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Authors	Cowan, Caitlin S. M.;Callaghan, Bridget L.;Kan, Janice M.;Richardson, Rick
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**The lasting impact of early-life adversity on individuals and their descendants:  
Potential mechanisms and hope for intervention**

Caitlin S. M. Cowan<sup>1\*</sup>, Bridget L. Callaghan<sup>2^</sup>, Janice M. Kan<sup>1^</sup>, and Rick  
Richardson<sup>1</sup>

<sup>1</sup> School of Psychology, The University of New South Wales, Sydney, Australia 2052

<sup>2</sup> Psychology Department, Columbia University, New York City USA

Email addresses: [c.cowan@unsw.edu.au](mailto:c.cowan@unsw.edu.au); [bridgetcallaghan281@gmail.com](mailto:bridgetcallaghan281@gmail.com);  
[janice.kan@unsw.edu.au](mailto:janice.kan@unsw.edu.au); [r.richardson@unsw.edu.au](mailto:r.richardson@unsw.edu.au)

\* indicates the corresponding author; email: [c.cowan@unsw.edu.au](mailto:c.cowan@unsw.edu.au)

^ indicates authors that made equal contributions

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treatment

## **Abstract**

The adverse effects of early-life stress are pervasive, with well-established mental and physical health consequences for exposed individuals. The impact of early adverse experiences is also highly persistent, with documented increases in risk for mental illness across the lifespan that are accompanied by stable alterations in neural function and hormonal responses to stress. Here, we review some of these “stress phenotypes”, with a focus on long-term mental health outcomes and potential intermediary factors such as altered development of the fear regulation system. Intriguingly, recent research suggests that such stress phenotypes may persist even beyond the lifespan of the individual, with consequences for their offspring and grand-offspring. Phenotypic characteristics may be transmitted to future generations via either the matriline or the patriline, a phenomenon that has been demonstrated in both human and animal studies. In this review, we highlight behavioral and epigenetic factors that may contribute to this multigenerational transmission and discuss the potential of various treatment approaches that may halt the cycle of stress phenotypes.

It is widely recognized that early experiences often have a profound impact on an individual's functioning across their lifespan. For example, the median age of onset for mental health disorders as a whole is 14 years, with the majority of adult disorders emerging during childhood to adolescent years (Jones, 2013, Kessler *et al.*, 2007, Lee *et al.*, 2014). Further, certain factors within the home (such as parental mental illness, criminality, violence, neglect) are strongly associated with the onset of child/adolescent mental illness, accounting for nearly half of all childhood-onset, and nearly a third of later-onset, mental health disorders (Green *et al.*, 2010). These data suggest that the experience of early life adversity renders individuals prone to psychopathology across the lifespan. While nearly all studies on mental health (across species) focus on end-state outcomes, that is, when disordered functioning is already present (Thompson & Levitt, 2010), understanding the developmental pathway towards aberrant outcomes is highly clinically relevant in order to establish early markers for identifying and treating at-risk individuals. For this reason, the focus of the first section of the current paper is to review some potential early development markers of risk associated with early life stress. This is followed by a discussion of the persistence of stress-induced phenotypes across generations as well as potential mechanisms for these generational effects. Finally, given the clinically significant outcomes associated with early adversity, some potential targets for intervention are discussed.

## **Early Adversity & the Development of the Individual**

### *Animal Models*

The importance of animal models is accentuated in adversity research, a field that is fraught with logistical and ethical complications when considering studies of human populations. As such, animal models have been highly informative in establishing some of the immediate, short-term, and longer-term consequences of early adversity, and in providing insights into the developmental pathways that might increase risk for mental illness across the

lifespan (see Meyer & Hamel, 2014, Sanchez *et al.*, 2001 for reviews). Here, we focus on work (conducted mainly in rodents) suggesting that early adversity has specific effects on the development of emotion learning systems (such as fear and extinction) that may place affected individuals at risk for dysfunction later in life.

In some early rodent studies demonstrating the effect of stress on the developing fear system, it was shown that exposure to stress or glucocorticoids within the first few weeks of postnatal life resulted in an early transition from approach to avoidance responses towards a shock-paired odor (e.g., Moriceau *et al.*, 2009). That is, seminal studies by Sullivan and her colleagues demonstrated that while post-natal day (P) 12 rats avoid an odor previously paired with shock, P6 rats exhibit a paradoxical approach response towards such an odor (Sullivan, 2001). However, this transition to odor-avoidance learning occurred at a younger age in infant rats exposed to a stressor (abusive care from the mother as a result of insufficient bedding) or those injected with corticosterone (Moriceau *et al.*, 2009). We have reported other changes in learned fear behaviors in infant rats (i.e., P17) exposed to either rearing stress or glucocorticoid exposure. Specifically, stressed or glucocorticoid-exposed infants exhibited much longer retention of learned fear responses than their same-aged, typically reared counterparts (the non-stressed peers exhibited rapid forgetting known as infantile amnesia; Callaghan & Richardson, 2012, Cowan *et al.*, 2013, also see Haroutunian & Riccio, 1979). In addition to these changes in fear retention, stressed/corticosterone-exposed infant rats also exhibit greater fear relapse after extinction training, a finding normally only observed later in development (Callaghan & Richardson, 2011, 2014, Cowan *et al.*, 2013). Taken together, these outcomes suggest that the effects of adversity on the developing fear system may lead to increased risk for mental health problems via an early switch into fear systems characterized by greater emotionality (i.e., stronger aversions), longer lasting fear

associations, and greater relapse after attempts to inhibit the fear through a process like extinction.

The use of animal models also allows for greater exploration of the neural correlates of early adversity. There is substantial evidence that early life stress alters the brain, both structurally (e.g., reductions in hippocampal volume) and functionally (e.g. alterations in neuroendocrine HPA-axis activity, neurotransmitter levels, and cellular signaling; for reviews, see Gunnar & Quevedo, 2007, Maccari *et al.*, 2014, Marco *et al.*, 2011, Tyrka *et al.*, 2013). As with studies of behavioral changes in the context of early adversity, the majority of research on these neural effects has been conducted in adult individuals, when pathological outcomes have already emerged. However, there is growing evidence that early life stress affects neuroendocrine and brain functioning during development. For example, increased secretion of the stress hormone cortisol, or its rodent equivalent corticosterone, has been reported in infant rats, guinea pigs, and non-human primates as a result of maternal separation (Gareau *et al.*, 2006, Hennessy & Moorman, 1989, Levine & Wiener, 1988, Nishi *et al.*, 2014). Similar elevations in the neuropeptide corticotrophin-releasing factor (CRF), which is also critically involved in the mammalian stress response, have also been observed in juvenile and adolescent primates (Coplan *et al.*, 2005). Interestingly, elevation of CRF in adolescent bonnet macaques as a result of early exposure to stress (variable foraging demand) was associated with increased volume of the left amygdala, which in turn associated with increased anxiety-like responses in adulthood (Coplan *et al.*, 2014). Likewise, in the Moriceau *et al.* (2009) study described earlier, the precocious development of odor-avoidance learning in stressed animals was associated with enhanced amygdala activity (measured by 2 DG uptake), which is normally only observed in older animals (Moriceau *et al.*, 2006). Such precocious maturation of the neural circuitry involved in emotion regulation is reminiscent of, and may underpin, the accelerated developmental trajectories observed in infant rats'

learned fear behaviors (Callaghan & Richardson, 2011, 2012, Cowan *et al.*, 2013, Moriceau *et al.*, 2009).

### *Humans*

The current evidence supports the suggestion that neural development in children is similarly affected by early stress exposure. As one illustration, alterations in cortisol secretion are observed in children exposed to early adversity (for review, see Hunter *et al.*, 2011), paralleling the findings of the animal literature. Although the direction of these effects in humans has been inconsistent (likely due to variation in selection criteria and experimental methodology), it is clear that early life stress alters the neuroendocrine system in children. The precocious maturation of emotion-related neural circuitry observed in rodents is similarly reflected in studies of children using brain-imaging techniques. For example, Tottenham and her colleagues have examined previously-institutionalized individuals using fMRI, finding that these children exhibit a more mature pattern of PFC-amygdala connectivity when viewing fearful faces (Gee *et al.*, 2013). Also, in an EEG study of children 2-4 years of age, McLaughlin *et al.* (2011) reported that institutional rearing alters the developmental trajectory of frontal asymmetry. In that study, deviations from the normal developmental trajectory predicted negative behavioral outcomes (higher internalizing symptoms) at 54 months of age.

### **Early Adversity across Generations**

Considering that stress has such profound effects on the biology and behavior of exposed individuals across their lifespan, it is possible that these alterations may have adverse consequences for future generations (see Figure 1 for an illustration of how this has been investigated in animals and humans).

## *Animal Studies*

There are a growing number of examples of multigenerational transmission of experience-dependent characteristics in controlled laboratory studies. In one such example, Dias and Ressler (2014) used two distinctive odors (acetophenone and propanol) that elicit non-overlapping patterns of activity in the main olfactory epithelium and olfactory bulb. Adult male mice received pairings of one of these odors with an aversive shock. Subsequently, these males were mated with odor-naïve females, and their male offspring (termed the F1 generation; see Figure 1 for further explanation of this terminology) were tested when they reached adulthood. Critically, these F1 offspring had never encountered either of these odors, nor had they ever been shocked. The results showed that the F1 males exhibited a potentiated startle response in the presence of the odor that their father had experienced with shock; further, these males had a lower detection threshold for this odor (i.e., they detected the odor at lower concentrations). Along with these behavioral effects, Dias and Ressler (2014) reported increases in the size of the glomeruli activated by the odor that the F1 male's father had experienced with shock. These effects, both behavioral and neural, were also observed when the F1 males were conceived via IVF and persisted in the offspring of the F1 males (termed the F2 generation). Similar results were obtained when it was the mother who experienced the odor-shock pairings, an effect that persisted following cross-fostering procedures. That is, regardless of whether the odor-shock pairings were experienced by their mothers, fathers, or grandfathers, subsequent generations of mice exhibited the same neuroanatomical changes and differential responses to an odor that they had never experienced prior to testing. All in all, this study provides an amazing illustration of intergenerational transmission of learned behavior and neuroanatomical structure. Multigenerational transmission of stress phenotypes has also been reported for the behavior and cortisol levels of nonhuman primates exposed to higher stress environments (Fairbanks *et*

*al.*, 2011, Kinnally *et al.*, 2013) and for the behavior and metabolic responses of rodents exposed to maternal separation and unpredictable maternal stress (in a complex, sex-specific manner; Franklin *et al.*, 2010, Gapp *et al.*, 2014). It will be interesting to determine whether the early emergence of long-term, and relapse prone, fear memory in infants exposed to early life stress (as described earlier) are also transmitted across generations.

### *Humans*

Significant events in human history have provided researchers with natural “experiments” to explore whether intergenerational transmission of stress occurs in humans. One such event is the Dutch famine (1944-1945), a five month period of extreme food shortage that placed significant stress on an otherwise well-nourished population (Painter *et al.*, 2005). It was found that prenatal maternal malnutrition during later or mid gestation resulted in physical underdevelopment (e.g., lower birth weights, heights, and smaller head circumferences), whereas famine exposure during early gestation resulted in a three-fold increase in coronary heart disease. Males of this generation went on to have offspring (i.e., grandchildren of the malnourished individual) who were more prone to obesity (Veenendaal *et al.*, 2013). In addition to these findings on offspring physical health, there is also evidence that the Dutch famine impacted offspring *mental* health. Specifically, offspring exposed to the famine in-utero had a heightened risk for schizophrenia spectrum disorders (Hoek *et al.*, 1998), while prenatal exposure (preconception or gestational) increased symptoms of depression in adulthood (Stein *et al.*, 2009). The intergenerational effect of trauma on offspring emotional development has also been demonstrated in studies of witnesses and survivors of natural disasters or acts of violence. For example, a study of the 1976 Tangshan earthquake in China found that prenatally-exposed individuals exhibited higher rates of severe depression at 18 years of age compared to a cohort born one year later (and also assessed at 18 years; Watson *et al.*, 1999), while risk for schizophrenia is consistently

elevated in the offspring of mothers exposed to war during pregnancy (see Babenko *et al.*, 2015 for review). As another example, Yehuda *et al.* (2005) examined women who were pregnant during the 2001 World Trade Centre attacks and later developed post-traumatic stress disorder (PTSD). Soon after the attacks these women exhibited lower salivary cortisol levels, considered a biological precursor to PTSD, as compared to a similar group of women who also witnessed the attacks during pregnancy but did not go on to develop PTSD. Importantly, the infant offspring of mothers with PTSD also exhibited low levels of salivary cortisol in the first year of life. These studies of maternal stress during gestation have provided important insights into how parental experiences persist beyond trauma exposure to impact offspring development.

The intergenerational transmission of the effects of stress is also evident when trauma exposure occurs outside of gestation. For example, offspring of combat veterans exhibit increased risk for psychological dysfunction, which appears to be exacerbated when the offspring themselves serve in combat (for a review, see Dekel & Goldblatt, 2008). Adult children of Holocaust survivors born after the war, or after their parents had escaped to safety, exhibit higher lifetime prevalence rates of depression, PTSD, and other anxiety disorders compared to Jewish individuals who did not have a parent who was a Holocaust survivor (Yehuda *et al.*, 2008). In this sample, depressive disorders in adult offspring were significantly associated with paternal and/or maternal PTSD, while PTSD risk in adult offspring was uniquely associated with maternal PTSD (Yehuda *et al.*, 2008). The presence of other anxiety disorders, such as Generalized Anxiety Disorder and Specific Phobia, was elevated among all Holocaust survivor offspring regardless of parental PTSD status. Similarly, a longitudinal study conducted in post-conflict Sierra Leone found significant co-variation between caregiver and child mental health, such that caregiver depression and anxiety were associated with an increase in internalizing symptoms in their adolescent

offspring (Betancourt *et al.*, in press), while in a general population sample PTSD incidence was elevated in the offspring of women with PTSD (Roberts *et al.*, 2012). Together, these studies highlight the inflated risk of psychiatric disorders in offspring of parents exposed to trauma. Further, they provide evidence for intergenerational transmission of stress in humans, such that individuals can acquire biological and behavioral phenotypes that match their parent's risky environment.

### **Potential Mechanisms**

The literature reviewed above demonstrates the relationship between stressful experiences (especially those occurring early in development) and later behavior, not just for directly affected individuals but also for their offspring. Exploration of the underlying mechanisms for such effects is necessary not just to deepen our understanding but also to illuminate potential avenues for intervention where the behavioral phenotype is maladaptive. *Parental behavior.*

One potential mechanism for transmission of parental experiences to the offspring is changes in the behavior of the affected parent. Indeed, parental styles of caregivers have been shown to be atypical post-trauma (Betancourt, 2015). For example, studies of caregivers who survived the Khmer Rouge regime in Cambodia found that a role-reversing parental style mediated the relationship between maternal PTSD and offspring anxiety (Field *et al.*, 2013, Field *et al.*, 2011). In role-reversing parenting, the parent relies on the child to meet their emotional needs (Macfie *et al.*, 2005). This style of parenting is thought to interfere with a child's development of autonomy and is often evident in at-risk samples (Cummings *et al.*, 1994). There is also some evidence that stress and trauma can result in more overtly destructive parenting approaches, such as the increased violence and poorer parental adjustment observed in families of male Vietnam veterans with PTSD (Jordan *et al.*, 1992). Further, adult offspring of Holocaust survivors report higher levels of childhood trauma,

especially emotional abuse and neglect (Yehuda *et al.*, 2001), and the relationship between child mental health and parental war trauma in the Gaza region was found to be mediated by psychological maltreatment of the child by the parent (Palosaari *et al.*, 2013). Maladaptive parental styles have been shown to persist beyond the second generation. In one study, adolescent grandchildren of Holocaust survivors perceived their parents as less accepting and overprotective in comparison to adolescents of similar cultural heritage with no family Holocaust background (Scharf, 2007). More specific to early-life trauma, there is evidence that childhood abuse is associated with dysfunctional parenting practices and attitudes (Ehrensaft *et al.*, 2015). Parents with a history of childhood abuse exhibit higher levels of emotional disengagement (i.e., lower availability, less time spent with child, and higher levels of neutral affect during interactions with their child) and harsh discipline or physical punishment, as well as higher perceived ineffectiveness as a parent (Banyard, 1997, DiLillo & Damashek, 2003, Ehrensaft *et al.*, 2015, Juul *et al.*, in press). Together, these studies demonstrate that environmental insults to parents may exert enduring effects of secondary traumatization on their offspring via parental interactions, resulting in increased risk of mental health problems and difficulties in parenting their own children (Scharf, 2007).

These observations in humans are supported by animal studies showing parental care can mediate the effects of environmental adversity on offspring development. The pioneering work of Michael Meaney highlighted the importance of maternal care for offspring development, demonstrating the links between low levels of maternal care and heightened stress reactivity, as well as the transmission of maternal behaviors across multiple generations (e.g., Francis *et al.*, 1999, Liu *et al.*, 1997, see also Fairbanks, 1996 for review of the transmission of maternal behavior in nonhuman primates). In addition, maternal care has been implicated in the multigenerational effect of stressful social environments (Champagne & Meaney, 2007). Specifically, female rodents exposed to social isolation showed decreased

levels of maternal care towards their offspring, and this effect was transmitted to a next generation of females. Importantly, the offspring raised by this next generation of females showed reduced exploratory behavior, an indicator of heightened anxiety-like behavior in animals.

As well as studying the “vertical transmission” of stress across multiple generations in rodents (i.e., the effects of parental stress exposure on offspring and grand-offspring), research can also be conducted within generations. This can be modeled by breeding rodent mothers multiple times following an initial and single period of stress to examine the persistent effects in caregivers (so called “horizontal transmission”). Using this method, one study found a direct effect of gestational stress on maternal behavior such that stressed mothers became less nurturing than control mothers (Champagne & Meaney, 2006). Those offspring raised with reduced maternal care had heightened anxiety later in adulthood. Strikingly, this style of maternal behavior and the anxious phenotype found in the offspring persisted to a subsequent set of offspring, despite the absence of any environmental stress imposed on the mother. This line of parental transmission across generations is also currently being examined by our group using a postnatal stressor, maternal separation. The results so far suggest that the effects of maternal separation on the maturation of emotion regulation observed in infants directly exposed to that stressor are also observed in infants from a mother’s subsequent litter (who were not directly exposed to any stressor; Kan *et al.*, 2015).

#### *Mating strategies.*

In rodent studies of paternal transmission of early-life stress (e.g., Dias & Ressler, 2014, Franklin *et al.*, 2010), contact between the affected parent and the offspring can be eliminated, apparently ruling out the possibility that any observed behavioral effect in the offspring is due to stress-induced changes in parental style. Nonetheless, it remains possible that there are alterations in maternal behavior that are dependent on paternal characteristics

and behavior during mating (see Curley *et al.*, 2011, for review). For instance, in a mate preference task, female rats exhibit a preference against males exposed to epigenome-modifying toxins (Crews *et al.*, 2007). Remarkably, this preference was observed in the F3 generation (i.e., the males were not directly exposed, but rather their great-grandfathers received in utero exposure to the toxin) and prior to the onset of any disease phenotype. It has also been shown that females alter their investment in pups in response to the male's social housing experience, and that maternal investment is negatively correlated with paternal anxiety (Mashoodh *et al.*, 2012). In other words, female rodents are able to detect altered epigenetic profiles and dysfunctional behavior in potential mating partners and adjust their behavior accordingly, being less likely to reproduce with these males and reducing investment in shared offspring when mating does occur.

Another way that mating behavior may be affected by stressful experiences is via alterations in sexual maturation. Girls exposed to early-life stress exhibit earlier onset of menarche and are more likely to reproduce at an earlier age (Chisholm *et al.*, 2005, Quinlivan *et al.*, 2004). Young motherhood, particularly during teenage years, is associated with poorer health outcomes for children and financial instability, with the implication that offspring are themselves exposed to a stressful early environment (Quinlivan *et al.*, 2004). Thus, stress phenotypes may be perpetuated through the generations by a cycle of early motherhood and resource-deficient environments.

### *Epigenetic Programming.*

The dynamic nature of changes to both brain and behavior in response to early life stress, in addition to the observed transmission of these changes across generations, suggests that epigenetic mechanisms may be involved. Alterations in epigenetic regulation of gene transcription have been implicated in learning and memory, as well as the expression and multi-generational transmission of psychiatric disorders (Bale, 2015, Morris & Monteggia,

2014, Peña *et al.*, 2014, Rodgers & Bale, 2015). Kundakovic and Champagne (2014) provide an excellent review of potential epigenetic mechanisms for the transmission of phenotypes associated with early-life adversity, including alterations in DNA methylation, posttranslational histone modifications, and non-coding RNAs. Briefly, they suggest that epigenetic changes may occur either through germ-line inheritance or through experience-dependent changes that are reiterated through generations via alterations in parental behavior. For example, abusive or inattentive caregiving is associated with epigenetic alterations of the neural circuitry that regulates caregiving behavior in the offspring (e.g., hypermethylation of the estrogen receptor *Esr1*), increasing the likelihood that offspring of abusive carers will imitate such behavior with their own offspring, thus perpetuating the cycle. Such a cyclic mechanism may at least partially account for the effects of parental behavior described above.

As noted by Szyf (2015), our current understanding of epigenetic mechanisms is still limited and there remains some skepticism regarding the stable inheritance of epigenetic marks through the germ-line. This skepticism is based on a long-standing principle of “epigenetic resetting” whereby global demethylation during primordial germ cell differentiation was thought to lead to complete erasure of methylation patterns (Reik *et al.*, 2001). However, emerging evidence suggests that demethylation is extensive but not complete and that experience-induced alterations in DNA methylation may indeed be inherited through the germ-line (for review, see Dias *et al.*, 2015, Szyf, 2015). For example, in Dias and Ressler’s (2014) olfactory fear learning study, described earlier, there was significant hypomethylation of the *Olf151* genetic locus, which specifically mapped to the receptors for the conditioned odor, in the sperm of both F0 and F1 generations. In the context of early-life adversity, Franklin *et al.* (2010) reported increased methylation of genes coding for methyl CpG-binding protein 2 (MeCP2) and the cannabinoid receptor-1 (CB1) and

decreased methylation for the genes coding for corticotrophin-releasing factor receptor 2 (CRFR2) in the sperm cells of adult males directly exposed to maternal separation.

Interestingly, similar changes in methylation patterns of these genes was observed in the brains of female offspring and the sperm cells of male offspring (with the exception of CB1, which was not significantly altered in offspring sperm). Other studies (e.g., Gapp *et al.*, 2014, Rodgers *et al.*, 2013) have shown that paternal stress can alter sperm microRNA content and that microinjection of sperm RNA from stressed individuals into fertilized oocytes can reproduce stress phenotypes, providing another potential route for the occurrence of epigenetic inheritance through the germ-line.

Regardless of whether epigenetic marks are transmitted via pure germ-line inheritance, experience-driven patterns, or some combination of the two, there are a number of potential sites that could be of functional importance for the behavioral and neural changes observed in the case of early-life adversity. Both animal and human studies have identified a broad range of genes that undergo epigenetic modification (often in a complex, brain region or tissue-specific manner) following early-life stress, including but not limited to the serotonin transporter (5-HTT; Beach *et al.*, 2011, Kinnally *et al.*, 2010), brain-derived neurotrophic factor (BDNF; Roth & Sweatt, 2011), and MeCP2 (Franklin *et al.*, 2010). Adversity-induced epigenetic regulation has been reviewed in detail elsewhere (e.g., Lutz & Turecki, 2014), so for the purposes of this paper we will limit our discussion to one potential epigenetic pathway of particular relevance for the specific behavioral effects focused on herein, that being methylation of promoter regions of the glucocorticoid receptor gene (*NR3C1*). Here, it is worth noting that studies of the *NR3C1* gene have thus far not addressed the question of whether direct germ-line inheritance of epigenetic changes to this gene occur. However, it presents as a strong candidate for further investigation. The *NR3C1* gene is structurally conserved across humans and rodents, regulating expression of the glucocorticoid

receptor, which provides negative feedback on the concentration of glucocorticoids to regulate their production by the HPA axis (the neuroendocrine stress response system; Suderman *et al.*, 2012). HPA axis dysregulation is a commonly reported consequence of early adversity that is also considered a risk factor for various forms of psychopathology, including PTSD, depression, and anxiety disorders (Faravelli *et al.*, 2012, Pesonen *et al.*, 2010, Sanchez *et al.*, 2001, Shea *et al.*, 2005). It has been hypothesized that this dysregulation (which is known to persist into adulthood) is driven by changes to the DNA methylation status of promoter regions of *NR3C1*.

The first studies to demonstrate altered methylation of *NR3C1* were conducted in infant rats exposed to differing levels of maternal care (Weaver *et al.*, 2004). Those rats that experienced low maternal care exhibited increased DNA methylation in the exon 1<sub>7</sub> promoter region of *NR3C1* in the hippocampus. In humans, hypermethylation of the exon 1<sub>F</sub> promoter region of *NR3C1* (the human analog of exon 1<sub>7</sub> in rodents) has consistently been observed in individuals exposed to early life stress and in those with associated disorders such as borderline personality disorder, major depression, and PTSD (for review, see Daskalakis & Yehuda, 2014). These changes are observed as early as 3-5 years of age and remain stable in adulthood (Perroud *et al.*, 2011, Tyrka *et al.*, 2015, Tyrka *et al.*, 2012). Further, there is some preliminary evidence that this pattern of hypermethylation is passed on to the offspring of traumatized parents. For example, Yehuda *et al.* (2014) reported a complex relationship between parental PTSD and *NR3C1*-1<sub>F</sub> methylation, which was in turn associated with neuroendocrine responsiveness, in the offspring of Holocaust survivors. In addition, maternal war stress exposure correlated with maternal and infant *NR3C1* methylation levels in a Congolese sample (Mulligan *et al.*, 2012). Given that exposure to the glucocorticoid corticosterone is sufficient to mimic the effects of early adversity on learned fear behaviors during development in rodents (Callaghan & Richardson, 2012, 2014, Moriceau *et al.*, 2006),

it is tempting to speculate that heightened glucocorticoid exposure as a result of epigenetic changes to GR expression in the offspring may contribute to the passing down of vulnerable phenotypes and subsequent risk for psychopathology.

### **Potential for Intervention**

Regardless of the mechanism of transmission, it is clear that there are altered behavioral and neural outcomes for individuals exposed to early-life adversity and that these outcomes can be passed down to future generations (see Figure 2). In many cases this may prepare an individual for survival in a stressful environment, but when the environment no longer matches the adaptation it can be problematic, which may explain why individuals exposed to early adversity are more vulnerable to mental health problems. Attempts to reverse or interrupt the transmission of stress-induced phenotypes across generations have been limited. Yet the nature of the stress-induced phenotype, as a response to environmental conditions, suggests that the system is inherently flexible and therefore potentially amenable to intervention.

### *Traditional Mental Health Treatments*

As reviewed above, exposure to adversity during development renders individuals vulnerable to mental health issues throughout the lifespan. It follows that individuals with a history of early adversity are likely to benefit from effective treatments for these illnesses. Unfortunately, some concerning evidence suggests that individuals with a history of early trauma may not only exhibit a higher incidence of psychopathology but are also less likely to respond to existing treatments. Research from our lab shows that rodents with a history of early life stress are more vulnerable to fear relapse following extinction (Callaghan & Richardson, 2011, Cowan *et al.*, 2013), which is in keeping with epidemiological evidence that childhood maltreatment is associated with lack of treatment response in clinical trials of psychological therapy, antidepressant medication, or combined treatment for depression

(Nanni *et al.*, 2012). Similarly, both pharmacological and psychological treatments have been found to be less effective in populations exposed to early adversity across ADHD (Sugimoto *et al.*, 2015) and substance abuse (Boles *et al.*, 2005, Sacks *et al.*, 2008), although not in social anxiety (Bruce *et al.*, 2013).

Turning to generational effects, to our knowledge there have been no studies directly examining the impact of early adversity on offspring treatment outcomes. However, some research has examined the impact of parental mental illness. For example, early results from an international collaborative study (the Genes for Treatment study) suggest that parental psychopathology interferes with children's recovery from anxiety disorders, predicting a significantly poorer response to treatment on follow-up (Hudson *et al.*, 2015). On a more positive note, there are also indications that cognitive-behavioral therapy (CBT) can be effective as either a prevention or intervention technique to reduce psychological risk in the offspring of individuals with a mental illness. A number of early intervention programs have targeted these individuals with promising results (for review, see Reupert *et al.*, 2013, Siegenthaler *et al.*, 2012). For example, a family-based CBT program significantly reduced anxiety disorder onset in the 7-12 year old offspring of anxious parents at 12-month follow-up (from 30% in the waitlist control to zero in the treatment group; Ginsburg, 2009). Several studies have also found CBT-based programs to effectively reduce internalizing and externalizing symptomology and rates of illness onset in the offspring of depressed parents (Reupert *et al.*, 2013). While these studies have generally examined relatively small samples, more definitive support for the efficacy of intervention in the F1 generation may be provided by a large-scale clinical RCT currently being conducted on the effects of a CBT intervention in children of depressed and anxious parents (described in the study protocol published by Nauta *et al.*, 2012). Further research is also needed to identify the lifetime effectiveness of

such programs and to ascertain whether effective psychosocial or pharmacological treatment of mental illness can reduce the risk of psychopathology in subsequent generations.

### *Parenting Interventions*

While therapeutic interventions that address psychopathology in affected parents and their offspring appear promising, the dysfunctional parenting practices discussed previously point to the need to consider specific targeting of parenting skills. Such parenting skills training has been shown to be effective in the treatment of children's psychopathology, particularly for childhood externalizing disorders (e.g., Triple P Positive Parenting Program, Parent-Child Interaction Therapy; see Kazdin, 1997, Thomas & Zimmer-Gembeck, 2007, for reviews), but also for childhood anxiety and depression (Eckshtain *et al.*, in press, Manassis *et al.*, 2014). In parents exhibiting psychopathology (e.g., depression, substance abuse) or exposed to various forms of stress (e.g., poverty, divorce, foster care), the evidence suggests that parenting competence is amenable to intervention (Ajilchi & Kargar, 2013, Guttentag *et al.*, 2014, Lowell *et al.*, 2011, Oriana Linares *et al.*, 2006, Suchman *et al.*, 2011, Wolchik *et al.*, 1993). Importantly, treatment-induced enhancements of parenting attitudes have been shown to be transmitted to a subsequent generation, indicating the potential generational impact of parenting interventions (Mahrer *et al.*, 2014). Furthermore, parenting interventions have also proven to be effective at reducing or preventing negative child mental health outcomes in these at-risk samples, with reductions in child internalizing and externalizing problems reported across different forms of parental psychopathology and stress (Bywater *et al.*, 2009, Guttentag *et al.*, 2014, Lam *et al.*, 2008, Lowell *et al.*, 2011, Malmberg & Field, 2013, Self-Brown *et al.*, 2011). However, there is evidence that parenting interventions are less effective for the offspring of individuals with greater symptoms of psychopathology (e.g., van Loon *et al.* 2011, Webster-Stratton & Hammond, 1990, but see also Gardner *et al.* 2010, Timmer *et al.* 2011). This suggests the need to concurrently target parental

psychopathology, a strategy which has been shown to be more effective in the treatment of depressed mothers and their disruptive children (Sanders & McFarland, 2000).

Interestingly, an even simpler behavioral parenting technique has recently been shown to reduce epigenetic aberrations in children exposed to perinatal maternal depression (Murgatroyd *et al.*, 2015). Specifically, Murgatroyd and colleagues assessed DNA methylation patterns in 181 infants born to women with varying levels of depression symptomatology. They found associations between postnatal depression and increased methylation of the *NR3C1-1<sub>F</sub>* promoter in infants. Strikingly, in a higher risk subgroup whose mothers reported post- but not pre-natal depression levels above the median there was a protective effect of self-reported maternal stroking. That is, there was a negative correlation between maternal tactile stimulation of infants and the DNA methylation patterns in those infants, speaking to the importance of parent-child interactions in the modulation of epigenetic risk factors and transmission of stress phenotypes.

### *Targeting Epigenetic Changes*

Psychopathological responses to early adversity are complex, with no discernible specificity of relationships between particular types of trauma and particular mental illnesses within nor across generations (Starr *et al.*, 2014). As such, an approach that targets underlying mechanisms rather than symptomatology might prove more effective, particularly when considering long-term, heritable changes. Such an approach also fits well with the Research Domain Criteria (RDoC) framework proposed by the National Institute of Mental Health (e.g., Insel, 2014). Given our discussion of the potential role of epigenetic alterations in the maintenance and transmission of stress-induced behavioral changes, targeting the epigenome seems a logical starting point. In this regard, there has been some use of epigenetic modifiers to reverse anxiety and stress-responsiveness phenotypes associated with different early experiences. Meaney, Szyf, Weaver, and colleagues conducted the

groundbreaking studies in this area (Weaver *et al.*, 2004, Weaver *et al.*, 2005, Weaver *et al.*, 2006). They demonstrated that central infusion of either the histone deacetylase (HDAC) inhibitor trichostatin A or the methyl donor methionine could reverse the effects of low or high maternal care, respectively, on DNA methylation patterns, glucocorticoid receptor expression, HPA stress responses, and behavioral expressions of depression and anxiety in adult mice.

Similar effects of epigenetic modulators have since been demonstrated in animals exposed to early stress (Kao *et al.*, 2012). Specifically, systemic injection with an HDAC inhibitor (valproic acid) immediately prior to maternal separation and isolation reversed the effects of isolation on DNA methylation in the frontal cortex and fear-potentiated startle behavior in adulthood. This study is valuable in that it demonstrates that epigenetic treatments can confer long-lasting protection against the effects of early stress when delivered during the stressful period. However, it isn't known whether these benefits might be transmitted to future generations, let alone whether such hypothetical transmission might occur through the germ-line or through changes to parenting behavior. Research on the use of epigenetic modifiers in humans is even more limited. However, the effect of valproic acid (a pharmacological agent widely used to treat epilepsy and bipolar disorder) on epigenetic profiles has been proposed as the mechanism for its efficacy in management of bipolar disorder (Phiel *et al.*, 2001). Ideally, manipulation of the epigenome would be targeted at the key gene promoter regions in specific brain regions that are susceptible to stress-induced epigenetic changes (e.g., *NR3C1* in the hippocampus). However, at this stage we are far from achieving that level of specificity, suggesting that alternative treatment options need to be considered.

### *Micronutrients: A More Palatable Approach?*

An alternative, and definitely more practicable, strategy for intervention involves alterations in dietary intake. As observed in the aforementioned studies of the Dutch famine, nutrition can play a key role in the overall health of individuals and their offspring. It has also been hypothesized that one driver of epigenetic changes in the context of stress is altered maternal intake of nutrients, including dietary methyl donors (Lucassen *et al.*, 2013). Folic acid, vitamins B2, B6, and B12 are involved in DNA methylation and changes in maternal or paternal micronutrient intake can alter offspring, grand-offspring, and great-grand-offspring outcomes (see Vanhees *et al.*, 2014, for review). Further, recent evidence from studies of the effects of the Christchurch earthquake indicate that micronutrients administered in the aftermath of trauma conferred a protective effect, resulting in a long-lasting reduction in the incidence of PTSD symptoms (Rucklidge *et al.*, 2012, Rucklidge *et al.*, 2014). It would be interesting to investigate whether this stress-protective effect also occurs in developing populations exposed to trauma.

### *Modification of the Microbiome*

Another potential source of epigenetic modifiers is the large population of commensal bacteria that reside in the gastrointestinal tract. Known collectively as the microbiome, these bacteria produce large quantities of short-chain fatty acids (e.g., butyrate, acetate, and propionate), which can act as HDAC inhibitors (Licciardi *et al.*, 2010). There is mounting evidence to suggest that the microbiome plays an important role in the expression of emotional behavior and the function of associated brain systems across species (Christian *et al.*, 2015, Foster & McVey Neufeld, 2013, Jašarević *et al.*, 2015). For example, the composition of the microbiome has been shown to be important for the development of normal emotional behavior and stress responses in studies of both germ-free mice (raised in the absence of a microbiome; De Palma *et al.*, 2015, Heijtz *et al.*, 2011, Sudo *et al.*, 2004)

and mice with differing emotional and gastrointestinal profiles (Bercik *et al.*, 2011). Further, recent evidence has shown that the gut microbiota produce metabolites with known importance for mental health (e.g. serotonin; Yano *et al.*, 2015). Manipulation of the microbiome via administration of probiotic microorganisms, which colonize the gastrointestinal tract to bring benefits to the host organism, has been shown to improve mood and reduce anxiety in both rodents and humans (Dinan *et al.*, 2013, Messaoudi *et al.*, 2011, Neufeld *et al.*, 2011, Rao *et al.*, 2009). In addition, probiotics have been shown to alter functional network activation in healthy women completing an emotional attention task (Tillisch *et al.*, 2013).

Importantly, there is also recent evidence to show that probiotic treatments can alter outcomes in animal models of early-life stress. Probiotic treatment of maternally-separated animals has been shown to dampen corticosterone responses to separation and reduce adult expression of depressive behaviors (Desbonnet *et al.*, 2010, Gareau *et al.*, 2007). In addition, research from our group (Cowan *et al.*, 2015) shows that the effects of maternal separation on development of fear-related learning and extinction are reversed by treatment with a probiotic supplement to dams' drinking water. These findings suggest that both the precocious development of fear systems and the long-term emotional dysregulation associated with early-life adversity can be reversed by targeting the microbiome. It will be important to determine the mechanism for these actions (e.g., alterations in epigenetic signals, changes in maternal behavior), as well as testing whether these results can be translated to human populations or whether such treatments would alter outcomes for the offspring of individuals exposed to early adversity.

Unfortunately, there are many places around the world where children are exposed to "extreme environments" which adversely impact on physical health and brain development (Nelson, 2015). Treatments that target either nutritional intake or microbial balance in the gut

would be particularly appealing in such settings for a number of reasons (for further commentary, see Knight, 2015). For example, individuals in these settings are often malnourished and have a number of issues in regard to gut functioning (Ashbolt, 2004). Further, these sorts of interventions are relatively inexpensive to implement, in part because they don't require large numbers of individuals with advanced training. Finally, such treatment approaches are more likely to be accepted, and adhered to, than more time-demanding or invasive treatments.

#### *Other Considerations for Intervention*

The above discussion is by no means comprehensive in the consideration of potential interventions for the effects of early-life stress across generations. There are a variety of other promising methods that have been described in the literature, including exercise and anti-inflammatory medication (Harrison & Baune, 2014). One factor that will need to be considered in future studies is the timing of the intervention, whatever it might be. Just as there are critical periods during which stress can have a particularly deleterious effect, timing is likely to play an important role in the effectiveness of treatment (Gee & Casey, 2015). This will be a particularly complex process when dealing with multigenerational effects of stress, and Figure 2 highlights some potential key points for intervention drawn from the research discussed above. Similarly, the dose and duration of exposure to both stress and treatment are likely to impact on the success of any intervention. At this stage, the parameters for successful intervention remain unclear, particularly with respect to the intergenerational impact of treatment for early stress. This leaves the door open for an exciting period in the advancement of our understanding of both the mechanisms that underpin multigenerational transmission of stress phenotypes and the clinical applications of this knowledge.

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## References

- Ajilchi, B. & Kargar, F.R. (2013) The impact of a parenting skills training program on stressed mothers and their children's depression level. *Procedia Soc Behav Sci*, **84**, 450-456.
- Ashbolt, N.J. (2004) Microbial contamination of drinking water and disease outcomes in developing regions. *Toxicology*, **198**, 229-238.
- Babenko, O., Kovalchuk, I. & Metz, G.A.S. (2015) Stress-induced perinatal and transgenerational epigenetic programming of brain development and mental health. *Neurosci Biobehav Rev*, **48**, 70-91.
- Bale, T.L. (2015) Epigenetic and transgenerational reprogramming of brain development. *Nat Rev Neurosci*, **16**, 332-344.
- Banyard, V.L. (1997) The impact of childhood sexual abuse and family functioning on four dimensions of women's later parenting. *Child Abuse Negl*, **21**, 1095-1107.
- Beach, S.R.H., Brody, G.H., Todorov, A.A., Gunter, T.D. & Philibert, R.A. (2011) Methylation at 5HTT mediates the impact of child sex abuse on women's antisocial behavior: An examination of the iowa adoptee sample. *Psychosom Med*, **73**, 83-87.
- Bercik, P., Denou, E., Collins, J., Jackson, W.P., Lu, J., Jury, J., Deng, Y., Blennerhassett, P., Macri, J., McCoy, K.D., Verdu, E.F. & Collins, S.M. (2011) The intestinal microbiota affect central levels of brain-derived neurotropic factor and behavior in mice. *Gastroenterology*, **141**, 599-609.
- Betancourt, T.S. (2015) The intergenerational effect of war. *JAMA Psychiatry*, **72**, 199-200.
- Betancourt, T.S., McBain, R.K., Newnham, E.A. & Brennan, R.T. (in press) The intergenerational impact of war: Longitudinal relationships between caregiver and child mental health in postconflict Sierra Leone. *J Child Psychol Psychiatry*.

- Boles, S.M., Joshi, V., Grella, C. & Wellisch, J. (2005) Childhood sexual abuse patterns, psychosocial correlates, and treatment outcomes among adults in drug abuse treatment. *J Child Sex Abus*, **14**, 39-55.
- Bruce, L.C., Heimberg, R.G., Goldin, P.R. & Gross, J.J. (2013) Childhood maltreatment and response to cognitive behavioral therapy among individuals with social anxiety disorder. *Depress Anxiety*, **30**, 662-669.
- Bywater, T., Hutchings, J., Daley, D., Whitaker, C., Yeo, S.T., Jones, K., Eames, C. & Edwards, R.T. (2009) Long-term effectiveness of a parenting intervention for children at risk of developing conduct disorder. *Br J Psychiatry*, **195**, 318-324.
- Callaghan, B.L. & Richardson, R. (2011) Maternal separation results in early emergence of adult-like fear and extinction learning in infant rats. *Behav Neurosci*, **125**, 20-28.
- Callaghan, B.L. & Richardson, R. (2012) Adverse rearing environments and persistent memories in rats: Removing the brakes on infant fear memory. *Transl Psychiatry*, **2**, e138.
- Callaghan, B.L. & Richardson, R. (2014) Early emergence of adult-like fear renewal in the developing rat after chronic corticosterone treatment of the dam or the pups. *Behav Neurosci*, **128**, 594-602.
- Champagne, F.A. & Meaney, M.J. (2006) Stress during gestation alters postpartum maternal care and the development of the offspring in a rodent model. *Biol Psychiatry*, **59**, 1227-1235.
- Champagne, F.A. & Meaney, M.J. (2007) Transgenerational effects of social environment on variations in maternal care and behavioral response to novelty. *Behav Neurosci*, **121**, 1353-1363.

- Chisholm, J.S., Quinlivan, J.A., Petersen, R.W. & Coall, D.A. (2005) Early stress predicts age at menarche and first birth, adult attachment, and expected lifespan. *Hum Nat*, **16**, 233-265.
- Christian, L.M., Galley, J.D., Hade, E.M., Schoppe-Sullivan, S., Kamp Dush, C. & Bailey, M.T. (2015) Gut microbiome composition is associated with temperament during early childhood. *Brain Behav Immun*, **45**, 118-127.
- Coplan, J.D., Altemus, M., Mathew, S.J., Smith, E.L.P., Scharf, B., Coplan, P.M., Kral, J.G., Gorman, J.M., Owens, M.J., Nemeroff, C.B. & Rosenblum, L.A. (2005) Synchronized maternal-infant elevations of primate CSF CRF concentrations in response to variable foraging demand. *CNS Spectr*, **10**, 530-536.
- Coplan, J.D., Fathy, H.M., Jackowski, A.P., Tang, C.Y., Perera, T.D., Mathew, S.J., Martinez, J., Abdallah, C.G., Dwork, A.J., Pantol, G., Carpenter, D., Gorman, J.M., Nemeroff, C.B., Owens, M.J., Kaffman, A. & Kaufman, J. (2014) Early life stress and macaque amygdala hypertrophy: Preliminary evidence for a role for the serotonin transporter gene. *Front Behav Neurosci*, **8**, 342.
- Cowan, C.S.M., Callaghan, B.L. & Richardson, R. (2013) Acute early-life stress results in premature emergence of adult-like fear retention and extinction relapse in infant rats. *Behav Neurosci*, **127**, 703-711.
- Cowan, C.S.M., Callaghan, B.L. & Richardson, R. (2015) A probiotic formulation (Lactobacillus rhamnosus and L. helveticus) reverses abnormal developmental trajectories of emotional learning in stressed infant rats.
- Crews, D., Gore, A.C., Hsu, T.S., Dangleben, N.L., Spinetta, M., Schallert, T., Anway, M.D. & Skinner, M.K. (2007) Transgenerational epigenetic imprints on mate preference. *Proc Natl Acad Sci U S A*, **104**, 5942-5946.

- Cummings, E.M., Hennessy, K.D., Rabideau, G.J. & Cicchetti, D. (1994) Responses of physically abused boys to interadult anger involving their mothers. *Dev Psychopathol*, **6**, 31-41.
- Curley, J.P., Mashoodh, R. & Champagne, F.A. (2011) Epigenetics and the origins of paternal effects. *Horm Behav*, **59**, 306-314.
- Daskalakis, N.P. & Yehuda, R. (2014) Site-specific methylation changes in the glucocorticoid receptor exon 1F promoter in relation to life adversity: Systematic review of contributing factors. *Front Neurosci*, **8**, 369.
- De Palma, G., Blennerhassett, P., Lu, J., Deng, Y., Park, A.J., Green, W., Denou, E., Silva, M.A., Santacruz, A., Sanz, Y., Surette, M.G., Verdu, E.F., Collins, S.M. & Bercik, P. (2015) Microbiota and host determinants of behavioural phenotype in maternally separated mice. *Nat Commun*, **6**, 7735.
- Dekel, R. & Goldblatt, H. (2008) Is there intergenerational transmission of trauma? The case of combat veterans' children. *Am J Orthopsychiatry*, **78**, 281-289.
- Desbonnet, L., Garrett, L., Clarke, G., Kiely, B., Cryan, J.F. & Dinan, T.G. (2010) Effects of the probiotic *Bifidobacterium infantis* in the maternal separation model of depression. *Neuroscience*, **170**, 1179-1188.
- Dias, B.G., Maddox, S.A., Klengel, T. & Ressler, K.J. (2015) Epigenetic mechanisms underlying learning and the inheritance of learned behaviors. *Trends Neurosci*, **38**, 96-107.
- Dias, B.G. & Ressler, K.J. (2014) Parental olfactory experience influences behavior and neural structure in subsequent generations. *Nat Neurosci*, **17**, 89-96.
- DiLillo, D. & Damashek, A. (2003) Parenting characteristics of women reporting a history of childhood sexual abuse. *Child Maltreat*, **8**, 319-333.

- Dinan, T.G., Stanton, C. & Cryan, J.F. (2013) Psychobiotics: A novel class of psychotropic. *Biol Psychiatry*, **74**, 720-726.
- Eckshtain, D., Kuppens, S. & Weisz, J.R. (in press) Amelioration of child depression through behavioral parent training: A preliminary study. *J Clin Child Adolesc Psychol*.
- Ehrensaft, M.K., Knous-Westfall, H.M., Cohen, P. & Chen, H. (2015) How does child abuse history influence parenting of the next generation? *Psychol Violence*, **5**, 16-25.
- Fairbanks, L.A. (1996) Individual differences in maternal style: Causes and consequences for mothers and offspring. *Adv Study Behav*. pp. 579-611.
- Fairbanks, L.A., Jorgensen, M.J., Bailey, J.N., Breidenthal, S.E., Grzywa, R. & Laudenslager, M.L. (2011) Heritability and genetic correlation of hair cortisol in vervet monkeys in low and higher stress environments. *Psychoneuroendocrinology*, **36**, 1201-1208.
- Faravelli, C., Lo Sauro, C., Lelli, L., Pietrini, F., Lazzaretti, L., Godini, L., Benni, L., Fioravanti, G., Talamba, G.A., Castellini, G. & Ricca, V. (2012) The role of life events and HPA axis in anxiety disorders: A review. *Curr Pharm Des*, **18**, 5663-5674.
- Field, N.P., Muong, S. & Sochanvimean, V. (2013) Parental styles in the intergenerational transmission of trauma stemming from the Khmer Rouge regime in Cambodia. *Am J Orthopsychiatry*, **83**, 483-494.
- Field, N.P., Om, C., Kim, T. & Vorn, S. (2011) Parental styles in second generation effects of genocide stemming from the Khmer Rouge regime in Cambodia. *Attach Hum Dev*, **13**, 611-628.
- Foster, J.A. & McVey Neufeld, K.-A. (2013) Gut–brain axis: How the microbiome influences anxiety and depression. *Trends Neurosci*, **36**, 305-312.

- Francis, D., Diorio, J., Liu, D. & Meaney, M.J. (1999) Nongenomic transmission across generations of maternal behavior and stress responses in the rat. *Science*, **286**, 1155-1158.
- Franklin, T.B., Russig, H., Weiss, I.C., Grff, J., Linder, N., Michalon, A., Vizi, S. & Mansuy, I.M. (2010) Epigenetic transmission of the impact of early stress across generations. *Biol Psychiatry*, **68**, 408-415.
- Gapp, K., Jawaid, A., Sarkies, P., Bohacek, J., Pelczar, P., Prados, J., Farinelli, L., Miska, E. & Mansuy, I.M. (2014) Implication of sperm RNAs in transgenerational inheritance of the effects of early trauma in mice. *Nat Neurosci*, **17**, 667-669.
- Gardner, F., Hutchings, J., Bywater, T. & Whitaker, C. (2010) Who benefits and how does it work? Moderators and mediators of outcome in an effectiveness trial of a parenting intervention. *J Clin Child Adolesc Psychol*, **39**, 568-580.
- Gareau, M.G., Jury, J., MacQueen, G., Sherman, P.M. & Perdue, M.H. (2007) Probiotic treatment of rat pups normalises corticosterone release and ameliorates colonic dysfunction induced by maternal separation. *Gut*, **56**, 1522-1528.
- Gareau, M.G., Jury, J., Yang, P.C., MacQueen, G. & Perdue, M.H. (2006) Neonatal maternal separation causes colonic dysfunction in rat pups including impaired host resistance. *Pediatr Res*, **59**, 83-88.
- Gee, D.G. & Casey, B.J. (2015) The impact of developmental timing for stress and recovery. *Neurobiol Stress*, **1**, 184-194.
- Gee, D.G., Gabard-Durnam, L.J., Flannery, J., Goff, B., Humphreys, K.L., Telzer, E.H., Hare, T.A., Bookheimer, S.Y. & Tottenham, N. (2013) Early developmental emergence of human amygdala-prefrontal connectivity after maternal deprivation. *Proc Natl Acad Sci U S A*, **110**, 15638-15643.

- Ginsburg, G.S. (2009) The child anxiety prevention study: Intervention model and primary outcomes. *J Consult Clin Psychol*, **77**, 580-587.
- Green, J.G., McLaughlin, K.A., Berglund, P.A., Gruber, M.J., Sampson, N.A., Zaslavsky, A.M. & Kessler, R.C. (2010) Childhood adversities and adult psychiatric disorders in the national comorbidity survey replication I: Associations with first onset of DSM-IV disorders. *Arch Gen Psychiatry*, **67**, 113-123.
- Gunnar, M. & Quevedo, K. (2007) The neurobiology of stress and development. *Annu Rev Psychol*, **58**, 145-173.
- Guttentag, C.L., Landry, S.H., Williams, J.M., Baggett, K.M., Noria, C.W., Borkowski, J.G., Swank, P.R., Farris, J.R., Crawford, A., Lanzi, R.G., Carta, J.J., Warren, S.F. & Ramey, S.L. (2014) "My baby & me": Effects of an early, comprehensive parenting intervention on at-risk mothers and their children. *Dev Psychol*, **50**, 1482-1496.
- Haroutunian, V. & Riccio, D.C. (1979) Age-dependent effects of preconditioning aversive experiences on the retention of conditioned fear in weanling rats. *Behav Neural Biol*, **26**, 248-253.
- Harrison, E.L. & Baune, B.T. (2014) Modulation of early stress-induced neurobiological changes: A review of behavioural and pharmacological interventions in animal models. *Transl Psychiatry*, **4**, e390.
- Heijtz, R.D., Wang, S., Anuar, F., Qian, Y., Björkholm, B., Samuelsson, A., Hibberd, M.L., Forssberg, H. & Pettersson, S. (2011) Normal gut microbiota modulates brain development and behavior. *Proc Natl Acad Sci U S A*, **108**, 3047-3052.
- Hennessy, M.B. & Moorman, L. (1989) Factors influencing cortisol and behavioral responses to maternal separation in guinea pigs. *Behav Neurosci*, **103**, 378-385.
- Hoek, H.W., Brown, A.S. & Susser, E. (1998) The Dutch famine and schizophrenia spectrum disorders. *Soc Psychiatry Psychiatr Epidemiol*, **33**, 373-379.

- Hudson, J.L., Keers, R., Roberts, S., Coleman, J.R.I., Breen, G., Arendt, K., Bögels, S., Cooper, P., Creswell, C., Hartman, C., Heiervang, E.R., Hötzel, K., In-Albon, T., Lavalley, K., Lyneham, H.J., Marin, C.E., McKinnon, A., Meiser-Stedman, R., Morris, T., Nauta, M., Rapee, R.M., Schneider, S., Schneider, S.C., Silverman, W.K., Thastum, M., Thirlwall, K., Waite, P., Wergeland, G.J., Lester, K.J. & Eley, T.C. (2015) Clinical predictors of response to cognitive-behavioral therapy in pediatric anxiety disorders: The genes for treatment (GxT) study. *J Am Acad Child Adolesc Psychiatry*, **54**, 454-463.
- Hunter, A.L., Minnis, H. & Wilson, P. (2011) Altered stress responses in children exposed to early adversity: A systematic review of salivary cortisol studies. *Stress*, **14**, 614-626.
- Insel, T.R. (2014) The NIMH research domain criteria (RDoC) project: Precision medicine for psychiatry. *Am J Psychiatry*, **171**, 395-397.
- Jašarević, E., Rodgers, A.B. & Bale, T.L. (2015) A novel role for maternal stress and microbial transmission in early life programming and neurodevelopment. *Neurobiol Stress*, **1**, 81-88.
- Jones, P.B. (2013) Adult mental health disorders and their age at onset. *Br J Psychiatry*, **202**, s5-s10.
- Jordan, B.K., Marmar, C.R., Fairbank, J.A., Schlenger, W.E., Kulka, R.A., Hough, R.L. & Weiss, D.S. (1992) Problems in families of male Vietnam veterans with posttraumatic stress disorder. *J Consult Clin Psychol*, **60**, 916-926.
- Juul, S.H., Hendrix, C., Robinson, B., Stowe, Z.N., Newport, D.J., Brennan, P.A. & Johnson, K.C. (in press) Maternal early-life trauma and affective parenting style: The mediating role of HPA-axis function. *Arch Womens Ment Health*.

- Kan, J.M., Callaghan, B.L. & Richardson, R. (2015) A mother's past can predict her offspring's future: Previous maternal separation leads to the early emergence of adult-like fear behavior in subsequent infant rat offspring.
- Kao, G.S., Cheng, L.Y., Chen, L.H., Tzeng, W.Y., Cherng, C.G., Su, C.C., Wang, C.Y. & Yu, L. (2012) Neonatal isolation decreases cued fear conditioning and frontal cortical histone 3 lysine 9 methylation in adult female rats. *Eur J Pharmacol*, **697**, 65-72.
- Kazdin, A.E. (1997) Parent management training: Evidence, outcomes, and issues. *J Am Acad Child Adolesc Psychiatry*, **36**, 1349-1356.
- Kessler, R.C., Amminger, G.P., Aguilar-Gaxiola, S., Alonso, J., Lee, S. & Üstün, T.B. (2007) Age of onset of mental disorders: A review of recent literature. *Curr Opin Psychiatry*, **20**, 359-364.
- Kinnally, E.L., Capitanio, J.P., Leibel, R., Deng, L., Leduc, C., Haghghi, F. & Mann, J.J. (2010) Epigenetic regulation of serotonin transporter expression and behavior in infant rhesus macaques. *Genes Brain Behav*, **9**, 575-582.
- Kinnally, E.L., Feinberg, C., Kim, D., Ferguson, K., Coplan, J.D. & Mann, J.J. (2013) Transgenerational effects of variable foraging demand stress in female bonnet macaques. *Am J Primatol*, **75**, 509-517.
- Knight, R. (2015) Why microbiome treatments could pay off soon. *Nature*, **518**, S5.
- Kundakovic, M. & Champagne, F.A. (2014) Early-life experience, epigenetics, and the developing brain. *Neuropsychopharmacology*, **40**, 141-153.
- Lam, W.K., Fals-Stewart, W. & Kelley, M.L. (2008) Effects of parent skills training with behavioral couples therapy for alcoholism on children: A randomized clinical pilot trial. *Addict Behav*, **33**, 1076-1080.

- Lee, F.S., Heimer, H., Giedd, J.N., Lein, E.S., Šestan, N., Weinberger, D.R. & Casey, B.J. (2014) Adolescent mental health - Opportunity and obligation: Emerging neuroscience offers hope for treatments. *Science*, **346**, 547-549.
- Levine, S. & Wiener, S.G. (1988) Psychoendocrine aspects of mother-infant relationships in nonhuman primates. *Psychoneuroendocrinology*, **13**, 143-154.
- Licciardi, P.V., Wong, S.S., Tang, M.L.K. & Karagiannis, T.C. (2010) Epigenome targeting by probiotic metabolites. *Gut Pathog*, **2**, 24.
- Liu, D., Diorio, J., Tannenbaum, B., Caldji, C., Francis, D., Freedman, A., Sharma, S., Pearson, D., Plotsky, P.M. & Meaney, M.J. (1997) Maternal care, hippocampal glucocorticoid receptors, and hypothalamic-pituitary-adrenal responses to stress. *Science*, **277**, 1659-1662.
- Lowell, D.I., Carter, A.S., Godoy, L., Paulicin, B. & Briggs-Gowan, M.J. (2011) A randomized controlled trial of Child FIRST: A comprehensive home-based intervention translating research into early childhood practice. *Child Dev*, **82**, 193-208.
- Lucassen, P.J., Naninck, E.F.G., van Goudoever, J.B., Fitzsimons, C., Joels, M. & Korosi, A. (2013) Perinatal programming of adult hippocampal structure and function: Emerging roles of stress, nutrition and epigenetics. *Trends Neurosci*, **36**, 621-631.
- Lutz, P.E. & Turecki, G. (2014) DNA methylation and childhood maltreatment: From animal models to human studies. *Neuroscience*, **264**, 142-156.
- Maccari, S., Krugers, H.J., Morley-Fletcher, S., Szyf, M. & Brunton, P.J. (2014) The consequences of early-life adversity: Neurobiological, behavioural and epigenetic adaptations. *J Neuroendocrinol*, **26**, 707-723.

- Macfie, J., McElwain, N.L., Houts, R.M. & Cox, M.J. (2005) Intergenerational transmission of role reversal between parent and child: Dyadic and family systems internal working models. *Attach Hum Dev*, **7**, 51-65.
- Mahrer, N.E., Winslow, E., Wolchik, S.A., Tein, J.Y. & Sandler, I.N. (2014) Effects of a preventive parenting intervention for divorced families on the intergenerational transmission of parenting attitudes in young adult offspring. *Child Dev*, **85**, 2091-2105.
- Malmberg, J.L. & Field, C.E. (2013) Preventative behavioral parent training: A preliminary investigation of strategies for preventing at-risk children from developing later conduct problems. *Child Fam Behav Ther*, **35**, 212-227.
- Manassis, K., Changgun Le, T., Bennett, K., Zhao, X.Y., Mendlowitz, S., Duda, S., Saini, M., Wilansky, P., Baer, S., Barrett, P., Bodden, D., Cobham, V.E., Dadds, M.R., Flannery-Schroeder, E., Ginsburg, G., Heyne, D., Hudson, J.L., Kendall, P.C., Liber, J., Masia-Warner, C., Nauta, M.H., Rapee, R.M., Silverman, W., Siqueland, L., Spence, S.H., Utens, E. & Wood, J.J. (2014) Types of parental involvement in CBT with anxious youth: A preliminary meta-analysis. *J Consult Clin Psychol*, **82**, 1163-1172.
- Marco, E.M., Macrì, S. & Laviola, G. (2011) Critical age windows for neurodevelopmental psychiatric disorders: Evidence from animal models. *Neurotox Res*, **19**, 286-307.
- Mashoodh, R., Franks, B., Curley, J.P. & Champagne, F.A. (2012) Paternal social enrichment effects on maternal behavior and offspring growth. *Proc Natl Acad Sci U S A*, **109**, 17232-17238.
- McLaughlin, K.A., Fox, N.A., Zeanah, C.H. & Nelson, C.A. (2011) Adverse rearing environments and neural development in children: The development of frontal electroencephalogram asymmetry. *Biol Psychiatry*, **70**, 1008-1015.

- Messaoudi, M., Lalonde, R., Violle, N., Javelot, H., Desor, D., Nejdi, A., Bisson, J.F., Rougeot, C., Pichelin, M., Cazaubiel, M. & Cazaubiel, J.M. (2011) Assessment of psychotropic-like properties of a probiotic formulation (*Lactobacillus helveticus* R0052 and *Bifidobacterium longum* R0175) in rats and human subjects. *Br J Nutr*, **105**, 755-764.
- Meyer, J.S. & Hamel, A.F. (2014) Models of stress in nonhuman primates and their relevance for human psychopathology and endocrine dysfunction. *ILAR J*, **55**, 347-360.
- Moriceau, S., Shionoya, K., Jakubs, K. & Sullivan, R.M. (2009) Early-life stress disrupts attachment learning: The role of amygdala corticosterone, locus ceruleus corticotropin releasing hormone, and olfactory bulb norepinephrine. *J Neurosci*, **29**, 15745-15755.
- Moriceau, S., Wilson, D.A., Levine, S. & Sullivan, R.M. (2006) Dual circuitry for odor-shock conditioning during infancy: Corticosterone switches between fear and attraction via amygdala. *J Neurosci*, **26**, 6737-6748.
- Morris, M.J.P. & Monteggia, L.M.P. (2014) Role of DNA methylation and the DNA methyltransferases in learning and memory. *Dialogues Clin Neurosci*, **16**, 359-371.
- Mulligan, C.J., D'Errico, N.C., Stees, J. & Hughes, D.A. (2012) Methylation changes at NR3C1 in newborns associate with maternal prenatal stress exposure and newborn birth weight. *Epigenetics*, **7**, 853-857.
- Murgatroyd, C., Quinn, J.P., Sharp, H.M., Pickles, A. & Hill, J. (2015) Effects of prenatal and postnatal depression, and maternal stroking, at the glucocorticoid receptor gene. *Transl Psychiatry*, **5**, e560.
- Nanni, V., Uher, R. & Danese, A. (2012) Childhood maltreatment predicts unfavorable course of illness and treatment outcome in depression: A meta-analysis. *Am J Psychiatry*, **169**, 141-151.

- Nauta, M.H., Festen, H., Reichart, C.G., Nolen, W.A., Stant, A.D., Bockting, C.L.H., van der Wee, N.J.A., Beekman, A., Doreleijers, T.A.H., Hartman, C.A., de Jong, P.J. & de Vries, S.O. (2012) Preventing mood and anxiety disorders in youth: A multi-centre RCT in the high risk offspring of depressed and anxious patients. *BMC Psychiatry*, **12**, 31.
- Nelson, C.A. (2015) An international approach to research on brain development. *Trends Cogn Sci*, **19**, 424-426.
- Neufeld, K.-A.M., Kang, N., Bienenstock, J. & Foster, J.A. (2011) Effects of intestinal microbiota on anxiety-like behavior. *Commun Integr Biol*, **4**, 492-494.
- Nishi, M., Horii-Hayashi, N. & Sasagawa, T. (2014) Effects of early life adverse experiences on brain activity: Implications from maternal separation models in rodents. *Front Neurosci*, **8**, 166.
- Oriana Linares, L., Montalto, D., Li, M. & Oza, V.S. (2006) A promising parenting intervention in foster care. *J Consult Clin Psychol*, **74**, 32-41.
- Painter, R.C., Roseboom, T.J. & Bleker, O.P. (2005) Prenatal exposure to the Dutch famine and disease in later life: An overview. *Reproductive Toxicology*, **20**, 345-352.
- Palosaari, E., Punamäki, R.-L., Qouta, S. & Diab, M. (2013) Intergenerational effects of war trauma among Palestinian families mediated via psychological maltreatment. *Child Abuse Negl*, **37**, 955-968.
- Peña, C.J., Bagot, R.C., Labonté, B. & Nestler, E.J. (2014) Epigenetic signaling in psychiatric disorders. *J Mol Biol*, **426**, 3389-3412.
- Perroud, N., Paoloni-Giacobino, A., Prada, P., Olie, E., Salzman, A., Nicastro, R., Guillaume, S., Mouthon, D., Stouder, C., Dieben, K., Huguelet, P., Courtet, P. & Malafosse, A. (2011) Increased methylation of glucocorticoid receptor gene (NR3C1)

- in adults with a history of childhood maltreatment: a link with the severity and type of trauma. *Transl Psychiatry*, **1**, e59.
- Pesonen, A.K., Räikkönen, K., Feldt, K., Heinonen, K., Osmond, C., Phillips, D.I.W., Barker, D.J.P., Eriksson, J.G. & Kajantie, E. (2010) Childhood separation experience predicts HPA axis hormonal responses in late adulthood: A natural experiment of World War II. *Psychoneuroendocrinology*, **35**, 758-767.
- Phiel, C.J., Zhang, F., Huang, E.Y., Guenther, M.G., Lazar, M.A. & Klein, P.S. (2001) Histone deacetylase is a direct target of valproic acid, a potent anticonvulsant, mood stabilizer, and teratogen. *J Biol Chem*, **276**, 36734-36741.
- Quinlivan, J.A., Tan, L.H., Steele, A. & Black, K. (2004) Impact of demographic factors, early family relationships and depressive symptomatology in teenage pregnancy. *Aust N Z J Psychiatry*, **38**, 197-203.
- Rao, A.V., Basted, A.C., Beaulne, T.M., Katzman, M.A., Iorio, C., Berardi, J.M. & Logan, A.C. (2009) A randomized, double-blind, placebo-controlled pilot study of a probiotic in emotional symptoms of chronic fatigue syndrome. *Gut Pathog*, **1**, 6.
- Reik, W., Dean, W. & Walter, J. (2001) Epigenetic reprogramming in mammalian development. *Science*, **293**, 1089-1093.
- Reupert, A.E., Cuff, R., Drost, L., Foster, K., van Doesum, K.T. & van Santvoort, F. (2013) Intervention programs for children whose parents have a mental illness: A review. *Med J Aust*, **199**, S18-S22.
- Roberts, A.L., Galea, S., Austin, S.B., Cerda, M., Wright, R.J., Rich-Edwards, J.W. & Koenen, K.C. (2012) Posttraumatic stress disorder across two generations: Concordance and mechanisms in a population-based sample. *Biol Psychiatry*, **72**, 505-511.

- Rodgers, A.B. & Bale, T.L. (2015) Germ Cell Origins of Posttraumatic Stress Disorder Risk: The Transgenerational Impact of Parental Stress Experience. *Biol Psychiatry*, **78**, 307-314.
- Rodgers, A.B., Morgan, C.P., Bronson, S.L., Revello, S. & Bale, T.L. (2013) Paternal stress exposure alters sperm microRNA content and reprograms offspring HPA stress axis regulation. *J Neurosci*, **33**, 9003-9012.
- Roth, T.L. & Sweatt, J.D. (2011) Epigenetic marking of the BDNF gene by early-life adverse experiences. *Horm Behav*, **59**, 315-320.
- Rucklidge, J.J., Andridge, R., Gorman, B., Blampied, N., Gordon, H. & Boggis, A. (2012) Shaken but unstirred? Effects of micronutrients on stress and trauma after an earthquake: RCT evidence comparing formulas and doses. *Hum Psychopharmacol*, **27**, 440-454.
- Rucklidge, J.J., Blampied, N., Gorman, B., Gordon, H.A. & Sole, E. (2014) Psychological functioning 1 year after a brief intervention using micronutrients to treat stress and anxiety related to the 2011 Christchurch earthquakes: A naturalistic follow-up. *Hum Psychopharmacol*, **29**, 230-243.
- Sacks, J.Y., McKendrick, K. & Banks, S. (2008) The impact of early trauma and abuse on residential substance abuse treatment outcomes for women. *J Subst Abuse Treat*, **34**, 90-100.
- Sanchez, M.M., Ladd, C.O. & Plotsky, P.M. (2001) Early adverse experience as a developmental risk factor for later psychopathology: Evidence from rodent and primate models. *Dev Psychopathol*, **13**, 419-449.
- Sanders, M.R. & McFarland, M. (2000) Treatment of depressed mothers with disruptive children: A controlled evaluation of cognitive behavioral family intervention. *Behav Ther*, **31**, 89-112.

- Scharf, M. (2007) Long-term effects of trauma: Psychosocial functioning of the second and third generation of Holocaust survivors. *Dev Psychopathol*, **19**, 603-622.
- Self-Brown, S., Frederick, K., Binder, S., Whitaker, D., Lutzker, J., Edwards, A. & Blankenship, J. (2011) Examining the need for cultural adaptations to an evidence-based parent training program targeting the prevention of child maltreatment. *Child Youth Serv Rev*, **33**, 1166-1172.
- Shea, A., Walsh, C., MacMillan, H. & Steiner, M. (2005) Child maltreatment and HPA axis dysregulation: Relationship to major depressive disorder and post traumatic stress disorder in females. *Psychoneuroendocrinology*, **30**, 162-178.
- Siegenthaler, E., Munder, T. & Egger, M. (2012) Effect of preventive interventions in mentally ill parents on the mental health of the offspring: Systematic review and meta-analysis. *J Am Acad Child Adolesc Psychiatry*, **51**, 8-17.
- Starr, L.R., Conway, C.C., Hammen, C.L. & Brennan, P.A. (2014) Transdiagnostic and disorder-specific models of intergenerational transmission of internalizing pathology. *Psychol Med*, **44**, 161-172.
- Stein, A.D., Pierik, F.H., Verrips, G.H.W., Susser, E.S. & Lurney, L.H. (2009) Maternal exposure to the dutch famine before conception and during pregnancy quality of life and depressive symptoms in adult offspring. *Epidemiology*, **20**, 909-915.
- Suchman, N.E., Decoste, C., McMahon, T.J., Rounsaville, B. & Mayes, L. (2011) The Mothers and Toddlers program, an attachment-based parenting intervention for substance-using women: Results at 6-week follow-up in a randomized clinical pilot. *Infant Ment Health J*, **32**, 427-449.
- Suderman, M., McGowan, P.O., Sasaki, A., Huang, T.C.T., Hallett, M.T., Meaney, M.J., Turecki, G. & Szyf, M. (2012) Conserved epigenetic sensitivity to early life

- experience in the rat and human hippocampus. *Proceedings of the National Academy of Sciences*, **109**, 17266-17272.
- Sudo, N., Chida, Y., Aiba, Y., Sonoda, J., Oyama, N., Yu, X., Kubo, C. & Koga, Y. (2004) Postnatal microbial colonization programs the hypothalamic-pituitary-adrenal system for stress response in mice. *J Physiol*, **558**, 263-275.
- Sugimoto, A., Suzuki, Y., Endo, T., Matsumoto, K., Sugiyama, T. & Someya, T. (2015) Efficacy of atomoxetine for symptoms of attention-deficit/hyperactivity disorder in children with a history of child abuse. *J Child Adolesc Psychopharmacol*, **25**, 269-271.
- Sullivan, R.M. (2001) Unique characteristics of neonatal classical conditioning: The role of the amygdala and locus coeruleus. *Integr Physiol Behav Sci*, **36**, 293-307.
- Szyf, M. (2015) Nongenetic inheritance and transgenerational epigenetics. *Trends Mol Med*, **21**, 134-144.
- Thomas, R. & Zimmer-Gembeck, M.J. (2007) Behavioral outcomes of parent-child interaction therapy and triple P-positive parenting program: A review and meta-analysis. *J Abnorm Child Psychol*, **35**, 475-495.
- Thompson, B.L. & Levitt, P. (2010) The clinical-basic interface in defining pathogenesis in disorders of neurodevelopmental origin. *Neuron*, **67**, 702-712.
- Tillisch, K., Labus, J., Kilpatrick, L., Jiang, Z., Stains, J., Ebrat, B., Guyonnet, D., Legrain-Raspaud, S., Trotin, B., Naliboff, B. & Mayer, E.A. (2013) Consumption of fermented milk product with probiotic modulates brain activity. *Gastroenterology*, **144**, 1394-1401.
- Timmer, S.G., Ho, L.K.L., Urquiza, A.J., Zebell, N.M., Fernandez Y Garcia, E. & Boys, D. (2011) The effectiveness of parent-child interaction therapy with depressive mothers:

- The changing relationship as the agent of individual change. *Child Psychiatry Hum Dev*, **42**, 406-423.
- Tyrka, A.R., Burgers, D.E., Philip, N.S., Price, L.H. & Carpenter, L.L. (2013) The neurobiological correlates of childhood adversity and implications for treatment. *Acta Psychiatr Scand*, **128**, 434-447.
- Tyrka, A.R., Parade, S.H., Eslinger, N.M., Marsit, C.J., Lesueur, C., Armstrong, D.A., Philip, N.S., Josefson, B. & Seifer, R. (2015) Methylation of exons 1D, 1F, and 1H of the glucocorticoid receptor gene promoter and exposure to adversity in preschool-aged children. *Dev Psychopathol*, **27**, 577-585.
- Tyrka, A.R., Price, L.H., Marsit, C., Walters, O.C. & Carpenter, L.L. (2012) Childhood adversity and epigenetic modulation of the leukocyte glucocorticoid receptor: Preliminary findings in healthy adults. *PLoS One*, **7**, e30148.
- Van Loon, L.M.A., Granic, I. & Engels, R.C.M.E. (2011) The role of maternal depression on treatment outcome for children with externalizing behavior problems. *J Psychopathol Behav Assess*, **33**, 178-186.
- Vanhees, K., Vonhögen, I.G.C., Van Schooten, F.J. & Godschalk, R.W.L. (2014) You are what you eat, and so are your children: The impact of micronutrients on the epigenetic programming of offspring. *Cell Mol Life Sci*, **71**, 271-285.
- Veenendaal, M.V.E., Painter, R.C., De Rooij, S.R., Bossuyt, P.M.M., Van Der Post, J.A.M., Gluckman, P.D., Hanson, M.A. & Roseboom, T.J. (2013) Transgenerational effects of prenatal exposure to the 1944-45 Dutch famine. *BJOG*, **120**, 548-553.
- Watson, J.B., Mednick, S.A., Huttunen, M. & Wang, X. (1999) Prenatal teratogens and the development of adult mental illness. *Dev Psychopathol*, **11**, 457-466.

- Weaver, I.C.G., Cervoni, N., Champagne, F.A., D'Alessio, A.C., Sharma, S., Seckl, J.R., Dymov, S., Szyf, M. & Meaney, M.J. (2004) Epigenetic programming by maternal behavior. *Nat Neurosci*, **7**, 847-854.
- Weaver, I.C.G., Champagne, F.A., Brown, S.E., Dymov, S., Sharma, S., Meaney, M.J. & Szyf, M. (2005) Reversal of maternal programming of stress responses in adult offspring through methyl supplementation: Altering epigenetic marking later in life. *J Neurosci*, **25**, 11045-11054.
- Weaver, I.C.G., Meaney, M.J. & Szyf, M. (2006) Maternal care effects on the hippocampal transcriptome and anxiety-mediated behaviors in the offspring that are reversible in adulthood. *Proc Natl Acad Sci U S A*, **103**, 3480-3485.
- Webster-Stratton, C. & Hammond, M. (1990) Predictors of treatment outcome in parent training for families with conduct problem children. *Behav Ther*, **21**, 319-337.
- Wolchik, S.A., West, S.G., Westover, S., Sandler, I.N., Martin, A., Lustig, J., Tein, J.Y. & Fisher, J. (1993) The children of divorce parenting intervention: Outcome evaluation of an empirically based program. *Am J Community Psychol*, **21**, 293-331.
- Yano, J.M., Yu, K., Donaldson, G.P., Shastri, G.G., Ann, P., Ma, L., Nagler, C.R., Ismagilov, R.F., Mazmanian, S.K. & Hsiao, E.Y. (2015) Indigenous bacteria from the gut microbiota regulate host serotonin biosynthesis. *Cell*, **161**, 264-276.
- Yehuda, R., Bell, A., Bierer, L.M. & Schmeidler, J. (2008) Maternal, not paternal, PTSD is related to increased risk for PTSD in offspring of Holocaust survivors. *J Psychiatr Res*, **42**, 1104-1111.
- Yehuda, R., Daskalakis, N.P., Lehrner, A., Desarnaud, F., Bader, H.N., Makotkine, I., Flory, J.D., Bierer, L.M. & Meaney, M.J. (2014) Influences of maternal and paternal PTSD on epigenetic regulation of the glucocorticoid receptor gene in Holocaust survivor offspring. *Am J Psychiatry*, **171**, 872-880.

Yehuda, R., Engel, S.M., Brand, S.R., Seckl, J., Marcus, S.M. & Berkowitz, G.S. (2005)

Transgenerational effects of posttraumatic stress disorder in babies of mothers exposed to the World Trade Center attacks during pregnancy. *J Clin Endocrinol Metab*, **90**, 4115-4118.

Yehuda, R., Halligan, S.L. & Grossman, R. (2001) Childhood trauma and risk for PTSD:

Relationship to intergenerational effects of trauma, parental PTSD, and cortisol excretion. *Dev Psychopathol*, **13**, 733-753.