An almost fearless brain

Wouldn’t it be great to live without fear? Or would it? Research is showing just how important fear can be.

By Jose Viosca

Living a fearless life is one of the favourite topics of fiction novels and films. In the 1932 novel Brave New World, happiness is easy: just take a pill and your problems will disappear, together with any fear. For me, a neuroscientist and science-fiction lover, it is amazing to see some of Aldous Huxley’s fiction becoming reality in laboratories that investigate the neural basis of fear.

Despite its varied causes, fear results in remarkably similar symptoms: your heart races, you start to sweat, you are suffocating, dizzy and nauseous. You are terrified of losing control or of dying.

The selective removal of specific fears is a key goal in neuroscience, as we search for treatments for phobias, post-traumatic stress disorder and other anxiety disorders. In Europe, more than 43 million people are affected by such conditions (see table 1), and current treatments include psychotherapy and antidepressants.

These are helpful approaches, but in some cases they do not work or they have unpleasant side effects. To provide more effective treatments, we need to understand how the brain processes fear.

Terrifying, isn’t it? What can be harmful usually also looks scary: that is one way dangers are avoided in nature.
Neurons (green) in the brain connect with thousands of other neurons, creating millions of neuronal circuits.

### Table 1: Europeans incidences of the most common fear and anxiety disorders

<table>
<thead>
<tr>
<th>Condition</th>
<th>Number affected (millions)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Panic disorder</td>
<td>5.3</td>
</tr>
<tr>
<td>Agoraphobia</td>
<td>4.0</td>
</tr>
<tr>
<td>Social phobia</td>
<td>6.7</td>
</tr>
<tr>
<td>Generalised anxiety disorder</td>
<td>5.9</td>
</tr>
<tr>
<td>Specific phobias</td>
<td>18.5</td>
</tr>
<tr>
<td>Obsessive compulsive disorder</td>
<td>2.7</td>
</tr>
</tbody>
</table>

Data source: Wittchen & Jacobi (2005)

To this end, researchers are now exploring one region of the brain: the hypothalamus. Located deep within the brain, the hypothalamus is a kind of ancient brain centre for motivations and internal drives such as hunger. Although human impulses such as the desire for happiness or love are not limited to the hypothalamus, they relate to basic motivations that have their roots in the need to survive and reproduce. If the energy levels in your body run low, for example, certain neurons in the hypothalamus will scream out, triggering the sensation of hunger. Similarly, a chain of neurobiological reactions underlies the sensation of thirst when your body is short of water; anger if someone invades your territory; and fear if you encounter a fierce dog.

How does this work? We know that the hypothalamus consists of more than 15 groups of nerve cells – neurons – which all use a variety of chemical messengers and receptors to communicate with each other. We know very little, however, about which of these types of neuron are involved in each survival motivation. Nor do we yet understand how the hypothalamus co-ordinates such...
an array of diverse functions. Are the same neurons involved? Or do distinct but neighbouring cells rapidly communicate to determine which response to trigger?

To answer these questions, scientists need to systematically deconstruct the hypothalamus into its component parts, like thousands of pieces of a large jigsaw puzzle. This can be done by switching specific neurons on or off using nothing but a flash of light.

Does this sound like the stuff of science fiction? In fact, it is one of the techniques used in the research group that I worked in at the European Molecular Biology Laboratory (EMBL) \(^1\). A photoactive protein, like those present in our eyes, is introduced into a specific type of neuron in the brain. The protein can then trigger or silence the activity of the neuron in response to a flash of laser light. When researchers in our group used light to switch off certain neurons in the hypothalamus of mice, the mice lost all sense of fear. Instead of avoiding a rat – a natural predator of mice – they approached it, forcing us to stop that particular experiment to protect the mice.

Intriguingly, a similar behaviour has been observed in humans – which first suggested to neuroscientists that fear is controlled by different areas of the brain. Before we knew the hypothalamus was involved in fear, we thought perhaps the only area of the brain involved was another small region known as the amygdala. A woman known in the scientific literature only as ‘SM’ had a genetic condition that destroyed neurons in the amygdala, which is known as the centre of emotions and an essential component in processing fear. Over the years, scientists exposed SM to all kinds of terrible stimuli including horror movies, poisonous snakes and spiders, but she showed no fear. One day, however, they asked her to breathe in air with a high concentration of carbon dioxide. This raises the carbon dioxide levels in the blood and would normally be associated with suffocation. This, finally, caused SM to panic – she screamed for help while pulling the mask away. Scientists observing her concluded that other brain regions besides the amygdala must be able to trigger and process fear.

For the first time, SM had experienced fear. What a wonderful idea, you might think. This is what scientists are searching for, to help treat anxiety disorders and other conditions. SM’s life, however, demonstrates the importance of fear. Over the years, her lack of fear had led her into many dangerous situations – including many assaults – because she did not recognise the early signs of danger, just like the mouse that carelessly approached the rat. Fear is not only unpleasant and disturbing but is also there to warn us of dangers.

This research illustrates the importance of striking a balance, which scientists continue to investigate. Basic emotions can be vital but become harmful in the extreme. In *Brave New World*, for example, people paid a very high price for happiness and fearless-ness: they lost their freedom. Perhaps the purpose of fear is also to point out the need to constantly evaluate and react, to make appropriate choices, and to seek answers about our emotions and behaviours at home, work or school.

**Reference**

Web reference

w1 – To learn more about the European Molecular Biology Laboratory (EMBL), see: www.embl.org

Resources

The fearless mouse approaching a rat was first observed in this research:
To learn more about the case of SM, see:

To find an explanation of optogenetics, the technique to control neuronal activity with light, see:
To learn how optogenetics can be used to investigate brain circuits, see:

To learn about previous EMBL research into the role of the amygdala in fear, see:

Jose Viosca is a neuroscientist turned science communicator. He investigated the neural circuits of fear as a postdoctoral researcher at the European Molecular Biology Laboratory. Now he is an editorial intern for Science in School.

The brain is a large puzzle. Neuroscientists try to understand each piece to grasp the whole.