A Study on the Ocular Infection with Rabies Virus in Mouse

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INTRODUCTION

Rabies is a zoonotic disease, which causes more than 60,000 human deaths around the world annually [1]. Rabies virus (RABV), belongs to the genus Lyssavirus of the Rhabdoviridae family. The most common mode of rabies virus transmission is through a bite wound or contact of broken skin with saliva of a rabid animal. Various other routes of virus transmission include exposure of mucous membranes (i.e. eyes, nose, and mouth) to infected saliva of a rabid animal, aerosol transmission, and corneal transplantation. Laboratory workers during work with rabies virus and veterinarians during examination and surgery of rabid animals may be at risk for exposure to saliva or other infectious fluids splashing into their eyes. The aim of this study was to investigate the possibility of ocular rabies pathogenesis in mice as an animal model. Our results will determine if rabies virus strains challenge virus standard (CVS) and street rabies virus (SRV) are able to infect the central nervous system (CNS) of mice through the ocular route. Methods: This study was performed in two experiments. In experiment 1, different lethal doses of fixed rabies virus strain CVS were made and instilled into both eyes of test mice. In experiment 2, concentrated rabies virus strains CVS and SRV were instilled into both eyes of the test mice. Main results: In both experiments, we showed that CVS and Street rabies virus (SRV) strains produced infection in mice following ocular instillation of fixed and street rabies viruses developed no clinical symptoms of rabies and all were healthy and alive during the 3 month observation period. The results were negative in both experiments. Conclusion: Our results suggest that CVS and SRV viruses are not able to infect the CNS of mice via intact conjunctiva and cornea. J Med Microbiol Infec Dis, 2014, 2 (2): 61-65.

Keywords: Eye, Infection, Mouse, Rabies Virus.

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MATERIAL AND METHODS

Viruses. In this study, two strains of rabies virus were used: fixed virus (CVS) and SRV. CVS stock (CVS-11) was a 20% viral suspension made by serial passage of mice brain and SRV was isolated from brain sample of a rabid wolf, which was confirmed by fluorescent antibody test (FAT) at WHO collaborating center (WHOCC) for reference and research on Rabies, Pasteur Institute of Iran.

Titration of virus strains. Serial tenfold dilution of CVS stock from $10^{-8}$ to $10^{-7}$, was made in diluents (deionized water with 2% horse serum) and 0.03 mL of each dilution was inoculated intracerebrally (IC) into 5 mice. Mice were euthanized after 5 days or more post-infection at the paralytic stage. Their brain was removed and rabies infection was confirmed by FAT. The LD$_{50}$ was calculated according to the Spearman-Kärber method [8].

A 10% suspension of SRV was prepared by homogenizing the wolf brain sample in an isotonic buffered solution containing 1560 IU/ml penicillin and 500 IU/ml streptomycin antibiotics. A Groups of 10 mice aged 21 days, weighing 12-14 g were inoculated IC with 0.03 ml of each dilution was inoculated intracerebrally (IC) into 5 mice. Mice were euthanized after 5 days or more post-infection at the paralytic stage. Their brain was removed and rabies infection was confirmed by FAT. The LD$_{50}$ was calculated according to the Spearman-Kärber method [8].

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Table 1. Results of experiment 1

<table>
<thead>
<tr>
<th>Group</th>
<th>No. of mice</th>
<th>Virus</th>
<th>LD$_{50}$/0.03 ml</th>
<th>Deaths/Exposed</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>5</td>
<td>CVS</td>
<td>1</td>
<td>0/5 (0%)</td>
</tr>
<tr>
<td>2</td>
<td>5</td>
<td>CVS</td>
<td>25</td>
<td>0/5 (0%)</td>
</tr>
<tr>
<td>3</td>
<td>5</td>
<td>CVS</td>
<td>50</td>
<td>0/5 (0%)</td>
</tr>
<tr>
<td>4</td>
<td>5</td>
<td>Normal Saline</td>
<td>-</td>
<td>0/5 (0%)</td>
</tr>
</tbody>
</table>

Note: Various Lethal Doses of fixed rabies virus CVS were used to infect mice by ocular instillation. In control group, normal saline was used. No deaths were observed either due to ravies or accident during surveillance period.

Table 2. Results of experiment 2

<table>
<thead>
<tr>
<th>Group</th>
<th>No. of mice</th>
<th>Virus</th>
<th>LD$_{50}$/0.03 ml</th>
<th>Deaths/Exposed</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>23</td>
<td>CVS Stock (20% suspension)</td>
<td>$10^6$</td>
<td>0/23 (0%)</td>
</tr>
<tr>
<td>2</td>
<td>23</td>
<td>SRV Stock (10% suspension)</td>
<td>$10^6$</td>
<td>0/23 (0%)</td>
</tr>
<tr>
<td>3</td>
<td>5</td>
<td>Normal Saline</td>
<td>-</td>
<td>0/5 (0%)</td>
</tr>
</tbody>
</table>

Note: In both groups of mice under study, no deaths were observed due to ravies or accidental during three months of observation.
Fluorescent Antibody Test (FAT). After 3 months of observation, 2 mice of each group in experiment 1, and 5 mice of each group in experiment 2 were selected randomly and euthanized. FAT was performed on brain samples of mice based on a technique previously described by DJ Dean et al. [8]. Briefly, smears of mice brain sample were prepared by the impression method on a clean slide and fixed for 30 min in cold acetone. The slides were covered with anti-rabies nucleocapsid rabbit immunoglobulin G conjugated with fluorescein isothiocyanate (BIO-RAD, Marnes-La-Coquette-France) and incubated at 37°C in a humid chamber for 1 h. Then, slides were washed twice with phosphate-buffered saline and observed by fluorescent microscope (Nikon Eclipse TE200; Nikon Corp, Tokyo, Japan).

RESULTS

Virus titration experiments determined the LD50s of CVS and SRV suspensions as 10^6.5 and 10^4.7, respectively. In experiment 1, none of the infected mice developed rhabies following ocular instillation of different LD50s of CVS (Table 1). During 3 months of observation, no signs of rhabies were observed in mice inoculated either with CVS or SRV. The second experiment gave similar results comparable to the first experiment. All mice in both experiments, including control and inoculated groups were alive and healthy. FAT results were negative for mice brains in both experiments. No viral antigens were detected in brain samples of mice.

DISCUSSION

Salivary glands of experimentally infected dogs and cats with detectable virus which contain geometric mean titers ranging from 3,400 to 386,000 mouse LD50/g. Saliva of rhabies animal could be highly infectious if it comes to contact with sensitive area, such as mucosa or broken skin [9]. Therefore, it could be potentially dangerous to cause rhabies infection through these routes as it has been mentioned previously. In this study, we have designed an experiment to test whether contamination of eye with rhabies virus could be a route of infection in mouse model. We hypothesized whether rhabies virus can enter and penetrate into eye neurons through any part of intact eye, including cornea or mucosa. In experiment 2, certain conditions of the experiment 1 were applied to repeat the research with SRV and check the dose dependency of the CVS. The results of this study showed that the suspension containing 10^6.7 LD50/0.03 ml of SRV from a wild type strain originated from a rhabid wolf or 10^6.5 LD50/0.03 ml of the fixed virus strain CVS could not infect mice by absorption via ocular instillation. Previous reports have indicated that high concentrations of virus are necessary to infect mouse through oral or nasal routes [7]. However, the results of the present study showed that ocular instillation, even with high concentrations of CVS, did not cause clinical manifestations of rhabies, even after 3 months. These findings are in agreement with unpublished data from Hanlon experiments, which showed that ocular instillation by drops of a virulent canine isolate of virus did not cause rhabies in Syrian hamsters [5]. Although, there are studies that show the possibility of rhabies infection through oral route. Charlton and Casey indicated that absorption of CVS through the oral mucosa is minimal in mice and skunks [3]. It has been shown that the lyophilized SAG2 oral rhabies vaccine is effective in immunizing captive arctic foxes [10]. There are different studies showing that there is no viral amplification or penetration in animal tissues following oral vaccination, which implies the safety of this live rhabies vaccine. However, SAG2 carries a double mutant attenuated virus strain that could not totally repeal the possibility of street virus penetration through oral rout [11]. In our study, both fixed and street viruses were tested, and the results in mice suggest that contamination of ocular mucosa could not cause rhabies infection. Lafay et al. demonstrated that olfactory neuroreceptors can be directly infected by CVS following its instillation in the nasal cavity, and CVS spreads into brain along the olfactory pathway [4]. So, in our study, probably nerve endings in intact cornea and conjunctiva are not as easily accessible to the virus as intranasal route.
In CVS-infected weanling mice and hamsters that were infected orally, rabies virus antigen was not observed in intestinal mucosal cells, but was found in neurons in Auerbach’s and Meissner’s plexuses of the stomach and intestine [5]. These findings suggest that viral entry through the oral route likely occurs via breaks in the integrity of the gastrointestinal mucosa [5]. Conjunctiva and the cornea epithelium are important barriers against viral entry [12, 13]. In this study, it is thought that the native immunity of auto clearance of eye or intact cornea and conjunctiva could result in the removal of viruses before entrance into the inner layers and neurons. Kucera et al. indicated that inoculation of CVS strain into the anterior chamber of the eye could result in rabies; however, the instillation needle needs to pass through the layers of cornea [14]. The cornea has a rich nerve-supply and corneal epithelial cells are in close contact with the nervous system and the nerve connection between the cornea and its corresponding CNS segments are short. Although, infection of the cornea with the rabies virus has been well indicated in the centrifugal spread of the virus [15], the results of this study revealed that the intact cornea does not serve as an entry site for the virus.

Yoned et al. showed that anti-rabies virus antibody titer in mice intranasally immunized with concentrated rabies virus antigen (CRV) plus cholera toxin (CT), was comparable to that of mice intraperitoneally immunized twice with the same amount of CRV. High levels of IgA and IgG were detected in mice immunized intranasally with CRV/CT [16]. Mucosal vaccination stimulates IgA production in mucosal tissues and IgG antibodies in serum [16]. The mucosal immunization has been proven to be efficient against various infectious pathogens, such as influenza virus, Newcastle disease virus, foot and mouth disease virus, Aujeszky’s disease virus, HIV and Ascaris suum [17-21]. In the present study, rabies virus was instilled into eye. Conjunctiva, is a part of eye mucosa, which its underlying structures are known as conjunctiva-associated lymphoid tissue (CALT). In the CALT, antigens are taken up by the follicles and presented to lymphocytes by antigen presenting cells. This leads to activation of B and T cells, which carry out the immune reaction [12, 22, 23]. In this study, CALT might have an interfering function in neutralization of rabies virus and prevention of virus entrance to the CNS of mice, since contact of conjunctiva with virus containing suspension is inevitable.

Davis et al. showed the presence of rabies virus neutralizing antibody (VNA) in bats and mice exposed through aerosol, to 3 variants of bat rabies virus. In this exposure, all bats and certain numbers of mice survived and produced detectable rabies VNA [24]. In our study, all mice survived, which were exposed to rabies virus by ocular route. Further independent experiments are necessary to demonstrate whether or not survival of mice would be related to immune response and VNA production.

Age, immune status of the host, and factors involved in species spillover have been proved to be important factors in viral neurovirulence [5]. Further investigations would be necessary to rule out the ocular mucosa as a site of virus penetration. As the mouse eye is too small and difficult to instill virus in it, rabbit eye could act as a better model for ophthalmic research, due to its more anatomical similarity to human eye [25]. The experiment can be repeated with rabbits as animal model in order to find more clear description of rabies infection via ocular route. Ocular transmission of rabies virus through abraded cornea could provide similar condition to a real ocular exposure, which could be considered for future studies.

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CONFLICT OF INTEREST

The authors declare that there are no conflicts of interest associated with this manuscript.

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