

WHAT WE DO

Written by Administrator

Saturday, 10 November 2012 00:00 - Last Updated Sunday, 02 March 2014 17:37

- DISSEMINATE ME/CFS and FM literature to medical and legal professionals and the general public;
- INTERVENE in court cases of benefit to the ME/CFS and FM population;
- COLLABORATE with support Networks in both Canada, U.S. and elsewhere;
- REPRESENT the ME/CFS and FM population at meetings with government officials and politicians;
- PUBLISH a quarterly newsletter QUEST to keep the ME/CFS and FM population informed;
- PROVIDE resource materials for disability issues;
- KEEP ABREAST of research on ME/CFS and FM around the world;
- SUPPORT scientific endeavors;
- EDUCATE the school systems on youth with ME/CFS and FM;
- KEEP a roster of Lawyers who are experienced in ME/CFS and FM issues;
- LIAISE with all forms of media on ME/CFS and FM issues;

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- WORK with Networks of related diseases;
- DO Special Projects that develop;
- HOLD conferences on ME/CFS and FM;
- ACCEPT opportunities to work with doctors, researchers and legal professionals;
- INFORM people who contact us on ME/CFS and FM;
- REFER people to the best help available;
- FOLLOW-UP leads on ME/CFS and FM to its source;
- USE website, Facebook and Twitter;
- SEND announcements to those on our email roster;
- [ADMINISTER the Dr. Alison Bested Scholarship Fund](#) ; and
- DEAL with issues as they arise.

FOR DETAILS ON THE ABOVE, PLEASE VIEW THE HISTORY &

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ACHIEVEMENTS OF THE NATIONAL ME/FM ACTION NETWORK.