Definition and refinement of the 7q36.3 duplication region associated with schizophrenia

Branko Aleksic1*, Itaru Kushima1*, Tamae Ohye2*, Masashi Ikeda3, Shohko Kunimoto1, Yukako Nakamura1, Akira Yoshimi1, Takayoshi Koide1, Shuji Iritani1, Hiroki Kurahashi2, Nakao Iwata3 & Norio Ozaki1

1Nagoya University, Graduate School of Medicine, Department of Psychiatry, 2Fujita Health University, Division of Molecular Genetics, Institute for Comprehensive Medical Science, 3Fujita Health University, Department of Psychiatry.

Using a very high-resolution oligonucleotide array for copy number variant (CNV) screening of samples comprising schizophrenic patients, we detected a novel CNV within the critical region (NCBI36/hg18, Chr7: 158,630,410–158,719,410) previously shown to be associated with schizophrenia. We investigated the association between the novel CNV identified in the current study and schizophrenia. Three independent samples were used: (1) Screening set, 300 Japanese schizophrenic patients (53.28 ± 14.66 years); (2) Confirmation set, 531 schizophrenic patients (46.03 ± 12.15 years); and (3) 711 healthy controls (47.12 ± 11.03 years). All subjects enrolled in the study were Japanese. Chromosomal position was determined using fluorescence in situ hybridization. We identified a novel duplication within the region associated with schizophrenia identified on 7q36.3 that is adjacent to VIPR2 and is not associated with schizophrenia. In the Japanese population, the 35-kb region that harbors the common, novel CNV should be excluded from the region associated with schizophrenia on 7q36.3.

Schizophrenia is a chronic, debilitating illness characterized by impairments in cognition, affect and behaviour1. The Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM IV-TR)2 defines the essential features of schizophrenia as a mixture of characteristic signs and symptoms (both positive and negative) that have been present for a significant portion of time during a 1-month period (or for a shorter time if successfully treated), with some signs of the disorder persisting for at least 6 months. In this regard, positive refers to the presence of active symptoms including delusions and hallucinations. Negative symptoms refer to a loss, typically of emotions, speech, or motivation. Schizophrenic disorders exist on a continuum from mild to severe. The DSM IV-TR2 recognizes a number of different types, which include disorganized, catatonic, paranoid, schizophreniform, residual, schizoaffective, undifferentiated and not otherwise specified3. Schizophrenia is a relatively common disorder, with a lifetime prevalence of about 1%1. Although the overall sex ratio is almost equal, males tend to have an earlier onset than females, a finding accounted for by the later age of onset in those females who lack a family history of the disease4. Family history is the most important risk factor for schizophrenia, consistent with a genetic contribution to its etiology4. However, as with most mental disorders, the origins and mechanisms of schizophrenia are not fully understood.

Genetic factors influence human disorders by determining disease susceptibility or resistance5. Therefore, genetic studies can help pinpoint the exact molecular mechanism of a disease. Recent successes in the genetic mapping and molecular mechanism of the Mendelian traits have been remarkable, owing to the development of genome wide screening techniques6. As such, attention has been gradually shifting towards more complex, common, genetic disorders and traits that involve multiple genes and environmental effects, such as celiac disease7, diabetes8, rheumatoid arthritis9 and psychiatric disorders10. In this context, recurrent microdeletions at 1q21.111, 15q13.312, and 15q11.212, microduplications at 16p11.213, and copy number variations (CNVs) at other genomic loci14 have been shown to be associated with schizophrenia in large cohorts examined by CNV analyses and other molecular studies. Furthermore, duplication at chromosome 7q36.3, encompassing VIPR2, was implicated in schizophrenia for the first time in a recent report15. In a specific genome-wide association study of 8,290 patients with schizophrenia performed by Vacic et al.15, the authors found that 0.35% of these patients carry rare CNVs in the chromosomal locus 7q36.3. In contrast, these microduplications were much less frequent.

OPEN

SUBJECT AREAS:
SCHIZOPHRENIA
MEDICAL GENETICS
MOLECULAR BIOLOGY
GENETICS OF THE NERVOUS SYSTEM

Received 16 January 2013
Accepted 20 August 2013
Published 4 September 2013

Correspondence and requests for materials should be addressed to B.A. (branko@med.nagoya-u.ac.jp)

* These authors contributed equally to this work.

SCIENTIFIC REPORTS | 3 : 2587 | DOI: 10.1038/srep02587
among the 7,431 healthy controls. All variants overlap with \textit{VIPR2} or lie within the noncoding subtelomeric region, <89 kb (NCBI36/hg18, Chr7: 158,630,410–158,719,410) from the transcriptional start site of \textit{VIPR2}. This gene encodes the vasoactive intestinal peptide (VIP) receptor VPAC2, which is a G-protein-coupled receptor that is expressed in the suprachiasmatic nucleus, hippocampus, amygdala and hypothalamus\textsuperscript{16}. VPAC2 binds VIP, activates cyclic AMP (cAMP)-signalling and PKA, regulates synaptic transmission in the hippocampus, and promotes the proliferation of neural progenitor cells in the dentate gyrus\textsuperscript{17}. Moreover, it has been shown that alteration in synaptic plasticity of hippocampal neurons may contribute to the symptoms observed in schizophrenic patients\textsuperscript{18}. The aforementioned lines of evidence provide support for the role of \textit{VIPR2} as a candidate gene for schizophrenia from a biological point of view.

In the present study, by using a very high-resolution oligonucleotide array for CNV screening of samples from schizophrenic patients, we were able to detect CNV within the critical region (NCBI36/hg18, Chr7: 158,630,410–158,719,410) on 7q36.3 that was shown to be associated with schizophrenia by the Vacic et al. study\textsuperscript{15}. Thus, the goal of the present study was to follow-up on the novel CNV that was previously detected in schizophrenic patients and further investigate any association between this CNV and schizophrenia.

**Results**

In the present study, we detected a smaller (35 kb) duplication (NCBI36/hg18, Chr7: 158,658,128–158,693,128) within the critical region identified by Vacic et al.\textsuperscript{15} (Figure 1). The observed frequency of the CNVs was ~2% and we did not detect any statistically significant difference between the patients and controls (Table 1). There was a 100% concordance rate between the custom NimbleGen 12 × 135,000 CGH arrays and the NimbleGen 3 × 720,000 CGH arrays or custom TaqMan copy number assay for the detection of the smaller (35 kb) duplication (NCBI36/hg18, Chr7: 158,658,128–158,693,128) within the critical region (NCBI36/hg18, Chr7: 158,630,410–158,719,410) previously identified by Vacic et al.\textsuperscript{15}.

In metaphase cells, all duplication-specific FISH signals localize to the subtelomeric region of 7q, confirming that the duplications lie adjacent to each other in the 7q36.3 region (Figure 2). In addition, NS102 exhibited two signals, one of which had a higher intensity compared to that of the other. This suggests that there is unilocus duplication in the \textit{VIPR2} promoter region. During the orientation analysis, an amplicon was detected by electrophoresis only in samples with duplication, which indicates that there is a head-to-tail orientation of the repeated DNA fragment (Supplementary Figure 1). Additionally, sequence analysis of the repeat junction revealed that all samples with duplication shared exactly the same sequence within the junction region (Supplementary Figure 2). Based on the

![Figure 1 | High-resolution aCGH data.](https://www.nature.com/scientificreports/)
Figure 2 | Tandem duplications of 7q36.3 confirmed in two patients by fluorescence in situ hybridization (FISH). 7p-green (arrowheads) and 7q-red (arrows, CNV specific). Left NS102 (3 copies); Right NS004 (2 copies). Cytogenetic confirmation was obtained for two samples with and without duplication of VIPR2. Probes for duplicated region were produced by long range PCR. The subtelomeric probe, 7p-green (Abbott Molecular), was used as a reference. Hybridizations were performed according to the manufacturer’s protocols.

Table 1 | Frequency distribution (confirmation set)

| Set      | Sample size | 0  | 1  | 2  | 3  | 4  | >2 copies | P-Value |
|----------|-------------|----|----|----|----|----|-----------|---------|
| Cases    | 531         | 0  | 0  | 520 | 0  | 0  | 10        | 0.96    |
| Controls | 711         | 0  | 0  | 696 | 0  | 0  | 13        |         |

The main limitation of the current study was that we examined a much smaller number of samples as compared to the Vacic et al. study\(^5\). It is of note that the frequency of the common 35-kb CNV detected in the current study was 2%, and thus with our current sample size of 300 schizophrenic patients, it was large enough to capture the variation. Regarding the individual with 4 copies, we do not have any data indicating whether the individual is a homozygote of duplication or is a carrier of triplication. This point should be considered as another limitation of the current study. The results of the current study do suggest that in case of a duplication event on 7q36.3, the relevant region is not the VIPR2 promoter (as has been suggested by Vacic et al.\(^5\)), but rather suggest that it is the VIPR2 gene region. In addition, the 2-bp microhomology in the promoter region of VIPR2 may be associated with the relative meiotic instability of the region harboring the common CNV that is adjacent to the VIPR2 gene\(^5\). This in turn may give rise to the larger VIPR2 duplications that were shown to be associated with schizophrenia with an odds ratio of 4.0\(^4\).

It is interesting that our findings demonstrated that CNV was detected in our study in contrast to the previous reports by both Vacic et al.\(^5\) and Beri et al.\(^6\). Moreover, CNV has not been listed in the database of genomic variants\(^27\). Although the CNV detected by our group may be specific to Japanese populations, further studies should be undertaken to ensure comprehensive characterization of the region surrounding the VIPR2 gene. In addition, to address the question regarding the origin of the CNV detected in the current study, it is necessary to perform family analysis of the carriers and determine whether CNV is a de novo event, or if it is transmitted from the parents. In conclusion, the 35-kb region that harbors the common CNV in the Japanese population should be excluded from the region of the association peak in the schizophrenia group reported in the Vacic et al. study\(^5\).

Methods

Three independent samples were used in the current study: (1) Screening set, 300 Japanese patients suffering from schizophrenia (53.28 ± 14.66 years); (2) Confirmation set, 531 patients suffering from schizophrenia (46.03 ± 12.15 years); and (3) 711 adult control subjects (47.12 ± 11.03 years). All schizophrenic patients met the current DSM IV-TR criteria, which was reflected by consensus diagnosis of two experienced psychiatrists. Prior to inclusion in the control set, subjects were screened on the basis of a brief diagnostic interview. Detailed characterization and psychiatric assessment of the subjects is available elsewhere\(^27\). All subjects enrolled in the study were Japanese and provided written informed consent prior to the study. Venous blood was drawn from each subject and genomic DNA was extracted according to the standard phenol/chloroform method. Comparative genomic hybridization of DNA was performed using the high-resolution NimbleGen (Roche NimbleGen, Inc., U.S.) CGH array (3 × 720,000 or 12 × 135,000). Labeling and hybridization of patient (test) and sex-matched commercial (Promega Corporation, U.S.) reference DNA was performed according to the manufacturer’s protocols. Test and reference DNA were labeled by Cy3- and Cy5-labeled random primers, respectively, and were combined and hybridized to the array for 40–72 h. Arrays were washed in four steps, as indicated in the protocol. Two-color scanning was performed using a NimbleGen MS 200 microarray scanner. Acquisition of the microarray images was performed with NimbleGen MS 200 software. Data extraction, analysis and visualization were done using NimbleScan version 2.4 software. CNV calling was performed using NEXUS software. The FAST2 Segmentation Algorithm, a Hidden Markov Model (HMM) based approach, was used to make copy number calls. The FAST2 algorithm, unlike other common HMM methods for copy number estimation, does not aim to estimate the copy number state at each probe, but uses many states to cover more possibilities, such as mosaic events. These state values are then used to make calls based on a log ratio threshold. The significance threshold for
segmentation was set at 10⁻³ and also required a minimum of three probes per segment. The log ratio thresholds for single copy gain and single copy loss were set at 0.3 and -0.3, respectively. The log ratio thresholds for the gain of two or more copies and homozygous loss were set at 0.9 and -0.9 respectively.

Custom TaqMan copy number assay was specifically designed to interrogate a duplication region (NCBI36/hg18, Chr7: 158,630,410–158,719,410) without interspersed repeats, low complexity or a homologous DNA sequence. A TaqMan copy number assay for RNase P was used as a reference. Experiments were carried out on four technical replicates according to the manufacturer’s protocol. CNV typing of the screening sample was performed using Roche NimbleGen, Inc. CGH array 3 × 720,000, while confirmation of the sample was performed using the TaqMan copy number assay. Sixteen randomly selected duplication events (both in the screening and confirmation samples) were validated using custom NimbleGen 12 × 135,000 CGH arrays (Roche NimbleGen, Inc., U.S.) covering the region (NCBI36/hg18, Chr7: 158,630,410–158,719,410) implicated in the Vacci et al. study14, with an average of one probe per 500 bp. P values derived from association analysis were based on Fisher’s exact test.

We performed PCR based analysis to determine the orientation of the detected duplications. We designed forward and reversed primers to align with the region of the duplication junction (F: 5’-TGTTGAGTCCTCCAGAGGCAC-3’, R: 5’-CATCTCCAGGCTAGTGCATC-3’) (Supplementary Figure 1). Cytogenetic confirmation was obtained for two samples with and without duplication of VIPR2. Probes for duplicated region were produced by long range PCR (NCBI36/hg18, Chr7: 158,658,128–158,693,128). Subtelomeric probe, 7p-green (Abbott Molecular, U.S.), was used as a reference. Hybridizations were performed according to the manufacturer’s protocols.

Haplotypes were estimated using the statistical software package PHASE version 3.4.1 (http://www.stat.washington.edu/ph2nph/). This program is based on a Bayesian statistical method using coalescent-based models that infers phases at loci from unphased genotype data for a sample of unrelated individuals20. The algorithm uses a flexible model for the decay of linkage disequilibrium with distance and explicitly incorporates an assumption about the recombination rate variation. PHASE uses Gibbs sampling, a Markov-Chain Monte Carlo algorithm for the estimation of the posterior distribution. Hence, the individual haplotype can be estimated from the posterior distribution by choosing the most likely haplotype reconstruction for each individual. Using the extension for unrelated individuals, we used the default settings to infer the haplotypes from the genotype data of the 8 SNPs (Supplementary Table 1) surrounding the duplication in sample that comprised 517 subjects (7 with a structural variant detected in the current study). Estimates of the sample haplotype frequencies together with their standard deviation, a list of the most likely pairs of haplotypes for each individual together with their probability, and the estimates of recombination parameters in the region, were calculated using the same software.

Acknowledgements
Funding for this study was provided by research grants from the Ministry of Education, Culture, Sports and Technology of Japan; the Ministry of Health, Labor and Welfare of Japan; Grant-in-Aid for “Integrated research on neuropsychiatric disorders” carried out under the Strategic Research Program for Brain Sciences by the Ministry of Education, Culture, Sports, Science and Technology of Japan; Grant-in-Aid for Scientific Research on Innovative Areas (Comprehensive Brain Science Network) from the Ministry of Education, Science, Sports and Culture of Japan; The Academic Frontier Project for Private Universities, Comparative Cognitive Science Institutes, Meijo University; the Core Research for Evolutional Science and Technology and SENSHIN Medical Research.

Author contributions
B.A., I.K.O. and N.O. designed the study and wrote the protocol. B.A., I.K.O., T.O., N.I., H.K. and N.O. performed the literature review. B.A., I.K.O. and T.O. made and managed the sample database. B.A., I.K.O., T.O., M.I., S.K., Y.N., A.Y., T.K., S.I., H.K., N.I. and N.O. collected and managed the genome samples. B.A., I.K.O., T.O., M.I., S.K., Y.N., A.Y., T.K. and S.I. conducted the statistical analysis. B.A., I.K.O., T.O., M.I., S.K., Y.N., A.Y., T.K., S.I., H.K., N.I. and N.O. interpreted and discussed the results. B.A., I.K.O., T.O., M.I., H.K., N.I. and N.O. wrote the manuscript and edited the final manuscript.

Additional information
Supplementary information accompanies this paper at http://www.nature.com/scientificreports

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Alekseev, B. et al. Discovery and refinement of the 7q36.3 duplication region associated with schizophrenia. Sci. Rep. 3, 2587; DOI:10.1038/srep02587 (2013). This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivs 3.0 Unported license. To view a copy of this license, visit http://creativecommons.org/licenses/by-nc-nd/3.0/