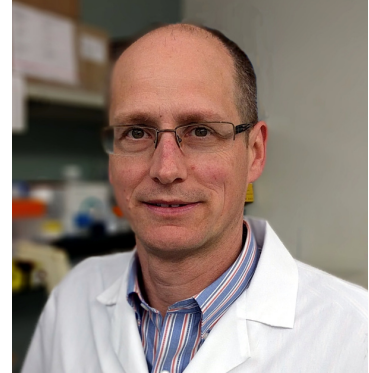


## セミナーのお知らせ

# ***A Flexible VLP Platform To Elicit Protective Responses For a Range of Targets***

**Dr. David Whitacre**  
**Senior Vice President**  
**VLP Biotech, Inc.,**  
**San Diego, California**



Date: January 6<sup>th</sup> Monday

Time: 3pm to 4 pm

Location: Biken Hall, 1<sup>st</sup> Floor, Main building,  
Research Institute for Microbial Diseases

Traditional vaccines have been based on live-attenuated or killed pathogens and newer vaccines have targeted critical pathogen subunits. Further research has identified specific peptide targets that enable neutralizing a virus or bacterial toxin, sequestering an allergen, blocking an undesired signaling event or even stopping parasitic infection when antibodies bind the correct sequence. The concept of epitope-focused vaccines is not new, but peptides alone are poorly immunogenic and require a carrier to elicit potent antibody responses. We have developed a fully-recombinant virus-like particle (VLP) carrier that elicits potent antibody responses to precise peptidic targets by presentation in a matrix array on a highly-immunogenic particle. Our VLP is fully compatible with microbial expression, is self-assembling and highly stable, enabling a low cost-of-goods compatible with vaccines intended for worldwide use. We have proof-of-concept for a variety of pathogens and advanced preclinical programs are for a malaria vaccine to block sporozoite liver infection; an immune therapeutic for chronic hepatitis B infection; and a lyophilizable vaccine for anthrax.

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連絡先: マラリアワクチン開発寄附研究部門 堀井俊宏

horii@biken.osaka-u.ac.jp

Tel: 06-6879-8280