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Meta-Taxonomic Analysis of Poultry and Slaughterhouse Microbiota: A Comprehensive Examination of Resident Microbial Communities

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INTRODUCTION & AIMS

Poultry contaminations occur during the processing chain and are found from farm to table.

The final microbial profile of poultry meat is influenced by each step of the process.

The consumption of contaminated poultry poses a significant threat to public health. It is associated to infections caused by pathogens such as: Campylobacter, Salmonella, Listeria monocytogenes and Arcobacter.

BROILER CARCASSESES FINAL MICROBIAL PROFILE **PRODUCTION STAGES FARM: LIVE BIRDS** FXE

The **objective** is to assess the contribution of broiler skin and gut to the indigenous slaughterhouse microbiota. In addition, the study aims to clarify how the persistence and environmental distribution of Arcobacter spp. are influenced by the crosscontamination patterns associated with these specific sources.

MATERIALS & METHODS

1. BROILERS AND ENVIRONMENTAL SAMPLING

154 samples were collected from a local poultry abattoir



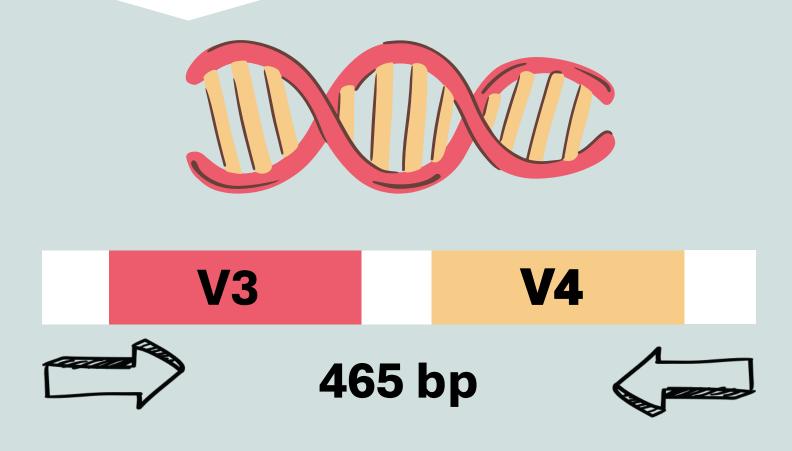
49 samples of broiler neck skin (BNS)

49 samples of broiler caecum (BC)

from samples slaughterhouse equipment (SE)

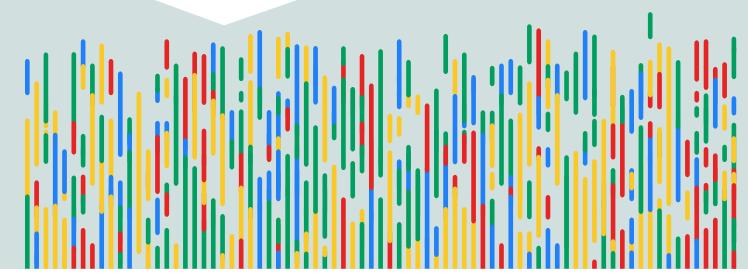
2. DNA extraction

3. V3-V4 region library constructed from the 16S rRNA gene region



4. BIOINFORMATIC ANALYSIS:

taxonomy was assigned with a 99% of sequence similarity by matching ASVs to the Silva prokaryotic reference database



RESULTS & DISCUSSIONS

North-Italy

The BC microbial community is significantly distinct from the BNS and SE community (Fig. 1, 2).

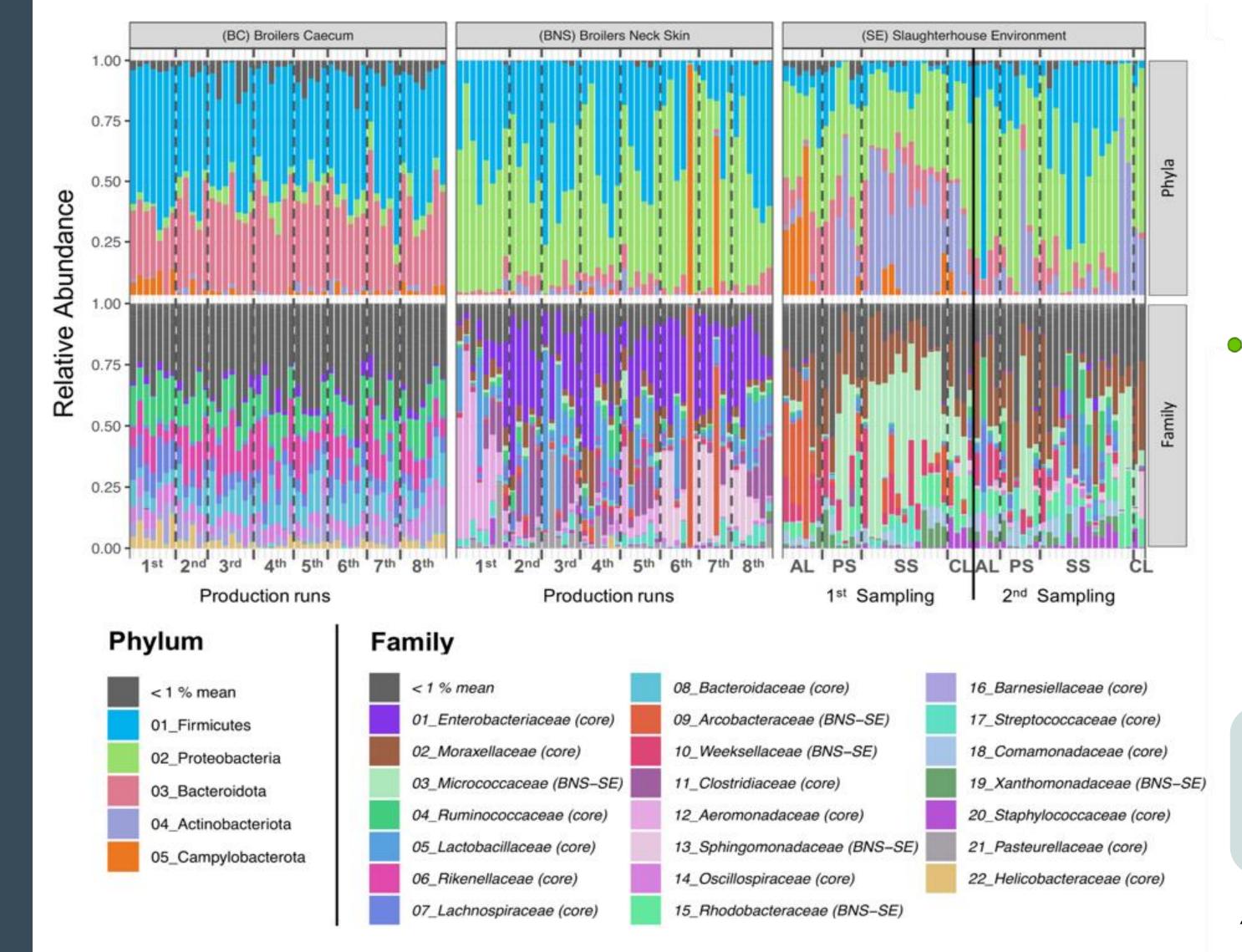


Figure 2. Microbiota composition and distribution in phylum and family taxa. The samples are grouped according to the temporal order of the sampling and according to the order of the slaughter of the flock and the processing stage, BC-BNS and SE. Slaughterhouse sectors and transport lines are indicated: FL = first row; PS = plucking sector; SS = slaughter sector; SL = second row. Taxa are ordered in the legend from most to least abundant (mean >1%).

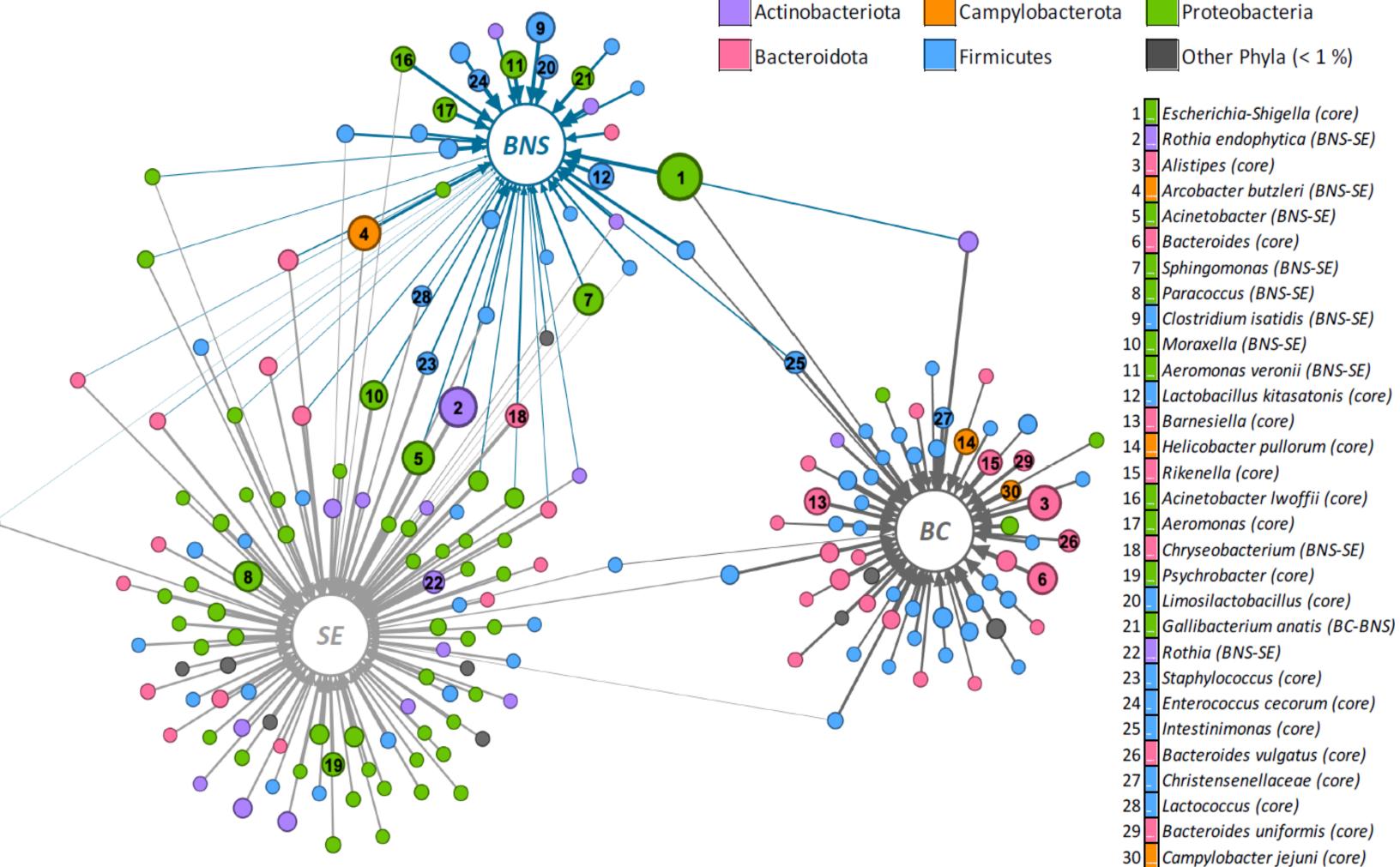
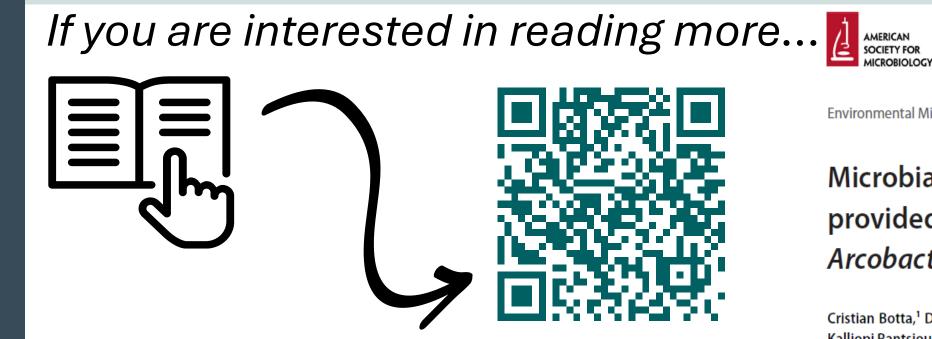


Figure 1. Graphic rapresentation of the network revealing the taxa associated to the three sampling sources. Taxa (colored nodes) are unidirectionally connected with arrows to the sampling sources if significant associations were detected (Indicator Species Analysis: multipatt statistics; R > 0.4and *P* value < 0.001).

Arcobacter butzleri emerged as one of the most abundant species and was detected throughout the slaughterhouse, showing a higher prevalence compared to other Campylobacterota (Fig. 1, 2). It was uniquely and significantly associated with BNS and SE, while Helicobacter pullorum and Campylobacter jejuni were indicators of BC (Fig. 1). A. butzleri was constantly present on BNS along production runs, but significantly more abundant the first sampling day in SE (Fig.2).

CONCLUSIONS & IMPORTANCE

Our findings have emphasised the **persistence of A. butzleri** in a modern poultry abattoir and its establishment as part of the resident microbiota in specific environmental niches. The analysis conducted underlines the significance of early monitoring of food pathogens in the production chain, supported by meta-taxonomic analysis. Using these detection approaches, the presence of these pathogens could soon be considered an indicator of food safety and quality in slaughtered poultry.



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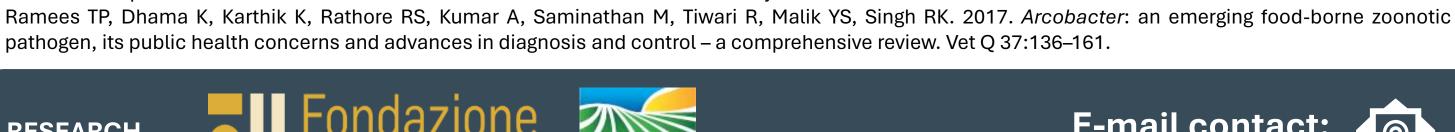
Microbial contamination pathways in a poultry abattoir provided clues on the distribution and persistence of *Arcobacter* spp.

Cristian Botta, Davide Buzzanca, Elisabetta Chiarini, Francesco Chiesa, Selene Rubiola, Ilario Ferrocino, Edoardo Fontanella,



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