

VISIBILITY THAT INSPIRES HOPE



TOGETHER AGAINST SARCOMA.

The ASARGA Association is a non-profit organization dedicated to providing comprehensive support to patients diagnosed with sarcoma and their families in Spain, promoting research, education, and the defense of rights related to this rare disease.



Our mission

To improve the quality of life of people with sarcoma through:

- Information and support.
- Promotion of research.
- Health and social training.
- Political advocacy and public awareness.



Areas of action

- Support for patients and families.
- Counseling, support groups, emotional guidance.
- Training and awareness.
- Conferences, campaigns, and workshops aimed at professionals and the general public.



Research

- Grants and projects in collaboration with scientific societies and universities.
- Carolina Cerezo Grant – the first Galician research on ultra-rare sarcomas, in collaboration with CINBIO – University of Vigo.
- Project “Pre-surgical differential diagnosis of uterine sarcoma,” endorsed by GEIS and SEEM.
- Buesa Grant – in collaboration with GEIS (Spanish Sarcoma Research Group).



Our community

Awards

SPAGN – *Advocacy in action 2025.*

GEPAC – *Minimizing the Risk of Spreading Hidden Uterine Sarcoma.*

FEDER – *Rare Disease Research Grant Program 2025.*

ADVOCACY AND SOCIAL IMPACT

We strengthen the rights of sarcoma patients through initiatives such as the **Non-Legislative Proposal** approved by the Spanish Congress (2023) and the **Sarcoma Manifesto**. At the same time, we foster a strong support network connecting patients, families, and sarcoma specialists. Our work has made a tangible difference, with **over 200 patients and families supported**, collaborations with **CSUR centers and national reference hospitals**, and a direct contribution to earlier diagnosis and comprehensive care.



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