

Towards malaria vaccine development -how basic researchers open the door to product development?-



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Session 5 2022.3.24

The first malaria vaccine to reduce infection: RTS,S



40% reduction in malaria episode. The effects are affected by polymorphisms in CSP.

An improved infection inhibition strategy is urgently required!



Basic research to elucidate parasite infection



Long road for product development



PDPs: Product development partners



The history of collaboration with PDPs



Global Health Innovative Technology Fund

e	愛媛大学 EHIME UNIVERSITY	University at Buffalo The State University of New York	Sumitomo Dainippon Pharma
	Year	Partners	Objectives/Approach
	2014	PATH-MVI	Evaluating Pf75 as a transmission blocking vaccine target
	2016	PATH-MVI	Dissecting Pfs230 for functional subdomain for transmission blocking vaccine
	2019	PATH-MVI, SUNY	Evaluating codelivery of Pfs230 and CSP with CoPoP for transmission & infection blocking vaccine
	2020	PATH-MVI, Sumitomo Dainippon Pharm	Evaluate full-length CSP formulated with SA-1, in comparison with RTS,S/AS01 as a benchmark
	2019	Mahidol University, University of Pennsylvania	Development of a novel Pvs25 nucleoside-modified mRNA vaccine that induces potent and long-lasting transmission blocking immunity



Dissecting Pfs230 for functional subdomain (T2016-207)



Co-delivery of Pfs230C1 and CSP with CoPoP for multistage malaria vaccine (G2019-111)

70-80% reduction in the parasite burden in the liver!!

Outcome of Partnership

- Huang *et al.*, under review
- Continuation to next GHIT project



Lessons learned, challenges and impact to our organization from the partnership

Important for the partnership

- Common goal, vision, and passion
- Complementary skill sets and strengths
- Clear partner role and responsibility
- Respect each partner



Impact from the partnership

- Meet more collaborators with similar aims
- Start new projects with new approaches
- Find new demands of basic research for vaccine development



Thank you!







