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Citation for published version:

Digital Object Identifier (DOI):
10.1093/schbul/sbq135

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published In:
Schizophrenia Bulletin

Publisher Rights Statement:
The Author 2010. Europe PMC open access link

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DISC1 in Schizophrenia: Genetic Mouse Models and Human Genomic Imaging

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Schizophrenia and related disorders have a major genetic component. Several large-scale studies have uncovered a number of possible candidate genes, but these have yet to be consistently replicated and their underlying biological function remains elusive. One exception is ‘Disrupted in schizophrenia 1’ (\textit{DISC1}), a gene locus originally identified in a large Scottish family, showing a heavy burden of major mental illnesses associated with a balanced t(1;11)(q42.1;q14.3) chromosome translocation. Substantial genetic and biological research on \textit{DISC1} has been reported in the intervening 10 years: \textit{DISC1} is now recognized as a genetic risk factor for a spectrum of psychiatric disorders and \textit{DISC1} impacts on many aspects of central nervous system (CNS) function, including neurodevelopment, neurosignaling, and synaptic functioning. Evidence has emerged from genetic studies showing a relationship between \textit{DISC1} and quantitative traits, including working memory, cognitive aging, gray matter volume in the prefrontal cortex, and abnormalities in hippocampal structures and function. \textit{DISC1} interacts with numerous proteins also involved in neuronal migration, neurite outgrowth, cytoskeletal modulation, and signal transduction, some of which have been reported as independent genetic susceptibility factors for psychiatric morbidity. Here, we focus on the growing literature relating genetic variation in the \textit{DISC1} pathway to functional and structural studies of the brain in humans and in the mouse.

\textit{Key words:} genetics/schizophrenia/mouse models/neurodevelopment/neuroimaging

Introduction

Schizophrenia, like many other common diseases, is complex and multifactorial, with contributions from multiple susceptibility genes, epigenetic, stochastic, and environmental factors.\textsuperscript{1} Family and twin studies have shown that genetic factors play a major role in the development of schizophrenia, and a number of candidate risk genes have associated with schizophrenia\textsuperscript{2–4} of which \textit{DTNBP1, NRG1, and DISC1} are particularly relevant to this review. \textit{DISC1} has captured much attention, not only being associated with schizophrenia but also predisposing individuals to a wide range of major mental disorders. Through genetics, cell biology, animal modeling, and neuroimaging, the \textit{DISC1} pathway is emerging as a pivotal mediator of quantitative and pathological brain dysfunction.

The Scottish \textit{DISC1} Family and Further Independent Genetic Evidence

\textit{DISC1} was first identified in a large Scottish family with a high loading of major mental illness.\textsuperscript{5} Specifically, the t(1; 11)(q42.1;q14.3) translocation allele of the \textit{DISC1} gene associates with schizophrenia, bipolar affective disorder, and recurrent major depression\textsuperscript{6} (log of the odds ratio = 7.1) in this large family. The first independent evidence for the involvement of the \textit{DISC1} locus in psychiatric illness came from studies in the Finnish population\textsuperscript{7,8} in which evidence for linkage of schizophrenia and schizoaffective disorder to the 1q32.2–q41 region, proximal to the \textit{DISC1} gene, was reported. Further population studies of Taiwan, Scotland, and Britain/Iceland have also shown linkage of chromosome 1q32–42 to major psychiatric illness.\textsuperscript{8}

\textit{DISC1} Genetic Mouse Models

Although it is difficult to model a psychiatric disorder like schizophrenia in animals (eg, hallucinations and delusions are likely to be human specific), anatomical, behavioral, and cognitive characteristics have been described in patients, and it is feasible to study these in...
There has been much debate about the functional consequence of the t(1;11) on DISC1. It may be simply the effect of haploinsufficiency (half normal levels of DISC1 protein) from conception onwards, but others have proposed a dominant negative effect from expressing a hypothetical truncated DISC1 protein. On this premise, several transgenic mouse models expressing an N-terminal or C-terminal truncated protein have been generated. Reduced cortical thickness and partial agenesis of the corpus callosum have been observed in transgenic mice expressing truncated DISC1 generated using a bacterial yeast artificial chromosome vector. Enlarged lateral and third ventricles have been shown by magnetic resonance imaging (MRI), in 2 other types of transgenic mice in which human DISC1 cDNA transgenes have been expressed postnatally in forebrain cells. These are similar to features seen in imaging studies of chronic schizophrenic patients. Furthermore, the L100P mutant mice showed behaviors, including profound PPI and latent inhibition (LI) deficits, akin to those seen in some with schizophrenia, and these could be reversed by antipsychotic treatment. In contrast, the Q31L mutant mice showed LI deficits, but only modest PPI deficits and exhibited a more depressive-like behavior, partially reversed by the antidepressant bupropion. Thus, as in humans, DISC1 mutations in mice can lead to 2 very different phenotypes. Despite the fact that none of these mouse models exactly matches all features characteristically seen in human patients, their phenotypes are consistent with human genetic studies, suggesting extensive locus and allelic genetic heterogeneity.

DISC1 Imaging Genetics in Humans

**DISC1 in Schizophrenia**

**DISC1 Imaging Genetics in Humans**

**DISC1 Expression, Cognitive, and Behavioral Phenotypes Reported to be Associated With Genetic Variants of DISC1 in Mice.** These are representative of endophenotypes of major psychiatric illness and likely to be due to widespread DISC1 expression.
of the hippocampus. For example, the Ser allele has been associated with reduced hippocampal gray matter volume and altered engagement of the hippocampus during cognitive tasks (see figure 2). Furthermore, the Cys704 allele has been associated with major depressive disorder, a reduction in gray matter volume in the cingulate cortex, and with decreased fractional anisotropy in prefrontal white matter.

There is robust evidence for prefrontal cortex (PFC) abnormalities in psychosis. During verbal fluency tasks, a classic test of prefrontal function, among healthy volunteers, activation of the PFC in Ser704/Ser704 subjects was greater, especially of the left hemisphere, a finding that can be interpreted as a manifestation of less efficient prefrontal function, with more activation needed to achieve the same behavioral output. These findings are consistent with reports of association between Ser704, schizophrenia, and impaired declarative memory in schizophrenic patients.

A Japanese MRI study recently reported that healthy carriers of Cys704 showed significantly larger volumes of the medial superior frontal gyrus and short insular cortex than the Ser704 homozygotes, whereas schizophrenic patients who were carriers of Cys704 tended to have a smaller supramarginal gyrus than the Ser704 homozygotes. There was also noted to be a significant correlation between the right medial superior frontal gyrus volume and daily dose of antipsychotic medication in schizophrenic patients who were Ser704 homozygotes.

Another study investigated the relationship between the DISC1 Leu607Phe polymorphism and prefrontal gray matter volumes using MRI. Among patients and healthy volunteers, carriers of Phe607 had significantly less gray matter in the superior frontal gyrus and anterior cingulate gyrus compared with Leu/Leu homozygotes. Patients who were Phe607 carriers also showed greater severity of positive symptoms suggesting that this polymorphism is clinically significant in terms of psychopathology, as suggested by others.

Raznahan et al have recently shown that the developmental rate of cortical thinning (CT) is dependent on DISC1 genotype. Consistent with other results, Phe607 carriers showed attenuated rates of CT compared with Leu/Leu homozygotes in bilateral superior frontal and left angular gyri between the ages of 9 and 23 years. The Ser704 homozygotes in turn showed accelerated CT compared with Cys704 carriers in the left anterior cingulate and temporal cortices. Interestingly, the Ser704Cys SNP affects DISC1-NDEL1 association, which also has a well-established role in cortical maturation. The next section of this review focuses on DISC1 interacting partners that we have genetic data from mouse models and from imaging genetic studies.

**DISC1 Interacting Partners: Structural and Anatomical Changes in Genetic Mouse Models and Imaging Genetic Studies**

DISC1 interacts with numerous neuronal proteins, reflecting the diversity of potential roles that it plays in brain function. These interactors can be broadly classified into...
Neuregulin-1 and Epidermal Growth Factor Receptor B4

Neuregulin-1 (NRG1) is a large multiexon gene on chromosome 8p. It produces multiple isoforms grouped into types I–VI according to their 5′ exon, through alternate promoters and splicing. NRG1 has multiple roles in the central nervous system (CNS) and in processes of possible relevance to schizophrenia including: myelination, glial cell development, migration of radial glial cells during cortical development, neuronal plasticity, development of GABAergic interneurons, and expression of dopamine and serotonin receptors and monoamine transporters. Complete knockout of NRG1 expression is lethal, standing potential roles for NRG1 in schizophrenia.

Our group in Edinburgh was the first to relate NRG1 status to brain imaging measures. Among 79 subjects from the Edinburgh High Risk Study, none of whom were receiving treatment at the time of the study, there was a highly significant association between the risk allele at SNP8NRG243177 and psychotic symptoms, across the course of the study. During the Hayling sentence completion task (known to activate frontal and temporal brain regions), functional magnetic resonance imaging (fMRI) revealed decreased activation of right medial PFC in the subjects with the risk (T/T) genotype at SNP8NRG243177 compared with those without the risk allele in the contrast of sentence completion vs rest. We extended this study to also investigate the National Adult Reading Test, a measure of premorbid intelligence quotient (IQ), and found that the T/T group had a significantly decreased IQ compared with the C/T and C/C groups. A similar pattern of deficits was seen using the Wechsler Adult Intelligence Scale, a measure of current IQ, although this effect failed to reach statistical significance.

In a follow on study, subjects with the risk-associated SNP8NRG243177 T/T genotype showed reduced white matter density in the anterior limb of the internal capsule and evidence of reduced structural connectivity in the same region using DTI, suggesting that NRG1 may increase susceptibility to psychosis by altering connections between PFC and other brain regions. These findings further implicate white matter disconnectivity in psychosis. Our group also investigated DTI tractography of NRG1 and white matter integrity and showed that the 2 risk-associated variants of NRG1 (T allele at SNP8NRG243177 and C allele at SNP8NRG221533) are associated with reduced white matter integrity in the anterior thalamic radiation, which connects the dorsomedial and anterior thalamic nuclei with the PFC. If this connection is indeed disrupted by psychosis-related NRG1 variants, as shown here, this would provide a possible explanation of the abnormal prefrontal function and connectivity often reported in psychosis.

Collectively, these findings suggest that NRG1, Erbs, AKT1, and DISC1 connect in a common pathway that regulates neurodevelopment, which may also contribute to schizophrenia susceptibility. Impaired AKT1 signaling has also been associated with schizophrenia. This study, therefore, brings together 3 prominent schizophrenia candidate genes (NRG1, AKT1, and DISC1).

The large number of NRG1 signaling mechanisms and isoforms parallels the many roles it has in the CNS, many of which could be involved in schizophrenia. A 5′ haplotype (HapICE) in the NRG1 gene has been identified that associates with schizophrenia, a finding supported by a number of subsequent studies. The NRG1 risk–associated variants are primarily in noncoding intronic and promoter regions, leading to the suggestion that the causative variant may be mediated by altering gene expression rather than protein levels. This hypothesis has been strengthened by a study, which demonstrated that genetic variation at SNP8NRG243177 from HapICE is associated with altered NRG1 expression.

Mouse studies have also been informative in understanding potential roles for NRG1 in schizophrenia. Complete knockout of NRG1 expression is lethal, whereas animals with heterozygous disruption of NRG1 or of Erb4 receptor function have been shown to have reduced numbers of N-methyl-D-aspartate acid receptors and increased levels of dopamine receptors in the PFC, as well as exhibiting behavioral abnormalities consistent with schizophrenia.

Pericentriolar Material 1

Pericentriolar material 1 (PCMI) is another putative schizophrenia susceptibility gene. It has also been
shown to interact directly with DISC1.\textsuperscript{35} DISC1 regulates PCM1 localization to the centrosome,\textsuperscript{36} an organelle that serves as the main microtubule organizing centre. Microtubule assembly is in turn PCM1 dependent.\textsuperscript{37} Recent data show that the Leu607Phe polymorphism of DISC1 affects both centrosomal PCM1 localization and spontaneous neurotransmitter (noradrenaline) release but interestingly not microtubule dynamics.\textsuperscript{38} The assembly of other centrosomal proteins and microtubule anchorage and organization are dependent on PCM1,\textsuperscript{36} suggesting that the DISC1 Leu607Phe polymorphism could impinge on microtubule function in the absence of changes in microtubule dynamics. The fact that microtubules act as railroads along which molecular motors transport intracellular cargoes, such as precursors of synaptic vesicles to axon terminals, provides a possible explanation for why the Leu607Phe polymorphism alters neurotransmitter release via changes in PCM1-related changes in microtubule stability and organization.

The PCM1 gene has also been associated with orbitofrontal gray matter volumetric deficits,\textsuperscript{39} and there are a number of haplotypes at the PCM1 locus found to associate with schizophrenia.\textsuperscript{40} Voxel-based morphometry of cases from a UK case-control sample showed a significant relative reduction in the volume of orbitofrontal gray matter in comparison with patients with non-PCM1-associated schizophrenia, who by contrast showed gray matter volume reduction in the temporal pole, hippocampus, and interior temporal cortex. PCM1 is part of a multicomponent complex with DISC1 so it will be interesting to consider other components as putative contributors to risk and brain pathology.

Serine-threonine protein kinase AKT1

AKT1 (also known as protein kinase B), a serine/threonine kinase with multiple cellular functions including metabolism, cell stress, cell-cycle regulation, and apoptosis, also regulates actin organization and cell motility via Girdin (Girders of actin filaments), which directly binds actin at the leading edge of migrating cells.\textsuperscript{41} Girdin has been shown to be a DISC1 interactor.\textsuperscript{41} Its expression in postnatal rodent brain is localized in the dentate gyrus and pyramidal cell layers CA1 and CA3 of the hippocampus, mirroring DISC1 expression.\textsuperscript{41}

There is evidence implicating the AKT pathway in schizophrenia. Genetic variants in AKT have been reported to associate with schizophrenia.\textsuperscript{9} A transgenic mouse has been generated with defective neuronal AKT1 signaling, which exhibits neurochemical and behavioral phenotypes relevant to schizophrenia, including decreases in prefrontal dopamine signaling and deficits in sensorimotor gating.\textsuperscript{42} Impaired cortical AKT1 activity in these mice significantly enhanced norepinephrine transporter function, blockade of which reversed the cortical hypodopaminergia and behavioral deficits in these mice, further supporting the potential of AKT1 and nor-ephinephrine transporter as drug targets.

Decreased AKT1 activity and decreased AKT-dependent phosphorylation of glycogen synthase kinase-3 (GSK-3β), a protein that mediates glucose metabolism and is also thought to regulate synaptic plasticity, have been reported in postmortem schizophrenic brains.\textsuperscript{43} Furthermore, AKT1 knockout mice show impaired PPI of the startle response. Both typical and atypical antipsychotic drugs enhance AKT1 signaling by activating AKT1 or by increasing phosphorylation of GSK-3β.\textsuperscript{44} The mechanism of action is through D2 receptor activation, which results in AKT1 being inactivated by protein phosphatase 2A (PP2A), in an arrestin-dependent complex, thereby increasing GSK-3β activity. AKT1 is activated by phosphorylation, whereas GSK-3β is inactivated if phosphorylated by AKT1. Levels of the phosphorylated GSK-3 isoform, GSK-3β, have been shown to be diminished in the frontal cortex of people with schizophrenia.\textsuperscript{27} GSK-3β has recently been identified as a novel DISC1 interactor, bringing the wingless (wnt) pathway and β-catenin into the arena for DISC researchers.\textsuperscript{45} DISC1 and Girdin dimerize and bind NDEL1, which in turn binds NDE1 and LIS1, other key proteins in neurodevelopment. DISC1 therefore forms higher-order multimers, in a process that alters the binding of NDEL1 and is sensitive to polymorphic variation at the Ser704Cys position,\textsuperscript{46} which has previously been related by fMRI and working memory tasks to differential hippocampal engagement in normal human subjects and also to altered brain expression of DISC1 partners.\textsuperscript{8} Like AKT1, DISC1 signaling also inhibits GSK-3β, and interestingly, from a therapeutic viewpoint, the mood stabilizer lithium chloride also targets GSK-3β.

Dysbindin/DTNBP1

DTNBP1, located at 6p22.3, has been identified as a putative schizophrenia susceptibility gene.\textsuperscript{37} It has also been shown to interact with DISC1.\textsuperscript{48} The molecular mechanism of action of DTNBP1 is not completely understood, but it is widely distributed in the CNS and appears to be involved in a range of processes including: postsynaptic density function and presynaptic glutamatergic transmission, neuron viability via glutamate signaling resulting in modulation of dopamine and other neurotransmitter activity.\textsuperscript{25} ‘Sandy’ mice, which have a spontaneous mutation in dysbindin, show cytoarchitectural changes in the hippocampus\textsuperscript{49} and also exhibit various abnormal behaviors similar to schizophrenia.\textsuperscript{50} Genetic variation in DTNBP1 has been associated with general cognitive ability,\textsuperscript{51} while carriers of a putative schizophrenia risk haplotype have reported to have a significantly greater decline in IQ compared with noncarriers suggesting that DTNBP1 influences the severity of intellectual
decline seen in schizophrenia. Furthermore, DTNBPI has also been associated with imaging phenotypes in schizophrenia, carriers of a risk genotype exhibiting reduced brain volume and regional CT. It has been suggested from the results of these imaging phenotypes that DTNBPI may be interacting with other schizophrenia-related risk factors to affect laminar thickness.

Conclusions
There is an abundance of genetic and biological evidence supporting a role for DISC1 in risk of psychiatric illness and in neurodevelopment and signaling. Here, we have focused on evidence from human brain imaging studies and from genetically engineered mouse models. We have extended this review to include comparable studies of DISC1 interactors to provide an overview of the role of the DISC1 pathway. Collectively, a picture is emerging of how DISC1, through a multiplicity of protein-protein interactions, including with other known risk genes, regulates 2 core etiological concepts in schizophrenia, namely neurodevelopment and neurotransmission. Further studies in both humans and rodents of defined genetic status for DISC1 and its many partners are likely to improve our understanding of schizophrenia etiology at the molecular and systems level, a prerequisite to the effective drug therapies, and the ability to predict the course of illness and treatment responses.

Funding
The Academy of Medical Sciences/Wellcome Trust (R41455 to M.J.); The RS MacDonald Trust (D21419, together with J.H.); Research Councils UK fellowship (GR/T227983/01 to P.A.T.).

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