Estrous Cycle in Mice with Intraperitonial Injected-CSE
(Mutiati et al.)

Estrous Cycle Response in Mice (Mus musculus) with CSE (Crude Sperm Extract) Injected Intraperitoneally

Enni Mutiati¹, Sri Rahayu², Gatot Ciptadi³, Moch. Nasich³

¹Master Program of Biology, Faculty of Mathematics and Natural Sciences, University of Brawijaya, Malang, Indonesia
²Department of Biology, Faculty of Mathematics and Natural Sciences, University of Brawijaya, Malang, Indonesia
³Department of Animal Production, Faculty of Animal Husbandry, University of Brawijaya, Malang, Indonesia

Abstract
Sperm protein has an important role in fertilization process. It becomes antigenic when it is injected to body and can increase TNF-α secretion. TNF-α in blood vessels can inhibit estradiol synthesis. Estradiol has a significant role in reproduction cycle, especially in estrous cycle. The study aims to understand the influence of Crude Sperm Extract (CSE) to mice (Mus musculus) estrous cycle. 16 fertile mice strain Balb-C aged 3-4 months, weighed 20-30 g was divided into 4 groups. P₀ is a control group injected by PBS, group P₁, P₂, and P₃ were injected by CSE with 1.5 mg.ml⁻¹, 2 mg.ml⁻¹, and 2.5 mg.ml⁻¹. CSE is injected intraperitoneally during mice’s diestrus phase. CSE is injected in day 0, day 12, day 24 and observed daily. The data are descriptively analyzed. The results show that CSE with molecule weight between 26.8-176.8 kDa influences estrous cycle.

Keywords: Crude Sperm Extract, estrous cycle, folliculogenesis, Mus musculus

INTRODUCTION
Sperm protein plays an important role in the fertilization process. It becomes antigenic when injected to body and causes immune responses [1,2,3]. Sperm protein will increase the secretion of TNF-α by macrophages-activated CD4 T cells [4,5].

TNF-α will bind to TNF-α receptor in the ovary [6], thus inhibits the synthesis of estradiol through cAMP (Adenosine-3,5'-Cyclic Monophosphate) and PKA (protein kinase A) pathway. cAMP and PKA are involved in cytochrome P450scx excretion that convert cholesterol to pregnenolone, and eventually estradiol biosynthesis and metabolism [7,8]. Estradiol has a significant role in estrous cycle [9]. Based on the mentioned facts and reasons, the present study aims to understand the influence of Crude Sperm Extract (CSE) in mice estrous cycle.

MATERIALS AND METHODS
Isolation and Charaterization of Bull’s Sperm Protein
We obtained 2 ml (900-2000×10⁶ cells.ml⁻¹) of bull’s sperm from BBIB (Balai Besar Inseminasi Buatan – Center for Artificial Insemination). The sperm was washed with 6 ml PBS (phosphate buffer saline), vortexed and centrifuged (2500rpm, 10 minutes). The pellet was resuspended with 3 ml TCM (Tissue Culture Medium) then vortexed and centrifuged once more (2500rpm, 10 minutes). The pellet was resuspended with 0.5 ml extract buffer and cold-sonicated (50% amplitude, 20 minutes). Total 1 ml suspension was centrifuged (9000rpm, 30 minutes, 4°C) and result’s 0.5 ml supernatant was further centrifuged (13,000 rpm, 45 minutes, 4°C) and then resuspended with KCL-HEPES buffer (1:1) and stored in -80°C [10]. The isolated sperm protein was characterized with 12.5% separating gel SDS-PAGE protocol [11].

Animal Treatment
Total of 16 fertile female mice (Mus musculus) strain Balb-C, aged 3-4 months and weighed 20-30 g were used as animal model in this study. They were divided into 4 groups: control group (P₀) without CSE injection; P₁ which injected with 1.5 mg.ml⁻¹ CSE; P₂ 2 mg.ml⁻¹ CSE injection; and P₃ 2.5 mg.ml⁻¹ CSE injection. All injections were all intraperitoneal and administered at 0th day, 12th day and 24th day of experiment. CSE is injected intraperitoneally during mice’s diestrus phase.

Estrous cycle observation
The observation of estrous cycle was conducted since 0th until 24th days that covers proestrus, estrous, metestrus dan diestrus periods. Vaginal smears were made by using pipette
with NaCl 0.9% solution. The length of estrous cycle was descriptively analyzed.

**RESULT AND DISCUSSION**

SDS-PAGE characterization reveals that Bull’s sperm CSE have 11 protein bands with molecule size 176.8, 63.2, 58.4, 55.3, 52.4, 49.7, 44.6, 38.02, 36.03, 34.1 and 26.8 kDa [12]. These protein bands were all used for treatment in mice.

We identified the phases of mice estrous cycle by its general criteria (Fig. 1). In proestrus phase, vaginal cytology dominated by parabasal cells and estrous phase vaginal cytology dominated by superficial cells. Intermediate and parabasal cells predominate in smears taken during metestrus. The onset of diestrus is marked by a precipitous decline in the number of superficial cells and reappearance of intermediate and parabasal cells [13].

![Figure 1. Vaginal cytology representing each stage of estrous cycle](image)

**Description:**
A. Proestrus, nucleated epithelial cells (e)
B. Estrous, cornification cell (k)
C. Metestrus, cornification cell (k) and leukocyte cell (l)
D. Diestrus, dominated by leukocytes (l).

Estrous cycle signifies sexual activity and organ function, like the function of the ovary, along with follicle development. We found irregularity in CSE injected group. It characterized by elongation of time in one phase of the estrous cycle, which is diestrus (Table 1, Fig. 2).

<table>
<thead>
<tr>
<th>Phase (day)</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Proestrus</td>
<td>P₀ 1.0 ± 0.0 1.0 ± 0.0 1.0 ± 0.0 1.0 ± 0.0</td>
</tr>
<tr>
<td>Estrous</td>
<td>P₀ 1.0 ± 0.0 1.0 ± 0.0 1.0 ± 0.0 1.0 ± 0.0</td>
</tr>
<tr>
<td>Metestrus</td>
<td>P₀ 1.0 ± 0.0 1.0 ± 0.0 1.0 ± 0.0 1.5± 0.58</td>
</tr>
<tr>
<td>Diestrus</td>
<td>P₀ 2.8 ± 0.7 3.1 ± 0.6 4.1 ± 0.8 4.3 ± 1.6</td>
</tr>
</tbody>
</table>

The data signify that CSE could influence the estrous cycle length, which have elongation in diestrus phase compared to control (P₀) (Fig. 2). The elongation tendencies of diestrus phase in each treatment are 3 days longer (P₁), 4 days (P₂), and 4-5 days (P₃) (Fig. 3). The average duration of normal estrous cycle in mice and rats is 4-5 days [14]. CSE injection in the body may cause immune responses and eventually increase TNF-α secretion in blood by CD4 T cells which are activated by macrophages [2-5]. TNF-α through the blood vessels will bind to TNF-α receptor in the ovary [6].

The increase of cytokine TNF-α in the ovarian inhibit synthesis of estradiol through inhibition of cAMP and PKA pathway. cAMP and PKA involved in regulating the expression of the enzyme cytochrome P450scc. P450scc cytochrome enzymes will convert cholesterol to pregnenolone. Pregnenolone be converted to estradiol [7].

Inhibition of cAMP and PKA will inhibit the enzyme P450scc, so it will inhibit the biosynthesis and metabolism of Estradiol [8]. Inhibition of estradiol can delay the replacement of diestrus phase to the proestrus phase, because the proestrus phase has estradiol in highest levels and the diestrus phase has estradiol is low levels [15]. It will affect on hormone regulation.

![Figure 3. Length of Estrous Cycle for Each Stage](image)
CONCLUSION
The protein in the bull’s sperm CSE have molecule size ranging from 26.8 to 176.8 kDa. Our findings suggested that bull sperm CSE influences the mice estrous cycle, especially in diestrus period elongation. This provides vital preliminary information for the usage of CSE as the candidate of immunocontraception for human. We infer that it is necessary to measure the estrogen blood level, the antibody formation in the blood and flowcytometry analysis to elucidate if there a TNF-a present in the blood.

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REFERENCES