Fluency (CF) and the CANTAB Reaction time (RTI) to assess speed of processing; the CANTAB Rapid Visual Processing (RVP) to assess attention; the Wechsler Adult Intelligence Scale (WAIS) III digit backward (DB) and the CANTAB Spatial Working Memory (SWM) to assess working memory; the MCCB Hopkins Verbal Learning Task (HVLT) to assess verbal learning; the MCCB Brief Visuospatial Memory Test (BVMT) to assess visual learning; the CANTAB Stocking Of Cambridge (SOC) to assess reasoning and problem-solving; and the MCCB Mayer-Salovey-Caruso Emotional Intelligence Test - Managing emotions (MSCEIT) to assess social cognition. A detailed explanation of these tests and their outcome measures can be found in previous literature (Massardo et al., 2015; Scoriels et al., 2012). We used parallel versions of the tests that were sensitive to

practice effects (i.e. HVLT, and BVMT) at 20 and 40 h of training. Global cognition was calculated from the z-scores of speed of processing, attention, working memory, verbal learning, visual learning, reasoning and problem-solving, and social cognition.

2.4. Cognitive training exercises

We designed the auditory and visual computerized cognitive trainings by selecting pairs of auditory and visual exercises from Posit Science, Inc. (www.brainhq.com) that had equivalent task dynamics, each in its perceptive modality. The auditory training included Sound Sweeps, Fine Tuning, Memory Grid, Auditory Ace, Syllable Stacks, and In the Know. The visual training included Visual Sweeps, Eye for Detail, Target Tracker, Card Shark, Juggle Factor, Recognition, and Face Facts. See Fig. S1 from the supplementary material for details. In both trainings, participants were driven to make progressively more accurate distinctions about the spectrotemporal fine structure of auditory or visual stimuli under conditions of increasing attention, memory and difficulty load (Fisher et al., 2009). Task difficulty increased systematically as performance improved and an algorithm was used to maintain individual rates of success in the exercises at around 80%. Correct performance was rewarded through entertaining visual and auditory embellishments and the accumulation of stars. To monitor the magnitude of progress in the training exercises, we evaluated subjects improvement in performance for each individual exercise using T-scores.

2.5. Planned analyses

The main outcome, i.e. the differential effect of the auditory and visual trainings on global cognition in schizophrenia participants, was analyzed using Intent-To-Treat (ITT) Repeated-Measures Analysis of Variance (RM-ANOVA) models, with last-observation-carried forward. This type of analysis allows incorporating data from the 79 participants, even if not all of them have finished the entire protocol study. Time (baseline, 20 h, and 40 h) was the within-subject factor and training type was the between-subject factor. Secondary outcomes, which included the individual cognitive domains, clinical measurements and quality of life, were also analyzed using RM-ANOVA. Normality and homogeneity of data distributions were tested using the Shapiro–Wilk and Levine tests. We calculated participants' neuropsychological tests z-scores using the formula $(x - \mu)/\sigma$ where x is the measure of interest, μ the mean and σ the standard deviation of the normative data of healthy participants available in the CANTAB and MCCB repository. Z-score change - or delta z-score - was also used where we wanted to show the difference between the final and initial z-scores (Delta z-score = Z-score after 40 h of training - z-score at baseline). t -tests, Chi squared test and Mann-Whitney U test were used to assess

differences in socio-demographic and cognitive data at baseline. Post hoc correlations were applied for statistically significant results from the main and secondary outcome measures' change scores (post 40 h minus baseline). We applied Bonferroni corrections for data with multiple dependent variables. Data were analyzed using SPSS.

3. Results

3.1. Participants

Two hundred and two subjects were screened and 123 (61%) were excluded because of ineligibility or lack of interest in the study. Thus, 79 schizophrenia subjects performed the first assessment and were randomized to either training program: 40 to the auditory training and 39 to the visual training. Fifty-three participants finished the training (Fig. S2). Participants were predominantly male, had an average of 42 years of age, 12 years of education and were chronically ill (all except two were taking antipsychotics over the course of the study). There were no significant socio-demographic and clinical differences between the two randomized groups (Table 1). Sixty-four participants trained in the laboratory and 15 participants trained at home, assessing the exercises on the internet. The training site did not affect the cognitive or the clinical outcomes when used as a covariate. The magnitudes of progress in the cognitive training exercises were similar between the auditory and the visual training groups.

3.2. Cognitive outcomes

There were no statistically significant differences in baseline cognition between the auditory training and the visual training groups. The Intent-To-Treat analysis on global cognition showed a significant group by time interaction: when compared to the auditory training group, participants who received the visual training showed significant improvements in global cognition from baseline to after 40 h of training ($F_{(77,2)} = 3.35, p = .04, d = 0.42$; Table 2). Furthermore, there was a significant main effect of time on global cognition ($F_{(77,2)} = 4.66, p = .01, d = 0.49$). Post hoc one-way RM-ANOVA for the two groups separately revealed significant improvement over time in global cognition in the visual training ($F_{(37,2)} = 6.02, p = .004, d = 0.80$), but not in the auditory training group (Fig. 1A and Table 2).

The analysis of individual cognitive domains, controlling for multiple comparisons, revealed a main effect of time for attention ($F_{(72,2)} = 6.80, p = .002, d = 0.65$), and reasoning and problem-solving ($F_{(73,2)} = 15.11, p = .000001, d = 0.97$), with no significant group by time interactions (Fig. 1B, 1C and Table 2). Post hoc one-way RM-ANOVA analysis, also controlling for multiple comparisons, revealed a significant effect of time on reasoning and problem-solving in the visual and auditory

Table 1
Baseline demographic and clinical information.

	All participants within inclusion criteria (n = 79)	Visual training group (n = 39)	Auditory training group (n = 40)	Differences between visual and auditory training groups (p-value)
Gender (men/women)	55/24	29/10	26/14	0.47
Age (mean (SD) ^b)	40 (12)	39 (12)	40 (12)	0.68
IQ (mean (SD))	101 (13)	100 (14)	100 (13)	0.96
Education (mean (SD))	12 (3)	12 (3)	12 (3)	0.69
Diagnostic (schizophrenia/schizoaffective)	72/7	36/3	36/4	0.69
Years of illness (mean (SD))	18 (11)	18 (11)	17 (10)	0.99
PANSS ^a score (mean (SD))	59 (14)	59 (16)	59 (12)	0.85
Antipsychotics generation (1st/2nd /multiple/none)	28/43/6/2	14/19/5/1	14/24/1/1	0.64
Anticholinergic/benzodiazepine/mood stabilizer/antidepressant	44/38/22/15	20/19/14/9	24/19/8/6	0.44
Chlorpromazine equivalent (mean (SD))	508 (417)	549 (371)	454 (429)	0.34
Anticholinergic equivalent (mean (SD))	3 (5)	6 (4)	5 (5)	0.54
Global progress in the exercises (mean t-scores (SD))	0.78 (0.70)	0.72 (0.59)	0.84 (0.85)	0.79

^a PANSS: Positive and Negative Syndrome Scale.

[‡] SD: Standard deviation.

Table 2
Effect of visual and auditory cognitive training on global cognition and the seven cognitive domains defined by MATRICS in schizophrenia subjects.

	Visual training (n = 39)					Auditory training (n = 40)					ANOVA comparing the 2 groups											
	Baseline (mean (SD))	20 h (mean (SD))	40 Hours (mean (SD))	F ANOVA (time)	p-value	Effect size (Cohen's d*)	Baseline (mean (SD))	20 h (mean (SD))	40 Hours (mean (SD))	F ANOVA (time)	p-value	Effect size (Cohen's d*)	F ANOVA (group)	p-value	Effect size (Cohen's d*)							
Global cognition	-0.75 (0.75)	-0.70 (0.69)	-0.52 (0.56)	6.02	0.004	0.80	-0.66 (0.83)	-0.58 (0.84)	-0.62 (0.86)	0.82	0.44	0.29	4.66	0.01	0.49	0.06	0.82	0.06	0.06	3.35	0.04	0.42
Speed of processing	-1.31 (2.18)	-1.36 (1.65)	-1.18 (1.67)	0.34	0.72	0.20	-1.19 (2.06)	-1.21 (1.74)	-1.53 (2.25)	0.34	0.71	0.20	0.03	0.97	0.04	0.01	0.94	0.02	0.02	0.66	0.52	0.20
Attention	-1.35 (1.44)	-1.03 (1.21)	-0.88 (1.20)	3.97	0.02	0.70	-1.26 (1.31)	-1.16 (1.21)	-0.94 (1.21)	2.96	0.06	0.63	6.80	0.002	0.65	0.01	0.93	0.02	0.02	0.21	0.81	0.11
Working memory	-0.30 (0.87)	-0.33 (0.99)	-0.24 (0.93)	1.63	0.20	0.46	-0.12 (1.19)	-0.05 (1.07)	-0.24 (0.96)	1.28	0.29	0.41	0.04	0.97	0.06	0.34	0.56	0.04	0.04	2.87	0.06	0.41
Verbal memory & learning	-0.21 (1.16)	-0.22 (1.08)	-0.21 (0.94)	0.25	0.78	0.20	-0.05 (1.05)	-0.08 (1.15)	-0.06 (1.12)	0.43	0.66	0.20	0.05	0.95	0.41	0.57	0.46	0.19	0.60	0.60	0.55	0.20
Visual memory & learning	-1.70 (1.49)	-1.47 (1.53)	-1.25 (1.32)	2.89	0.06	0.59	-1.32 (1.47)	-0.99 (1.34)	-1.30 (1.78)	1.14	0.33	0.41	2.44	0.09	0.41	0.96	0.33	0.29	1.41	0.66	0.52	0.20
Reasoning & problem solving	-0.17 (0.73)	0.08 (0.46)	0.18 (0.44)	5.75	0.005	0.84	0.02 (0.58)	0.11 (0.50)	0.29 (0.47)	12.74	0.00002	1.28	15.11	1.10 ⁻⁶	0.97	2.49	0.12	0.41	0.66	0.64	0.43	0.20
Social cognition	-0.26 (0.64)	N/A	-0.05 (0.71)	0.62	0.44	0.29	-0.38 (0.73)	N/A	-0.23 (0.71)	3.72	0.06	0.67	3.54	0.07	0.51	0.08	0.78	0.06	0.64	0.64	0.43	0.20

Means represent z-scores. **Bold** represents statistically significant results ($p < .05$). *Italic* represents trends for statistical significance ($0.05 < p < .1$). * Cohen's d converted from Eta².

training groups ($F_{(37,2)} = 5.75, p = .005, d = 0.84$ and $F_{(38,2)} = 12.74, p = .00002, d = 1.28$ respectively), and on attention in the visual training group ($F_{(37,2)} = 3.97, p = .02, d = 0.70$). These results showed medium to large effects in the direction of an improvement of the cognitive domains evaluated (Fig. 1 and Table 2).

Additionally, we performed an exploratory analysis of individual cognitive tests separated by the sensory modality used in the neuropsychological assessment: visual or auditory (Tables S1 and S2, respectively). The visual training group improved performance over time in two visual tests: the Stockings of Cambridge (that assessed reasoning and problem-solving), and the Rapid Visual Processing (that assessed attention) (Table S1). On the other hand, the auditory training group improved performance in the visual Stockings of Cambridge test and in the auditory Digit Backward test (that assessed working memory) (Table S1 and S2).

3.3. Symptoms and quality of life

The RM-ANOVA showed an interaction of time by type of training for PANSS negative symptoms scores ($F_{(75,1)} = 3.90, p = .05, d = 0.45$; Table S3 and Fig. 2C). Overall participants had a significant decrease of PANSS total score with time ($F_{(75,1)} = 14.39, p = .0003, d = 0.87$), and in particular in positive and general psychopathology symptoms ($F_{(75,1)} = 18.92, p < .0001, d = 1$ and $F_{(75,1)} = 8.07, p = .006, d = 0.65$ respectively), which reflected improvement in symptomatology (Table S3 and Fig. 2A-D).

Next, we performed a post hoc one-way RM-ANOVA of PANSS symptoms scores for the two groups separately, controlling for multiple comparisons. The visual training showed statistically significant decreases over time for total ($F_{(37,1)} = 12.91, p = .001, d = 1.17$), positive ($F_{(37,1)} = 13.38, p = .001, d = 1.19$), negative ($F_{(37,1)} = 4.19, p = .05, d = 0.66$), and general psychopathology ($F_{(37,1)} = 7.43, p = .01, d = 0.89$) symptoms, whereas the auditory training showed significant decrease in positive symptoms ($F_{(38,1)} = 6.4, p = .02, d = 0.82$) only (Table S3 and Fig. 2A, 2B, and 2D).

We found no significant effect of time, type of training or interaction of time by type of training in the quality of life (Table S3).

3.4. Association between changes in cognition and in clinical symptoms

We used Pearson correlations to test the association between the post-intervention changes in global cognition, attention, and reasoning and problem-solving, and changes in PANSS symptoms scores. Interestingly, we found a statistically significant association between the improvement in global cognition and the decrease in positive symptoms ($r = -0.37, p = .02$) in the schizophrenia subjects who performed 40 h of the visual training (Fig. 3A). We also found in the visual training group significant associations between the improvements in attention and the decreases in positive, negative and total symptoms ($r = -0.42, p = .01; r = -0.36, p = .03, \text{ and } r = -0.49, p = .002$ respectively, Fig. 3B-D). We did not observe these associations in the auditory training group.

4. Discussion

This study provides evidence that visual and auditory exercises of neuroscience-informed cognitive trainings are differentially effective at improving global cognition in schizophrenia subjects. On one hand, the visual training group notably improved in global cognition, with specific improvement observed in both low-order (attention) and high-order (reasoning and problem-solving) cognitive processes. On the other hand, the auditory training group improved over time in reasoning and problem-solving. However, only the training of visual processes was able to significantly improve global cognition.

The higher efficiency of the visual training may be explained by the predominance of the visual over the auditory processing system. Vision

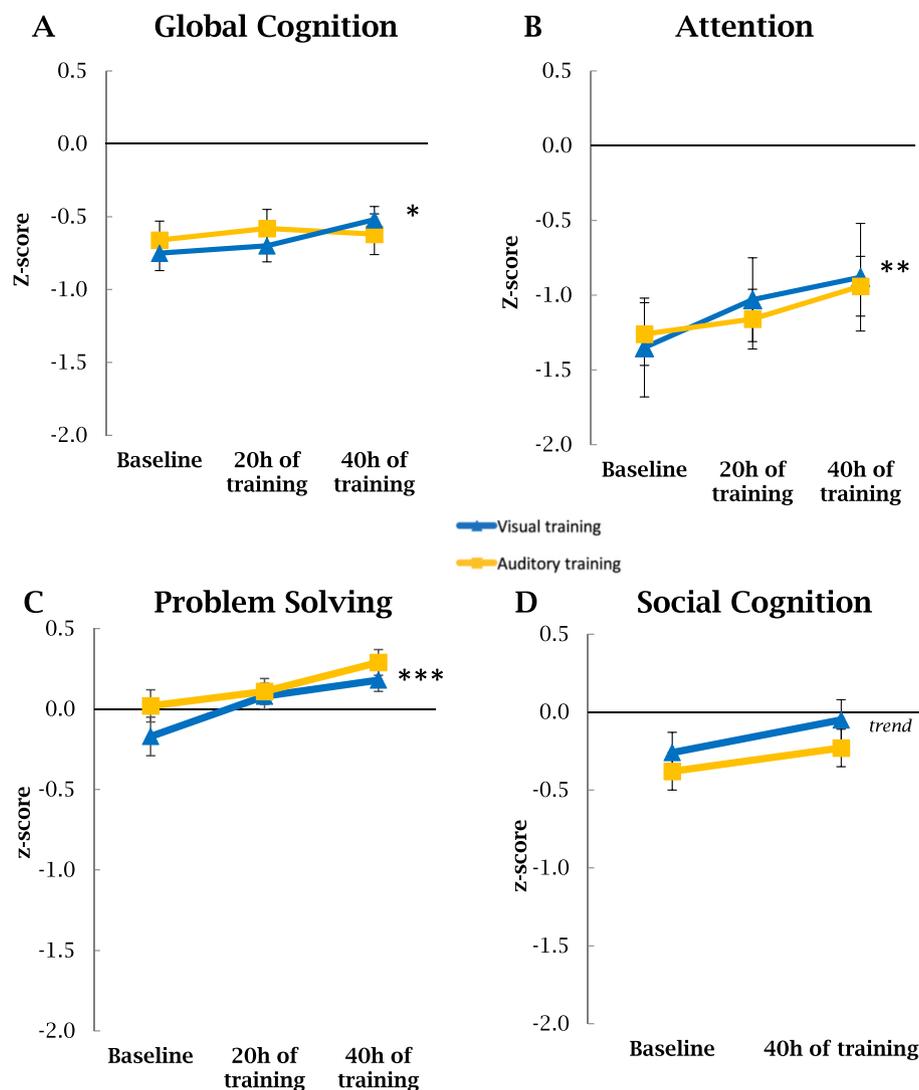


Fig. 1. Cognitive Performance with the Auditory and Visual training. **A.** Global cognition scores were significantly higher for patients who performed the visual training compared to the auditory training ($p = .04$). **B.** Attention scores increased significantly over time for patients who performed the visual training and at trend levels for the auditory training ($p = .02$ and $p = .06$ respectively). **C.** Reasoning and problem-solving scores increased significantly over time for patients who performed both types of trainings ($p = .005$ and $p = .00002$ respectively). **D.** Social cognition scores increased at the limit of significance over time for patients who performed the auditory training ($p = .06$). Error bars represent SEM. * $p < .05$, ** $p < .01$, *** $p < .001$.

plays a predominant role over audition, and we are generally more prone to attend to visual stimuli over auditory stimuli (Godfroy-Cooper et al., 2015). The McGurk effect is a great demonstration that the visual system takes over the auditory system, when conflicting information is observed between the two perceptual modalities (McGurk and Macdonald, 1976). Our results also suggested that the perceptive modality of the tests may be sensitive to the perceptive modality of the training. Indeed, both the visual and auditory trainings improved visual tests whereas only the auditory training improved auditory tests.

In a previous study, 50 h of auditory exercises of neuroscience-informed cognitive training improved verbal learning and memory and cognitive control in schizophrenia subjects that lasted 6 months beyond the intervention (Fisher et al., 2010). The addition of 50 more hours of visual and cognitive control training drove enduring gains also in speed of processing and global cognition (with a z-score change of 0.4). These findings emphasize that impairments in visual processes are important targets for improving global cognition in schizophrenia.

Yet, the moderate effect of the auditory training was surprising, given that previous studies using similar auditory training exercises had found significant improvement in several cognitive domains (Fisher et al., 2015, 2009). These studies have shown that cognition is

more impaired in the chronic phases than the early phases of schizophrenia, and that auditory training is more efficient in the chronic phases (Fisher et al., 2016) versus the early phases of schizophrenia (Fisher et al., 2015) or high risk for psychosis (Loewy et al., 2016). In our study, chronic schizophrenia participants in the auditory training group showed, at baseline, average performance in 4 cognitive domains, and low average performance in 3 domains, reflecting more intact cognition, and improvement in reasoning and problem solving only. This pattern of results indicates that less impairment at baseline leads to improvement in fewer cognitive domains after auditory training.

Feasibility of implementing the cognitive training program in schizophrenia subjects in different sites was verified in a multi-site, multinational study (Murthy et al., 2012). The results on cognitive outcomes were also mixed, and could be the result of the studied population, including the level of cognitive deficits, and differences in the measures used.

We found a significant decrease in symptoms of schizophrenia in both training groups. Interestingly, the visual training group showed significant improvements over time in a broad range of symptoms, which correlated significantly with improvements in attention. The literature has shown that impairments in attention are associated with

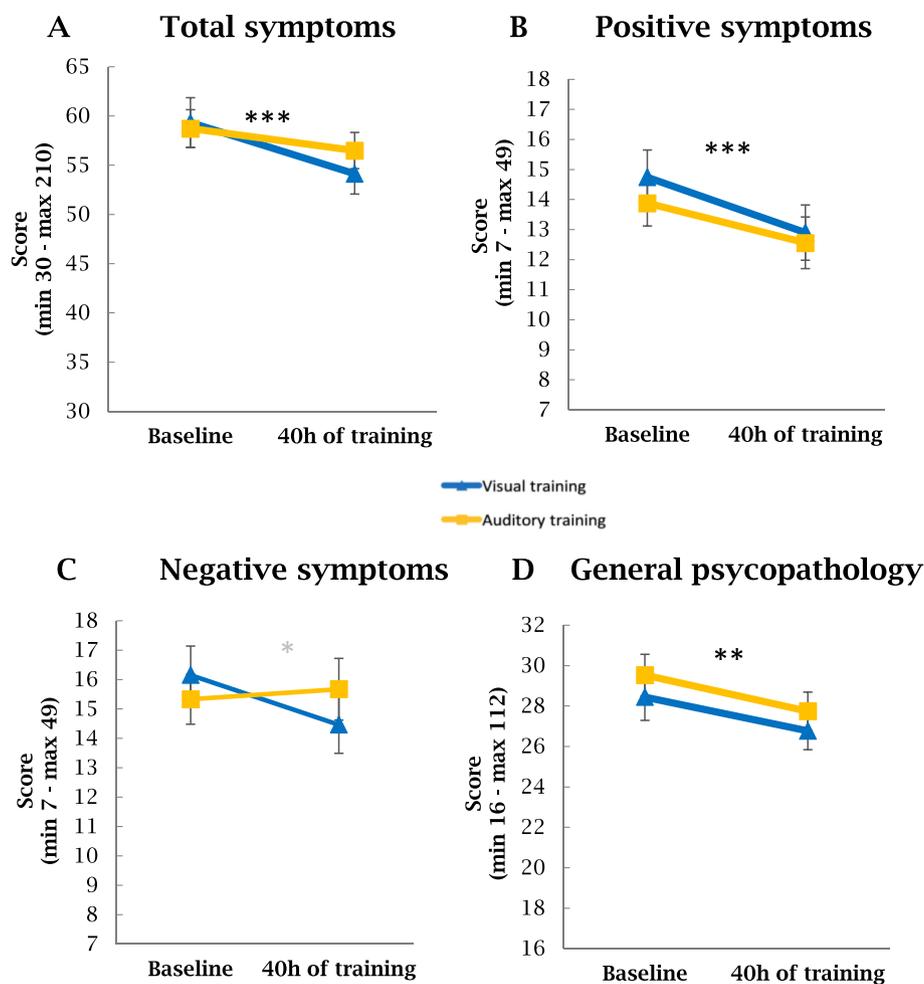


Fig. 2. PANSS symptoms evolution with 40 h of auditory or visual cognitive training in schizophrenia subjects. **A.** The overall effect of time was significant for PANSS total symptoms. Post hoc analyses showed that the visual training group (but not the auditory group) improved significantly over time for PANSS total symptoms ($p < .001$). **B.** The overall effect of time was significant for PANSS positive symptoms. Post hoc analyses showed that the visual training group (but not the auditory group) improved significantly over time for PANSS positive symptoms ($p < .001$). **C.** The interaction between time and type of training was significant for PANSS negative symptoms ($p < .05$). **D.** The overall effect of time was significant for PANSS general psychopathology. Post hoc analyses showed that the visual training group (but not the auditory group) improved significantly over time for general psychopathology ($p < .01$).

both negative and positive symptoms of schizophrenia (Berman et al., 1997; Mazza et al., 2013; Sanz et al., 2012) and the correlation found in our study suggests a pathway between the improvements in attention and the decrease in the clinical symptoms of schizophrenia. This hypothesis is supported by data from a meta-analysis that found that cognitive remediation is often related to small to moderate beneficial effects in symptoms (Cella et al., 2017). Therefore, we could speculate that even small changes in cognition, and more specifically in basic cognition (i.e. attention), can make a significant difference in patients' symptoms.

The sample size may have limited our study results, as we had a high attrition rate, with several gains in cognition very close to significance. One could also argue that the improvements we observed in cognition may be due to a test re-test effect. However, we evaluated a pilot group of schizophrenia subjects at baseline with the same battery of tests twice (one week apart), and there were no significant differences between the two sessions, demonstrating that a practice effect was improbable (unpublished data). A study by Pietrzak and colleagues also shows stable cognitive impairments in schizophrenia subjects when assessed within weeks or months (Pietrzak et al., 2009). Further, whenever possible, we used parallel versions of the same tests, and we avoided tests known to have high practice effects. Also, our study was designed such that the visual training and the auditory training would be the control of one another, thus it did not include an independent control group such as treatment as usual. However, we acknowledge

that it would have been beneficial to include such a group to assess the specificity of the gains in symptomatology.

This study shows for the first time that a neuroscience-informed computerized cognitive training can be applied effectively in the Brazilian population. We conclude that the visual training and the auditory training are differentially efficient at remediating cognitive deficits and symptoms of clinically stable schizophrenia subjects. A follow-up after twelve months of training is currently in progress to evaluate the long-term effects of the training in cognition and symptoms. Should the effects persist, we would contemplate implementing this type of intervention for patients with schizophrenia in our clinical facilities.

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.schres.2020.05.017>.

Role of the funding source

This work was supported by the National Institutes of Health – Fogarty International Center (Grant R03TW009002 to SV and RP); Fundação de Amparo a Pesquisa do Estado do Rio de Janeiro (FAPERJ) (Grant E-26/110.305/2014 to RP) and Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq) (Grant 400455/2012-9 to RP). RP is an Atlantic Fellow of the Global Brain Health Institute. LG, LM, SK, AG, PR, FT and CN were supported by fellowships from FAPERJ and CNPq. The cognitive training software used in this study and all technical support were provided to us free of charge by Posit Science, Inc.

Contributors

LS, LTG, AIF, NG, BJS, MTC, MF, SV, and RP formulated the research questions and designed the study; LS, LTG, LGCM, SK, ALDVG, PVSR, FMT, CN carried the research out; LS,

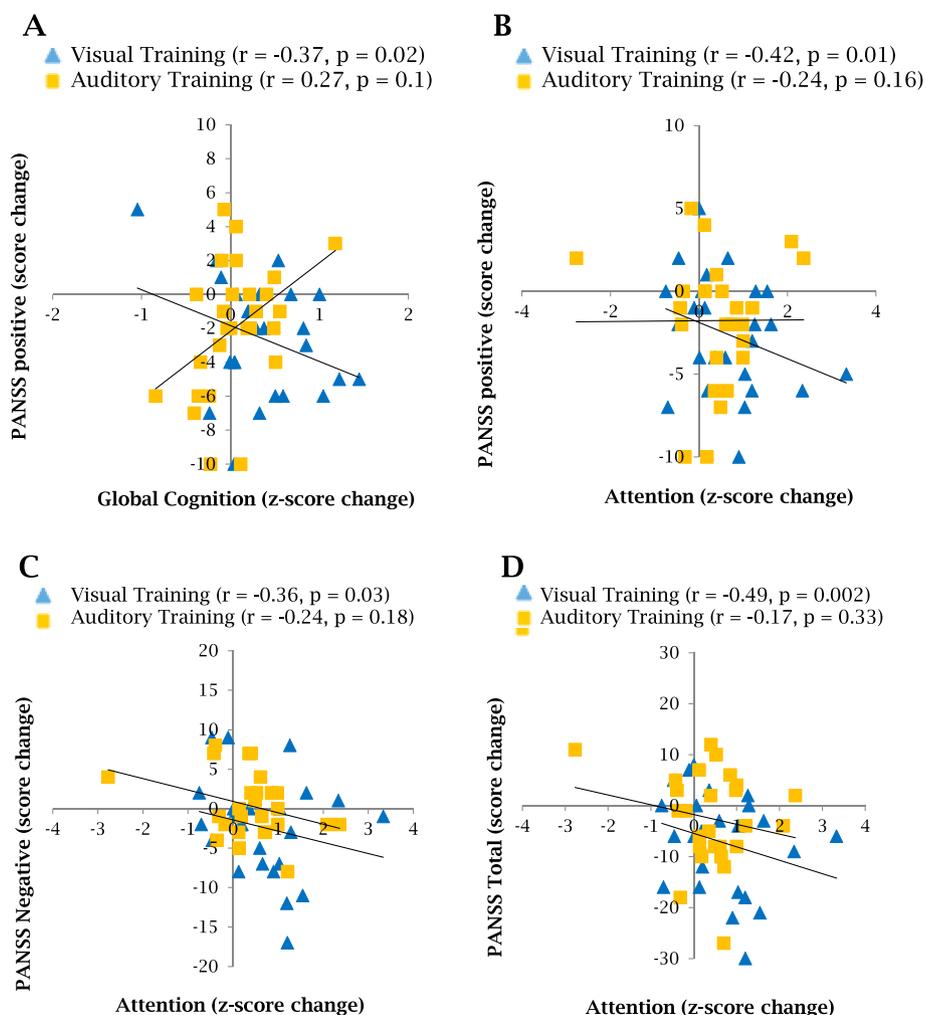


Fig. 3. Association between PANSS symptoms and cognitive changes after 40 h of auditory or visual cognitive training in schizophrenia subjects. **A.** Pearson correlation shows a significant association between global cognition and PANSS positive symptoms improvements for subjects who performed the visual training. **B.** Pearson correlation shows a significant association between attention and PANSS positive symptoms improvements for subjects who performed the visual training. **C.** Pearson correlation shows a significant association between attention and PANSS negative symptoms improvements for subjects who performed the visual training. **D.** Pearson correlation shows a significant association between attention and PANSS total symptoms improvements for subjects who performed the visual training.

LTG, LGCM, SK, ALDVG, PVS, and RP analyzed the data and all authors contributed to writing the article.

Declaration of competing interest

Dr. Panizzutti is the founder of NeuroForma LTDA, a company with a financial interest in cognitive training. The remaining authors have no conflict of interest to disclose.

Acknowledgements

The authors gratefully acknowledge participants and their families; and thank Ana Claudia Rangel and Andrea Fantinatti for the excellent technical support.

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