

May 22, 2024



## Melatonin ELISA

DOI

#### dx.doi.org/10.17504/protocols.io.81wgbx35ylpk/v1

daniel.dautan daniel<sup>1</sup>, Per Svenningsson<sup>1</sup>

<sup>1</sup>Karolinska Institute Stockholm

ASAP Collaborative Rese...

Kaplitt Protocols



### Eileen Ruth Torres

Weill Cornell Medicine





DOI: dx.doi.org/10.17504/protocols.io.81wgbx35ylpk/v1

Protocol Citation: daniel.dautan daniel, Per Svenningsson 2024. Melatonin ELISA. protocols.io

https://dx.doi.org/10.17504/protocols.io.81wgbx35ylpk/v1

License: This is an open access protocol distributed under the terms of the Creative Commons Attribution License, which permits

unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Protocol status: Working We use this protocol and it's

working

Created: February 15, 2024

Last Modified: May 22, 2024

Protocol Integer ID: 95309

Keywords: ASAPCRN, melatonin, sleep

**Funders Acknowledgement: Aligning Science Across** 

Parkinson's Grant ID: 020608



# **Abstract**

Measurement of mouse plasma melatonin using ELISA Kit (Enzo Life Sciences, ENZ-KIT150-0001, NY, US) according to manufacturer instructions.

# Materials

Enzo Life Sciences, ENZ-KIT150-0001



#### Melatonin Extraction

- 1 Mix 4 200 µL of plasma with an equal volume of cold Ethyl Acetate. Vortex gently.
- Allow layers to separate on ice for 00:03:00. Vortex again and incubated on ice for 00:02:00.
- Spin samples at 1000g for 00:10:00 at 4 °C. Transfer the organic layer to a new tube.
- 4 Dry samples and resuspend in Δ 220 μL of 1X stabilizer.

### **ELISA**

2h

1h

5m

10m

- Add  $\perp$  100  $\mu$ L of standards' working solutions and samples to provided 96-well plate in duplicates.
- Immediately after add  $\Delta 50 \, \mu$  of melatonin tracer to each well (except blanks) followed by  $\Delta 50 \, \mu$  of 1X melatonin antibody (except blanks).
- Cover plates with the provided plate sealer. Incubate for 01:00:00 at 37 °C with 500 rpm shaking.
- 8 Decant the solution from each well. Add  $\perp$  400  $\mu$ L of wash solution to each well.
- 9 Decant the solution from each well and pat dry against clean absorbent paper.
- 10 Repeat wash step 3 times.
- 11 Add Δ 200 μL of melatonin conjugate solution to each well (except blanks). Cover plate with the sealer. Incubate for 00:30:00 at 8 Room temperature.



- 12 30m Incubate for 👏 00:30:00 at 🖁 37 °C protected from light.
- 13 Add  $\perp$  50  $\mu$ L of stop solution to each well. Measure the optical density using a micro-plate reader with absorbance set to 450 nm.