

A patient led charity, raising awareness and supporting people with CRPS through experience, understanding and education

Facts about CRPS

Complex Regional Pain Syndrome (CRPS) is a debilitating and disabling neurological disorder causing chronic pain. It most commonly occurs in a limb following a minor injury (such as a sprain or broken bone), surgery, or can appear spontaneously without known cause. CRPS is believed to be due to a problem in the central and peripheral nervous systems, resulting in some or all of the symptoms below:

- Feeling of burning or freezing sensation
- Hypersensitivity of the skin
- Changes in skin colour and temperature
- Changes in hair or nail growth
- Swelling or stiffness in affected limbs
- Difficulty moving

CRPS is poorly understood and people often fail to receive the information and treatment they need. This puts a strain on relationships with family and friends and can affect a person's ability to work. It is therefore, not uncommon for patients to require psychological intervention and support, alongside medical intervention, to deal with the particularly distressing nature of the condition.

CRPS can go into remission and prognosis is good when treatment begins early. However, for a small percentage of people it causes lifelong pain and disability.

CRPS is **NOT** a psychological illness.

CRPS can strike any age and affects men, women and children but statistics show it is more common in women.

There is no single test for CRPS, it is usually diagnosed by ruling out other conditions. Patients must have a particular set of symptoms which doctors and health professionals identify as Complex Regional Pain Syndrome.

CRPS is difficult to treat but physiotherapy, occupational therapy, hydrotherapy, psychological interventions and medications may be used to reduce symptoms.

Very little is known or understood about CRPS. Around the world, research is taking place to improve understanding of the condition in order to discover more effective treatments.

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