

# **Neurocriminology and Serial Violence: A Meta-Analysis of Biological and Neurological Correlates of Criminal Behavior**

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## **Abstract**

This meta-analysis synthesizes recent findings in neurocriminology to examine the biological and neurological underpinnings of criminal behavior, with a specific focus on serial killers and individuals exhibiting extreme violence. Drawing from studies across neuroimaging, genetic research, and psychophysiological assessments, the review identifies consistent patterns of dysfunction in brain regions such as the prefrontal cortex, amygdala, and limbic system. Genetic markers, hormonal imbalances, and prenatal factors are also examined for their predictive role in antisocial behavior. Across reviewed studies, a common thread emerges: deficits in impulse control, emotional regulation, and moral reasoning are strongly associated with both structural and functional brain abnormalities. Environmental interactions—especially early trauma—appear to exacerbate biological vulnerabilities. By integrating multi-domain findings, this paper highlights the value of neuro criminological frameworks in informing criminal profiling, legal responsibility assessments, and rehabilitative strategies. Implications for mental health interventions and forensic evaluations are discussed.

**Keywords:** neurocriminology, serial killers, brain abnormalities, meta-analysis, violence, antisocial behavior, forensic psychology

## **Introduction**

The intersection between biology and criminal behavior has gained significant attention with the emergence of neurocriminology, an interdisciplinary field that merges neuroscience and criminology to examine the biological bases of violent and antisocial behavior. While criminology has traditionally focused on sociological and psychological explanations for crime, neurocriminology emphasizes the role of brain structure, function, and neurochemical pathways in shaping behavior that deviates from societal norms.

In recent decades, research using functional magnetic resonance imaging (fMRI), positron emission tomography (PET), and other neurological techniques has revealed compelling links between brain abnormalities and violent tendencies, particularly in individuals classified as serial killers or diagnosed with antisocial personality disorder (ASPD) and psychopathy. These findings are supported by evidence of genetic predispositions, hormonal dysregulation, and early developmental insults that may affect the brain's capacity for emotional regulation and impulse control.

This paper presents a meta-analytic review of empirical studies investigating the neurobiological underpinnings of violent criminal behavior. The review aims to identify consistent findings across multiple domains—including neuroanatomy, genetics, hormones, and psychophysiological responses—to better understand the convergence of biological vulnerabilities in violent offenders. By synthesizing the available literature, this paper seeks to highlight key patterns, identify gaps in existing research, and propose directions for more integrative approaches to criminal profiling and forensic intervention.

## **Method**

### **Search Strategy**

A systematic literature search was conducted using academic databases including PubMed, PsycINFO, Scopus, and Google Scholar to identify peer-reviewed studies examining biological, neurological, or genetic correlates of violent or antisocial behavior. The search spanned publications from 2000 to 2024, using combinations of the following keywords: neurocriminology, serial killers, antisocial personality disorder, brain abnormalities, psychopathy, genetics and crime, neuroimaging, violence, and hormones and aggression.

### **Inclusion Criteria**

Studies were included if they:

- Were published in English between 2000–2024
- Focused on violent criminal behavior (e.g., homicide, serial killing, psychopathy)
- Investigated biological, neurological, hormonal, or genetic factors
- Used empirical research methods (e.g., neuroimaging, genetic testing, physiological measurement)
- Provided full-text access to results

### **Exclusion Criteria**

Studies were excluded if they:

- Focused solely on non-violent or white-collar crime
- Were theoretical or editorial in nature
- Did not include biological or neurological components
- Had non-human samples (e.g., animal models)

### **Study Selection and Data Extraction**

Titles and abstracts of identified articles were screened for relevance. Eligible full-text studies were then reviewed in detail. Data was extracted on sample characteristics, methodology, neurobiological focus (e.g., brain region or system studied), and key findings. A total of X studies (you can fill in the number) were included in the final analysis.

## **Results and Thematic Analysis**

### **Brain Structure and Function in Violent Offenders: A Meta-Analytical Perspective**

Research across neurocriminology consistently points to abnormalities in prefrontal-limbic circuitry as a key factor in violent behavior. This meta-analysis identifies common neuroanatomical patterns from multiple studies focusing on individuals with antisocial personality disorder (ASPD), psychopathy, and homicidal behavior, particularly among serial offenders.

**Prefrontal Cortex**

Meta-analyses and neuroimaging studies frequently report reduced gray matter volume and hypoactivity in the prefrontal cortex, especially the orbitofrontal and dorsolateral regions (Yang & Raine, 2009; Brower & Price, 2001). These areas govern impulse control, moral judgment, and emotional regulation. Deficits here are associated with poor decision-making, lack of remorse, and disinhibited aggression—hallmarks of violent criminal behavior.

**Amygdala**

The amygdala, a core structure in emotional processing, shows structural and functional irregularities in violent individuals. Studies suggest both hypoactivity during fear processing (Blair, 2007) and volume reduction (Glenn et al., 2010), which may contribute to empathy deficits, emotional blunting, and lack of fear conditioning—traits observed in many serial killers and individuals with psychopathy.

**Limbic System Connectivity**

Disruptions in prefrontal-limbic connectivity have also been noted, indicating impaired regulation of emotional impulses (Raine et al., 1998). The hippocampus, cingulate cortex, and hypothalamus show variable activity, but many studies converge on a model where emotional dysregulation, paired with poor executive control, increases risk for violent action.

Overall, findings across the literature support the theory that violent criminal behavior is often underpinned by neurostructural deficits in regions essential for emotional regulation, inhibition, and moral decision-making. These findings offer potential utility in forensic assessment and early intervention programs.

**Genetic and Epigenetic Influences on Criminal Behavior**

A growing body of research supports the role of genetic and epigenetic factors in the predisposition toward violent and antisocial behavior. This section synthesizes findings from studies examining heritable traits, gene-environment interactions, and molecular mechanisms associated with aggression, impulsivity, and criminality.

**MAOA Gene and Violence**

One of the most frequently cited genetic markers in neurocriminology is the MAOA gene, often referred to as the "warrior gene." Variants of this gene, particularly the low-activity MAOA-L allele, have been associated with increased risk for impulsive aggression and antisocial behavior, especially when combined with early-life trauma (Caspi et al., 2002). Meta-analytic reviews confirm a significant gene-environment interaction, where the effect of MAOA-L is amplified in individuals exposed to childhood abuse or neglect (Byrd & Manuck, 2014).

**Other Candidate Genes**

Beyond MAOA, genes regulating dopaminergic (e.g., DRD2, DRD4) and serotonergic systems (e.g., 5-HTTLPR) have also been implicated. These genes influence neurotransmitter function, which affects emotional reactivity, risk-taking, and self-regulation—traits frequently observed in offenders with violent histories. However, findings across studies are mixed, suggesting that no single gene determines criminality; rather, polygenic models are more accurate.

### **Epigenetic Modifications**

Recent studies have begun to explore how epigenetic changes—modifications that affect gene expression without altering DNA sequences—may contribute to antisocial behavior. For example, methylation of genes associated with stress regulation (e.g., NR3C1) has been observed in individuals with traumatic backgrounds and conduct problems (Provencal et al., 2015). These findings highlight how environmental stressors can biologically embed into behaviorally relevant systems.

Together, these genetic and epigenetic insights suggest that biological predispositions, especially when paired with adverse environments, may significantly increase the risk for violent behavior. However, the field cautions against deterministic interpretations and emphasizes the need for integrated biosocial models.

### **Prenatal and Perinatal Risk Factors in Violent Behavior**

Meta-analytic reviews and longitudinal studies have revealed that early biological insults—beginning as early as prenatal development—can increase the likelihood of violent and antisocial outcomes later in life. These include birth complications, prenatal exposure to toxins, and malnutrition, all of which may affect brain development, particularly in areas related to emotional regulation and executive functioning.

#### **Birth Complications and Brain Injury**

Studies consistently link obstetric complications—such as hypoxia (oxygen deprivation), premature birth, and low birth weight—with increased risk of violent behavior in adolescence and adulthood. A prominent meta-analysis by Hodgins and colleagues (2002) found that individuals with a history of perinatal complications had significantly higher rates of aggressive conduct disorder and criminal conviction, especially when combined with psychosocial adversity.

#### **Prenatal Exposure to Toxins**

Exposure to alcohol, nicotine, and lead during pregnancy has also been implicated in the development of behavioral disorders. Fetal Alcohol Spectrum Disorders (FASD), in particular, are associated with reduced volume in the prefrontal cortex and impaired social cognition—traits commonly linked to conduct disorder and delinquency (Mattson et al., 2001).

#### **Maternal Stress and Malnutrition**

Emerging research also points to the role of prenatal maternal stress in altering fetal brain development through elevated cortisol levels. Malnutrition during pregnancy, particularly deficiencies in omega-3 fatty acids, iron, and zinc, may impair neural connectivity, increasing susceptibility to impulsivity and externalizing behaviors in childhood and adolescence (Raine et al., 2003).

Collectively, the literature suggests that prenatal and perinatal factors contribute significantly to the development of neurobiological vulnerabilities. When these vulnerabilities are coupled with negative postnatal environments, the risk of persistent violent and antisocial trajectories increases markedly.

### **Hormonal and Neurochemical Influences on Aggression and Violence**

A substantial body of research has investigated how hormonal imbalances and neurochemical dysregulation contribute to aggressive, impulsive, and antisocial behavior. This section synthesizes

findings from meta-analyses and neurochemical studies on the roles of testosterone, cortisol, dopamine, serotonin, and oxytocin in shaping violent tendencies.

### **Testosterone and Aggression**

Testosterone, a steroid hormone associated with dominance behavior, has been repeatedly linked to aggression and antisocial traits. Meta-analytical findings suggest a modest but consistent correlation between high testosterone levels and violent behavior, particularly in males (Book et al., 2001). However, the effect size is often moderated by social context, suggesting that testosterone may amplify pre existing behavioral tendencies rather than cause violence independently.

### **Cortisol and Stress Reactivity**

In contrast to testosterone, cortisol—the primary stress hormone—has shown an inverse relationship with aggression. Individuals with low basal cortisol levels often exhibit fearlessness, risk-taking, and reduced sensitivity to punishment, characteristics common among habitual offenders (McBurnett et al., 2000). This hypoarousal profile may reflect a blunted physiological response to stress, diminishing internal restraints on antisocial behavior.

### **Serotonin and Impulse Control**

Serotonin plays a key role in mood regulation and impulse control. Numerous studies link low serotonergic activity—measured via cerebrospinal fluid (CSF) or pharmacological challenge tests—to impulsivity and violent aggression, particularly in individuals with borderline or antisocial traits (Lesch & Merschdorf, 2000). These deficits may undermine the capacity for emotional regulation, contributing to reactive aggression.

### **Dopamine and Reward Sensitivity**

Dopaminergic systems are central to reward-seeking behavior. Abnormalities in dopamine transmission—particularly overactivation of the mesolimbic pathway—have been associated with sensation seeking, substance abuse, and criminal risk-taking. While not a direct cause of violence, these tendencies may increase vulnerability to antisocial lifestyles.

### **Oxytocin and Social Bonding**

While often considered a “pro-social” hormone, oxytocin’s role in aggression is context-dependent. Some studies show reduced oxytocin activity in individuals with psychopathic traits, suggesting a blunted capacity for empathy and attachment. However, other findings suggest that oxytocin may intensify in-group loyalty and out-group hostility, complicating its interpretation.

In summary, the literature supports a nuanced relationship between hormonal and neurochemical systems and violent behavior. These systems interact dynamically with environmental triggers and cognitive appraisals, reinforcing the need for multifactorial models in understanding and managing aggression.

### **Psychophysiological Correlates of Violent Behavior**

Psychophysiological studies offer insights into the autonomic nervous system (ANS) responses of individuals with antisocial and violent tendencies. This area of research focuses on measurable

biological signals such as heart rate, skin conductance, and electroencephalographic (EEG) activity, providing evidence of emotional underarousal, fearlessness, and impulse dysregulation in violent offenders.

### **Resting Heart Rate and Fearlessness**

One of the most consistent findings across meta-analyses is that low resting heart rate is significantly associated with aggressive and antisocial behavior, particularly in males (Ortiz & Raine, 2004). This physiological profile is theorized to reflect a fearlessness that reduces responsiveness to punishment or social consequences. It may also signal a need for external stimulation, making individuals more likely to engage in risky or thrill-seeking behavior, including violence.

### **Skin Conductance and Emotional Blunting**

Studies on skin conductance response (SCR)—a marker of sympathetic nervous system activity—indicate that individuals with psychopathic or callous-unemotional traits often display blunted SCRs when exposed to emotionally salient or threatening stimuli (Lorber, 2004). This under-reactivity suggests a lack of emotional arousal, contributing to reduced empathy, poor fear conditioning, and disinhibition.

### **Electroencephalography (EEG) and Impulsivity**

EEG studies show that violent individuals often exhibit abnormal brain wave patterns, particularly excessive slow-wave activity (theta and delta waves) in the frontal cortex. These patterns are commonly associated with attention deficits, impulsivity, and behavioral dysregulation—all of which are elevated in antisocial populations (Gao & Raine, 2010).

### **Startle Reflex and Emotional Deficits**

Deficits in the startle reflex—typically measured through eyeblink magnitude in response to sudden stimuli—have also been documented in individuals with psychopathy. Reduced startle potentiation in threatening or aversive contexts suggests emotional hyporesponsiveness, consistent with neuroimaging findings of amygdala dysfunction.

Collectively, these psychophysiological patterns reinforce the idea that biological underarousal and emotion-processing deficits are central features in violent and antisocial individuals. These markers may hold promise for early identification, risk assessment, and tailored intervention strategies in forensic contexts.

### **Brain Imaging and Neurological Evidence in Violent Offenders**

Neuroimaging techniques such as functional magnetic resonance imaging (fMRI), positron emission tomography (PET), and structural MRI have revolutionized our understanding of how brain abnormalities relate to violent and antisocial behavior. This section synthesizes findings from studies that utilize neuroimaging to explore structural and functional deficits in offenders, particularly those with psychopathy, antisocial personality disorder (ASPD), and histories of homicidal or serial violence.

### **Structural Abnormalities**

Meta-analyses of MRI studies have revealed consistent reductions in gray matter volume in areas associa



ted with impulse control, moral reasoning, and empathy—notably the prefrontal cortex, anterior cingulate cortex, and amygdala (Yang & Raine, 2009). For example, violent individuals often show reduced volume in the orbitofrontal cortex, which is crucial for integrating emotional signals into decision-making.

### **Functional Impairments**

fMRI and PET studies suggest that violent offenders exhibit hypoactivity in frontal and temporal regions during tasks involving moral judgment, emotional recognition, and inhibition control. For instance, research by Kiehl et al. (2001) found significantly decreased activity in the paralimbic system (including the insula and temporal poles) in incarcerated psychopaths during emotional processing tasks.

### **Connectivity Deficits**

Recent studies using diffusion tensor imaging (DTI) highlight disrupted connectivity between the prefrontal cortex and limbic regions, including the amygdala and hippocampus (Craig et al., 2009). These findings suggest a breakdown in the communication pathways needed to regulate affective responses, contributing to impulsivity, emotional detachment, and poor behavioral control.

### **Brain Abnormalities in Serial Killers**

Though sample sizes are limited, case studies of serial killers and mass murderers have frequently reported anomalies in frontal and temporal lobes, as well as lesions or damage from traumatic brain injury (TBI). Such damage may impair moral reasoning and empathic capacity, two hallmarks of psychopathic behavior.

Together, brain imaging findings underscore the neurobiological contributions to violent behavior and provide objective markers that may enhance forensic evaluation, risk assessment, and rehabilitation planning.

### **Challenges and Limitations in Neurocriminology**

While the field of neurocriminology has made substantial progress in identifying biological correlates of violent and antisocial behavior, several methodological and ethical challenges complicate the interpretation and application of these findings.

### **Methodological Variability**

One of the primary limitations across studies is the inconsistency in research design and measurement tools. Sample sizes often vary widely, and operational definitions of “violent behavior” can range from aggressive tendencies to serial homicide. This heterogeneity makes meta-analytical synthesis difficult and raises questions about generalizability. Additionally, many studies rely on cross-sectional designs, limiting causal inference between brain abnormalities and criminal behavior.

### **Overemphasis on Biological Determinism**

There is an ongoing risk of biological reductionism, where complex behaviors are attributed solely to genetic or neurological causes. While neurocriminology aims to highlight biological vulnerabilities, critics argue that it sometimes downplays sociocultural and environmental influences such as poverty,

trauma, or systemic oppression. This can lead to oversimplified narratives about “born criminals,” potentially reinforcing stigma or bias in forensic contexts.

### **Ethical Implications**

The application of neurocriminological findings in legal and clinical settings raises serious ethical concerns. For instance, if certain brain patterns are associated with violence, how should the justice system weigh this information in terms of culpability, sentencing, or rehabilitation eligibility? The potential for neuroscientific evidence to be misinterpreted or overvalued in courtrooms calls for clear guidelines and caution.

### **Lack of Diversity and Cultural Representation**

Most research in neurocriminology has been conducted in Western, industrialized contexts, often neglecting cultural variability in emotional expression, moral reasoning, and justice systems. The underrepresentation of marginalized groups in neurocriminological samples limits the universality of current findings and risks perpetuating bias.

In summary, while neurocriminology offers valuable tools and perspectives, its findings must be contextualized within a broader biopsychosocial framework. Future research should prioritize methodological rigor, ethical responsibility, and inclusive sampling to ensure that scientific advancements are applied with justice and equity.

### **Future Directions in Neurocriminology**

As neurocriminology continues to evolve, there is growing recognition of the need for more integrative, ethical, and culturally sensitive research. Building upon current findings, several key directions can guide the future of this interdisciplinary field.

### **Longitudinal and Developmental Studies**

Future research should prioritize longitudinal designs to better understand how neurobiological vulnerabilities interact with life experiences over time. By tracking individuals from childhood into adulthood, researchers can more clearly identify developmental trajectories that lead to persistent antisocial behavior and examine critical windows for intervention.

### **Multimodal Approaches**

Combining neuroimaging, genetic testing, psychophysiological measures, and behavioral assessments in a single study can provide a more nuanced understanding of the biological roots of violence. Multimodal approaches also allow for the development of individualized risk profiles, which can inform tailored rehabilitation strategies and preventive measures.

### **Ethical Integration into Forensic Practice**

The increasing use of neuroscientific evidence in criminal justice calls for standardized guidelines that balance scientific validity with legal fairness. This includes training legal professionals in the interpretation of brain-based evidence and ensuring it is used to enhance, not replace, human judgment in evaluating responsibility and sentencing.



### **Early Intervention and Prevention**

Findings from neurocriminology should be translated into early screening and prevention programs, particularly in high-risk youth populations. Identifying markers of impulsivity, emotional dysregulation, or physiological underarousal in early childhood may allow for proactive mental health interventions that reduce future antisocial trajectories.

### **Cultural and Global Expansion**

To ensure that neurocriminological insights are relevant and equitable, future studies must expand beyond Western samples and consider the role of culture in shaping brain-behavior relationships. This includes examining how societal norms, trauma exposure, and structural inequalities influence the expression and consequences of biological vulnerabilities.

By embracing complexity and interdisciplinarity, neurocriminology can move toward a future that is not only scientifically rigorous, but also ethically sound and socially responsible.

### **Conclusion**

This meta-analysis provides a comprehensive synthesis of the current literature on biological, neurological, and physiological correlates of violent and antisocial behavior, with particular emphasis on individuals exhibiting extreme patterns such as serial killing. Across domains—including brain structure and function, genetic predisposition, prenatal risk factors, hormonal influences, and psychophysiological patterns—findings converge on a shared insight: biological vulnerabilities play a significant role in shaping violent behavior, particularly when compounded by environmental risk factors such as trauma and neglect.

Structural and functional deficits in regions such as the prefrontal cortex, amygdala, and limbic system consistently emerge as biological hallmarks of impaired emotional regulation and poor impulse control. Genetic markers like MAOA-L, hormonal patterns involving testosterone and cortisol, and psychophysiological traits such as low resting heart rate or reduced skin conductance all reflect measurable biological tendencies that may increase the likelihood of violent behavior—especially when left unbuffered by protective social or psychological factors.

While the contributions of neurocriminology are substantial, this field must continue evolving in a direction that embraces methodological diversity, cultural inclusivity, and ethical reflection. Biological findings should not be used in isolation to determine criminal responsibility, but rather to enhance holistic, just, and humane approaches to understanding and preventing violence.

By situating biological risk within broader psychosocial frameworks, neurocriminology offers the potential not only to understand the roots of crime but to help create effective early interventions, rehabilitative strategies, and more compassionate criminal justice systems.

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