Rett Syndrome Europe is a non-profit organisation supporting a network of national associations who are the voice for people living with Rett Syndrome.

The aims of RSE are:

- To raise awareness and increase the understanding of this devastating disorder in every European country
- To improve communication within the European Rett Community
- To promote the interests of people with Rett syndrome and their families
- To extend the work of RSE to all European countries, assisting in the creation of national associations where none exist
- To promote research into Rett syndrome
- To advise and influence policy and legislation on all matters that affect people living with Rett syndrome and their carers

CAN YOU HELP?

www.rettsyndrome.eu
What is Rett Syndrome?

Rett syndrome is a complex neurological disorder that affects mainly girls. Although signs of Rett syndrome may not be initially obvious, it is present at birth and becomes more evident during the second year. People with Rett Syndrome are profoundly and multiply disabled and totally dependent on others for their needs throughout their lives. Approximately 1 in 10,000 girls are diagnosed with Rett syndrome in Europe.

Support us so that we can:
- Attend international conferences
- Give each child the same chances of support
- Improve international scientific collaboration for Rett syndrome

Your donation can make a difference...

To make a one off donation or set up a standing order:
Rett Syndrome Europe
Bank Account details: ING Bank
IBAN/ LU710141745029900000
BIC CODE CELLLULL

Thank you