# **ACCESS TO CARE EVENTS**

During 2021 the EMHA conducted a groundbreaking survey which allowed to learn about the access to care barriers that different European countries have when dealing with migraine. More than 11 countries participated, extracting then powerful insights that helped to address their national policy makers and other stakeholders during 2022 through the Access to Care events. These events helped the countries hosting them to reach politicians and

even achieve significant changes in their local policies addressing migraine treatments and patient's rights.

### In Italy, significant progress has been made in the recognition and management of chronic headaches and migraines, thanks to collaboration between various institutions and key stakeholders. A pivotal

Italy

event was organized by the EMHA in collaboration with the CIRNA Group Foundation, hosted by Member of European Parliament (MEP) Aldo Patricciello. This event brought together policymakers, including Senator Paola Boldrini, Honorable Celeste D'Arrando, and Honorable Nicola Provenza, who quickly mobilized to provide concrete solutions for the needs of chronic patients. During the event, Lara Merighi, a migraine patient and representative of the Alleanza Cefalalgici-CIRNA Foundation, delivered a compelling speech that led to two major outcomes. First, the Italian Congress approved the

creation of a Scientific Advisory Board within the Italian Health Cluster. Second, migraines were officially recognized as a social disease under Italian law. This recognition was supported by the publication of an expert consensus document on criteria for assessing disability and invalidity in

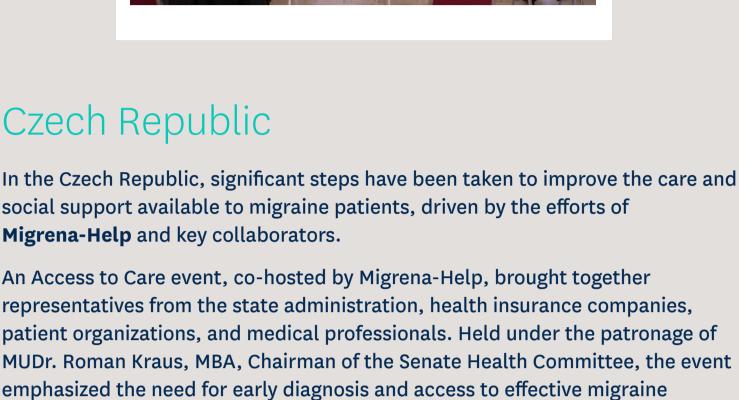
chronic primary headaches, developed by the Alleanza Cefalalgici-CIRNA Foundation in collaboration with EMHA. In March 2023, the implementing decree of Law 81/2020 was issued, officially recognizing chronic headaches and migraines as social diseases. This milestone encouraged Italian regions to develop innovative projects tailored to patients' specific needs, addressing shared priorities such as establishing diagnostic and treatment networks, improving access to innovative therapies, training general practitioners and pharmacists, and promoting public awareness campaigns. Efforts are also underway to

include chronic migraine in the list of officially recognized disabling

disability and providing benefits.

diseases, which would establish uniform national criteria for assessing

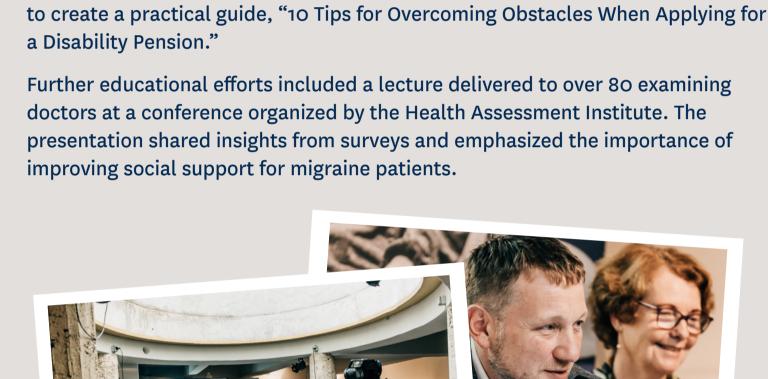
The Alleanza Cefalalgici Foundation has played a crucial role in advising regions on their project development and participating in the review of initiatives. At the national level, a technical working group was established, producing a policy brief outlining common tools and strategies to guide the implementation of innovative care pathways for chronic primary headache patients. However, the need for sustainable funding beyond the 2023-2024 biennium has been highlighted to ensure the long-term success of these initiatives. These advancements mark a significant shift in the care landscape for migraines and chronic headaches in Italy, improving access to comprehensive care and enhancing the quality of life for patients.



treatments while fostering ongoing cooperation with policymakers.

Building on this momentum, Migrena-Help engaged with the Ministry of Labor to advocate for increased access to disability pensions for individuals with chronic migraines. This advocacy led to an educational seminar for medical examiners, during which an expert delivered a lecture on the severity of migraines and the challenges faced by patients. Additionally, Migrena-Help

collaborated with Medical Tribune and the Czech Social Security Administration



On June 11, Migrena-Help and Medical Tribune will host a roundtable discussion in the Chamber of Deputies of the Czech Republic. This event will showcase the journey of a migraine patient through the Czech healthcare system, highlighting

obstacles encountered from general practitioners to specialized Headache

to participants, further enriching the discussion.

Centers. A comprehensive document summarizing this journey will be provided

These initiatives underline Migrena-Help's commitment to raising awareness,



In Spain, the treatment of migraine is typically handled by primary care

and not all of these cases are directed to headache units. This situation

Experts have highlighted the inadequate use of triptans and the limited

availability of preventive treatments, which are currently only provided

to less than 15% of eligible patients.

professionals who may not be knowledgeable about the correct treatment

for the condition. Only the most severe cases are referred to neurologists,

suggests that there is room for improvement in the treatment of migraine.



## event celebrated in the Spanish Parliament in 2022 government ministers and political party delegates agreed to meet with AEMICE to promote the development of a National Migraine Plan. These political party delegates

Spain

impulsed parliamentary questions to the rest but with little success.

As a novelty, during the 12th September and following AEMICE's strategy,

ministry a National Plan for migraine recognizing publicly the need of work

involved Health Minister Mr. Jose Manuel Miñones in boosting from his

in a new strategy involving AEMICE and Spanish Scientific Societies.

To address these issues, the EMHA organized together with the Spanish

in October 2022. The goal of this strategy would be to address all relevant

healthcare and social needs, minimize regional disparities, and ensure the

highest possible quality of care for patients. Following the Access to Care

patient organization, AEMICE, an event in the Spanish Parliament



## During the conference, the results of a study conducted in 2021, aimed at identifying challenges in healthcare access for people with migraine and

discuss this essential issue in Portugal.

other headaches, were presented. Following the event, a significant step was taken in advocating for change. On March 15, 2023, a meeting with a group of MPs from the Health Commission of the Portuguese Parliament took place. The purpose was to share the study findings and raise awareness among political decision-makers about the social, economic, and personal impact of migraine and headaches.

The campaign culminated in the Access to Care in Migraine and Headaches

Conference, which brought together diverse stakeholders from society to

Currently, Migra Portugal is in the process of writing a scientific article

establish meetings with the National Health Executive and the Government

to publish the study results. Additionally, efforts are being made to

Secretary responsible for Health Promotion, in the pursuit of driving

positive change in migraine and headache care.

ACHIEVING CHANGE IS POSSIBLE WHEN WE ALL WORK TOGETHER

TO IMPROVE THE LIVES OF ALL PATIENTS WHO SUFFER FROM MIGRAINE.

The Migraine Movement