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Bacteriophage: a potential performance aid for broiler chickens

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BACKGROUND

a growing need to address problems with flock performance. Feed will enhance broiler chicken performance. feed or extra time to achieve saleable weights, which impacts tissues, caecum, ileum and duodenum. processing smaller birds at the abattoir.



Fig 1. Same-age flock containing birds of poor performance. Poorly performing birds are identified with blue arrows.

Prior to 2006 in the EU and UK, subtherapeutic doses of antibiotics, known as antibiotic growth promoters (AGPs), were provided to flocks in order to enhance performance. Due to major concerns of antimicrobial resistance (AMR) and antibiotics entering the food chain (2), along with increased customer demand for poultry products from flocks raised without antibiotic use, nontherapeutic antibiotic use has subsequently been banned in the EU and UK. Further banning of all use of routine antibiotics in farming came into force in the EU in Jan 2022 (Regulation (EU) 2019/6 on Veterinary Medicines.) Alternatives such as herbal extracts, essential oils, prebiotics, probiotics, enzymes and phytochemicals exist but none alone are as effective as AGPs (3).

Bacteriophages (aka phages) are naturally occurring viruses which specifically parasitise bacteria and may regulate gut bacteria, thus potentially providing greater nutrition for the bird's growth. Most investigations into phage use focus on therapeutic use in humans and pigs (4, 5). In poultry, the therapeutic use of phage has been investigated and has seen success in controlling infections such as salmonella (6). However, the characterisation of phage communities (phageomes) in the gut and their relationship with constituent gut bacteria (bacteriomes or microbiomes) is not well understood and little work has been carried out on developing natural phage cocktails and investigating their impact on broiler performance.

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AIMS AND OBJECTIVES

The increasing consumption of poultry meat worldwide highlights. The aim of the project is to produce an innovative phage cocktail that

makes up the largest part of broiler chicken production costs and Objective 1 is to characterise phageomes from good and poorly poorer performing flocks (Fig 1.) may require greater amounts of performing broiler flocks at different lifecycle stages from several gut

farmers' profits (1). Further losses can occur due to difficulties Objective 2 is to culture selected phageomes identified from objective 1 and investigate their potential for controlling bacterial populations by in vitro methods.

> Objective 3 is to feed refined phageome products to broiler chicks to determine potential their performance enhancers. as

WORKFLOW

Longitudinal survey. 10 commercial flocks of various performance. Ilea, duodena and caeca from 5 birds pooled across 3 timepoints.

Bacterial enrichment. Centrifuge at 3k RPM for 15 mins.

Extraction via Roche MagNA Pure 96 system

Viral enrichment. Centrifuge at 3k RPM for 15 mins. 0.45µm and 0.22µm filter. Ultracentrifuge at 113k RCF for 4 hours.

(Roche, Basel, Switzerland).

Extraction and elution of RNA and DNA via Norgen RNA/DNA/Protein Purification Plus kit (Norgen Biotek, Thorold, Canada).

Outsourced for next generation sequencing (NGS).

Trimming and quality checking of NGS data (>4.27) billion reads) using Trim Galore (7).

Sequencing alignment of viral RNA and DNA NGS data via metaSPAdes (8).

Future classification of bacterial 16S and viral RNA and DNA data using appropriate software to identify key phage species and constituent bacteria.

Future culture study to produce phage cocktails based on classified data using Electrolab Fermac 320 bioreactor (Fig 2.) Pre and post culture NGS to confirm presence of species of interest.

Future feed trial experiment using commercially obtained broilers to investigate phage cocktail performance via recording of daily weights.

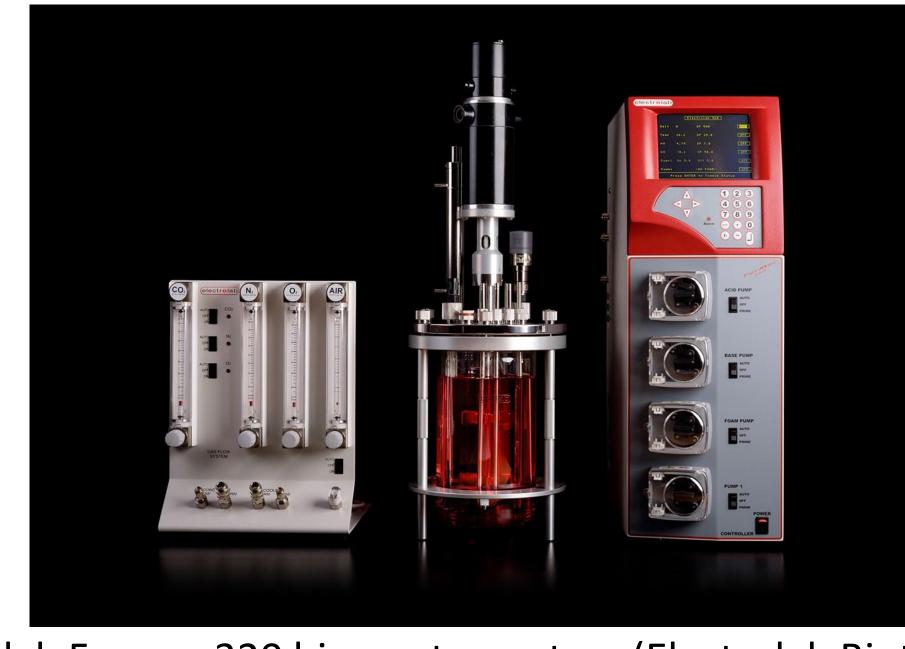


Fig 2. The Electrolab Fermac 320 bioreactor system (Electrolab Biotech, Tewkesbury, England) offers precise control of various environmental factors such as temperature and pH in the process of culturing. (9)

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