Our main Objectives

1. to collect and disseminate information on updated epidemiological indicators on the basis of 83 population-based Cancer Registries from 24 countries in Europe.
2. to describe the healthcare pathways for rare cancers.
3. to develop a clinical database on a sub-group of rare cancers.
4. to propose quality criteria for centres of expertise for rare cancers.
5. to list centres of treatment for rare cancers in Europe.
6. to support the identification of European Reference Networks.
7. to develop and disseminate current clinical approaches to rare cancers.
8. to increase awareness among general practitioners and pathologists about rare cancers.
9. to disseminate information tailored to the needs of patients and all concerned stakeholders.
10. to support patients’ associations empowerment.

In this general framework full of difficulties, there is always something useful that can be done

Visit us at www.rarecarenet.eu
RARECAREnet is working to put rare cancers on the map.