

ABOUT US

Written by Administrator

Monday, 16 April 2012 00:00 - Last Updated Sunday, 02 March 2014 17:35

The NATIONAL ME/FM ACTION NETWORK became a Canadian charitable organization on June 18, 1993 dedicated to Myalgic Encephalomyelitis, also known as Chronic Fatigue Syndrome (ME/CFS) and Fibromyalgia (FM) through support, advocacy, education and research. Since its founding, the organization has been an ACTION Network. Some of our accomplishments are:



- Providing knowledge and empowerment to the people who need information and help
- Establishing relationships with both the provincial and Canadian governments in Canada to better the lives of people ill with ME/CFS and FM
- Developing contacts with medical and legal professionals
- Working with National and International Support Groups
- Publishing a quarterly newsletter keeping people informed about the progress in research and related matters

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- Spearheading the development of the Consensus Documents, known as the Canadian Definitions
- Resource guides for disability matters
- TEACH-ME guide for Teachers and Parents of children and youth with ME/CFS and FM

The NATIONAL ME/FM ACTION NETWORK hosted the 10th International IACFS/ME research and clinical conference for ME/CFS and FM and related illnesses in Ottawa from September 22nd to 25th, 2011 consisting of professional workshops and meetings as well as a one-day meeting for the general public.

See the History and Achievements of the organization for more details. [Click here](#) .

Our Motto: People Helping People Helping Themselves

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