

## RARE DISEASE UK: THE CASE FOR A NATIONAL PLAN FOR RARE DISEASE

### What is the issue?

There are over 6000 rare diseases affecting over 3.5 million people (1 in 17) in the UK. Collectively, rare diseases are not rare.

NHS services to support people with rare diseases remain patchy and poorly integrated, meaning that hundreds and thousands of UK families with rare diseases struggle to access the help and support that they need.

### What is Rare Disease UK asking for?

We urge the Government to support the proposed Council Recommendation on a European action in the field of Rare Diseases, which is due to be ratified in June 2009.

### How would this benefit the UK?

A National Plan would aim to create efficiency in expenditure and service allocation. It would:

- help to make the most efficient use of the limited NHS resources available;
- support the Government's health research agenda, including strengthening translational research;
- facilitate early and correct diagnosis and timely treatment, which could offer improved health and quality of life for people with rare diseases;
- ensure the UK keeps pace with other EU countries that have adopted national plans on rare disease.

### What is Rare Disease UK?

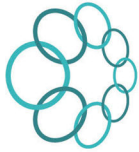
It is the National Alliance of key stakeholders brought together to develop a National Plan for Rare Disease in the UK. It is a joint initiative of the Genetic Interest Group (the UK alliance of patient organisations with a membership of over 130 charities which support children, families and individuals affected by genetic disorders) and others in response to unmet health care needs of hundreds and thousands of families who currently struggle to get access to integrated care and support from the NHS.

### Why is a National Plan for rare disease needed?

Much progress has been made in this field in the past decade, with innovative commissioning strategies and the establishment of centres of excellence delivering multidisciplinary care for some conditions. However, there is currently no coherent strategy for care and support for people affected by rare disease.

Many patients with rare conditions have difficulties accessing high quality care and services, in many cases due to the lack of communication and coordination of services that are sometimes already in existence. Due to the rarity of many conditions, health professionals often have little or no experience in supporting patients in order to find optimum care pathways which can often be difficult or impossible. This leaves patients and families feeling frustrated and 'stuck' in the system.

A National Plan is needed to improve the current fragmentation of services and enable patients and health professionals to provide and use best practice care. This will ensure that all patients with rare disease can not only be diagnosed quickly, but also have timely access to the care and support that they need.



## IMPROVING HEALTHCARE SERVICES

### How could a National Plan improve healthcare services for people with rare disease?

Care and services for families and patients with rare disease are currently patchy and fragmented. A National Plan could ensure that the hundreds and thousands of individuals and families in the UK who are affected by rare genetic disorders have ready access to integrated care and support from the NHS.

### How would this benefit the UK?

A National Plan would help the NHS make the most efficient use of scarce expertise by targeting health care resources to maximise the benefits for patients and families affected by rare disease.

Empowering patients and families by improving information on rare diseases would also contribute to the Government's vision of a patient-led NHS.

### A National Plan should include commitments to:

- identify national or regional centres of expertise, and create them where they do not exist;
- ensure clear healthcare pathways for patients with rare disease through the cooperation between relevant experts;
- provide coordinated care for patients with rare disease within a multidisciplinary team approach model of care, to counter fragmentation across organisational boundaries (this is important because many rare disease affect a number of systems within the body, so patients need to see healthcare professionals from several different specialties during their lifespan);
- improve information on rare diseases for patients and the public.

### How much would this cost?

Recommendations such as those outlined above are largely about reorganisation of resources, rather than an increase in resources. They would enable a more efficient use of resources and maximised benefits at minimum cost to Government.

## PROMOTING RESEARCH INTO RARE DISEASE

### How could a National Plan promote research into rare disease?

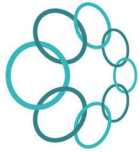
The creation of a National Plan is a timely opportunity to ensure:

- that the UK is an attractive location for rare disease research. That means providing the right framework and environment to support this research – this could include, for example, patient registries to facilitate clinical trials.
- better coordination of the rare disease research that is already taking place in the UK, for example by building research networks.

### How could this benefit the UK?

A clear strategy on rare disease research, embracing both the academic and private sectors, would help to ensure the UK's position as a world leader in biomedical research. Coordinating and fostering rare disease research would:

- mean more possibilities for diagnosis and treatment;
- make the UK a more attractive research location;
- lead to knock-on benefits for medical research in general, particularly for conditions with a genetic component, such as asthma and diabetes.



The National Plan would open up opportunities for greater international research cooperation, which is particularly important for rare disease with very low number of patients. This is an opportunity that the UK needs to grasp in order to keep pace with other countries and keep investment in the UK, and it is closely aligned with the Government's own health research agenda, for example in:

- supporting translational research;
- developing a thriving research culture;
- maintaining and attracting investment by the pharmaceutical and biotechnology industries.

### **A National Plan should include commitments to:**

- identify current research projects and existing research resources;
- identify needs and priorities for basic, clinical and translational research in the field of rare disease;
- foster research in the field of rare disease.

### **How much would this cost?**

The aim of the National Plan would be to create efficiency in current expenditure.

## **ENABLING HEALTH GAIN**

### **How would a National Plan lead to increased health gain?**

By enabling early diagnosis and timely access to appropriate treatment. This is particularly important for the many rare diseases that are progressive in nature.

Alstrom Syndrome is one of the rare conditions for which there is coordinated care, with specialised multi-disciplinary clinics which make a real difference to families' lives. However, many other patients with rare diseases do not have access to coordinated specialist care. This can result in misdiagnosis or non-diagnosis, which are barriers to improving quality of life. For example, it is very common for families with Gorlin Syndrome (of which one of the main characteristics are skin cancers called basal cell carcinomas, or BCCs), to remain undiagnosed for several generations. Some people with Gorlin Syndrome are still misdiagnosed and given radiotherapy, which accelerates the BCCs.

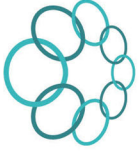
Where there are treatments for rare conditions, not all patients are able to access the right treatment, either at all or at the right time. Medicines for such small numbers of patients are unlikely to meet NICE's cost-effectiveness criteria, but even if they do, decisions to fund treatments are then often made on a case by case basis. As well as causing undue worry for patients, wasting NHS resources and leading to geographical variations, the subsequent delays in treating patients can reduce the health benefits of the treatment.

### **How would this benefit the UK?**

The more effective and efficient the diagnosis and treatment of rare disease, the more people with rare conditions can live healthier, longer lives and make a contribution to society and the economy.

### **A National Plan should include commitments to:**

- improve diagnostic facilities and genetic testing services;
- foster greater awareness of rare disease among the medical profession, to ensure early diagnosis;
- ensure that there is a systematic way of introducing new interventions for health gain;
- establish a robust, transparent, evidence-based system to assess new medicines for rare diseases, involving all stakeholders;
- commission treatment of rare diseases at regional or national level;
- ensure adequate access to cost-effective treatments.



## How much would this cost?

There is a misconception that the healthcare budget would be overwhelmed with a 'tidal wave' of new medicines for rare diseases. There are several reasons why that would not be the case:

- Each year around 100 medicines in development receive orphan designation (official recognition that they would be used to treat rare disease). However, only 10 per cent of these eventually become licensed medicines.
- At any one time it is unlikely that there would be more than 150 orphan medicinal products on the market in Europe under exclusivity arrangements (meaning that directly competitive similar products cannot normally be placed on the market at the same time).

## A EUROPEAN COMPARISON

The UK is in danger of slipping behind other countries who are taking action. France has implemented a comprehensive plan, the Netherlands has implemented a national strategy and Italy has adopted regional strategies. Bulgaria and Portugal have also adopted national plans. The UK now needs to consider this as a positive opportunity for patients in this country.

## NEXT STEPS

Rare Disease UK calls on the Government to support the proposed Council Recommendation on a European action in the field of Rare Diseases.

Rare Disease UK aims to create a draft National Plan for Rare Disease in the UK with input from expert stakeholders from research, patient organisations, clinicians, the pharmaceutical industry, commissioners and government. The plan will bring together research into all aspects of rare disease, prevention and diagnosis of rare disease, best practice in the treatment of rare disease, and the dissemination of information on rare disease to the public, patients and professionals.

## PATIENTS' STORIES

"Sixteen years on from the formation of the Gorlin Syndrome Group, we still have families contacting us with concerns about the lengthy delays in diagnosis despite many years of treatment. Some people have diagnosed themselves or their children by searching the internet and arriving at our website. Other patients tell us about the reluctance of GPs to refer to genetic services and delays in accessing timely treatment."

"When our baby daughter was diagnosed with a rare chromosome disorder, a doctor sat down with us and showed us the results of her genetics test which had the words 'Abnormal Result' stamped on them. The doctor said he had never seen another child with the condition and therefore was unable to answer any of the questions which were swimming around in our heads."

"Each time we see a medical professional we've never seen before (frequently!), we have to begin right at the beginning and go through our daughter's entire medical history, explaining everything we know about her condition. We have to explain what our daughter's condition is, how it affects her, the difficult delivery and post-natal experiences and the potential effects of her condition that she has not yet experienced. "

"We recently saw a consultant who asked 'why are you here?'"

"I took my son to an optician when he was six as he was experiencing difficulties with reading and I thought he might be dyslexic. The optician told my son he wasn't trying hard enough when he couldn't do the tests ... My son was eventually diagnosed with Batten disease five years after a previous diagnosis of cone and rod dystrophy. ... receiving the diagnosis earlier would have given us more time to come to terms with the situation, even though the prognosis was, and still is, bleak."