Innate immunity receptors in depression and suicide: upregulated NOD-like receptors containing pyrin (NLRPs) and hyperactive inflammasomes in the postmortem brains of people who were depressed and died by suicide

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**Background:** Abnormalities of inflammation have been implicated in the pathophysiology of depression and suicide, based on observations of increased levels of proinflammatory cytokines in the serum of people who were depressed and died by suicide. More recently, abnormalities in cytokines and innate immunity receptors such as toll-like receptors have also been observed in the postmortem brains of people who were depressed and died by suicide. In addition to toll-like receptors, another subfamily of innate immunity receptors known as NOD-like receptors containing pyrin (NLRPs) are the most widely present NOD-like receptors in the central nervous system. NLRPs also form inflammasomes that play an important role in brain function. We studied the role of NLRPs in depression and suicide. **Methods:** We determined the protein and mRNA expression of NLRP1, NLRP3 and NLRP6 and the components of their inflammasomes (i.e., adapter molecule apoptosis-associated speck-like protein [ASC], caspase1, caspase3, interleukin [IL]-1β and IL-18) postmortem in the prefrontal cortex of people who were depressed and died by suicide, and in healthy controls. We determined mRNA levels using quantitative polymerase chain reaction, and we determined protein expression using Western blot immunolabelling. **Results:** We found that the protein and mRNA expression levels of NLRP1, NLRP3, NLRP6, caspase3 and ASC were significantly increased in people who were depressed and died by suicide compared to healthy controls. **Limitations:** Some people who were depressed and died by suicide were taking antidepressant medication at the time of their death. **Conclusion:** Similar to toll-like receptors, NLRP and its inflammasomes were upregulated in people who were depressed and died by suicide compared to healthy controls. Innate immunity receptors in general — and NLRPs and inflammasomes in particular — may play an important role in the pathophysiology of depression and suicide.

**Introduction**

Suicide is a major public health concern. According to the National Center for Health Statistics (US Centers for Disease Control and Prevention),1 48,344 suicides took place in 2018 in the United States alone. Many clinical, demographic and psychological studies of suicide have been conducted, but there are relatively few studies of the neurobiology of suicide. In recent years, it has become evident that abnormal immune function may be associated with suicide and depression.3,4 Cytokines are major mediators of inflammation (see reviews by Black and Miller5 and Dowlati and colleagues6); the hypothesis that abnormal immune function is associated with the pathogenesis of suicide6 and depression5 is based primarily on the observation that levels of proinflammatory cytokines such as interleukin (IL)-1β, IL-6 and tumour necrosis factor (TNF) α are increased in the plasma, serum and cerebrospinal fluid of people who are depressed5 and suicidal.3 A role for cytokines in suicide and depression is also supported by the observations of Tonelli and colleagues,7 and by our observation8,9 that mRNA and protein levels of proinflammatory cytokines such as IL-1β, IL-6 and TNFα were increased in the prefrontal cortex (PFC) of people who were depressed and died by suicide. However, the mechanism of increased cytokines in the postmortem brains of such people is unclear.

One of the mechanisms of cytokine production in the brain may be related to the activation of innate immune receptors. The innate immune system plays an important role in the central nervous system, especially in pathological conditions.4 In the brain, several cell types (such as microglia, astrocytes and neurons) express these innate immune receptors,10 which include subfamilies of toll-like receptors (TLRs) and NOD-like receptors (NLRs; see review by Lampron and colleagues11). TLRs and NLRs are pattern-recognition receptors...
that recognize 2 types of ligands: pathogen-associated molecular patterns (PAMPs), linked to various microbes, and damage-associated molecular patterns (DAMPs), produced during tissue-based injury.12-14 After microbial recognition, both TLRs and NLRs activate signalling pathways that lead to the formation of cytokines and chemokines.15

PAMPs and DAMPs include proteins from bacterial membranes, intracellular proteins (such as heat-shock proteins) and protein products (such as adenosine triphosphate [ATP], urea and nucleic acid).16 In a previous study, we reported that the protein and mRNA expression of IL-1β, IL-6 and TNFα was increased in the PFC of adults9 and teens8 who were depressed and died by suicide. We have also found increased mRNA and protein expression of TLRs — particularly TLR2, TLR3, TLR4, TLR6 and TLR1017-18 — in the PFC of people who were depressed and died by suicide compared to healthy controls. This finding suggests that the increased IL-6 and TNFα we observed8,9 in the postmortem brains of people who were depressed and died by suicide may have been related to abnormalities in the components of NLRP inflammasomes.19 In general, the inflammasomes consist of an NOD-like receptors containing pyrin (NLRP) β, which is then processed by the main and the downstream effector enzyme procaspase1.20,21 Because the conversion of pro-IL-1β to active or mature IL-1β involves the activation of NLRP inflammasomes, our objective in this study was to examine whether increased protein and mRNA expression of IL-1β in the postmortem brains of people who were depressed and died by suicide19 was related to abnormalities in the components of NLRP inflammasomes.

The NLR family has 21 members; of these, NLRP1, NLRP3 and NLRP6 show the highest levels of expression in the central nervous system, mainly in microglia, but also in astrocytes and neurons,22 and they form inflammasomes. To examine whether increased IL-1β in people who were depressed and died by suicide was related to increased activity of NLRP inflammasomes, we determined the mRNA and protein expression of the following in the PFC of 24 people who were depressed and died by suicide and 24 healthy controls: NLRP1, NLRP3 and NLRP6; the components of NLRP3 inflammasomes (i.e., ASC and caspase1); and the components of the product of NLRP3 inflammasome activation (i.e., IL-1β and IL-18).

Methods

Participants and diagnoses

This study was performed in the PFC (Brodmann area 9) of 24 people who were depressed and died by suicide and 24 nonpsychiatric controls (healthy controls). Brain tissues were obtained from the Maryland Brain Collection at the Maryland Psychiatric Research Center in Baltimore. Tissues were collected only after a family member had provided informed consent. All tissue from healthy controls and people who were depressed and died by suicide was grossly examined by experienced neuropathologists. Toxicology data were obtained by analysis of urine and blood samples. All procedures were approved by the institutional review boards of the University of Maryland and University of Illinois at Chicago.

Diagnostic method

Diagnoses were based on the Structured Clinical Interview for DSM-IV (SCID I).23 At least 1 family member or a friend underwent an interview after providing written informed consent. Two psychiatrists made diagnoses by consensus based on data obtained in the interview, medical records from the case and records obtained from the medical examiner’s office. We verified that healthy controls lacked mental illnesses using the same consensus diagnostic procedures.

Determination of mRNA levels

Total RNA from 100 mg of frozen tissue was extracted using Trizol reagent (Invitrogen; Thermo Fisher Scientific Inc.), followed by DNase treatment according to the manufacturer’s protocol. Concentration of total RNA samples was measured using a NanoDrop 1000 (NanoDrop Technologies, LLC), and purity was determined as a 260/280 nm ratio of 1.8 to 2.0. The quality of the RNA was assessed using a 2100 Bioanalyzer (Agilent Technologies, Inc.) to determine the RNA integrity number (RIN). All samples had 28S/18S ratios greater than 1.2 and RINs greater than 7.0. Finally, RNA samples were stored at −80°C until required for further analysis.

The mRNA expression levels were determined using a 2-step, real-time, quantitative polymerase chain reaction (qPCR) method as follows: reverse transcription in duplicate (25°C, 10 minutes; 37°C, 60 minutes; 70°C, 15 minutes; then hold at 4°C); qPCR in duplicate from each reverse transcription (95°C, 10 minutes; then 95°C for 15 seconds and 60°C for 30 seconds, cycled 40 times). For reverse transcription, 1 µg of total RNA was reverse-transcribed using 50 ng of random hexamers, 2 mM dNTP mix, 10 units of ribonuclease inhibitor and 200 units of MMLV reverse transcriptase enzyme in a final reaction volume of 20 µL. Real-time qPCR was performed using predesigned TaqMan gene expression assays (Applied Biosystems; Thermo Fisher Scientific Inc.) for all target and housekeeping genes on a MX3005p sequence detection system (Agilent Technologies, Inc.). The TaqMan assay IDs for the housekeeping and target genes are listed in Table 1.

To determine the stability and optimal number of the housekeeping genes, we used geNORM version 3.4 (PrimerDesign Ltd.) according to the manufacturer’s instructions. The average gene-stability measure ranked β-actin and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) as the most stable genes in our samples. We tested qPCR efficiency over a 5-log dilution series and confirmed that all target genes and housekeeping genes had similar amplification efficiencies. For each primer–probe set, we performed qPCR using 10 µL of cDNA diluted 1:10-fold. Each qPCR plate included “no reverse transcriptase” and “no template” controls.
to eliminate nonspecific amplification. One sample from each target gene was run on a gel to confirm specificity, and all samples were run in triplicate. Target gene qPCR data were normalized to the geometric mean of β-actin and GAPDH and were expressed relative to the control samples using the 2^{-ΔΔCt} method.

Determination of protein levels

The cytosol fraction of the brain tissue samples was prepared by homogenizing 100 mg of tissue in 0.6 mL of buffer containing 100 mM Tris (pH 7.4); 150 mM NaCl; 1% Triton X-100; 1% sodium decanoate; 0.1% sodium dodecyl sulfate; 5 mM ethylenediaminetetraacetic acid; 1 mM phenylmethylsulfonyl fluoride; 5 μg/mL aprotinin, leupeptin and pepstatin; and 100 mM sodium orthovanadate. The homogenate was centrifuged at 15 000 g for 10 minutes at 4°C, and the supernatant was used for all assays. Protein concentration was determined according to the method of Lowry and colleagues.

Quantitation of NLRPs, ASC, caspase1 and caspase3

Immunolabelling of NLRP1, NLRP3, NLRP6, caspase1, caspase3 and ASC was performed as described previously. Briefly, 20 μg protein samples were loaded onto 7.5% (wt./vol.) acrylamide gel. The proteins were then transferred electrophoretically to an enhanced chemiluminescent nitrocellulose membrane. Blots were incubated with anti-NLRP1, NLRP3, NLRP6, caspase1, caspase3 and ASC antibodies overnight and with horseradish peroxidase-linked secondary antibody for 3 hours at room temperature, exposed to enhanced chemiluminescent film. Membranes were probed by monoclonal β-actin antibody (1:5000 for 2 hours), followed by secondary anti-mouse IgG antibody (1:5000 for 2 hours). The bands on the autoradiogram were quantified using the Loats Image Analysis System (Loats Associates, Inc.). The optical density of each band for NLRP1, NLRP3, NLRP6, caspase1, caspase3 and ASC was corrected using the optical density of the corresponding β-actin band. Values were presented as a percentage of the control. The NLRP1, NLRP3, caspase1, caspase3 and ASC polyclonal antibodies were purchased from Cell Signalling Technology, Inc.; the NLRP6 antibody was purchased from Abcam.

Determination of IL-18 protein levels

Levels of IL-18 were determined in aliquots of the cytosol fraction by enzyme-linked immunosorbent assay using commercially available Quantikine kits for human IL-18, purchased from R & D Systems.

Statistical analysis

We performed data analyses using SPSS statistical software (version 27; IBM). All data were embedded in a generalized linear model to compare the effects of the 2 groups (people who were depressed and died by suicide and healthy controls) on the mRNA and protein expression of NLRP and NLRP-associated inflammasome molecules by adjusting the effects of age, sex, postmortem interval and brain pH. To examine whether the protein and mRNA expression of NLRPs and other outcome measures were affected by post-mortem interval, age, or brain pH, we included these variables as covariates in the multivariate model. We analyzed the results for protein and mRNA expression individually using SPSS. We also tested each outcome measure for normality using the Kolmogorov–Smirnov test. We performed univariate comparisons between the 2 groups for each studied molecule using the Bonferroni post hoc test. Changes with p values of 0.05 or less were considered significant.

Results

Sample demographic and clinical characteristics

The demographic and clinical characteristics of the 2 study groups are summarized in Table 2, and detailed characteristics are provided in Appendix 1, Table S1, available at jpn.ca. This report describes observations for variables such as age, sex, race or ethnicity, postmortem interval and brain pH; Appendix 1, Table S1, also includes the method of suicide and the presence of antidepressants or ethanol at the time of death for people who were depressed and died by suicide.

| Primer/probe | TaqMan accession | Probe location (exon boundary) | Assay function |
|--------------|------------------|-------------------------------|---------------|
| ACTB         | Hs99999903_m1    | 1–1                           | Housekeeping gene |
| GAPDH        | Hs99999905_m1    | 3–3                           | Housekeeping gene |
| NLRP1        | Hs00248187_m1    | 6–7                           | Target gene |
| NLRP3        | Hs00918082_m1    | 1–2                           | Target gene |
| NLRP6        | Hs00373246_m1    | 5–6                           | Target gene |
| CASP1        | Hs00354836_m1    | 8–9                           | Target gene |
| CASP3        | Hs00234387_m1    | 4–5                           | Target gene |
| PYCARD/ASC   | Hs00203118_m1    | 2–3                           | Target gene |
| IL18         | Hs01038788_m1    | 3–4                           | Target gene |

PCR = polymerase chain reaction.
NLRP inflammasomes in depression and suicide

Table 2: Participant demographic and clinical characteristics

| Characteristic       | Healthy controls* | Depressed and died by suicide* |
|----------------------|-------------------|--------------------------------|
| Participants, n      | 24                | 24                             |
| Age, yr              | 42.08 ± 15.35     | 38.95 ± 15.39                  |
| Race or ethnicity    | 7 Black, 17 White | 2 Black, 22 White              |
| Sex                  | 20 male, 4 female | 14 male, 10 female             |
| Postmortem interval, h | 16.54 ± 6.4     | 18.91 ± 6.02                   |
| Brain pH             | 7.01 ± 0.15       | 6.96 ± 0.25                    |

*Values are presented as n or mean ± standard deviation.

Protein and mRNA expression of IL-1β, IL-6 and TNFα

In an earlier study, we determined the protein and mRNA expression of TNFα, IL-1β and IL-6 in the PFC of healthy controls and people who were depressed and died by suicide. Compared to healthy controls, we found a significant increase in the protein and mRNA expression of TNFα, IL-1β and IL-6 in the PFC of people who were depressed and died by suicide (Figure 1A and B).

mRNA expression

We determined the mRNA expression of 3 NLRPs (NLRP1, NLRP3 and NLRP6) using qPCR, followed by mRNA expression studies of some component genes of NLRP3-associated inflammasomes (ASC, caspase1, caspase3 and IL-18); results are shown as fold change (Figure 2A and B). Comparison of the mRNA expression of these genes in both groups using multivariate analysis showed overall significant differences in the mRNA expression of NLRP3 (F<sub>4,49</sub> = 1.331; p = 0.018) and caspase3 (F<sub>4,49</sub> = 2.654; p = 0.045). Univariate comparisons using the Bonferroni post hoc test suggested significant up-regulation of the mRNA expression of NLRP1 (F<sub>4,49</sub> = 5.127; p = 0.028), NLRP3 (F<sub>4,49</sub> = 4.656; p = 0.036) and NLRP6 (F<sub>4,49</sub> = 4.944; p = 0.031) in the PFC of people who were depressed and died by suicide compared to healthy controls (Figure 2A). Univariate comparison tests also revealed significant increases in the mRNA expression of ASC (F<sub>4,49</sub> = 4.377; p = 0.042) and caspase3 (F<sub>4,49</sub> = 6.187; p = 0.017) in the PFC of people who were depressed and died by suicide compared to healthy controls (Figure 2B). However, we observed no significant differences between the 2 groups in the mRNA expression of caspase1 and IL-18 (Figure 2B). Confounding variables such as age, postmortem interval and brain pH showed no significant effects on mRNA expression in either of the 2 study groups.

To examine whether mRNA expression results had a normal bimodal distribution, we performed the Kolmogorov-Smirnov test of normality in SPSS and observed that mRNA expression of NLRP1 in people who were depressed and died by suicide failed the normality test (p ≤ 0.05). NLRP1 in healthy controls and other measures in both groups passed the normality test.

Protein expression

We also studied the protein expression of NLRPs and NLRP3-associated inflammasome protein components using Western blotting; the results are shown as representative blots and histograms in Figure 3 and Figure 4. Statistical analysis of protein levels using multivariate analysis showed significant overall changes in the protein expression of NLRP1 (F<sub>4,49</sub> = 3.856; p = 0.013), NLRP3 (F<sub>4,49</sub> = 3.405; p = 0.016), NLRP6 (F<sub>4,49</sub> = 3.358; p = 0.018; Figure 3B) and ASC (F<sub>4,49</sub> = 4.987; p = 0.002; Figure 4B) in the PFC of people who were depressed and died by suicide compared to healthy controls. Univariate comparison using the Bonferroni post hoc test showed significant increases in the protein expression of NLRP1 (F<sub>4,49</sub> = 13.573; p = 0.001), NLRP3 (F<sub>4,49</sub> = 12.164; p = 0.001) and NLRP6 (F<sub>4,49</sub> = 5.193; p = 0.028) in people who were depressed and died by suicide (Figure 3B). Protein expression of ASC (F<sub>4,49</sub> = 18.796; p < 0.001) and caspase3 (F<sub>4,49</sub> = 3.987; p = 0.05; Figure 4) were also
upregulated significantly in the PFC of people who were depressed and died by suicide, as suggested by representative blots (Figure 4A) and pair-wise comparison testing (Figure 4B). Protein expression of caspase1 (Figure 4B) and IL-18 (Figure 4C) showed no significant differences between the 2 study groups. These results were similar to and consistent with our mRNA results (Figure 2B). We found no significant effects of confounding variables (age, postmortem interval or brain pH) on protein expression in either study group, except for an effect of age on the protein expression of IL-18 ($p = 0.045$) and an effect of postmortem interval on the protein expression of NLRP6 ($p = 0.046$).
Effect of suicide method and antidepressants on mRNA and protein expression

Because some studies have reported that suicide method is an important variable, we determined the effect of suicide method (i.e., gunshot wound, hanging or drug overdose) on our variables using SPSS.

In multivariate analysis, suicide method had no significant effect on the mRNA or protein expression of any of the molecular markers studied. We also determined the effect of 3 suicide methods (gunshot wound, hanging and drug overdose) separately for every marker in those who were depressed and died by suicide, and we observed no significant effects (p ≥ 0.45).

We also examined whether changes in the protein and mRNA expression of NLRPs and NLRP-associated inflammasome molecules that we observed in the PFC of people who were depressed and died by suicide were related to previous treatment, the presence of antidepressant drugs or both. Among those who were depressed and died by suicide 11 of 24 were taking antidepressants at the time of their death. We divided this group into 2 subgroups (those taking antidepressants and those not taking antidepressants) and performed t tests. We found no significant differences in the means of the 2 subgroups for mRNA expression (except NLRP1, p ≤ 0.01) or protein expression, suggesting that apart from NLRP1 mRNA expression, antidepressants had no significant effect on the gene or protein expression of NLRPs and NLRP-associated inflammasome molecules.

Discussion

In the present study we found that the mRNA and protein expression of NLRP1, NLRP3, NLRP6 and ASC were significantly increased in the PFC of people who were depressed and died by suicide compared to healthy controls. However, we found no significant differences between the 2 groups in the expression of caspase1 or IL-18.

Because caspase1 activation increases IL-18, and caspase3 inactivates IL-18 but not IL-1β, it is possible that our failure to find increased IL-18 may have been because of the increased caspase3 we observed. In a previous report, we found that the mRNA and protein expression of IL-1β was significantly increased in people who were depressed and...
died by suicide compared to healthy controls. We did not find significant changes in the expression of IL-18 between the 2 groups, suggesting that IL-18 abnormality is not involved in depression and suicide.

The NLRs comprise 3 different domains: the C-terminal domain, which consists of leucine-rich repeats and is thought to recognize microglial PAMPs and DAMPs or endogenous host molecules; a conserved central nucleotide binding and oligomerization domain (NOD or NACHT) and an N-terminal pyrin domain,72 and a pyrin domain, which is required for its activation.19,20 A unique feature of the NLRPs in general and NLRP3 in particular is that they are involved in forming a multiprotein complex known as an inflammasome.20,29 The NLRP inflammasome consists primarily of a cytosolic sensor molecule NLRP3 or other NLRPs, the adaptor protein ASC and a cytokine protease pro-caspase as the effector molecule.21 It is formed when NLRP3 associates with the adaptor protein ASC and pro-caspase1,19,30 upon activation by different stimuli, such as PAMPs or DAMPs. The activation of inflammasomes results in the cleavage of caspase1 from its pro-form to its enzymatically active form. The activated caspase1 then causes the cleavage of its several substrates, such as pro-IL-1β and pro-IL-18 to active IL-1β and IL-18.29

Although all 3 of the NLRPs we studied form inflammasomes, the NLRP3 inflammasome is the most widely studied at both the basic and clinical levels.19,31 It has also been studied in neurodegenerative disorders such as Alzheimer disease.31–33 To the best of our knowledge, neither the NLRP inflammasomes in people with bipolar disorder by Kim and colleagues,34 who found higher mRNA levels of mitochondrial NLRP3 and ASC in people with bipolar disorder. They also found increased levels of IL-1β, TNFα and caspase1 in the postmortem brains of people with bipolar disorder and schizophrenia.

Alcocer-Gomez and colleagues35 found higher mRNA expression of NLRP3 and ASC in the peripheral blood mononuclear cells and higher protein expression of IL-1β and IL-18 in the serum of patients with major depressive disorder compared to controls. Syed and colleagues36 evaluated the components of the NLRP3 inflammasome and found significantly increased levels of ASC, caspase1 and IL-18, but not IL-1β, in the plasma of patients with major depressive disorder compared to healthy controls.

A role for the NLRP3 inflammasome in depression is also supported by stress-induced depressive behaviour in animal models of depression.37 Mice who were subjected to immobilization or the forced swim test exhibited increased levels of NLRP3 compared to control mice. As well, when stressed, NLRP3-knockout mice showed no depressive behaviour, microglial activation or reduced hippocampal neurogenesis. Thus, NLRP3 inhibition or deletion of microglial activation impairs the stress-induced alterations associated with depression.27

Another piece of evidence suggesting a role for the NLRP3 inflammasome in depression is derived from the studies of Pan and colleagues38 in rats treated with chronic unpredictable mild stress, a model of depression. The rats exhibited depressive-like behaviour, and the authors found increased protein and mRNA levels of NLRP3 and IL-1β in their PFC, suggesting a role for NLRP3 in depression. Thus, both clinical and animal studies suggest the involvement of NLRP inflammasomes in major depressive disorder.

The activation of NLRP3 is a 2-step process of priming and activation, as shown in Figure 5. Priming is achieved by recognition of PAMPs or DAMPs via TLRs or cytokines (such as TNFα or IL-1β), or a combination of both.28,39 The second step consists of the activation of NLRPs by a wide variety of unrelated stimuli, including several DAMPs (such as ATP, urate or aluminum) and PAMPs (such as lipopolysaccharide, peptidoglycan or several toxins; see reviews by Swanson and colleagues39 and Vanaja and colleagues40).

Thus, there is distinct cooperation in the activation of NLRP3 inflammasomes between TLRs, cytokines and inflammasomes. The priming signal is provided by TLRs and cytokines such as TNFα with the formation of pro-IL-1β and pro-IL-18 (the substrates for the NLRP inflammasome),28 which are formed as a result of TLR interaction with PAMPs and DAMPs and the activation of nuclear factor xB (NF-xB). Our studies showing abnormalities of specific TLRs in the PFC of people with depression who died by suicide17,18 confirmed that the priming signal for the activation of NLRP inflammasomes is increased, leading increased activation of NLRP inflammasomes themselves. As well, the main function of priming is to induce pro-IL-1β and potentiate NLRP3 inflammasome activity through NF-xB-dependent induction of NLRP3, which may be a limiting component of the complex.40 Thus, it is primarily the upregulation of NLRPs (and probably ASC) that causes the IL-1β increase we have observed in the PFC of people who were depressed and died by suicide.5

Although the NLRP3 inflammasome has been more widely studied and characterized, NLRP1 and NLRP6 also form inflammasomes. In fact, NLRP1 inflammasomes were the first NLRP inflammasome to be characterized.20 Although NLRP1 does not need ASC to form the inflammasome and can interact directly with pro-caspase1 and bypass the ASC requirement, inclusion of ASC in the complex augments the activation of the human NLRP1 inflammasome.15,42 NLRP1 also interacts with caspase5, which may contribute to IL-1β processing in humans.20

NLRP6 forms an inflammasome as well, but the function of NLRP6 is not well known. This novel molecule was initially characterized as a regulator of NF-xB and caspase1.43 It functions as a positive activator of NF-xB when coexpressed with ASC, and it induces cytokine production. The increase in NLRP6 we found may not only cause the maturation of IL-1β but also lead to the production of other cytokines, such as TNFα.15,42,43

ASC plays a central role in the activation of caspase1 and the secretion of IL-1β and IL-18 in response to PAMPs and DAMPs. The activation of caspase1 and the production of IL-1β
and IL-18 is abolished in ASC-deficient macrophages, showing a crucial role for ASC in the IL-1β and IL-18 formation.

The other question is the cause and consequences of NLRP3 inflammasomes. The main causes of NLRP3 inflammasomes may be related to environmental factors, such as exposure to PAMPs and DAMPs. Exposure of TLRs to PAMP ligands, such as lipopolysaccharides produced by pathogens or mitochondria, may be a factor in activating NLRPs (see review by Vanaja and colleagues). The generation of ATP through DAMPs, as well as exposure to toxic materials such as potassium ATP or mitochondrial-derived reactive oxygen species or uric acid, may also be factors involved in the activation of NLRP3. Increased levels of reactive oxygen species and mitochondrial dysfunction have also been found in the post-mortem brains of people with bipolar disorder, another factor for the upregulation of NLRP inflammasomes.

Inflammasome activation has behavioural, clinical and biological consequences. For example, active inflammasomes and increased IL-1β have been associated with a loss of cognitive function. NLRP3 or caspase1 deficiency substantially attenuates the impairment of spatial memory. In addition to the behavioural abnormalities caused by NLRP inflammasomes and IL-1β, the latter also causes many biological effects. IL-1β plays an important role in the initiation and continuation of the inflammatory reaction in the central nervous system in response to various adverse stimuli. It also reduces the integrity of the blood–brain barrier, allowing the infiltration of peripheral immune cells into the central nervous system.

However, one of the specific effects of IL-1β is an increase in a process known as pyroptosis. Increased levels of caspase1 activated by the NLRP3 inflammasome can result in a highly inflammatory form of cell death known as pyroptosis. Pyroptosis occurs most frequently after infection with intracellular pathogens. The process of pyroptosis is exclusively mediated by caspase1, is different from apoptosis and occurs independently of proapoptotic caspases. The major consequences of increased inflammasome activity that we observed in people who were depressed and died by suicide may have been increased cell death mediated by pyroptosis in the brain.
Abnormalities in IL-1β, IL-18 and NLRP3 inflammasomes have been observed in neurodegenerative disorders such as Alzheimer disease. A recent report showed that protein and mRNA levels of IL-18 were significantly increased in the post-mortem brains of people with Alzheimer disease, especially in the frontal lobe. Heneka and colleagues reported that expression of NLRP3 and caspase1 was significantly increased in the post-mortem brains of people with Alzheimer disease.

The observation that NLRP3 inflammasomes and IL-1β in were increased the brains of people who were depressed and died by suicide provides a potential therapeutic target. NLRP3 may be a unique target in several ways. Abnormalities in cytokines and TLRs enhance the activation of NLRP3. Some studies have identified compounds that may block IL-1 signalling or cause the inhibition of NLRP3 inflammasome activation. Several compounds related to the IL-1 receptor and NLRP3 are being tested in conditions such as inflammatory and neurodegenerative disorders.

**Limitations**

This study had 2 main limitations. We did not have information about body mass index for the people in our study groups, and we were not certain if any of the observed changes in NLRP and its inflammasomes were related, at least in part, to body mass index. Some people were taking antidepressant treatment at the time of their death. Although we did not observe differences in the gene and protein expression of NLRPs and NLRP-associated inflammasome molecules between those who were taking antidepressants and those who were not, this is one of the limitations of a postmortem brain study.

**Conclusion**

The innate immunity system in the brain is mediated by subfamilies of innate immunity receptors such as TLRs and NLRPs — also known as pattern recognition receptors — which recognize PAMPs and DAMPs. Interaction of PAMPs and DAMPs with these receptors initiates a signalling mechanism, resulting in the activation of NF-kB and the production of cytokines such as pro-IL-1, pro-IL-18, IL-6 cytokines and other chemokines. Our studies have indicated increased NLRP1, NLRP3, NLRP6 and IL-1β signalling; the upregulation of NLRP1, NLRP3 and NLRP6 in general; and the upregulation of inflammasomes, consisting of ASC and caspase1 in particular. Activation of this multiprotein complex by PAMPs and DAMPs causes the activation and cleavage of caspase1; active caspase1 in turn cleaves pro-IL-1β and pro-IL-18, resulting in the formation of IL-1β and IL-18, which perform many important functions in the central nervous system, including pyroptosis (programmed cell death, which is different than apoptosis). These results not only provide a mechanism for the involvement of NLRPs in the pathophysiology of depression and suicide, but they also show great potential for developing specific therapeutics to treat depression and suicidal behaviour, especially in those who are unresponsive to antidepressants.

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**Contributors:** G. Pandey designed the study. H. Zhang and X. Ren acquired the data, which A. Sharma analyzed. G. Pandey wrote the article, which H. Zhang, A. Sharma and X. Ren reviewed. All authors approved the final version to be published and can certify that no other individuals not listed as authors have made substantial contributions to the paper.

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NLRP inflammasomes in depression and suicide

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