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Ketamine: The illicit party psychedelic that promises to heal depression

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Summary: Ketamine, an anesthetic which has been commonly used as an illicit party drug, is helping people overcome depression. The drug works rapidly and provides lasting relief from depressive symptoms for weeks after exposure. Researchers are examining how ketamine works in the brain to help positively influence behavior in those with mood disorders.

Source: The Conversation

It's been 50 years in the making, but the anaesthetic and illicit party drug ketamine is now having a clinical comeback. New studies show that this commonly used anaesthetic can provide quick relief of core symptoms associated with severe depression, including suicidal ideation.

Amazingly, ketamine works within hours and its effects are maintained for <u>at least one week</u>. Most strikingly, ketamine is effective in those patients who are <u>resistant to ordinary</u> antidepressants, and they make up around 30 to 50 percent of the depressed population.

MCFIP - Ketamine is a neuromodulator that functions as part of BDNF within the neuropeptide pancreatic polypeptide. Along with CBD and psilocybin, ketamine (esketamine) are all biosimilars that activate 5-HT serotonin receptors. DIY!

Now, efforts in <u>Mexico</u>, <u>Australia</u>, <u>France</u>, <u>Canada</u> and the <u>United States</u>, among others, are focusing on understanding exactly how ketamine does this, and to what extent it is safe and effective in a clinical setting. Together, these studies will increase our understanding of depression around the world and perhaps expand ketamine's potential to treat other forms of mental illness as well. MCFIP - Biosimilars for serotonin modulation of BDNF offset imbalances in the catecholamines within neuropeptide Y to regulate behavioral health. Regulation of these two activities are part of peptide YY; i.e. autophagy.

The focus of our lab at the University of Guelph is to understand how specific drugs, such as ketamine, work in the brain and influence behaviour.

My doctoral research, specifically, looks at the link between stress, inflammation and behaviour. I am studying how ketamine influences behaviour and can reduce the effects of stress, and what this means for mood disorders, such as major depression.

The first dissociative anaesthetic

Initially, ketamine was developed as an alternative to the well-known, illegal party drug, phencyclidine (PCP). In the late 1950s, PCP was the focus of Parke-Davis pharmaceuticals for its use as an anesthetic. However, the drug came with <u>uncomfortable side effects</u> such as delirium and a loss of feeling in the limbs, which lasted for several hours after the drug was taken.

To fix this issue, Cal Bratton of Parke-Davis encouraged scientists to look into possible ways to modify PCP, with a primary goal of reducing side effects. In 1962, the organic chemist, Calvin Stevens, made a PCP-like compound which he said had similar anaesthetic properties, with shorter acting psychoactive effects than PCP.

This compound, originally known as <u>CI-581</u>, was eventually named <u>ketamine</u> based on the ketone and amine group that formed its chemical structure.

Followings its discovery, ketamine was then used in the first human trials in the mid-1960s which included testing on volunteer inmates of Jackson Prison in Michigan, United States.

After consistent reports of feeling "disconnected" from the environment when given ketamine, it was classified as the first <u>dissociative anesthetic</u>.

In the years after its initial testing, the effects of ketamine rapidly gained popularity around the globe, and the approval as a human anaesthetic was passed by the U.S. Food and Drug Administration (FDA) in 1970 — to be sold as Ketalar.

Unique antidepressant effect

Recent studies describing ketamine's antidepressant properties have led to a dramatic shift in how we view the drug and treat mental illness.

Typical antidepressants work by controlling levels of neurotransmitters in the brain known as monoamines, such as serotonin and norepinephrine. Ketamine's antidepressant effect is unique, as it modifies the activity of glutamate, which is the main excitatory neurotransmitter in the brain and is not a monoamine. MCFIP - As neuromodulators, biosimilars for serotonin can modulate the activities of the amino acid neurotransmitters (AANTs); i.e. glutamate, glycine and aspartate.

One of the fascinating findings about ketamine is that it can rapidly reduce depressive symptoms in <u>patients who do not respond</u> to typical monoamine antidepressants. This suggests the role of glutamate in depression. MCFIP - The roles of the AANTs can be discussed for verification with qualified bioinformatics professionals.

In fact, studies have related ketamine's antidepressant effect with its ability to maintain connections between neurons (or nerve cells) in the brain. These connections are known to continuously change in response to our environment, which is a process known as plasticity. Interestingly, the ability of these neurons to change connections highly relies on normal glutamate activity. MCFIP - Our quantum biology modeling has identified, with near certainty, that serotonin is one of the three substances that function as interneurons to "modulate" neurons. Refer to the following for use as part of discussions: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4996712/

A combination of human and animal studies suggest that ketamine's antidepressant effect may involve regulating glutamate levels to <u>strengthen these connections</u> and/or restore them back to a pre-stressed state. MCFIP - Refer to previous comment relative to AANTs.

Further study of ketamine's ability to restore these neural connections and how glutamate relates to mood disorders will surely open new doors for understanding mental illness.

Nasal sprays and biomarkers

Current research has shown positive effects of ketamine for other forms of mental illness as well, such as post-traumatic stress disorder (PTSD) and <u>bipolar disorder</u>. Though the results seem quite positive, more studies are required to validate their use beyond depression.

Other studies using a repeated treatment method <u>over several weeks</u> showed that ketamine can produce long-term reductions in symptoms of treatment-resistant depression, lending potential to its safety and effectiveness over longer periods of treatment.

Recently, the FDA has approved esketamine (ketamine's "close cousin") in the U.S., sold as Spravato in the form of a nasal spray. Importantly, the spray is only prescribed to treatmentresistant patients who continue to take an oral antidepressant and can only be used under the supervision of a health-care provider. MCFIP - In our opinion, with biosimilars of ketamine currently available without prescription, the revenue potential of ketamine is minimal. Instead, the value of biosimilars of serotonin will be in the diagnostic testing of all activities associated with the neuropeptides including but not limited to BDNF and nerve growth factors. Additionally, another substantial stream of revenue will be behavioral health counseling to modify and maintain neurohormone reconfiguration. It should be noted that the physiological mechanisms regulated by the neuropeptides (e.g. formation of DNA, its repair and chirality of amino acids) will transform nearly all that is known about the causal paths of chronic diseases. Proof of concept for this assertion can been seen on the website <u>WWW.MCFIP.net</u> in various tabs for discussion; e.g. cancer issues, cardiovascular, neurodegenerative diseases, etc.

Finally, studies are also looking into biological markers that can predict response to treatment, also known as biomarkers. If successful, this research would allow for more accurate and effective treatment delivery in the form of personalised treatment plans.

Increasing accessibility to ketamine treatment for depression will be the next major milestone for this drug. It is sure to provide effective relief for those who continue to experience treatment-resistant and severe depression.

The promise of ketamine for treating mood disorders only applies in a carefully monitored clinical setting. The street drug Special K can be addictive, put users at high risk and cause long-term psychological harm.

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ABOUT THIS NEUROSCIENCE RESEARCH ARTICLE

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