

# ANALYSIS OF THE EXPRESSION OF A TUMOR SUPPRESSOR GENE IN TUMORAL AND NON-TUMORAL CELLS

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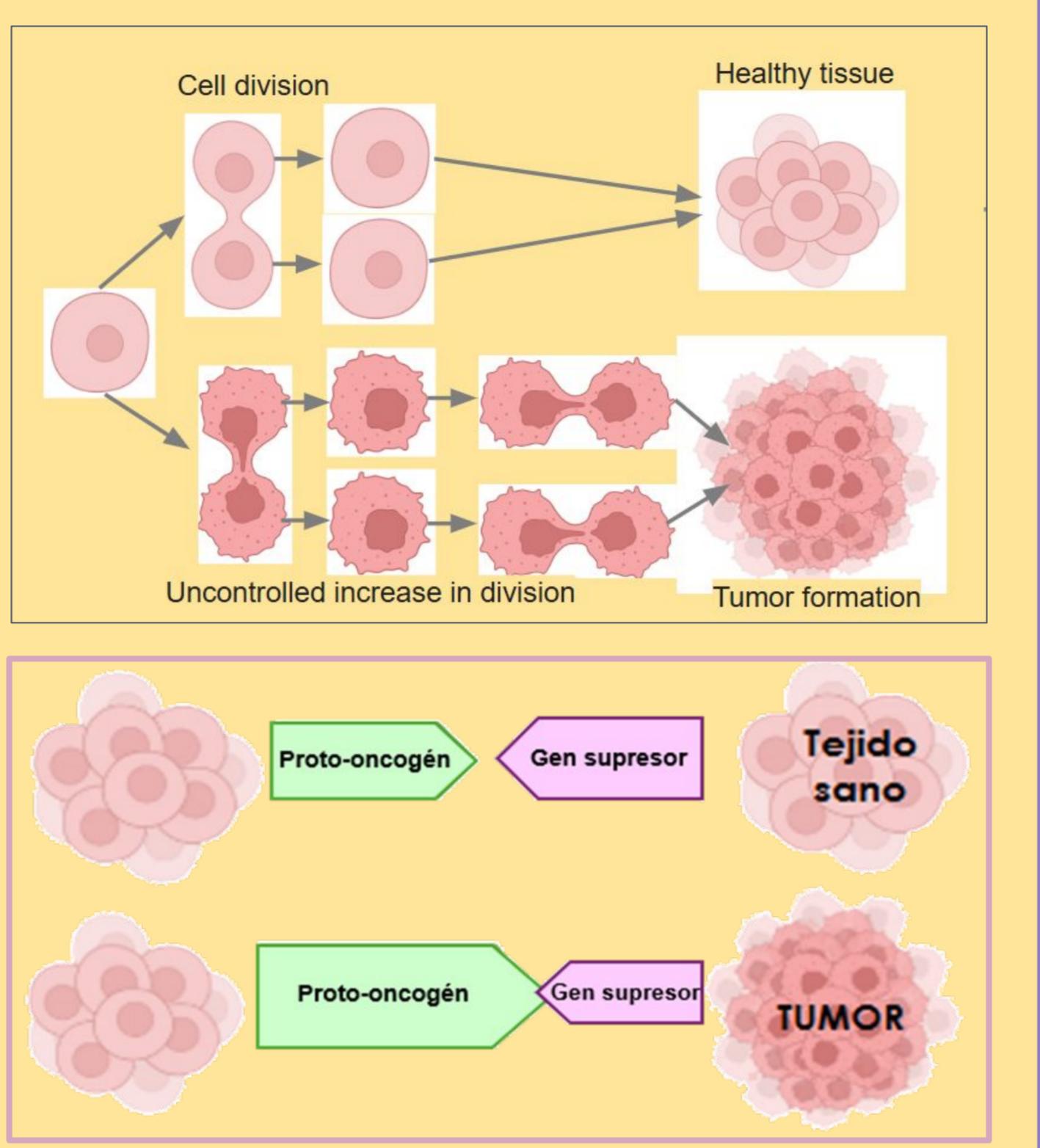
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## INTRODUCTION

- Cancer is a disease caused, among other things, by the **uncontrolled multiplication of cells** in a specific area of the body.
- In healthy cells, **tumor suppressor genes** are important, so its altered expression could result in a functional imbalance. This could lead to the development of tumor cells, characterized by a high division rate and poor regulation of apoptosis.



## OBJECTIVES

- Get RNA from cell pellets.
- Get DNA copy from RNA.
- Amplify by PCR (polymerase chain reaction) a control gene and a tumor suppressor gene.
- Observe the PCR results on an agarose gel.
- We want to study the expression of a tumor suppressor gene in healthy cells and tumour cells.
- To interpret, discuss and represent the most relevant project results and conclusions.

## HYPOTHESIS

The tumor suppressor gene will be expressed in healthy cells, but not in tumor cells.

## MATERIALS AND METHODS

### FIRST SESION



Presentation of the **theoretical background** and facilities by researchers. Familiarization with **laboratory materials and equipment**.

### SECOND SESION

1. Obtaining RNA from **cellular pellets** using a commercial kit.
2. Measurement of RNA concentration in **Nanodrop**.
3. Obtaining **DNA copy** by **retrotranscription**.
4. Tumor suppressor gene and control gene amplification by **PCR**.

### THIRD SESION

1. **Agarose gel** elaboration.
2. **Samples** loading into the **gel wells**.
3. Observation of **results** in a **transilluminator**.

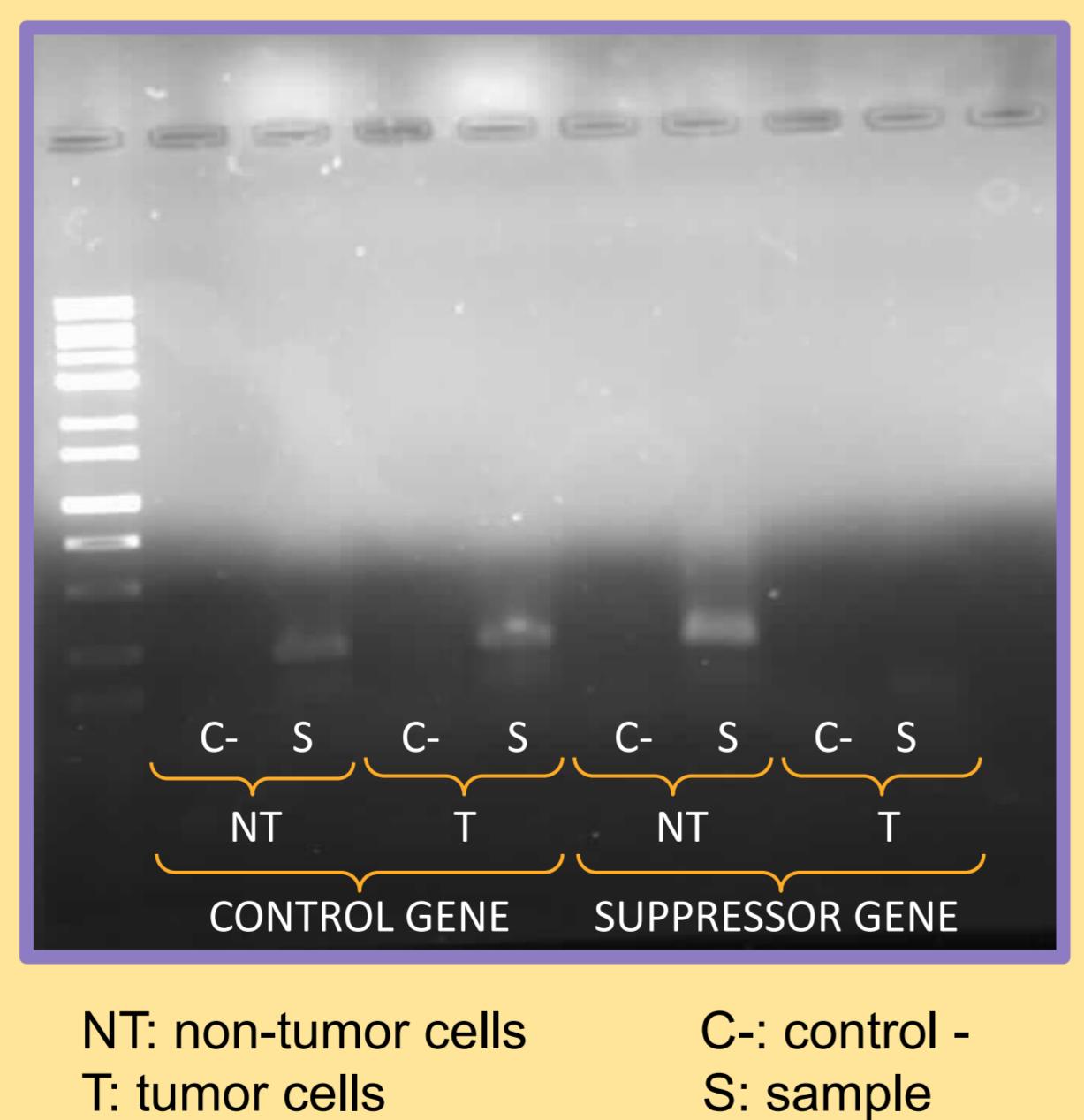
## RESULTS

### ARN QUANTIFICATION

CELL LINE	CONCENTRATION (ng/μl)
Non-tumoral	243,8
Tumoral	162,3

### VISUALIZATION OF THE PCR PRODUCT IN AGAROSE GEL

- In the **negative control** there is no amplification in any of the situations.
- Amplification of the **control gene** is observed in sample of both cell types.
- In the **tumor suppressor gene**, there is amplification in the non-tumor line sample, while there is no amplification in the tumor line sample.



## CONCLUSIONS

- RNA obtained has a **sufficient concentration and quality** to be used as a template for the DNA copy obtaining.
- PCR has been developed **correctly** for the **control gene**, since expression is observed in both healthy and tumour cells.
- Negative controls** for the control gene and for the tumor suppressor gene show **no expression**, so we consider the results to be adequate.
- The tumor suppressor gene studied is **expressed in healthy cells but not expressed in tumoral cells**.

## ACKNOWLEDGEMENTS

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