



# Comparison of positonium lifetime in living tissues for PALS detector and J-PET tomograph.

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Jagiellonian University

“Is Quantum Theory exact? From quantum foundations to quantum applications”

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# Motivation

Positron Annihilation Lifetime Spectroscopy (PALS) is a method based on analysis of the lifetime of positronium, used widely in material studies. This method has a potential in cancer diagnostics.

The main goal of this work was to examine the differences in o-Ps lifetime in tissues (healthy and cancer) measured with two detectors, The first setup, a standard one, consisting of two BaF<sub>2</sub> scintillators, and the second one, the J-PET detector consisting of around 200 plastic scintillators, arranged in three circles.

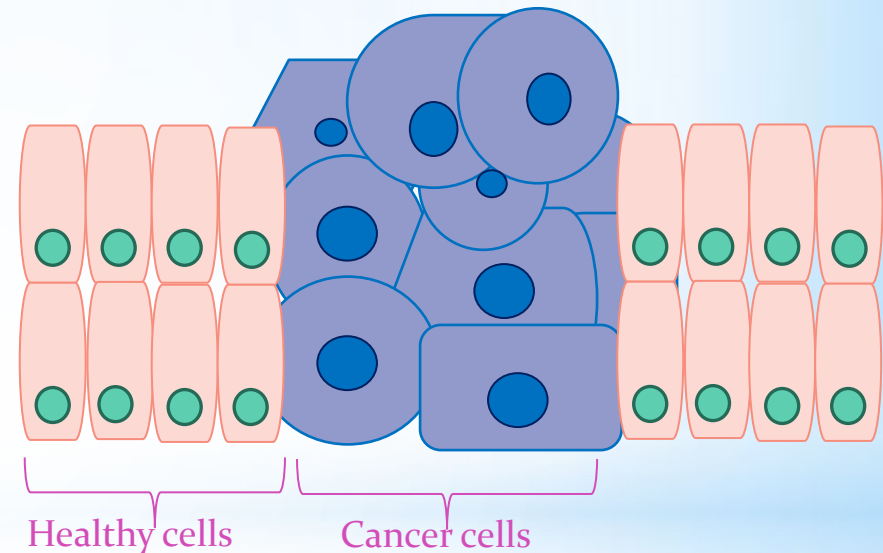
This research may help in establishing the precision of J-PET tomograph for PALS measurements.

# Introduction

Specific changes associated with cancer are connected with enhanced metabolism.

Cancer cells metabolise more glucose than normal cells

Metabolism of a human body can be visualized by the Positron Emission Tomography based on injection of a pharmaceutical labeled with a radioisotope, e.g.  $^{44}\text{Sc}$ .



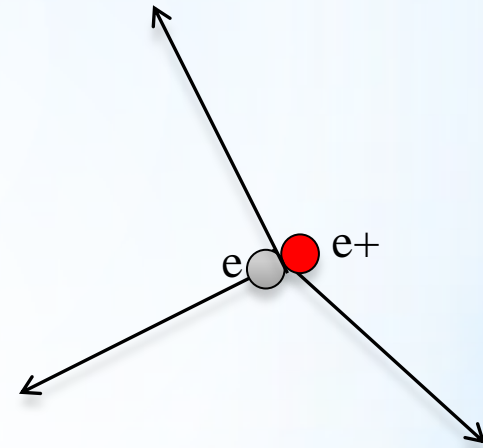
Picture 1. Cancer and Healthy cells

# Introduction

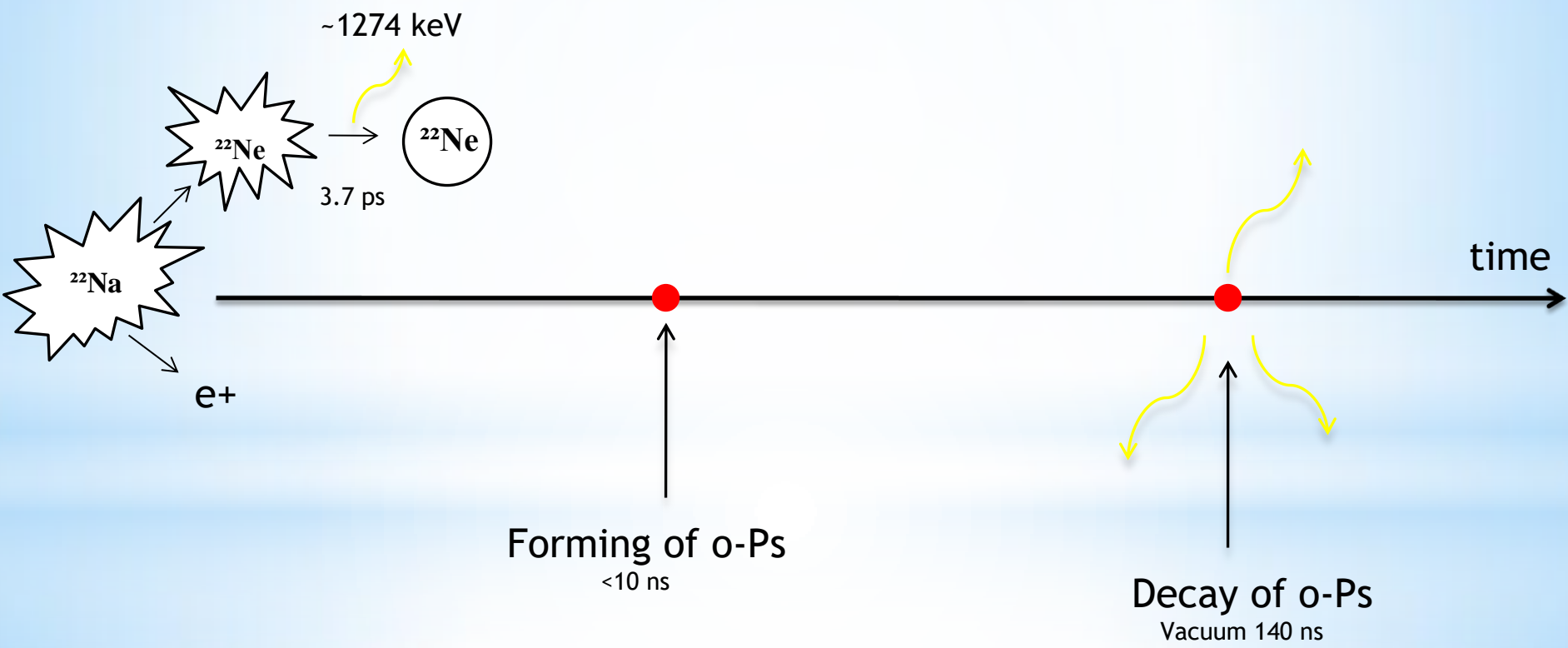
Radioisotopes are attached to biological carrier molecules that target components of cellular metabolism.

These isotopes undergo  $\beta^+$  decay emitting positrons, which may form a bound state with electron in the cell called positronium.

Properties of this atom, for example lifetime, may be different in cancerous and healthy tissues due to their different structure and activity.



Picture 2. o-Ps decay

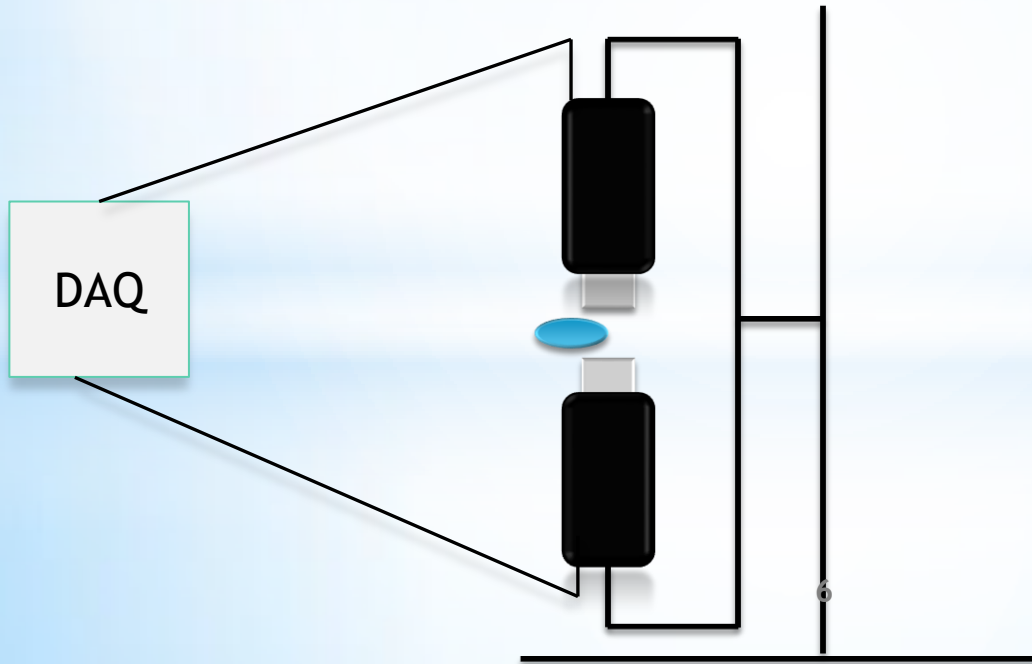


# Measurements with crystal detectors

In measurements we used 22-sodium radioisotope with activity 1 MBq.

Tested samples were delivered from the University Hospital in Krakow. Both normal and cancerous tissues come from large intestine.

The sample (containing two parts of the same tissue fixed in formaline) was inserted between two crystal detectors.



Picture 4. PALS detector

Picture 5. PALS detector

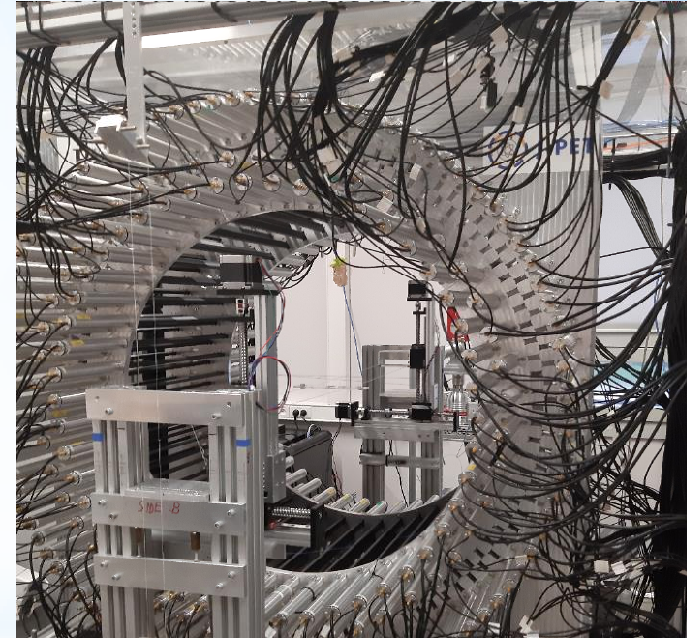


# Measurements with the J-PET detector

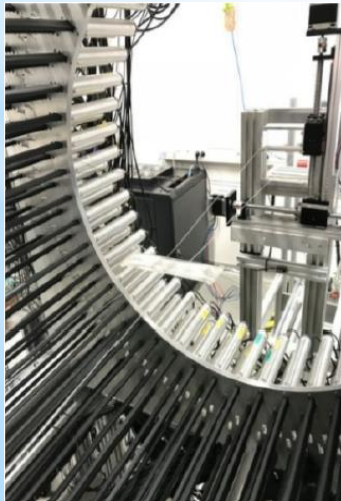
For second part of measurements the same samples were measured with the J-PET detector.

We have used  $^{22}\text{Na}$  source with activity of 0.77 MBq and 0.61 MBq for cancer and normal tissue, respectively.

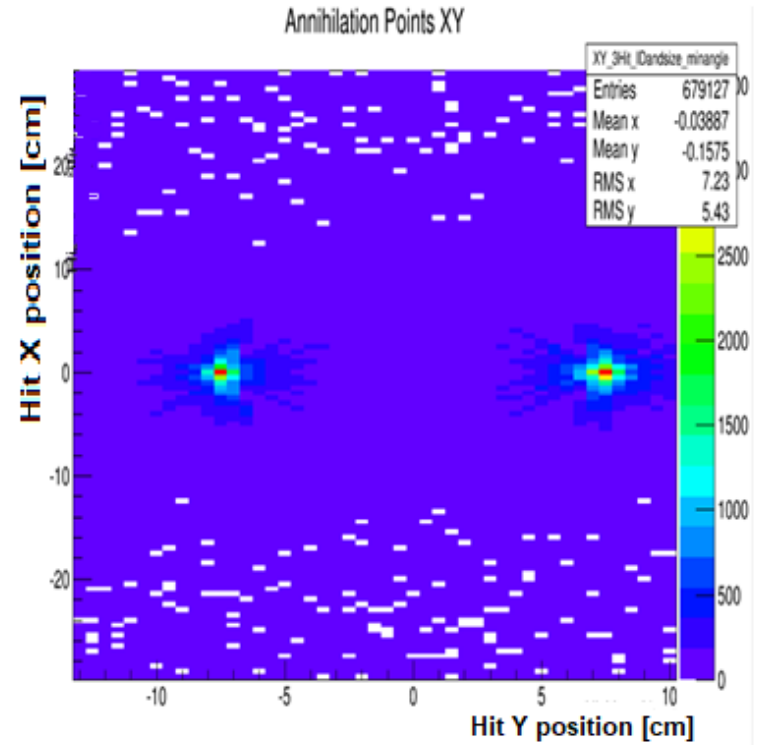
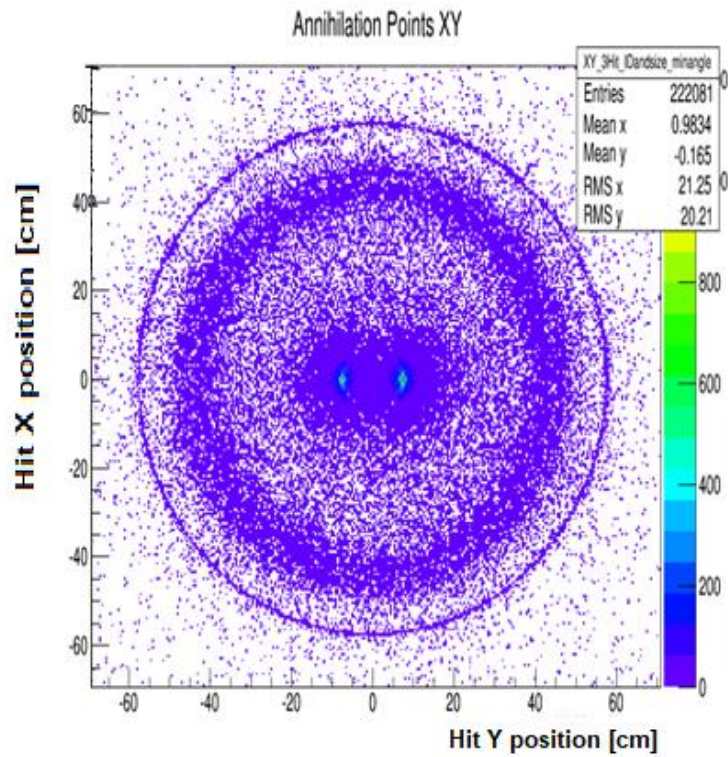
Both cancer and healthy tissues delivered from the same patient were measured at the same time but in different chambers. Getting results for one patient took half of the day.



Picture 6. J-PET

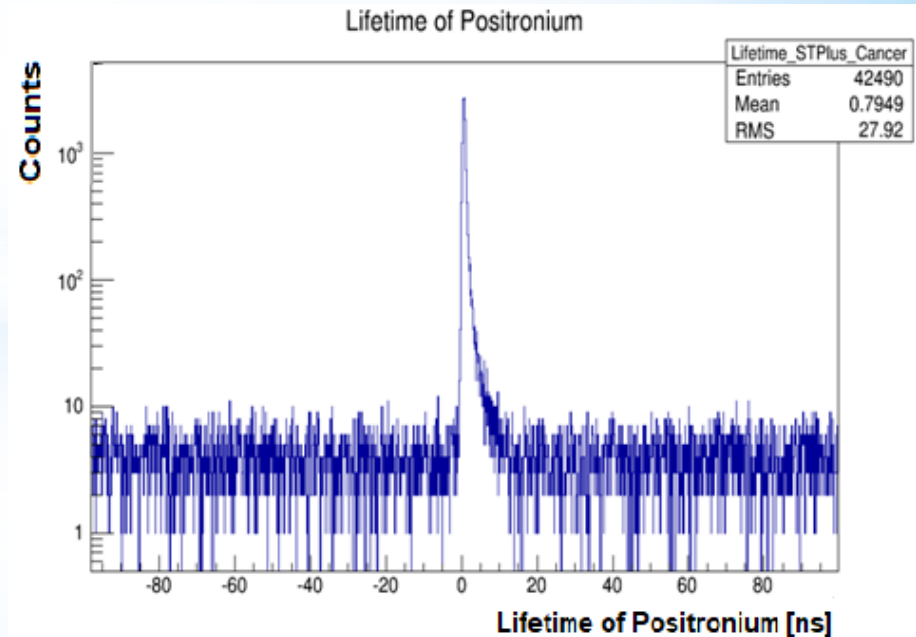
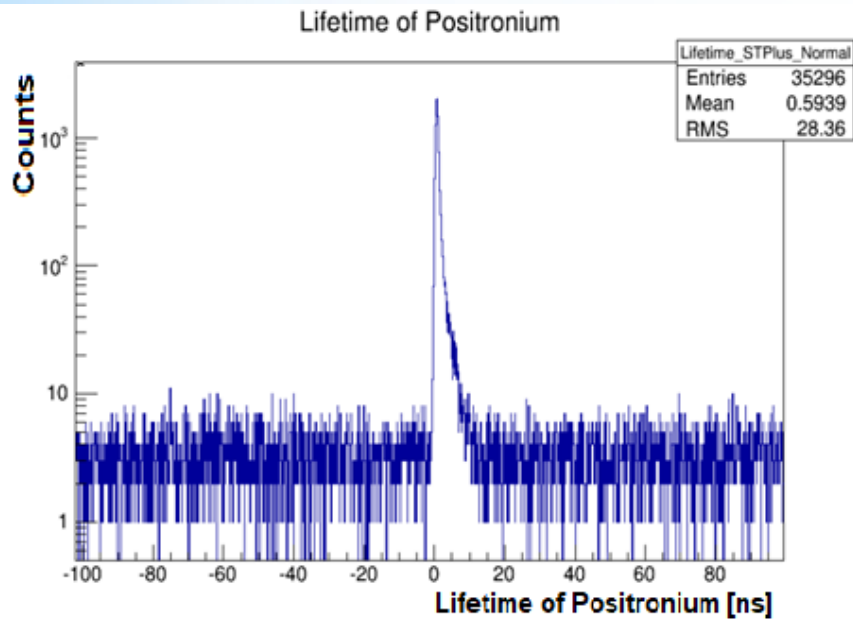
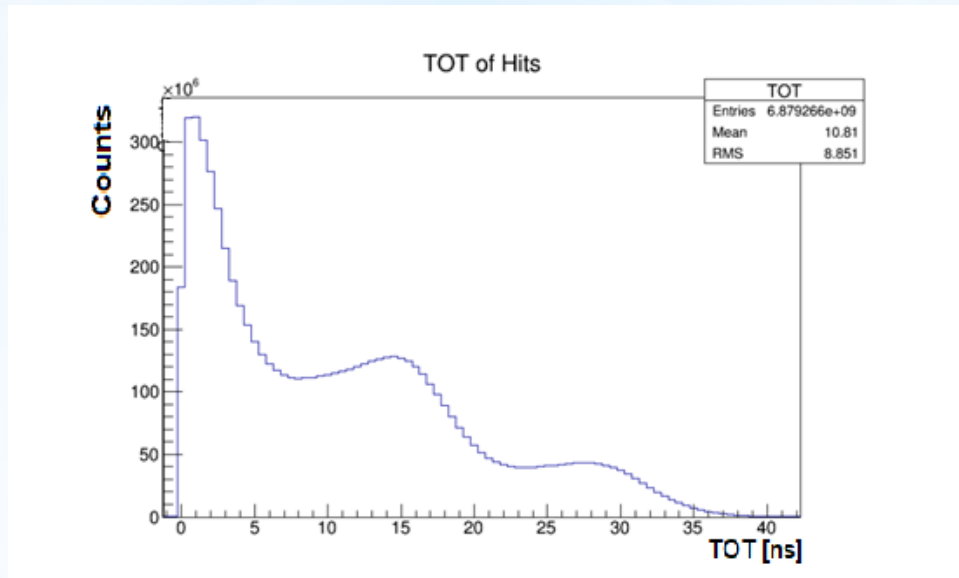


Picture 7 The way of holding samples in the detector

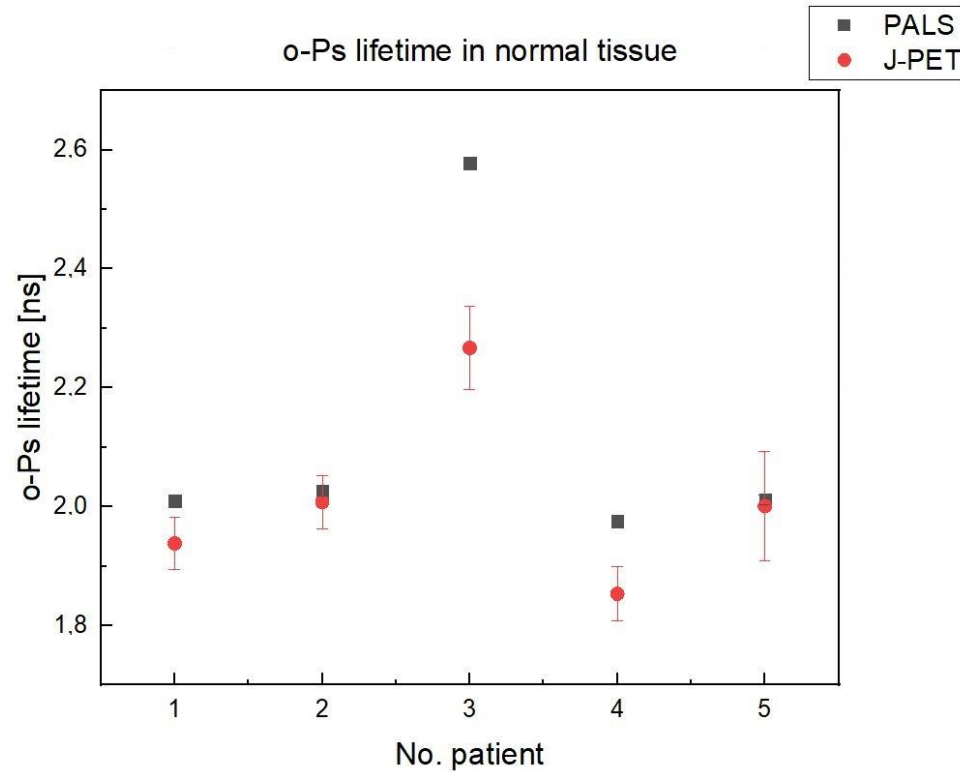
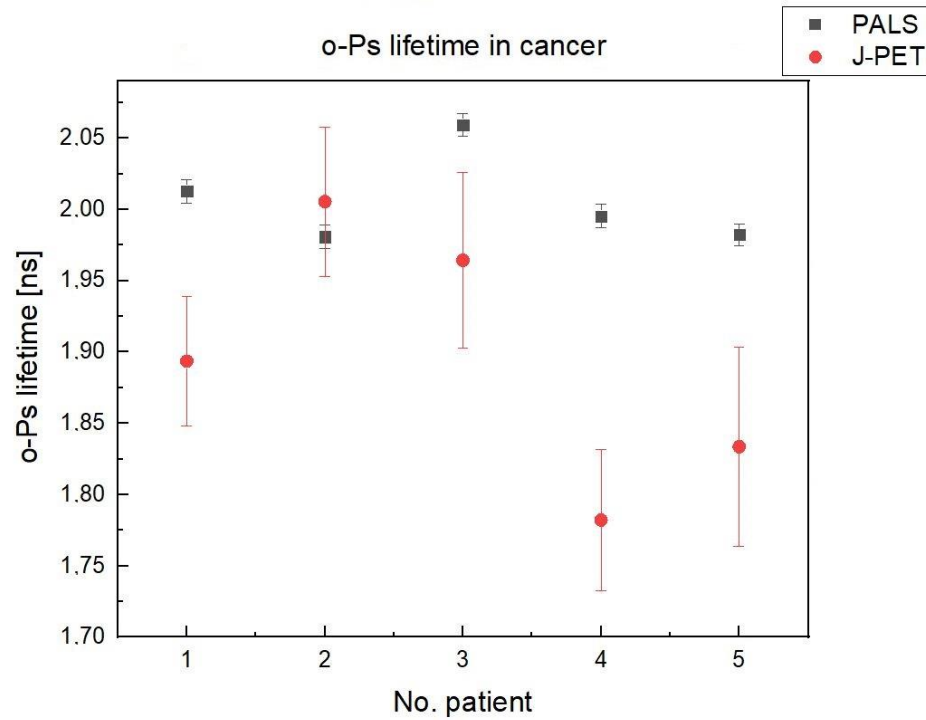




# Results- Control histograms

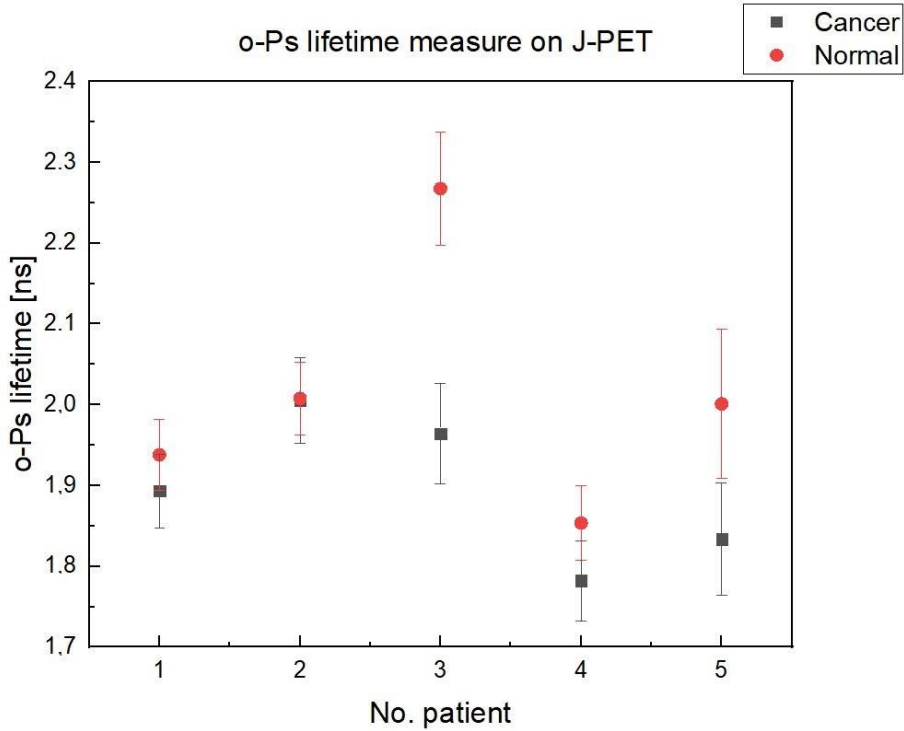


# Results

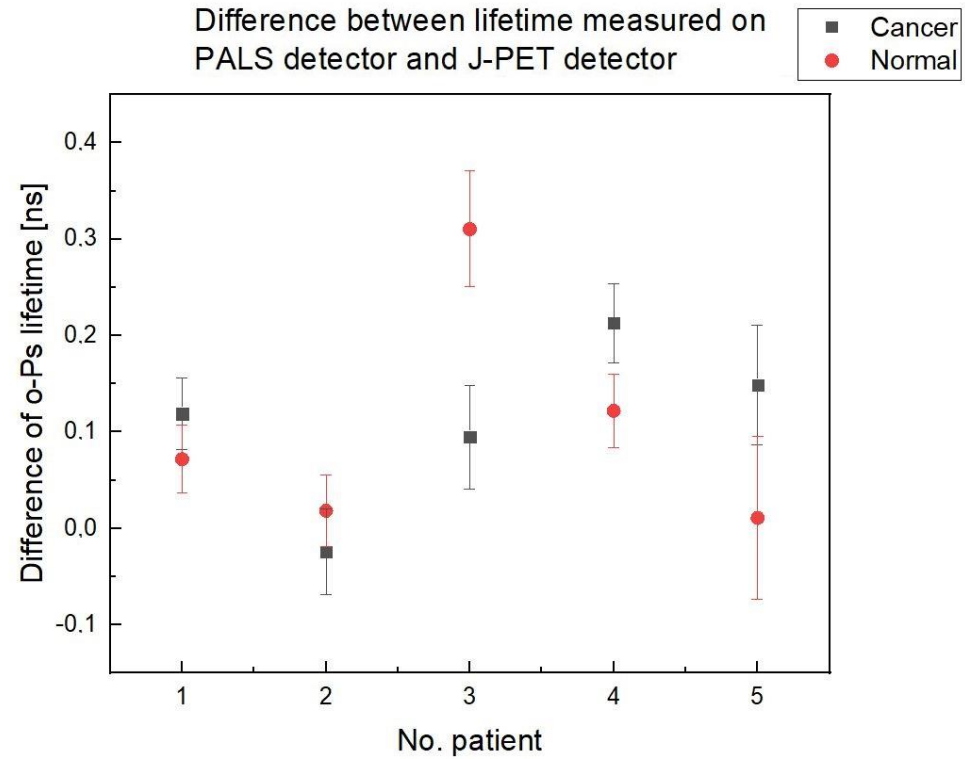


# Results

o-Ps lifetime measure on J-PET



Difference between lifetime measured on PALS detector and J-PET detector



# Summary and outlook

- Patient number 3-> lots of fat in normal tissue
- Time window used in analysis was too wide.
- The resolution of J-PET detector is worst than crystal BaF2 detectors for PALS measurement.

FWHM [ns]	
PALS	J-PET
0.29844(90)	0.638(12)
0.30104(90)	0.616(12)
0.29720(89)	0.675(15)

Amount of patients measured by:		Amount of data file for one person	Summary
day	14	700	9800
night	10	1300	13000
weekend	3	6500	19500
			42300

1 data file= 1 hour of analysis; Max 15 analysis run in 1 hour

42300 data file/15 data file/h=2820h=117.5days=3.9 month

In future we plan to end the analysis for 27 samples and start measuring new samples from Hospital. Also in addition we want to check the impact of the holder for cancer and healthy tissues o-Ps results.

# Bibliography

- [1] P. Moskal et al., Physic in Medicine and Biology 055017, 64 (2019)
- [2] P. Moskal et al., Phys. Med. Biol. 2025, 61 (2016)
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**Thank you**