Paramylon of Euglena

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<Motive>

- We researched euglena; which have properties similar to plants and animals.
- While we researched euglena, paramylon turned out to be only in euglena.
- Paramylon has many effects, for example, anticancer, detoxification, and protection from infection.
- We think paramylon has antibacterial properties because it has protection from infection.



<Purpose>

- We compare content of paramylon from euglena with that from white euglena to know which euglenas have more paramylon.
- We do experiments to know whether paramylon has antibacterial properties.



We used equipment which can operate brightness and temperature.

	Time	Brightness	Temperature
Daytime	12h	1314Lx	20 degrees
Night	12h	0Lx	20 degrees

<About raising>

- 1. Natural water and Hyponex, Chlorogonium culture fluid \rightarrow We failed because of mold.
- 2. Chlorogonium culture fluid
 - \rightarrow We succeeded and could get a lot of euglena.
- 3. K&H culture fluid
 - \rightarrow We could raise euglena best.

<Extraction of paramylon>

We extracted paramylon with this way.

- I. We separated euglena by using a cyclone separator and got a thick precipitate.
- II. We mashed I by using a homogenizer.
- III. We added II to a 1mass % SDS solution and kept it at $95^\circ\!\mathrm{C}$ for 1 hour.
- IV. We separated ■ by using a cyclone separator for 15 minutes.
- V. We washed precipitate with clean water.
- VI. We skimmed off the precipitate by using acetone.
- VII. We dried it and we got pure paramylon.

<Content of paramylon>

We extracted paramylon and compared the content.

Euglena 0.11 g from 200ml of K&H culture fluid.

White euglena 0.090 g from 200ml of K&H culture fluid.

Euglena has more paramylon than white euglena.

We think this may be because euglena does photosynthesis.

<Observation>

microbes.



We tried to observe euglena by using this property.

However, we couldn't observe euglena well, because as time went by , the water started to evapourate and the concentration of NiCl₂ got too high , so the euglena was squashed by osmotic pressure.

Now we use Vaseline and observe them.

<Precipitation of paramylon by neutralizing>

Paramylon was not aqueous but it dissolves into a basic solution or an organic solvent.

We dissolved paramylon into NaOH aqueous solution (1.0mol/L), and neutralized it with HCl aqueous solution (1.0mol/L), then we checked the precipitation of paramylon.

<Result>

We found something in the bottom of the beaker.

But we couldn't make the decision if it was paramylon or not by a using digital microscope.

We found that there was a little paramylon dissolved in the aqueous solution after neutralizing.

<Antibacterial action of paramylon>

- I. We spread out *Luteus* on the 4 petri dishes.
- II. We put 4 kinds of paperdisk on the petri dishes.
- III. We put petri dishes into 30.5 degrees' incubator.

<Result>



We could not admit inhibition ring in any petri dishes. We checked all, but we couldn't see any antibacterial action from the solutions.

<Consideration>

We thought 4 things from this experiment.

- 1. Paramylon doesn't have antibacterial action.
- 2. We didn't understand this experiment, so we failed.
- 3. Property of paramylon had changed because of the basic aqueous solution it was in.
- *4. Luteus* isn't influenced by antibacterial action of paramylon. So, we need to know about paramylon more and more.

<References>

Function of paramylon from *Euglena gracilis* as filer https://www.jstage.jst.go.jp/article/sptj/50/10/50_728/_pdf

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