# MONOCLONAL **ANTIBODIES CORE UNIT**

#### Giovanna Roncador Core Unit Head

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## **OVERVIEW**

The Monoclonal Antibodies Unit provides CNIO and other national and international research groups with the capability to generate "à la carte" monoclonal antibodies (mAbs) that are used as research tools to isolate, identify, and characterise new pathways relevant to cancer diagnosis, prevention, and treatment.

Our mAbs are useful tools to understand cancer biology and to diagnose neoplastic diseases, since they allow the identification of molecular markers that are selectively expressed by specific tumour subtypes.

We are particularly specialised in the production and validation of mAbs for immunohistochemistry (IHC), a technique that allows the localisation and study of proteins in tissue sections. This type of reagent allows for a more accurate diagnosis,

"The Monoclonal Antibodies Unit is highly specialised in mAbs production and characterisation, providing CNIO researchers with reliable and well-validated reagents that give added value to their research projects."

## resulting in a better classification of cancer and the selection of the most adequate cancer treatment.

The Unit also offers mAb characterisation and validation, medium-scale mAb production, as well as a service of Mycoplasma testing for the cell culture facility.

### **RESEARCH HIGHLIGHTS**

During the last 22 years, the Monoclonal Antibodies Unit has receptor superfamily. TACI, also known as CD267, promotes generated a large number of mAbs, directed against more T-independent antibody production, in part by facilitating than 180 different antigens, mostly targeting molecules for plasma cell differentiation. Since the distribution of CD267 which mAbs are not commercially available. Many of those in reactive and neoplastic lymphoid tissues has not been mAbs (63) have been licensed to external companies, investigated, we are currently evaluating its expression using generating royalties that represent an important source of a novel rat monoclonal antibody (CLOE240B) against the revenues for the CNIO. CD267 intracellular domain, which recognises its target in paraffin-embedded tissue sections. Large series of normal Each year we prepare and update a detailed CNIO mAbs tissues and B and T-cell lymphomas are being studied using Catalogue, which contains the datasheets of more than 100 whole sections and tissue microarrays. The aim is to determine thoroughly validated, high-quality mAbs (accessible at http:// the pathological diagnostic roles and clinical significance of www.cnio.es/ing/servicios/anticuerpos/default.aspx). This the CD267 receptor in B-cell neoplasms.

catalogue is offered to specialised companies looking for licensing opportunities.

#### **Research activities:**

National and international collaboration. In addition to our collaboration with the CNIO's Research Groups, during the last 22 years we have also developed many joint projects with groups from other national and international research institutions. In these collaborations, the scientists provide their extensive and profound knowledge of cancer research, generating fresh perspectives, diverse viewpoints, and innovative methodologies, which allow the targeting of proteins that play an important role in tumour transformation. We provide them with access to the generation of reliable tools (mAbs), useful both to confirm the results obtained, as well as to further investigate in their research field. In addition, we can develop and set up novel products that can lead to the generation of diagnostic tools for the prevention and diagnosis of cancer. Some of our most recent (last 2 years) and successful collaborations have been with the Spanish National Centre for Cardiovascular Research, CNIC (anti-ALDHl4 mAb), the Hospital Universitario Fundación Jiménez Diaz (anti-hPIGR mAb), and the Centre for Cooperative Research in Biosciences, CICbioGUNE (anti-IL4l1 mAbs).

TACI (CD267) in lymphomas. In 2022, we produced and characterised a novel mAb against TACI protein (encoded by TNFRSF13B gene) that belongs to the tumour necrosis factor

#### **PUBLICATIONS**

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Piris MÁ, Montes-Moreno S, Rodríguez-Justo M, Mena MP, Fernández de Larrea C, Engel P (2022). CD229 (Ly9) a novel biomarker for B-cell malignancies and multiple myeloma. Cancers (Basel) 14, 2154.

Carreras J, Roncador G, Hamoudi R (2022), Artificial intelligence predicted overall survival and classified mature B-cell neoplasms based on immuno-on-

EuroMAbNet. In 2008, in collaboration with Oxford University, we founded EuroMAbNet (www.euromabnet. com), a non-profit organisation that currently spans 13 European countries. EuroMAbNet's primary goal is to provide an arena for people working in the field of monoclonal antibody production and technology to exchange knowledge and updated methodologies, and to create common strategies to improve and standardise the production of properly validated antibodies.



FIGURE 1 Double immunofluorescence staining of PIGR mAb (red) and cytokeratin (green) in paraffin section of human epithelium

cology and immune checkpoint panels. Cancers (Basel) 14, 5318.

- Peña-Cardelles JF et al. (incl. Roncador G) (2022). Prognosis value of immunoregulatory molecules in oral cancer microen vironment: an immunohistochemical study, Biomedicines 10, 710.
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