

Evaluation of a *Campylobacter*-N-glycan presenting *Salmonella* Typhimurium for vaccination of broiler chickens against *Campylobacter*

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Key words

Campylobacteriosis, broiler chicken, live-attenuated vaccine, campylobacter-N-glycan, humoral immune response, maternal antibodies

Aim of the study

„Proof-of-concept“ study. A newly created live-attenuated *S. Typhimurium* vaccine strain, expressing the *Campylobacter*-N-glycan (a heptasaccharide) on its surface, was tested in broiler chickens.

Material and methods

102 commercial day-of-hatch broiler chickens were used in this study. Animals vaccinated on day-of-hatch were kept in isolators for 25 to 35 days with feed and water ad libitum. Blood samples for immunoblot analysis were collected on day 10, 15, 20, 25, (30 and 35). At the end of the experiment, chickens were killed and dissected. Caecal contents was used for quantitative *Salmonella* detection and in the challenge experiments also for quantitative *Campylobacter* counting. Real-time PCR standards were set for *C. jejuni* (Hong et al. 2007) and *S. Typhimurium* (Park et al. 2008) and used for quantitation of the vaccine strain and the challenge strain in liver, spleen and caecal contents.

Results and significance

The first four days after vaccination self-limiting diarrhoea was observed. The *S. Typhimurium* vaccine strain was found in liver and spleen 25 days after vaccination using bacteriology and real-time PCR. Up to 3×10^7 cfu/g *S. Typhimurium* were detected in caecal contents at the end of the experiments. In immunoblot analysis maternal anti-N-glycan IgY were present in sera of all animals until day 15. No anti-N-glycan IgM and IgY were detectable until day 25, thus the vaccine could not stimulate an active humoral immune response against the *Campylobacter* heptasaccharide. In sera from challenged animals specific IgM and IgY antibodies were detected on day 25 or 35, proving an elicited immune response presumptive to the challenge *C. jejuni* strain. In challenge experiments no biologically significant reduction of *C. jejuni* in caecal contents was observed. In conclusion, the vaccine *S. Typhimurium* was able to colonise the chicken without provoking severe disease. The commercial broiler chickens used had maternal IgY against the *Campylobacter*-N-glycan. Chickens did not produce an immune response to the vaccine, however did produce anti-N-glycan IgM and IgY following p.o. inoculation of *C. jejuni*. The experiments showed that the vaccine is safe in broiler chicken and that the *Campylobacter*-N-glycan is immunogenic in chicken. However presentation on the *Salmonella* surface and timepoint of vaccination needs to be adjusted.

Publications, posters and presentations

Presentation: *Campylobacter* Plattform 2010, Bern, 4. November 2010 (Schweizer Zoonosebericht 2011 Seite 9-11)

Presentation: Geflügelseminar der Fachsektion für Geflügelkrankheiten, Zürich, 23. März 2011

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