



FRAGILE X INTERNATIONAL

An Introduction to FraXI

A timeline of FraXI



**2012
INTERNATIONAL
FRAGILE X DAY**

The European Fragile X Network initiated International Fragile X Awareness Day on the 10th day of the 10th month: X.X.



**2018
NON-MEDICAL
INTERVENTIONS**

The European Fragile X Network helped modify how the treatment of Fragile X Syndrome is described in Orphanet to emphasise non-medical interventions "Speech, physical and sensory integration therapy as well as individualized educational plans and behavioral interventions".
(https://orpha.net/consor/cgi-bin/OC_Exp.php?Ing=EN&Expert=908)



**2020
FXPAC**

The country associations worked together to introduce the term Fragile X Premutation Associated Conditions (FXPAC).
<https://www.frontiersin.org/articles/10.3389/fped.2020.00266/full>



**2022
GENE NAME
CHANGE**

FraXI representatives led in changing the name of the FMR1 gene and of the FMRP protein.



**2022
FRAXI IS LAUNCHED**

FraXI formed the first ever internationally-registered Fragile X charity open to all FX organizations (either as full or associate members). We have a Vision! We are sharing our Vision, Mission and Values on our Website: www.fraxi.org. FraXI is working with partners across Europe to develop European guidelines for Fragile X Syndrome.



**2023
NEW FRAXI MEMBERS**

Fragile X Association Australia and Fragile X New Zealand are new FraXI Full Members. The National Fragile X Foundation (U.S.) is a new FraXI Associate Member.



**2024
ADVOCACY AND
AWARENESS**

- FraXI Congress 2024 in Barcelona, Spain, 7-10 November
- Growing our membership
- Raising awareness on International bodies and leading up to X.X. 2024
- Providing resources and webinars on best practices for our members