Binocular rivalry occurs when conflicting images are presented in corresponding locations of the two eyes. Perception alternates between the images at a rate that is relatively stable within individuals but that varies widely between individuals. The determinants of this variation are unknown. In addition, slow binocular rivalry has been demonstrated in bipolar disorder, a psychiatric condition with high heritability. The present study therefore examined whether there is a genetic contribution to individual variation in binocular rivalry rate. We employed the twin method and studied both monozygotic (MZ) twins (n = 128 pairs) who are genetically identical, and dizygotic (DZ) twins (n = 220 pairs) who share roughly half their genes. The binocular rivalry rate was signiﬁcantly higher for MZ than for DZ twins (0.51 versus 0.19, respectively). The best-ﬁtting genetic model showed 52% of the variance in binocular rivalry rate was accounted for by additive genetic factors. In contrast, nonshared environmental inﬂuences accounted for 18% of the variance, with the remainder attributed to measurement error. This study therefore demonstrates a substantial genetic contribution to individual variation in binocular rivalry rate. The results support the vigorous pursuit of genetic and molecular studies of binocular rivalry and further characterization of slow binocular rivalry as an endophenotype for bipolar disorder.

Genetic contribution to individual variation in binocular rivalry rate

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Results
Participants were adolescent twins (722 individuals; 48% male) with a mean age of 14 years. A subsample (97 individuals) was retested approximately 2 years later. Fig. 1 shows the binocular rivalry measures, data collection protocol, and genetic modeling procedure. The binocular rivalry stimuli comprised drifting vertical and horizontal square-wave gratings, viewed through liquid crystal display goggles, with no training in ﬁxation required. Participants pressed one key for the vertical percept, an adjacent key for the horizontal percept, and a third response option (“mixed”) for mixed percepts, unusual or uncertain percepts, or a previously incorrect response. Binocular rivalry rate was calculated by dividing the number of perceptual switches by the total viewing period in Blocks 2 and 3, excluding the periods immediately preceding and following a mixed response. The resulting value for binocular rivalry rate is the number of perceptual switches per second (expressed in Hz). Predominance was calculated by dividing the total time spent perceiving the vertical grating by the total time spent perceiving the horizontal grating in Blocks 2 and 3 (with the same mixed-response exclusions). The resulting ratio then was log transformed.

Exclusive binocular rivalry (i.e., minimal mixed percepts) was achieved successfully (Table 1). Binocular rivalry rates varied widely between individuals (0.08–1.32 Hz; mean, 0.54 ± 0.15 SD), as did predominance (Table 1). The main ﬁnding is that the twin correlation for binocular rivalry rate was signiﬁcantly higher for MZ (0.51) than for DZ twins (0.19), indicating genetic inﬂuence on this measure (Fig. 2C). In contrast, neither MZ or DZ twin correlations were signiﬁcant for predominance or nonexclusive (mixed) rivalry periods (Table 1 and Fig. 2C), so these measures were not included in genetic modeling analyses.

The within-test reliability for binocular rivalry rate was very high (Fig. 2C), and reliability over time (retest) was high (Fig. 2D). Within-test reliability for mixed hits and mixed time was very high; within-test reliability for predominance was lower but was still high (Table 1). Reliability over time was moderate for predominance, mixed hits, and mixed time (Table 1). Measures of between-block change in predominance showed poor reliability over time (retest correlations were nonsigniﬁcant) and therefore are not reported.


Conflict of interest statement: S.M.M. and J.D.P. are co-inventors on a granted University of Queensland, national and international patent concerning slow binocular rivalry in bipolar disorder. All other authors declare no conﬂict of interest.

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Binocular rivalry and genetic modeling. (A) Presenting a different image simultaneously, one to each eye, induces binocular rivalry (with occasional mixed percepts). In the present study, stimuli comprised drifting vertical and horizontal square-wave gratings. Binocular rivalry measures were rate (Hz), predominance (ratio of time spent perceiving one image relative to the other), number of mixed hits, and time associated with mixed hits. (B) Binocular rivalry data were collected for 21 min in three blocks. Block 1 was used for training and was discarded. Data from Blocks 2 and 3 were analyzed and were of most interest because binocular rivalry rates tend to stabilize with viewing time (B). (C) Path modeling of variance (12) into additive (A) and nonadditive (i.e., dominance/epistasis, D) genetic sources, unique environmental sources (E), and measurement error/unreliability (U). An ADEU model was chosen because the MZ twin correlation was more than twice the DZ twin correlation, indicating the importance of the A and D components over C components of variance. Reliable genetic and environmental variance is identified by equating pathways from A, D, and E components to data from the first and second test occasions. The remaining variance (U) is unshared between the two test occasions but represents an equal amount of variance for each variable on each test occasion and therefore is equated. Correlations between covtins for factors A and D are fixed at Mendelian expectations.

Preliminary analyses of binocular rivalry rate before genetic modeling showed homogeneity of sampling, with no birth order, zygosity, or sex effects for means or variances. Further, no mean effect was found for age. However, a significant mean effect was found for acuity (Δρ^2 = 1.70), such that 37 individuals with acuity of 6/9 in either eye had a marginally slower binocular rivalry rate than the rest of the sample for whom acuity was 6/6 or better in both eyes. Therefore, acuity was included as a covariate in all further analyses.

Figure 1. Binocular rivalry and genetic modeling. (A) Presenting a different image simultaneously, one to each eye, induces binocular rivalry (with occasional mixed percepts). In the present study, stimuli comprised drifting vertical and horizontal square-wave gratings. Binocular rivalry measures were rate (Hz), predominance (ratio of time spent perceiving one image relative to the other), number of mixed hits, and time associated with mixed hits. (B) Binocular rivalry data were collected for 21 min in three blocks. Block 1 was used for training and was discarded. Data from Blocks 2 and 3 were analyzed and were of most interest because binocular rivalry rates tend to stabilize with viewing time (B). (C) Path modeling of variance (12) into additive (A) and nonadditive (i.e., dominance/epistasis, D) genetic sources, unique environmental sources (E), and measurement error/unreliability (U). An ADEU model was chosen because the MZ twin correlation was more than twice the DZ twin correlation, indicating the importance of the A and D components over C components of variance. Reliable genetic and environmental variance is identified by equating pathways from A, D, and E components to data from the first and second test occasions. The remaining variance (U) is unshared between the two test occasions but represents an equal amount of variance for each variable on each test occasion and therefore is equated. Correlations between covtins for factors A and D are fixed at Mendelian expectations.

Table 1. Binocular rivalry summary statistics and correlations

<table>
<thead>
<tr>
<th>Variable</th>
<th>Test Mean (SD)</th>
<th>Test Range</th>
<th>Retest Mean (SD)</th>
<th>Retest Range</th>
<th>Reliability Within-test r (95% CI)</th>
<th>Between-test r (95% CI)</th>
<th>MZ r (95% CI)</th>
<th>DZ r (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Binocular rivalry rate, Hz</td>
<td>0.54 (0.15)</td>
<td>0.08–1.32</td>
<td>0.53 (0.16)</td>
<td>0.21–1.04</td>
<td>0.93 (0.92–0.94)</td>
<td>0.70 (0.58–0.78)</td>
<td>0.51 (0.37–0.62)</td>
<td>0.19 (0.07–0.31)</td>
</tr>
<tr>
<td>log predominance</td>
<td>0.17 (0.09)</td>
<td>−0.44–0.48</td>
<td>0.14 (0.08)</td>
<td>−0.24–0.22</td>
<td>0.71 (0.67–0.75)</td>
<td>0.43 (0.23–0.59)</td>
<td>0.08 (−0.10–0.25)</td>
<td>0.07 (−0.06–0.19)</td>
</tr>
<tr>
<td>Mixed hits</td>
<td>18.4 (25.4)</td>
<td>0–156</td>
<td>12.5 (18.0)</td>
<td>0–83</td>
<td>0.94 (0.92–0.95)</td>
<td>0.30 (0.03–0.53)</td>
<td>−0.10 (−0.32–0.13)</td>
<td>−0.11 (−0.27–0.05)</td>
</tr>
<tr>
<td>Mixed time, seconds</td>
<td>33.9 (50.2)</td>
<td>0–372.1</td>
<td>25.3 (42.8)</td>
<td>0–243.0</td>
<td>0.91 (0.89–0.93)</td>
<td>0.39 (0.13–0.60)</td>
<td>−0.14 (−0.34–0.08)</td>
<td>−0.12 (−0.28–0.04)</td>
</tr>
</tbody>
</table>

All mean, SD, and range values are before winzorization. Data were winzorized to ±3.3 SD before (i) reliability and twin maximum likelihood correlation analyses, (ii) preliminary analyses for genetic modeling, and (iii) genetic modeling. Mixed hits and mixed time were categorized for correlation analyses because of non-normal distribution (thresholds were set that divided each variable into three categories of approximately equal size based on scores from the first test occasion). Exclusion of binocular rivalry was achieved successfully, indicated by comparison of mixed hits/time with the hits/time associated with vertical (V) and horizontal (H) gratings percepts, for which the Test group means, SD, and ranges are, respectively: V hits, 189.0, 64.3, 19–516; H time, 43.5, 61.8, 109.3–354.8; H hits, 188.9, 64.3, 19–514; H time, 339.1, 61.8, 109.3–354.8.

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Population binocular rivalry and genetic modeling data. (A) MZ and DZ twin maximum likelihood correlations for binocular rivalry measures (error bars indicate 95% CI). BR, binocular rivalry; Predom, predominance. (B) The variance in binocular rivalry rate, estimated from the AEU (best-fitting) model, was accounted for by a substantial additive genetic component; plus unique environmental influences and measurement unreliability over a period of 2 years. The 30% unreliable variance splits 7% within tests and 23% between occasions. (C) Scatterplot showing very high within-test reliability of binocular rivalry rate (95% CI, 0.92–0.94; n = 722) in 14-year-old twins. (D) Scatterplot showing high retest reliability of binocular rivalry rate (95% CI, 0.58–0.78; n = 97) in twins retested at 16 years of age.

**Discussion**

This study represents the largest binocular rivalry population dataset yet published and provides confirmation of reports that binocular rivalry rate varies widely between individuals (3–8). The data also confirm reports of very high within-test (8) and high retest reliability of binocular rivalry rate (4, 5, 7, 8). We have demonstrated a substantial genetic contribution to individual variation in binocular rivalry rate. The mechanisms of this genetic influence remain to be determined.

Studies of binocular rivalry mechanistic models have focused on the level at which the phenomenon occurs in the brain, with perception-dependent neural activity reported as early as the lateral geniculate nucleus (13, 14), in the primary visual cortex (15), as late as the inferotemporal cortex (16), and also in high-level nonvisual regions (17). Similarly, modulation of binocular rivalry can occur with brain stimulation applied at low (18) or high (19, 20) levels. Proposed mechanistic models have included rivalry between specific populations of neurons (21), rivalry between stimulus representations at a high level of visual processing (22), hierarchical computational models (23, 24), and rivalry between independent attentional selection mechanisms in each cerebral hemisphere (19, 20, 25).

Early studies argued against an influence of peripheral factors, such as eye movements, on binocular rivalry (26, 27; notwithstanding central control of peripheral factors). More recently, an influence of eye movements (that induce retinal image shifts) has been reported (28; see also ref. 29 in regard to blinking). However, perceptual alternations also were shown to occur independently of peripheral factors, thus implicating a central binocular rivalry process (28; see also ref. 30). Further studies are required to identify the precise role of peripheral and central factors in determining binocular rivalry rate. An influence of voluntary attention on binocular rivalry has been reported also, but such influence is limited (31, 32) and is much less than the order of magnitude of individual variation reported here and in previous studies (3–8). Nonetheless, the emerging genetics of attentional networks (33) may be relevant to the present finding, because mechanisms of involuntary attention are thought to be engaged during binocular rivalry (1, 20, 25). Few studies have examined the role of neurotransmitter systems and the effects of pharmacological agents on binocular rivalry; however, recent reports suggest involvement of serotonergic (34, 35) and noradrenergic (30) systems. Such studies may provide clues to molecular mechanisms underlying individual variation in binocular rivalry rate. Indeed, the present finding warrants expanding the focus of binocular rivalry research from levels-based investigation to genetic and molecular aspects of the phenomenon.

Also warranted is a focus on models of binocular rivalry that accommodate individual variation in binocular rivalry rate. One such model (7, 19, 25, 36), which remains under investigation, suggested a role for cationic channel levels in determining binocular rivalry rate. This suggestion was based on demonstrating slow binocular rivalry as an endophenotype for bipolar disorder (7); a finding that has since been replicated (8, 37) and also has been shown with different types of perceptual rivalry (38–40). The present finding supports the use of slow binocular rivalry as an endophenotype for bipolar disorder because endophenotypes for heritable conditions must themselves be heritable traits (11). The heritability of binocular rivalry rate reported here is comparable to that reported for other neuropsychiatric endophenotypes, such as P300 event-related potential amplitude (heritability range, 0.3–0.8) in subjects with alcohol dependency (41, 42). The present finding has the potential to reveal not only mechanisms of binocular rivalry but also mechanisms of bipolar disorder and suggests gene-finding approaches to both. It also suggests use of the trait to help overcome challenges posed by the heterogeneity of the bipolar clinical phenotype (7–11, 37).

However, further characterization of the trait is required, including assessment of binocular rivalry rate in other psychiatric
conditions, particularly schizophrenia and major (unipolar) depression, to establish the trait’s specificity (8, 10). Also required is assessment of binocular rivalry rate in first-degree relatives of bipolar probands and of possible effects of medication and clinical state. However, effects of medication or clinical state are unlikely to account for slow binocular rivalry in bipolar disorder (7, 8, 37), a suggestion that is consistent with the present finding of a substantial genetic contribution to binocular rivalry rate. Similarly, eye movements are unlikely to account for the slowing of binocular rivalry in bipolar disorder because the limited number of studies of saccades during smooth pursuit in this clinical population (e.g., 43, 44) show either no significant difference from controls or an increased saccade frequency (which should cause a faster rather than slower binocular rivalry rate). Moreover, slow binocular rivalry in bipolar disorder also has been demonstrated with stationary gratings (8) that do not elicit pursuit eye movements.

If further studies confirm that endogenous binocular rivalry rate is fundamentally determined by central processing factors, the present finding also represents demonstration of, in a large sample, a substantial genetic contribution to individual variation in a postretinal visual processing phenomenon (45). Although there have been previous reports of genetic contributions to illusory movement (46), flicker fusion thresholds (47), and Rorschach indices (47), the sample sizes in those studies were small. A recent twin study, which did employ a large sample, assessed contrast sensitivity in middle-aged males and found only a modest heritability estimate (0.14–0.38) (48). Moreover, it is not known whether deficient contrast sensitivity occurs in the lens, retina, or postretinal processing (48).

Inspection time for line-length discrimination also has been examined in a large twin sample and was shown to have substantial genetic influence (heritability estimate, 0.57), but this perceptual task is thought to reflect attentional and decision processes used in response monitoring (49). Although a role for decision-making during binocular rivalry has been proposed (30, 50), the perceptual alternations cannot be prevented, and, as discussed above, voluntary attention has only a limited effect on the phenomenon (1, 31, 32).

Finally, the heritability and molecular basis of individual variation in human color vision—a retinal phenomenon—is well understood (51, 52). We therefore propose that if individual variation in binocular rivalry rate is indeed a predominantly central processing phenomenon, then, as for color vision and the retina, binocular rivalry may serve as a paradigm case to unravel the genetics, physiology, and pathophysiology of postretinal vision and perception. Although binocular rivalry is likely to be a complex trait with complex inheritance (unlike color vision), it is a phenomenon that has proven highly amenable to research in animals and humans using a wide variety of investigative methods (1). It is a phenomenon that also may provide a unique window into the science, and indeed genetics, of visual consciousness (53, 54).

Several hundred years since binocular rivalry was discovered, and more than a century since early reports that the rate of binocular rivalry varies widely between individuals, we have shown this aspect of the phenomenon to be under substantial genetic influence. This result suggests a range of novel approaches to investigate binocular rivalry further in both general and clinical populations, with a focus on genetic, molecular, and endophenotype studies.

Materials and Methods

Participants. A population sample of twins was recruited by the Queensland Institute of Medical Research (QIMR) for a genetic study of melanocortin nevi (moles) (55) through mailings to schools in South-East Queensland between 2000 and 2009. Binocular rivalry data were collected during a routine mole-count visit scheduled when twins turned 14 years of age. The sample included 722 individuals (48% male; 128 MZ pairs, 220 DZ pairs, and 5 unpaired cotwins; mean age, 14.1 ± 0.51 SD; range, 14–15 years). Zygosity was determined by typing nine independent polymorphic DNA markers using the AmpFLSTR Profiler PCR Amplification Kit (Applied Biosystems) and checked with ABO, MN, and Rh blood groups and/or phenotypic information (hair, skin, and eye color), with an extremely low probability of error (<10⁻⁴). At ≥16 years of age, twins were invited to participate in a study of cognition (56), and a subset (n = 97; 53% male) was retested 1.9–2.8 years (mean, 2.1 ± 0.2 SD) after the first test. The retest sample comprised 11 MZ pairs, 35 DZ pairs, and 5 unpaired cotwins aged 16.0–16.9 years (mean, 16.1 ± 0.2 SD). Individuals at age 14 or 16 years were excluded if they (i) reported a history of, or medication for, bipolar disorder, depression, or attention deficit hyperactivity disorder, (ii) reported a history of brain injury or other neurological condition, (iii) reported a history of uncorrected strabismus, (iv) had visual acuity worse than 0.6 in either eye (acuity was measured using a Snellen chart at 3 m), or (v) there were problems with data collection. There were 26 exclusions at the first test and 11 exclusions at retest. Written, informed consent was obtained from all participants and a parent or guardian. The study was conformed to the National Statement on Ethical Conduct in Human Research (2007) issued by the National Health and Medical Research Council (NHMRC) of Australia and was approved by the QIMR Human Research Ethics Committee.

Binocular Rivalry Stimuli, Recording Procedure, and Measures. Binocular rivalry stimuli were presented on a monochrome (green) computer monitor situated 3 m from the participant, in a dimly lit room. The orthogonal stimuli were vertical gratings drifting left-to-right, always presented to the left eye, and horizontal gratings drifting downward, always presented to the right eye. The gratings had a spatial frequency of 8 cycles/degree, were drifting at 4 cycles/second and were presented in a circular patch subtending 1.5° of visual angle. Contrast of the gratings was 0.9. Participants were instructed not to consume caffeinated beverages for 2 hours before the binocular rivalry test session. To assist with training, they received an explanatory sheet showing the various possible perceptions and explaining how to respond in each scenario. Participants were instructed to view the stimuli passively rather than to attempt to influence their perceptions. They were not aware of a research support that was using the data collection. Block 1 recording was used to train the participant, checking that the patient understood the instructions and was performing the task correctly. Questions could be asked during this period. Participants used the left hand to press a raised key in response to the vertical percept and the right hand to press an adjacent raised key in response to the horizontal percept. Mixed perceptions were indicated by pressing the space bar, using either hand, and this response option also was used to indicate unusual percepts and incorrect or undecided responses. Although the mixed hits and mixed time values provide an approximate indication of nonexclusive rivalry (i.e., mixed perceptions), this value is likely to be an overestimation because the mixed response option also was used to indicate unusual percepts and incorrect or undecided responses. Analyses were performed with specialized software (BiReme Systems) and PASW Statistics 17.0 (SPSS Inc). Within- and between-test reliability and genetic analyses used the structural equation modeling package Mx (www.vcu.edu/mx) which employs maximum likelihood estimation from raw data observations.

Genetic Model Fitting. Genetic modeling (56) can estimate additive (A) and nonadditive (i.e., dominance/epistasis, D) genetic effects, common (C) and unique (E) environmental effects and, when and if present, a suggestion of the importance of the A component and of the D component over C (57), so an ADE model was fitted (Fig. 1). Further con-
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