New Frontiers in Ophthalmology



Opinion ISSN: 2397-2092

Vaccination of corneal herpes (HSK)

Kozaburo Hayashi*

Laboratory of Immunology, National Eye Institute, National Institutes of Health, Bethesda, Maryland, USA

Corneal herpes is not a major concern anymore after effective antivirals have been introduced in ophthalmology clinics. However, immune mediated herpetic stromal keratitis (HSK) still remains the major infection related visual disturbances.

To prevent stromal haze, extended period of combined use of steroid +antivirals is mandatory. Obviously prolonged steroid treatment should be avoided. Up to now, several vaccines trials in murine genital herpes caused by HSV-2 have been reported.

Attenuated HSV-2 (HSV TK-) has been inoculated in the deep nasal mucosa where the attenuated virus proliferated. Subsequently, it elicited educated CD4T+ cells jn the nasal mucosa but not in local dorsal lymph nodes. These CD4 T cells migrated to the vaginal mucosa and educated cervical and/or vaginal mucosal B cells to secrete secretary IgA. However, application of these protocols to the corneal herpes (HSK), there are many pit falls in these projects. Olfactory nerve

is directly extended from the olfactory bulb, therefore it may serve the main entry route of HSV to the brain. Another likely way of HSV to the brain may through tentory nerve via dorsal root ganglion.

Spontaneous shedding of the HSV can be detected quite frequently in tears, therefore bilateral nasolacrimal ducts continually carries numerous virus particles.

We have to pay enough considerations to the HSV-1 vaccination in the deep nasal mucosal epithelium. Attenuated live vaccine is far better than killed/component vaccine to cope with complicated immune mediated HSK.

Clinical HSK in human population results long after prolonged repeated recurrent events. Although we have learned details of immune mediated characteristics of corneal herpes in mouse (C57/Bl or Balb/C) models, we need more studies before safe effective HSV vaccine could be in common usage.

Copyright: ©2020 Hayashi K. This is an open-access article 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.

*Correspondence to: Kozaburo Hayashi, 1-26-8 Chofugaoka, Chofu, Tokyo, Japan 182-002; E-mail: kozaburo@yc4.so-net. ne.jp

Received: November 11, 2020; **Accepted:** December 03, 2020; **Published:** December 10, 2020

New Front Ophthalmol, 2020 doi: 10.15761/NFO.1000253 Volume 6: 1-1