Major depression (M D) is one of the most costly diseases in the world (Wasser- man, 2006). Untreated affective disorders tend to be chronic and also life-threaten- ing, since depression commonly pre- cedes suicide. In the clinic, these patients must be assessed for their risk of suicide, as it is estimated to be the cause of death in 10-15% of individuals with M D (Wasser- man, 2006). To expand the possi- bilities of intervention and therapy of suicidality, particularly in connection to the treatment of M D, its neurobiological causality is being explored (M ann, 2003; W asserman, 2001).

In a stress-vulnerability model, genetic set-up, as well as environmental exposure to psychological stress, contributes to a person’s predisposition for suicidality, as well as to M D (H asler et al., 2004; M ann, 2003; W asserman, 2001). The main neuro- chemical findings on suicidality have suggested alterations in neurosystems which are usually implicated in M D; a lowered serotonergic (5HT) activity, depletion of the noradrenergic (NA) system and dysregulation of the hypothalamic-pituitary-adrenal (HPA) axis. Whereas the genes of e.g. the 5HT system and of the key NA -biosynthesis’ enzyme, tyrosine hydroxylase, have been studied extensively in this context (Bondy et al., 2006; Rujescu et al., 2007), the genes in the HPA axis have only begun to be investigated recently.

A non-overflow HPA axis is also the most consistent neurobiological indicator of M D (Bale, 2005; H asler et al., 2004; N emeroff & Vale, 2005; R oy et al., 1987; Swaab et al., 2005). This is true also true for some suicidal individuals, as an over- active HPA axis, particularly in connection to depression, is often preceding suicide (C oryell & Schlesser, 2001; Young, 2005). The HPA axis is primarially attributed to the action of corticotropin releasing hormone (CRH) and by altered feedback regulation of the axis through the glucocorticoid and mineralocorticoid receptors, which is manifested as hypercortisolism and a failure to sup- press cortisol. CRH alterations have also been observed in extrahypothalamic brain regions, e.g. the frontal cortex, limbic regions and cerebro spinal fluid (CSF). CRH acts as the key mediator of behavioral, cognitive, autonomic, neuro- endocrine and immunologic responses to aversive stimuli (emotional and physical stress or inflammation), acting mainly through the CRH receptor 1 (CRH R1).

In the HPA axis, CRH binds to CRH R1 in the anterior pituitary. This results in production of proopiomelanocortin (POMC)-derived peptides, such as adrenocorticotropic hormone (ACTH), which in turn stimulate the adrenal gland to secrete glucocorticoids (but also mineralocorticoids and androgens). The steroid acts on G Rs and M Rs in many tissues, including the brain, particularly parts of the brain stem and limbic regions. A functional response in the feedback regulation, occuring through the balanced activation of G Rs and M Rs, as well as by downregulating the expression of CRH R1 in the pituitary (A guiera et al., 2004), is crucial for returning to a normal state. A chronic state of CRH-mediated activity, e.g. due to adverse environmental stress or defective coping with low levels of stress, may result in disruption of the HPA axis feedback with consequences for e.g. the innervating limbic functions. Monoaminergic neurotransmission is affected, by e.g. suppression of the 5-HT receptor 1a in the dorsal raphe nuclei (M ejer & de Kloet, 1998) or by affecting NA in locus coeruleus (LC). Parallel effects of high levels of CRH (by action through its receptors), includes the increased release of pro-inflammatory cytokines by immune cells, activating the indoleamine 2,3-dioxygenase (IDO) pathway with subsequent consequences of tryptophan (5HT-) depletion and neurotoxicity, as well as glutamatergic hyperfunction (Leonard, 2005; Muller & Schwarz, 2007). Thus, disturbances in HPA axis responsivity in connection to CRH, are relevant in M D, as well as in suicidality.

The effects of CRH dysregulation are primarily mediated through CRH R1. Consequently, antagonists active against CRH R1 are being developed as a novel type of antidepressants (Nielsen, 2006). From a pharmacogenomic perspective, it is of importance to have access to the level of individual variation in this gene in context to its functionality, which can be needed for evaluation of treatment efficacy as well as being used in diagnostic tools.

Knowledge about dysfunctional CRH R1 variants may also bring about new, highly specific treatment possibilities with inhibitory RNA s (Sah, 2006). Surprisingly little is known about influence of genetic variations in the CRH R1 gene in the context of stress and depression, since only a few such studies have been reported up to date (Licinio et al., 2004; Liu et al., 2007; Liu et al., 2006; Papiol et al., 2007; Wasserman et al., 2007). In fact, our group has been the first to study the genetic variation in the CRH R1 gene in connection to depression and stress among suicidal individuals (Wasserman et al., 2007). Genetic variation in CRH R1 showed association and linkage to suicide attempt in depressed males exposed to low levels of stress. This association to low stress was true also for female suicide attempters, but without relation to depression. This may reflect the genetics behind the fact that a higher proportion of the depressed males have been shown to be suicidal, compared with depressed females (Wasser- man, 2001). Although there is need of further confirmation of our novel finding and analysis of the CRH R1 locus, we hypothesized that these individuals may be genetically predisposed to react with dysregulated HPA activity upon a low threshold of stress, through some alterations of the CRH R1 gene expression or functionality, possibly experiencing a chronic state of HPA activation and/or a reduced feedback mechanism, which partly substitute the environmental influence of high levels of stressful life events in the suicidal process (Wasserman et al., 2007). Interestingly our previous novel findings demonstrated that genetic

1) This review was first published August 16th, 2007 in the WPA Electronic Bulletin (World Psychiatric Association), http://wpanet.org/bulletin/eb54.html
variation in a transcription factor of the PO/MC gene, TBX19, which is regulated by CRH, showed association and linkage to the anger/hostility personality trait and suicidality (Wasserman et al., 2006). Taken together, our result suggest that genetic variation in the CRH-mediated regulation of the HPA axis is a factor of importance in suicidality and, as other have shown as well, for MD.

A thorough understanding of the dysregulation occurring at the HPA axis in suicidality is however only in its infancy. Whereas the dexamethasone suppression test (DST) is proving itself as an increasingly more viable prognostic test for the suicidality among certain depressed individuals, there is a substantial need for inclusion of additional measurement variables (Coryell & Schlesser, 2007; Coryell et al., 2006; Mann et al., 2006). For example, there is a need to explain the observed reduction in HPA axis activity in connection to suicidality (Pfenning et al., 2005), as well as to be able to differentiate from non-suicidal patients with MD. Since feedback in the HPA axis occurs both at the level of glucocorticoid receptors, as well as CRH R1 in the pituitary, it is likely that genetic variation in several of these genes system must taken into account in evaluating such results. Moreover, other suicidal individuals show lowered 5HT activity and yet others may show variation in further neurobiological systems (Mann, 2003; Wasserman, 2001). This is of interest in light of that the HPA axis is complexly interconnected with many of these systems, e.g. monoaminergic and glutamatergic (Leonard, 2005; Meijer & de Kloet, 1998; Müller & Schwarz, 2007), and that the understanding of this network is likely to require a more detailed dissection of the variabilities in these genes, as well as a consideration of interacting or redundant effects between genes, in the context of environmental stress (Kendler, 2005).

Thus, the exploration of the HPA axis at the level of genetic variability hold promises of delivering significant explanations as to why certain individuals are at risk for suicidality, in connection to MD or other conditions of stress, as well as being seemingly necessary for the construction of reliable diagnostic and treatment tools, with sufficient sensitivity and specificity for the suicidal aspect of these patients.

References


