Elucidating the Black Box From Stress To Paranoia

Steffen Moritz*,1, Pia Burnette1, Sabine Sperber1, Ulf Köther1, Marion Hagemann-Goebel1, Maike Hartmann2, and Tania M. Lincoln2

1Department of Psychiatry and Psychotherapy, University Medical Center Hamburg-Eppendorf, Martinistr. 52, D-20246 Hamburg, Germany; 2Section for Clinical Psychology and Psychotherapy, Faculty of Psychology, Philipps-Universität Marburg, Gutenbergstr. 18, D-35037 Marburg, Germany

*To whom correspondence should be addressed; tel: +49-40-7410-56565, fax: +49-40-7410-57566, e-mail: moritz@uke.uni-hamburg.de.

Sensitivity to stress has long been implicated in the pathogenesis of schizophrenia. It remains unclear, however, which exact mechanisms underlie the progression from vulnerability to psychotic breakdown. For the present study, we hypothesized that the induction of stress would aggravate cognitive biases in schizophrenia. A total of 20 acute and remitted schizophrenia patients and 15 healthy controls were tested with parallel versions of cognitive biases paradigms under 2 laboratory conditions: stress (loud noise, 75 dB) vs no-stress. In the course of both conditions, participants had to fill out a questionnaire that assessed depressive, obsessive–compulsive, and paranoid symptoms. For the patients with acute psychotic symptoms, paranoid but not other psychiatric symptoms were elevated under stress in comparison with no-stress. In contrast, stress somewhat diminished subclinical paranoid symptoms in healthy participants. Jumping to conclusions was evident in schizophrenia under both conditions but significantly more pronounced when stress was applied first in the acute group. A tendency emerged in both acute and remitted patients to attribute events to other people under stress which was not seen in healthy subjects. The present study may serve as a starting point for further research investigating how stress translates vulnerability into acute paranoia and to pinpoint cognitive risk factors that can be modified by treatment.

Key words: schizophrenia/stress/cognitive biases/jumping to conclusions/noise

Introduction

It is one of the most well-established findings in schizophrenia research that stress triggers or aggravates psychosis. Daily hassles as well as sustained levels of stress, as exerted, for example, by urbanicity, migration, and increased stress sensitivity, easily translate into symptoms such as paranoia in vulnerable individuals. The vulnerability–stress conceptualization of schizophrenia is picked up in almost all psychosocial interventions and is at the core of psychoeducation programs: However, while valuable as a heuristic model, many of its constituents remain unexamined, especially the modulating mechanisms on the way from stress to psychosis.

Given the large literature suggesting that cognitive distortions, such as reasoning and attributional biases, especially jumping to conclusions (JTC) and blaming others for failure, are present in the disorder and are presumably linked with the positive symptoms of schizophrenia, it is tempting to speculate that these biases are especially sensitive to stress and pave the way to breakdown. Starting with a study by Mujica-Parodi and coworkers, this assumption has obtained some empirical support. For example, our group found that healthy participants scoring high on psychosis-prone items showed a direct increase of paranoia under stress, which was mediated by negative emotion. However, elevated paranoia was not accompanied by an increase in reasoning biases in this sample. In another study, healthy participants with higher baseline vulnerability for psychosis showed an increase of paranoia after the induction of anxiety, which was mediated by increased JTC. In yet another study, patients high on delusions showed pronounced hasty decision making under emotional music relative to controls lending further support to the assumption that stress and emotional arousal aggravate cognitive biases.

Although these studies have not produced entirely consistent findings, they underline the potential relevance of cognitive biases for the translation from stress to
psychosis. So far, most studies have focused on JTC and did not address other cognitive biases (eg, attributional biases) which tap independent cognitive functions. Furthermore, the aforementioned studies used very different induction paradigms and currently we do not know whether general fear, social anxiety, or stress per se impact on cognitive biases the most.

For the present study, we aimed to extend on previous research by investigating the cognitive performance of schizophrenia patients and controls on 2 laboratory conditions, either during sensory stress (noise) or without stress. We chose loud noise as it is a valid and potent stressor. We concurrently measured several cognitive biases (eg, JTC, bias against disconfirmatory evidence [BADE]) along with a neurocognitive parameter (nonverbal memory). We assumed that patients would display deviant responses on the no-stress condition for cognitive biases, especially JTC, which would be further aggra-
vated under stress. No changes were expected for cognitive deficits as these are not related to positive but rather to nonacute negative and disorganized symptoms which are less stress sensitive. In order to ascertain the specificity of the association between stress and psychotic symp-
toms, we assessed other clinical symptoms as well. We predicted that stress would have the strongest impact on paranoid ideation, particularly for the acute patient group.

Methods

Participants

Twenty schizophrenia patients were recruited from the psychosis units of the University Medical Center Hamburg. Patients met criteria for schizophrenia and schizo-
affective disorder as determined by the Mini
International Neuropsychiatric Interview (MINI). According to the MINI criteria, 13 of the patients were currently in remission. None of the patients had a macroscopic neurological disorder or substance dependence. All patients were on stable antipsychotic medication. We recruited 15 healthy participants via an established subject pool who did not have any axis I diagnosis as assessed with the MINI and had had no prior contact with mental health institutions (for sociodemo-
graphic characteristics, see table 1).

Design and Procedure

The experiment was conducted as a randomized repeated-measures design. All participants underwent an assessment under 2 conditions within a time interval of 1–7 days: stress vs no-stress. During each session, they had to complete parallel versions of various cognitive tasks as well as the symptom rating scales. In the stress condition, stress was induced by loud construction noise via headphones (75 dB) except for the respective instruc-
tion phase. The no-stress condition was identical in admin-
istration except that no noise was applied.

Parallel versions of the experiments and the clinical questionnaires (randomized item order) were random-
ized to the conditions. The order of the assessment was the same in each condition. Participants who first un-
derwent the stress condition will be referred to as “stress first” group, whereas participants who first underwent the no-stress condition second will be referred to as the “stress second” group (see table 2).

Instruments

Baseline Assessments. To assess the baseline stress level, the Perceived Stress Scale (PSS) was administered which measures the degree to which situations in one’s life are appraised as stressful. It has shown satisfactory reliability and concurrent as well as discriminate validity.

In addition, the Community Assessment of Psychic Experiences (CAPE) questionnaire was administered before the experimental procedure to assess the extent of psychotic, negative, and depressive symptoms.

Assessments during the Experimental Conditions. Symptomatology A combined questionnaire on para-


oid, obsessive–compulsive, and depressive symptoms was administered in random item order. Participants had to rate the extent to which each item applies “at the moment.” It consisted of the following 3 instruments:

1. The Paranoia Checklist is an 18-item scale capturing ideas of persecution and reference. The German version has good internal consistency and convergent val-

2. The Obsessive–Compulsive Inventory-Revised (OCI-
R) was administered to evaluate the frequency and distress experienced by OC symptoms across 6 sub-

3. The General Depression Scale (GDS) is a German 20-

item self-report questionnaire tapping depressive fea-

ures. The GDS shows high internal consistency and is strongly correlated with other indexes of depression.

Six times during each condition (see table 1), partici-

pants had to complete a Visual Analog Scale (VAS) assessing the subjective experience of stress by making a mark on a 10-cm line for 4 stress items.

Cognitive Biases JTC was determined using a variant of the so-called beads task. The revised version is easier to comprehend than the original task and controls for effects of motivation. Participants were shown 2 lakes with fish of 2 different colors (lake 1: 80% color A and 20% color B and vice versa for lake 2). Participants were told that fish would be caught successively from
only one lake and were asked to judge from which lake the fish were caught. In total, 10 fish were drawn for all subjects in a seemingly random but actually fixed order. Participants were asked to give an estimate for each fish. In one version, lake 1 was favored (sequence: A-A-A-B-A-A-A-A-B-A). In the other version, lake 2 was favored (sequence: B-B-B-A-B-B-B-B-A-B). A decision after only one fish indicated JTC. At each of the 10 stages, a decision could be made, withdrawn, or changed.

The BADE task was adapted from earlier versions and entailed 16 delusion-neutral scenarios each consisting of 3 successive statements each providing additional information that increasingly disambiguates the scenario. Along with the statements 4 interpretations appeared.

Table 2. Protocol of Investigation (Stress Test without baseline)

<table>
<thead>
<tr>
<th>Time (minutes)</th>
<th>Tasks</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>Instruction</td>
</tr>
<tr>
<td>8</td>
<td>VAS(1) &gt; instruction BADE &gt; [noise on—start] &gt; execution &gt; BADE</td>
</tr>
<tr>
<td>28</td>
<td>VAS(2) &gt; [noise out] &gt; complex figure instruction &gt; [noise on] &gt; execution complex copy and immediate recall</td>
</tr>
<tr>
<td>40</td>
<td>VAS(3) &gt; [noise out] &gt; instruction fish task &gt; [noise on] &gt; execution fish task</td>
</tr>
<tr>
<td>68</td>
<td>VAS(5) &gt; [noise out] &gt; instruction IPSAQ-R &gt; [noise on] &gt; execution IPSAQ-R</td>
</tr>
<tr>
<td>90</td>
<td>End of IPSAQ-R &gt; VAS (6) &gt; [noise out—end]</td>
</tr>
</tbody>
</table>

Note: BADE, bias against disconfirmatory evidence; IPSAQ-R, Internal, Personal, and SituationalAttributions Questionnaire; POD, Questionnaire on Paranoid, Obsessive–Compulsive, and Depressive Symptoms; VAS, Visual Analog Scale.
between-subject factors. These 3 factors will be referred to as the standard set of factors in the following and complemented by additional factors depending on the experimental variable. The report of significant analyses will be confined to the core effects involving the Group factor.

**Results**

**Sociodemographic and Clinical Variables**

As can be seen in table 1, the groups did not differ with respect to any of the background variables. Patients had elevated CAPE scores which were in the expected range according to prior studies. The baseline stress level, as assessed with the PSS, was elevated in patients.

**Effect of Stress on Psychopathology**

A 4-way mixed ANOVA was computed: Psychopathology (Paranoia Checklist, OCI-R, GDS) and Condition (stress vs no-stress) served as the within-subject factors and Group (schizophrenia, healthy) and Sequence (stress condition applied first or second) were the between-subject factors. As expected, the main effect of Group was significant, \( F(1,31) = 16.18, P < .001, \eta^2_{\text{partial}} = .34 \). As can be derived from figure 1, the schizophrenia group had overall higher scores than the healthy participants. In addition, the Group \( \times \) Sequence interaction was significant, \( F(1,31) = 4.62, P = .01, \eta^2_{\text{partial}} = .13 \): Patients in the stress first condition had overall higher psychopathology scores than in the stress second condition. This was qualified by a significant 3-way interaction of Psychopathology \( \times \) Condition \( \times \) Group, \( F(2,62) = 4.55, P = .01, \eta^2_{\text{partial}} = .13 \). Figure 1 shows that patients were more paranoid under stress than under no-stress, and this was slightly opposite for healthy participants. For depression, this response pattern was inverted: Stress led to an increase of depressive symptoms. Results are displayed as means and SEs.

For the BADE, none of the effects involving Group achieved significance (\( P > .3, \eta^2_{\text{partial}} < .04 \)). For the Internal, Personal, and Situational Attributions Questionnaire (IPSAQ-R), we added 2 within-subject factors to the standard set of factors: Scenario (positive, negative) and Attribution (‘‘I,’’ ‘‘others’’; the percentage for the response option ‘‘circumstances’’ was not entered as an additional step as it can be derived from the percentages of the 2 other steps). The 3-way interaction of Group \( \times \) Condition \( \times \) Attribution was significant, \( F(1,31) = 6.94, P = .01, \eta^2_{\text{partial}} = .18 \). Healthy participants attributed events (irrespective of valence) less often to others, whereas the patients considered others more responsible under stress than without stress. Results remained essentially unchanged when the patient sample was split into acute vs remitted participants.

**Nonverbal Memory**

For the memory test, we added another within-subject factor to the standard set of factors: Reproduction Trial (copy figure, immediate, and delayed recall). The main effect of Group achieved significance, \( F(1,31) = 0.18, P = .005, \eta^2_{\text{partial}} = .23 \), reflecting decreased overall performance by the patients. This was qualified by a Group \( \times \) Reproduction Trial interaction, \( F(2,62) = 6.91, P = .002, \eta^2_{\text{partial}} = .18 \). While the groups performed similar on the copy trial, patients were far worse for the 2 recall trials. Results remained essentially unchanged when the patient sample was split into acute vs remitted participants.

**Heart Rate**

For analyzing the heart rate data, we added Time (first vs final heart rate) as additional factor to the standard set of factors. Only the Group factor achieved significance,
patients with schizophrenia are hastier in their decision making in the stress condition to contextual factors (ie, noise). It appears as if the healthy participants personal attributed any psychological changes during the experimental situation than controls (M = 4.46 vs 1.91). Results remained essentially unchanged when the patient sample was split into acute vs remitted participants: acute patients were not significantly more stressed (M = 4.95) than remitted patients (M = 4.29, t(18) = 0.66, P > .5).

Discussion

The present study aimed to shed light on the interrelationships of paranoid ideation, stress, and cognitive biases in schizophrenia. As expected, stress increased paranoid symptoms in schizophrenia patients, irrespective of symptom status. In marked contrast, stress somewhat diminished subclinical paranoid symptoms in the healthy participants. It appears as if the healthy participants (correctly) attributed any psychological changes during the stress condition to contextual factors (ie, noise).

The study confirms prior findings indicating that patients with schizophrenia are hastier in their decision making than controls. As expected, this response pattern was increased under stress, particularly in acute patients and—in line with another study—when stress was applied first. Moreover, stress increased external-personal attributions in schizophrenia patients, as measured with the IPSAQ-R, which was not the case in healthy subjects.

Whether the observed effects of stress on JTC are primary or secondary due to stress regulation strategies (eg, hasty decision making in order to decrease cognitive load) needs further exploration. In a prior study, we inferred that the association between anxiety and paranoia was moderated by the extent of baseline symptoms and mediated by an increase in the tendency to jump to conclusions: Threat-related interpretations, evoked by state anxiety, will be more rapidly accepted in the presence of a tendency to jump to conclusions. The excess of external-personal responses in schizophrenia patients on the IPSAQ-R under stress could reflect source confusion: tension and arousal exerted by stress may have been misattributed as tension due to other persons. If this speculation holds true, potential misattributions may become even more prevalent in a real-life stressful situation, in which other people are directly involved in contrast to a laboratory situation. We also suspect that the impact of stress on cognitive biases will be most pronounced when stressors are less explicit. Stress obviously did not affect all cognitive parameters similarly: neither the BADE nor nonverbal memory was differentially affected by stress in the 2 groups.

Subjective stress appraisal at baseline and during the 2 conditions as well as heart rates revealed that patients felt generally more stressed than healthy participants. This result supports the clinical observation that patients experience even minor challenges as stressful which has been associated with enhanced emotional reactivity. Although the heart rate was significantly correlated with subjective stress parameters, it cannot entirely be ruled out that some heart rate increase was caused by antipsychotic medication.

Several shortcomings of the present study must be conceded. First, the sample was small although our within-subject design has advantages over cross-sectional designs regarding power. We are currently conducting a larger trial which will also allow to look for moderating and mediating variables as in a previous study on subclinical paranoia. Secondly, we only used one stressor. Thus, further investigations should clarify whether stress in general or specific stressors are pertinent for the emergence and/or maintenance of symptoms and the aggravation of cognitive biases.

To conclude, the results suggest that sensory stress selectively aggravates decision-making and attributional biases as well as psychotic symptoms in (acute) schizophrenia patients. In contrast, stress somewhat increased depressive responses and decreased external attributions in healthy subjects. If replicated, interventions aimed at relapse prevention might benefit from including emotion regulation strategies and teaching patients ways to counter cognitive biases under stress.

Acknowledgments

Declaration of interests: None. The present study was part of the diploma theses of P.B. and S.S.

References


39. Osterrieth PA. Le test de copie d’une figure complexe; contribution a l’étude de la perception et de la mémoire (Test of copying a complex figure; contribution to the study of perception and memory). Arch Psychiat. 1944;30:206–356.


