

# Novel Approach For An Evolution Proof Influenza Vaccine



Lauren Myburgh<sup>1</sup>, Ruben A. Walen<sup>3</sup>, Karlijn van Loon<sup>1</sup>, Elisabeth J.M. Huijbers<sup>1,2</sup>, Judy R. van Beijnum<sup>1,2</sup>, Colin A. Russell<sup>3</sup>, Arjan W. Griffioen<sup>1,2</sup>

<sup>1</sup> Angiogenesis Laboratory, Cancer Center Amsterdam, Department of Medical Oncology, Amsterdam UMC

<sup>2</sup> CimCure BV, The Netherlands



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<sup>3</sup> Laboratory of Applied Evolutionary Biology, Department of Medical Microbiology, Amsterdam UMC.



1:2700 -

1:8100

1:24300 -

1:72900 -

1:100

1:300

1:900

1:2700

1:8100

1:24300-

1:72900-

0.6

0.4

0.2

0.8

0.6

0.4

0.2

OD Levels (655 nm)

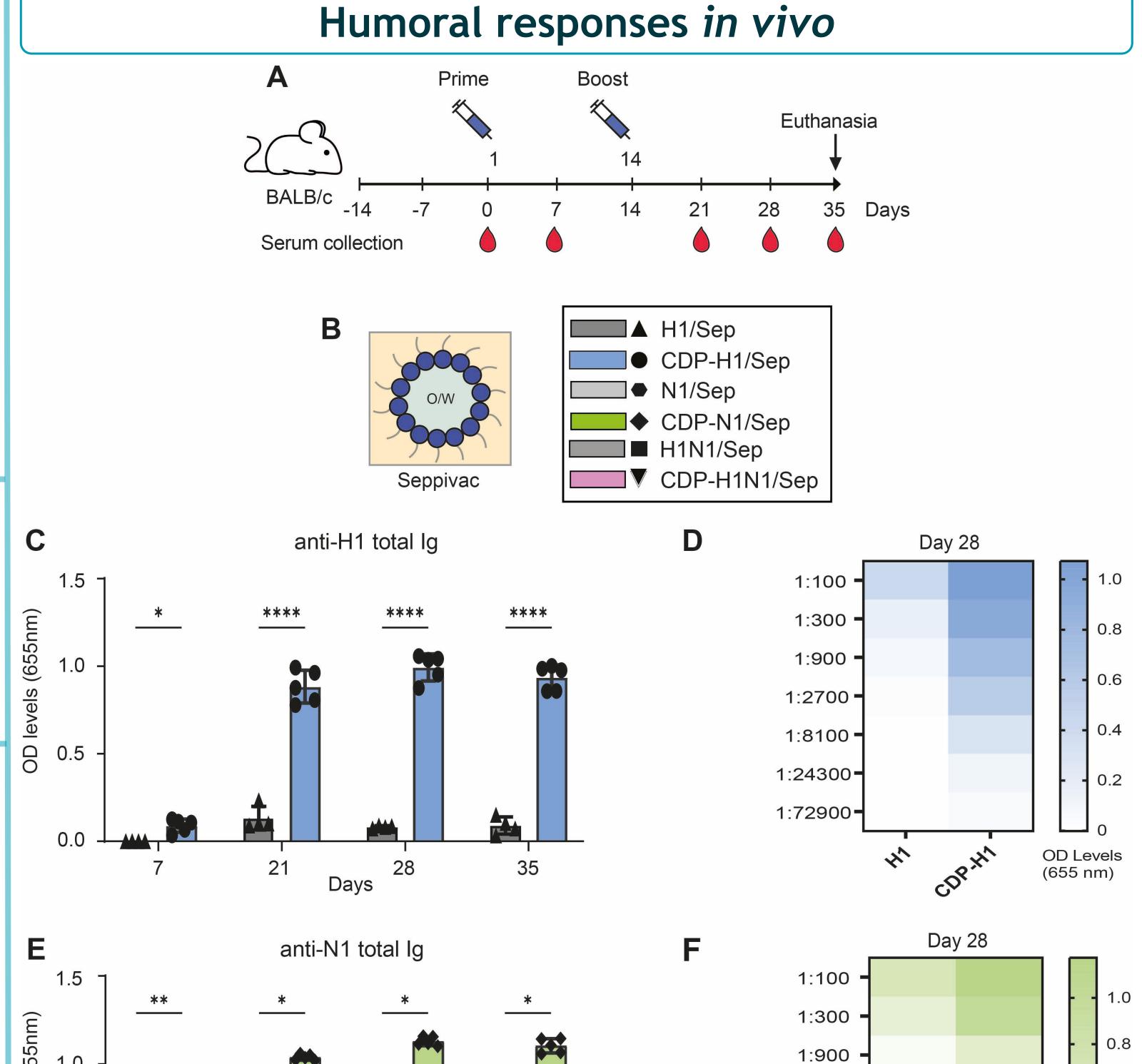
OD Levels (655 nm)

Day 28

HUM

## Background

Our research group developed a technology called Immuno-Boost (iBoost), which aims at increasing targeted immune recognition of non-or low-immunogenic epitopes with the use of a conjugate vaccine. This approach seems perfectly suited for vaccination against the influenza surface glycoproteins, which may allow for safe and scalable production while improving vaccine immunogenicity



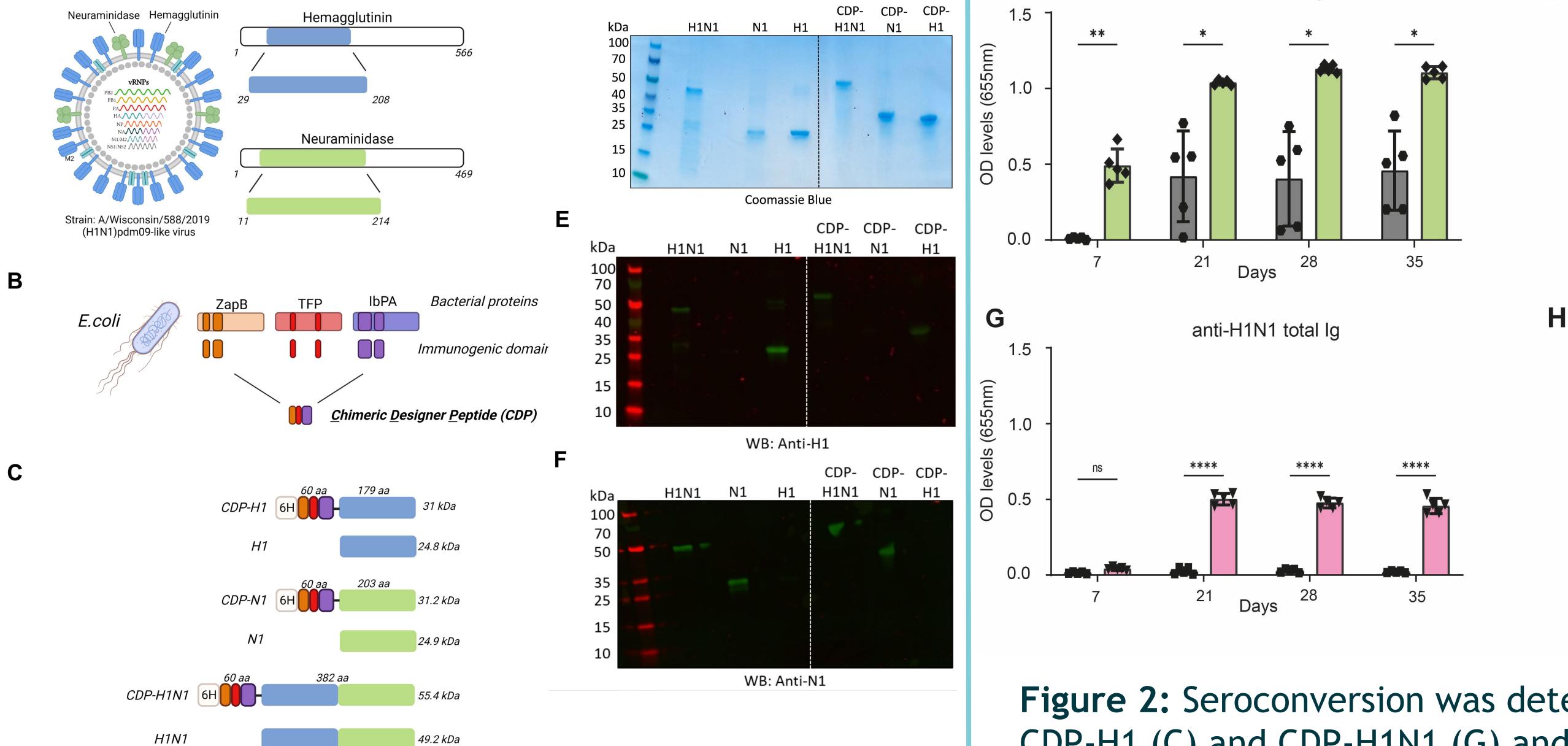
#### Aim

Development of an evolution-proof vaccine that targets both HA and NA using the **iBoost technology** 

## Methods & Results

D

Design of an iBoost-based vaccine against HA and/or NA



**Figure 1:** Constructs (C) were derived from the A/Wisconsin/67/2022 (H1N1) strain (A). These epitopes are then conjugated to CDP (B) and validated by SDS-page and Western Blot using commercial anti-H1 or anti-N1 antibodies (D-F)

**Figure 2:** Seroconversion was detectable by day 21 for CDP-H1 (C) and CDP-H1N1 (G) and by day 7 for CDP-N1 (E). By day 28 (D, F, H) all immunized with the CDP conjugated vaccines elicited a stronger antibody response compared to H1, N1 or H1N1 alone

### Conclusion

Using the iBoost platform for vaccination against HA and/or NA results in not only faster, but also stronger antibody responses compared to the unconjugated counterparts

## Future perspectives

The antibodies induced in this study most likely target major antigenic sites, which are susceptible to antigenic drift. Therefore, to identify suitable, conserved epitopes, we are developing a tailor-made computational pipeline taking into account multiple factors, such as surface exposure and residue-level mutability

E: <u>l.myburgh@amsterdamumc.nl</u>