The effects of stress on hypothalamic-pituitary-adrenal (HPA) axis function in subjects with schizophrenia

Os efeitos do estresse na função do eixo hipotalâmico-pituitário-adrenal em indivíduos com esquizofrenia

FRANCESCA L. GUEST1, DANIEL MARTINS-DE-SOUZA2,3, HASSAN RAHMOUNE2, SABINE BAHN2,4, PAUL C. GUEST2

1 Faculty of Medicine and Dentistry, University of Bristol, Bristol, UK.
2 Department of Chemical Engineering and Biotechnology, University of Cambridge, Cambridge, UK.
3 Max Planck Institute of Psychiatry & Ludwig Maximilians University (LMU), Munich, Germany; Laboratory of Neurosciences (LM-27), Institute of Psychiatry, School of Medicine, University of Sao Paulo (IPy-FMUSP), Sao Paulo, SP, Brazil.
4 Department of Neuroscience, Erasmus Medical Centre, The Netherlands.

Received: 9/23/2012 - Accepted: 11/7/2012

Abstract

Over the last few decades, evidence has been emerging that the pathogenesis of psychiatric disorders such as schizophrenia can involve perturbations of the hypothalamic-pituitary-adrenal (HPA) axis. Variations in the manifestation of these effects could be related to the differences in clinical symptoms between affected individuals as well as to differences in treatment response. Such effects can also arise from the complex interaction between genes and environmental factors. Here, we review the effects of maternal stress on abnormalities in HPA axis regulation and the development of psychiatric disorders including schizophrenia. Studies in this area may prove critical for increasing our understanding of the multi-dimensional nature of schizophrenia. Further research in this area could ultimately lead to the development of improved diagnostics and novel therapeutic approaches for treating this debilitating psychiatric condition.

Keywords: Psychiatric disorders, schizophrenia, HPA axis dysfunction, diagnosis, biomarkers.

Introdução

During a stressful event, chemicals such as adrenaline and glucocorticoids are released into the bloodstream leading to a chain reaction of events inside the body. The effect of these chemicals from stress situations experienced during pregnancy has also been investigated. David Barker proposed the “fetal programming hypothesis” – also known simply as the Barker hypothesis – in his paper of 1989, which details the importance of the intrauterine environment in the development of the fetal organs and tissues1. This hypothesis suggests that deviation from the ideal intrauterine environment may have lasting effects on organ structure and function1–2. Extrapolation of this concept to the brain suggests that the intrauterine environment during the first and second trimesters of pregnancy will have direct effects on the structure of the brain in the fetus, which may continue on into adulthood.

Correlations between prenatal maternal stress and behavioral and psychological abnormalities in animal offspring have been widely documented. It has been proposed that hormones released into the bloodstream in response to a stressful stimuli perceived by the mother, have a direct effect on development of the brain and other organs in the fetus. The correct regulation of hormone release in pregnancy is critical for correct fetal development and any deviation from the normal concentration of these hormones can produce microscopic and macroscopic changes in the brain, particularly in synaptic connectivity within and across distinct brain regions3. However, it is not one acute stressor, but chronic stress which predisposes to psychiatric disorders in the offspring. The development of behavioral and psychological problems including autism, attention-deficit hyperactivity disorder (ADHD), major depressive disorder, bipolar disorder and schizophrenia are thought to be linked to such perturbations in the hypothalamic-pituitary-adrenal (HPA) axis and in other organs of the diffuse neuroendocrine system.

From the research carried out by others it is clear that there is some correlation between increased maternal stress and behavioural and psychological problems in the offspring, but this is not thought to be purely hormonal. It also speculated that genetic links may exist for many of these diseases and that these may be, in some way, hereditary. It should also be considered that mothers with psychological problems or who have problems with stress management may be more likely to become stressed during pregnancy and thus induce a similar condition in their offspring. What is more likely is that these problems have both hormonal and genetic causes. The two hit hypothesis4 suggests that a genetic susceptibility exists in the fetus and then an anomalous hormonal environment in the uterus allows an abnormality to come to fruition.

This review aims to summarize and correlate the existing literature on the effect of prenatal stress on development of the fetal brain and how this may affect the behavior, psychology and overall health of the offspring later on in life. As much of the current literature...
reports the use of animal models to test these hypotheses, we will speculate on the extent to which these findings can be extrapolated to humans. We will also propose the likely effects of prenatal stress on the development of psychiatric disorders in humans and discuss the reported effects of this on the function of the HPA axis and other neuroendocrine systems. Finally, we will address the importance of the timing of the stress stimuli with respect to the gestational period in the mother as well as the impact of this on the existence and severity of behavioral problems in the offspring.

Stress: definitions

The book *Neuroendocrinology, an integrated approach* describes stress as "any event, whether real or perceived, that acts to disturb the homeostatic balance". When we think of stress in everyday life, we tend to think of emotional stress. This could mean the effects of big deadlines, crucial examinations, overbearing bosses and important decisions which can all be included as stressful stimuli. Not everyone responds to stress in the same way nor do they have the same stress threshold. Some people will become stressed only in extreme situations, whereas others could become overly-stressed over seemingly small matters. Stress can be evoked by fear of the unforeseen, the unknown or even possible bad outcomes. For example, the fear that a loved one will have a serious injury, even though there is no evidence to suggest that this may happen. The most intense emotional stressors occur in close relationships, such as those between family members or partners. Other types of stress include those of a mechanical or physical nature, such as impact injuries. Although this can lead to emotional stress in many circumstances, mechanical stress is not necessarily relevant to the subject of this review since this is focused on the physical nature, such as impact injuries. Although this can lead to emotional stress in many circumstances, mechanical stress is not necessarily relevant to the subject of this review since this is focused on the physical nature, such as impact injuries.

The stress response in humans

In reaction to a stress-producing stimulus, the body produces a response via activation of the HPA axis. This is a necessary mechanism as it prepares the body for the well-known "fight or flight" reaction to a potentially dangerous situation. However, the stress response that is produced does not differ greatly depending on the stressor. In humans the molecules greatly responsible for the stress response are adrenaline and glucocorticoids. There are three main phases of the generalized stress response that are set into motion when a stressor is detected. The first phase involves the production of the catecholamine molecules noradrenaline and adrenaline. These are examples of monoamine hormones, which are derived from the amino acid tyrosine. The conversion from tyrosine to dihydroxyphenylalanine (DOPA) takes place in chromaffin cells found in the adrenal medulla. Noradrenaline and adrenaline are released from the adrenal medulla almost instantaneously into the blood stream via the autonomic nervous system and affect a number of responses in the body. For example, the release of noradrenaline and adrenaline causes an increase in heart rate, constriction of blood vessels, widening of the pupils (midriasis), and an opening of respiratory passages. In addition insulin secretion is inhibited and glycojenolysis is stimulated in the muscles and liver, leading to a higher concentration of glucose in the blood stream.

Secondly, around 20 minutes afterwards, corticotrophin-releasing factor (CRF) is released from the paraventricular nucleus (PVN) of the hypothalamus into the portal vessel system and median eminence of the hypothalamus. This, in turn, stimulates the release of adrenocorticotropic hormone (ACTH) from the anterior pituitary directly into the blood stream. ACTH travels through the circulation to the adrenal glands, culminating in the release of glucocorticoids such as cortisol from the adrenal cortex. With respect to prenatal stress in humans, cortisol is the most important glucocorticoid. The elevation of cortisol levels in the blood results in the inhibition of the release of ACTH and CRF in a negative feedback loop mechanism.

In the third stage, glucocorticoids produce similar responses to those of the catecholamines, including increased gluconeogenesis and increased lipolysis, leading to increased volumes of glycerol and fatty acids in the blood stream. Glucocorticoids also stimulate the mobilization of fats from skeletal muscle and their deamination.

Fetal programming in response to stress

Chronic prenatal stress is linked with morphological changes in the fetal brain but the mechanisms underlying these effects are not well known. The maternal and fetal HPA axes and placenta have been studied to determine whether these can serve as conduits for such morphological effects. During pregnancy the mother's body undergoes many physical and chemical changes, including altered production of certain hormones. The levels of cortisol are typically found to be elevated in a pregnant mother. Such an elevation of cortisol is essential for the growth of the fetus and it stimulates the production of surfactant. However excessively elevated levels of cortisol can permanently modify the growth of the fetus. Stress alters the levels of many hormones and other molecules found in the blood and this can lead to changes in the fetus. Prenatal stress affects the placenta in particular, decreasing the function of this organ via decreased levels of nutrients and oxygen that reaches the fetus. The mechanism of this effect results from increased levels of adrenaline in the blood which induces increased vascular resistance, and restriction of the flow of blood to the placenta.

The fetus has a natural barrier enzyme to maternal cortisol called 11β-hydroxysteroid dehydrogenase-2 (11β-HSD2). This molecule is down-regulated in instances of maternal stress, leaving the fetus more susceptible to cortisol exposure. The reduction in 11β-HSD2 also leads to an increase in the production of other hormones such as prostaglandins, oestrogens, and placental lactogen. Also, placental CRH levels are subject to regulation by the maternal HPA axis. Studies have shown that stress-induced increases in CRH concentrations can lead to macroscopic and microscopic changes in the hippocampus of the offspring.

Behavioral and psychological problems

A number of different psychological problems have been shown to be linked to antenatal stress. An increased incidence in attention deficit hyperactivity disorder (ADHD) has been found in rhesus monkey offspring when mothers were exposed to loud noises during pregnancy. Studies have shown that the offspring of mothers exposed to long term stress showed greater responses to stressors, abnormalities in coping with stress and disturbed social behaviour. The incidence of ADHD was also found to be increased in offspring whose mothers experienced bereavement during pregnancy, as shown in a Danish study. In this case, unexpected bereavement resulted in a 72% increase in the likelihood of ADHD occurring in the offspring. Many other researchers have reported a greater prevalence of ADHD in the offspring of stressed mothers.

A number of reports have suggested that there is an increased incidence of schizophrenia in the offspring of prenatally-stressed mothers. The German army invasion of the Netherlands in May 1940, led to nationwide stress. A cohort consisting of the offspring...
of mothers exposed to this stress during the first, second and third trimesters of pregnancy were followed up and the prevalence of schizophrenia observed. This revealed a greater incidence of schizophrenia in these subjects, with the largest increase seen in those whose mothers were exposed in the first trimester. Similar findings resulted from studies of the Chinese Famine of 1958-1961. Again, in those individuals conceived at the peak of the famine showed an increased risk of schizophrenia and other conditions. Increased incidence of schizophrenia in the offspring of stressed mothers has also been reported by Huttenen et al., Kinney et al., and others.

Further evidence supporting the importance of intrauterine environment in the precipitation of psychiatric illnesses is seen in animal models. For example, studies involving the deprivation of protein in prenatal rats have shown that this can have long term effects on the brain in the offspring. These effects include changes in hippocampal morphology and glutamate and dopamine receptor binding. Other studies have shown that restriction of protein intake during pregnancy may have a negative effect on fetal brain development. This is thought to be due to changes in maternal metabolism of lipids, which are an essential component of cellular membranes. Interestingly, the behavioral abnormalities seen in these protein restricted models do not manifest until adulthood, including effects on brain-regulated activities such as decreased pre-pulse inhibition of the startle response and hyper-locomotion, which are traditionally used as a measure of schizophrenia-like behavior. A pre-pulse is a reduced version of a stimulus delivered to an individual shortly before a full version of the stimulus (pulse) is applied. The brain will normally down regulate its startle response to a stimulus if this is preceded by a pre-pulse. However, individuals with schizophrenia or Alzheimer’s disease will show a startle response of the same magnitude regardless of the presence of a pre-pulse.

Watson et al. reported a 13.3% incidence of depression in the offspring of mothers who were exposed to a severe earthquake compared to an incidence of 5.5% seen in the offspring on non-exposed mothers. Depression has also been linked to abnormalities seen in the HPA axis of rat dams suffering from prenatal stress. After Louisiana was hit by a bout of hurricanes and tropical storms, a study examining the difference between the incidences of autism in the offspring of mothers who were carriers was carried out. This showed an incidence of 26.5/10,000 in the high exposure group, compared with that of 3.72/10,000 in the low exposure group. Beversdorf et al. also established a link between prenatal stress and autism. A number of other noteworthy fetal abnormalities have been linked with prenatal stress seen in the mother including cognitive, behavioural and emotional issues. Interestingly, rat studies have shown that the offspring of mothers who were restrained in the last week of pregnancy are more likely to self-administer drugs, such as cocaine and amphetamines, than those from non-restrained mothers.

Microscopic and macroscopic changes in the brain

Specific regions of the brain have been shown to be affected by prenatal stress both macroscopically and microscopically. These regions include the hippocampus, amygdala, corpus callosum, cerebral cortex, cerebellum and hypothalamus. Since morphological changes in the above brain regions have been linked with certain psychological and behavioral problems, it is possible that prenatal stress affects the neurodevelopmental formation of these areas. This would, in turn, be expected to have behavioral effects, such as those seen in psychiatric disorders.

Changes in the size of the corpus callosum and the number of specific cell types within this brain structure have been linked with autism, ADHD and schizophrenia. Links of autism, ADHD and schizophrenia to prenatal stress have been detailed above. It may therefore be reasonable to extrapolate these findings to suggest that prenatal stress directly affects the structure of the corpus callosum. However, no studies in this area have been carried out in humans.

The hippocampus is known to be involved in the formation of memories and plays a major role in learning. Lemaire et al. reported that prenatal stress had a negative effect on the memory of rats. The hippocampal granule neurons continue to be created throughout life and are responsible for the formation of memories. However, the number of such granule neurons was reduced in adult rats with mothers that had been exposed to stress while pregnant. In addition, the authors showed that prenatal stress reduced hippocampal cell proliferation and survival rate of newborn cells, along with a reduced number of differentiated neurons. Interestingly, all of these deleterious effects could be countered by neonatal handling.

Although animal models have given us some insight into the connection between morphological changes in the brain, prenatal stress and behavioral disorders, it cannot be assumed that the same effects occur in humans. In fact, this is not likely to occur in some cases due to the fact that there are large differences with respect to the extent to which the brain undergoes development before, during and after birth in different animal species.

Timing and severity

In rats and non-human primates, the timing of the stressor in the pregnancy has been shown to be important. A study using rats has shown a 64% increase in the production of corticosterone – a similar hormone to aldosterone in humans – after handling pregnant rats in the last week of gestation and placing them in new, unfamiliar cages. Although some evidence has been put forward in the case of animal models, the timing and severity of a stressor which affects the human fetus is harder to determine. This difficulty is mainly due to the fact that it would be unethical to expose pregnant mothers to stress at different points during their pregnancy. However, natural disasters can be used for this purpose; such events can provide a large cohort of people affected by an identical stress. This allows retrospective assessment of those who were pregnant during the natural disaster and follow up studies of the offspring. In this way, we can assess whether any of the patterns seen in animal models can be extrapolated to humans.

A study termed “Project Ice Storm” attempted to determine the timing of the effects of maternal stress on the offspring using human subjects. In this case over 3 million people were unfortunately exposed to extreme cold due to power outages. This study revealed an abnormality in fingerprint formation in offspring of mothers exposed to the storm between weeks 14 and 22 of pregnancy. Interestingly, the period of fingerprint formation is known to overlap with the formation of the hippocampus. Therefore, it is possible that changes in the hippocampus may be seen in the same individuals. The project is still ongoing and is currently utilizing a magnetic resonance imaging (MRI) approach to determine whether behavioural and psychological problems seen in the offspring are linked to any changes in brain morphology. As the hippocampus is responsible for memory formation and cognitive reasoning, this region is of primary interest. It must be acknowledged, that the effects of a particular stressor on a mother depends not only on the nature of the stressor itself but also on the severity of the perceived threat of the stressor by the mother and on the mother’s stress tolerance and behavior in response to that stressor.

Potential advantages of stress

An interesting and potentially useful aspect of the stress response is that during acute stress the hippocampal dentate gyrus region is more active. This is the area of the brain responsible for learning, memory and neurogenesis. In response to a stressful stimulus the neural progenitor cells in the dentate gyrus are transformed into neurons and glia, which eventually lead to new synaptic connections. It is supposed that this is at least part of the mechanism of how memories are made and this process enables subjects to remember the object or event that caused the stress. An emotional input from the amygdala and the recognition of stress by the hippocampus cooperate to form a memory of fear and this will hopefully lead the individual to avoid similar dangerous situations in the future.
In contrast, chronic stress appears to decrease the ability to form new memories. Conrad et al. showed that chronically-stressed rats performed less well on a spatial maze test than their non-stressed counterparts. In addition, studies in humans have shown that years of stressful living can lead to increased incidence of depression and other psychiatric conditions. This leads to the possibility that acute stress can be beneficial and chronic stress detrimental.

**Effects of stress on insulin resistance and the HPA axis**

In humans, prenatal maternal stress during pregnancy has been shown to predict low birth weight and pre-term delivery. Studies have shown that this is, in turn, linked with metabolic dysfunctions such as decreased glucose tolerance and increased insulin resistance. Using an oral glucose tolerance test, a recent study analyzed the glucose levels of young adults whose mothers had experienced stressful life events during pregnancy such as relationship conflicts, death or severe illness of a close relative or friend, severe financial difficulties and car accidents, in comparison to control subjects whose mothers had relatively stress-free pregnancies. The results showed no significant difference between the glucose levels of the two groups, although the subjects with mothers that suffered from prenatal stress had significantly higher insulin levels at the 120 minute time point after administration of the glucose tolerance test.

Interestingly, another study by the same authors showed that adults whose mothers were stressed during pregnancy had increased cortisol levels in response to the Trier Social Stress Test (TSST) with a decreased cortisol response after administration of an ACTH stimulation test, indicating a possible dysregulation of the HPA axis. This provided evidence in humans of an association between prenatal stress exposure and alterations of HPA axis function in the offspring (Figure 1).

**Figure 1.** Schematic diagram showing interaction between genetic predisposition and early life environmental factors on neuronal function and later development of psychiatric illnesses.

A study using rat models showed that prenatal stress induced long-term changes in feeding behavior, glucose metabolism and insulin signalling. These deviations from normal glucose and insulin handling are similar to problems seen in type II diabetes. The authors speculated that this was linked to the increased levels of glucocorticoids in the intrauterine environment that accompanies prenatal stress. Another study tested the effects of administering stress hormones directly into sheep in the early stages of pregnancy, with particular interest in the effect that this had on the regulation of glucose and insulin signalling in adult male offspring. The study showed that the adult offspring of mothers which were administered stress hormones during pregnancy had an impaired glucose tolerance and hyperinsulinaemia. This suggested that glucocorticoid exposure in early pregnancy might lead to long-term pancreatic problems and diabetes, consistent with the fact that stress during early pregnancy contributes to such outcomes in humans.

**Effects on the insulin signalling and the HPA axis in schizophrenia**

Despite decades of research, the pathophysiology and aetiology of schizophrenia and other psychiatric disorders are not completely understood. The main hypotheses have focused on alterations in neurotransmitter systems such as the glutamatergic and dopaminergic pathways and current antipsychotic medications mainly target these systems. However, schizophrenia is often associated with peripheral manifestations including hyperinsulinaemia and type II diabetes mellitus. Although these effects can result from antipsychotic medications, they were also observed decades before the development and clinical use of antipsychotics. In addition, recent evidence has emerged that patients can show these effects at the first clinical presentation and prior to receiving medication and blood-based analyses have demonstrated hyperinsulinaemia and abnormalities in secretion of other endocrine factors at first presentation of symptoms.

In a recent study of 66 first onset schizophrenia patients and 68 matched control subjects, we used a series of immunoassays to measure the levels of insulin, proinsulin and des 31,32 proinsulin using two-site time-resolved fluorescence assays employing different combinations of monoclonal antibodies that discriminate between the specific forms of the molecule. Also, C-peptide and the insulin secretory granule protein chromogranin A were measured using commercially-available immunoassays. All of these molecules were found at significantly elevated levels in the circulation of the schizophrenia patients. In contrast, the glucose levels were relatively normal which was consistent with the possibility that at least some of these patients were insulin resistant at the onset of the disease. This could have important implications since elevated insulin can have deleterious effects on brain function.

Assuming that the finding of increased insulin levels in schizophrenia patients is associated with impaired insulin signalling, we also tested the possibility that secretion of other hormones of the diffuse neuroendocrine system and the HPA axis are affected in schizophrenia. We carried out multiplex immunoassay analysis of 21 hormones and hormone-related molecules using blood serum from 236 first onset schizophrenia patients and 230 matched controls using the Multi-Analyte Profiling (MAPTM) platform from Rules Based Medicine (Austin, TX, USA). Analysis using the multiplex immunoassay technology revealed that the serum concentrations of insulin and chromogranin A were increased in schizophrenia subjects, consistent with the above findings. In addition, we found elevated concentations of pancreatic polypeptide, prolactin, progesterone and cortisol, and decreased levels of growth hormone (Figure 2).

Other researchers have identified higher levels of CRH, arginine vasopressin (AVP), ACTH and cortisol in studies of schizophrenia and other psychiatric conditions. Functional magnetic resonance imaging (fMRI) studies have also identified larger pituitary volumes in first episode schizophrenia patients and pituitary abnormalities in chronic sufferers have been shown to be smaller, which may indicate adaptation, medication effects or desensitization to HPA hyperactivity.

Many hormones are affected by ultradian and circadian rhythms. Therefore, it is likely that the molecules identified in the above studies are co-regulated in a feedforward-feedback loop between the endocrine pancreas, pituitary and other tissues of the HPA and hypothalamic-pituitary-gonadal (HPG) systems. As a case in point, increased insulin levels have been associated with increased prolactin secretion and decreased growth hormone secretion pulses. Furthermore, studies using a diet-induced insulin resistance rat model have found increased insulin and progesterone levels. Also, we have previously reported the co-regulated expression levels of insulin, growth hormone, leptin and cortisol in first onset schizophrenia patients, using a targeted analyze cluster method which detects patterned behavior. The finding of increased cortisol levels using this method is consistent with other studies.
previous studies have linked abnormal AVP levels to the disease. The finding of increased ACTH and cortisol supports the hypothesis that these diverse methods we identified changes in cortisol, ACTH, AVP, agouti-related protein, growth hormone, prolactin and secretagogin.

We also analyzed post-mortem pituitaries from schizophrenia patients using a combination of liquid chromatography tandem mass spectrometry (LC-MS), two dimensional difference in-gel electrophoresis (2D-DIGE) and multiplex immunoassays. Using these diverse methods we identified changes in cortisol, ACTH, AVP, agouti-related protein, growth hormone, prolactin and secretagogin.

The finding of increased ACTH and cortisol supports the hypothesis that HPA axis hyperactivity may be involved in the pathophysiology of the disease. Previous studies have linked abnormal AVP levels to changes in mood and behaviour, and to psychotic disorders. In addition, AVP may affect HPA axis sensitivity since there appears to be a positive correlation between the circulating levels of AVP, ACTH and cortisol in schizophrenia patients. The finding that some of these changes were also detected in the circulation of living schizophrenia subjects (see above) suggests that they may play a role in the pathophysiology of the disease and could also lead to translation of these molecules as peripheral biomarkers for schizophrenia.

Therapeutic implications

The discovery that hyperinsulinaemia might play a role in late onset disorders in individuals with prenatally-stressed mothers, suggests that drugs which improve insulin signalling and glucose handling, may represent a potential novel treatment strategy. In the case of psychiatric disorders, such as schizophrenia, therapeutic strategies that target the underlying metabolic dysfunction could provide an effective alternative to traditional antipsychotic medications. This possibility is supported by the finding that the insulin-sensitizing agents metformin and rosiglitazone can correct the insulin resistance that seems to come hand in hand with antipsychotic treatment without inhibiting the antipsychotic drug efficacy in treating the disease.

Similar strategies are already proving fruitful for treatment of memory deficits in Alzheimer’s disease. Clinical trials are focusing on the use of gamma peroxisome proliferator-activated receptor (PPARγ) agonists such as rosiglitazone and pioglitazone as an alternative therapy to enhance cognition. PPARγ agonists induce the transcription of specific genes leading to the increase in the body’s sensitivity to insulin, amongst other effects, and were first used in clinical trials with the aim of treating diabetes mellitus and atherosclerosis. One research group conducted a 6-month, randomized controlled trial in patients suffering from mild Alzheimer’s disease and type II diabetes. Patients were assigned randomly to one of two groups. One group was treated with 15-30 mg pioglitazone once a day and the other was used as a control. The pioglitazone group showed improved cognitive function and increased blood flow in the parietal lobe of the brain and the control group showed no improvements. It was interesting that the effects of pioglitazone were accompanied by an increase in cerebral blood flow as this could potentially lead to an increase in the amount of glucose available in the brain.

The adrenal steroid dehydroepiandrosterone (DHEA) has antiglucocorticoid properties that may prove useful regulating high cortisol levels and glucocorticoid action in the brains of psychiatric patients. Studies using DHEA in addition to antipsychotic medication in already-medicated schizophrenic patients found a significant improvement in the negative, depressive and anxiety symptoms of the disease. DHEA treatment has also produced improvement in extrapyramidal side-effects such as involuntary tremors associated with antipsychotic treatment.

Conclusions and future prospects

Prenatal stress has been linked with many psychological and behavioral problems such as schizophrenia, ADHD, autism, and depression. Changes are seen in the brains of a variety of animal models in response to prenatal stress. In particular, changes are seen in the hippocampus, amygdala, corpus callosum, cerebral cortex, cerebellum and hypothalamus. These areas of the brain are responsible for the control of behavior and their alteration could explain the psychological problems witnessed, although this has not been proven to occur in humans. The mechanisms by which these changes in the brain are achieved most likely surround the maternal and fetal HPA axes and the effect of the intrauterine environment on fetal brain development. Regardless of the effect of environment on the developing fetus, it is likely that there is also a genetic element to the contraction of such problems.

It appears that the timing and severity of the stress experienced by mothers has an impact on the development of psychological and behavioural problems in the fetus. However, further research is needed before the exact effects of timing and severity of this can be determined. There is an ever-increasing amount of evidence for metabolic and hormonal components in conditions such as schizophrenia, which in some cases may be associated with prenatal stress. Abnormalities in the metabolism of glucose, insulin signalling and the HPA axis appear to be present in the early stages of these disorders and may provide the basis for the development of much-needed biomarkers for psychiatric conditions. The use of such biomarkers could lead to improved diagnosis and patient orientated personalized medicine strategies (the right drug for the right patient at the right time) as well as providing the possibility of pre-emptive treatment. Given the potential of this line of research to improve diagnosis and create alternative treatment strategies, more research is warranted.

It is clear however that the environmental trigger of stress in the intrauterine environment is not the only catalyst for development of the disease. It is likely that there is also a genetic predisposition that when combined with an environmental trigger leads to disease development. This is in accordance with the two hit hypothesis. We have the tendency to dismiss our “emotional” responses like sadness or stress as they are not tangible. However, everything that we feel is due to the movement of chemicals within us. These chemicals, neurotransmitters and hormones, produce the feeling of stress but also lead to a cascade of other reactions in the body, which can have very real and tangible outcomes.

Acknowledgements

This research was supported by the Stanley Medical Research Institute (SMRI) and the European Union FP7 SchizDX research programme (grant reference 223427).

References


