

EURACAN EUROPEAN REFERENCE NETWORK ON RARE SOLID ADULT CANCERS



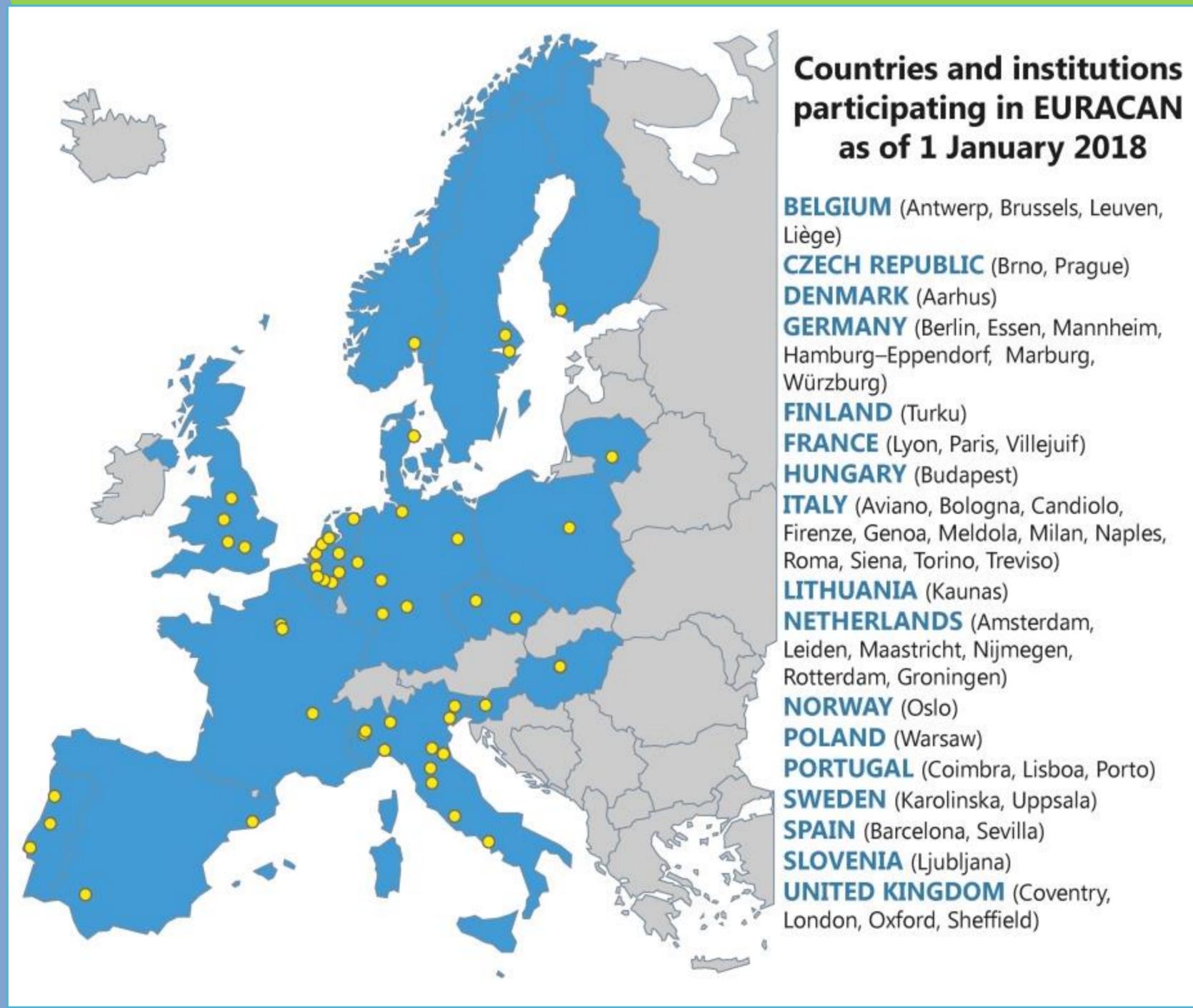
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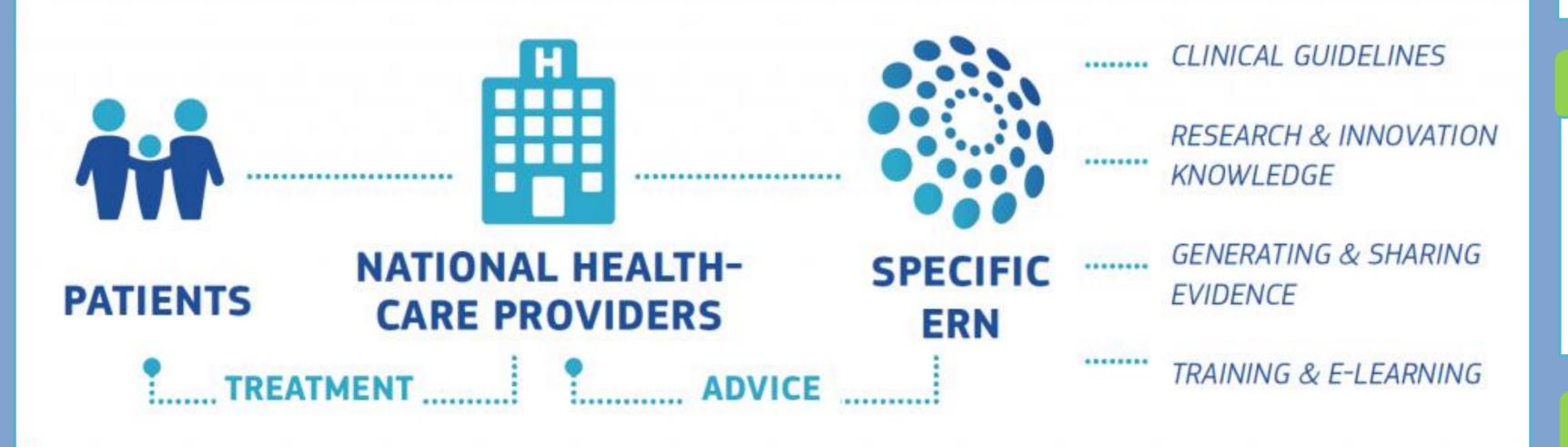
INTRODUCTION

- ERNs are virtual networks involving healthcare providers across Europe.
- EURACAN is the ERN on rare adult solid cancers and it gathers together 66 centres from 17 European countries. It aims to improve the quality of care and outcome of patients with rare adult solid cancers in the EU: diagnosis, treatment management, knowledge, research and communication on all adult solid rare cancers for patients, families, physicians and all stakeholders;

EURACAN EUROPEAN COVERAGE: Healthcare providers were identified on the basis of documented expertise, accrual in rare cancers, and endorsement by their Member State



METHODOLOGY: Patients across the EU will have access to the best expertise available on diagnosis and treatments through the Clinical Patient Management System (CPMS). The CPMS allows healthcare providers from all over the EU to organise virtual consultations and share experience together across national borders



EURACAN TARGETED RAC: EURACAN groups all rare cancers solid adult **into ten "domains"** corresponding to the RARECARE list of rare cancers based on the ICD-O



EURACAN invited Associated Partners from major scientific societies and leading patient advocacy organisations to participate in the different network bodies.



OUTCOMES

- ** Homogenisation of Clinical Practice Guidelines across all domains
- Development of a public **website** translated in ten languages and a Facebook page and a Twitter account.
- First **enrolment of patients** in the CPMS started in December 2017

CONCLUSION

EURACAN provides a unique network to European patients affected with rare cancers, from all member states, enabling access to optimal care, rare resources, education, innovation and research. Additional affiliated centres will join EURACAN in due course to further extend the network's outreach and increase coverage.