

## Occurrence and Antimicrobial-Resistant *Salmonella* Serovars Isolated from Turkey Carcasses and Broiler Turkey Farms in Meknès-Morocco

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### Commentary

Between November 2011 to November 2013 a total of 680 samples were collected: pools of droppings (n=600) and drinking water (n=80). The rate of insulation of *Salmonella* is important (35 %), and the isolated serotypes are worrying: 36 S. Kentucky, 15 S. Saintpaul, 8 S. Parkroyal and 3 S. Ruzizi.

They have relatively high rates of resistance to tetracycline (79 %) and streptomycin (72.5 %), followed by resistance to nalidixic acid (37.1 %), ciprofloxacin (33.9 %). Three strains of *S. Agona* Expanded Spectrum Betalactamase (ESBL) which have a high level of resistance to ceftriaxone with a minimum inhibitory concentration (CMI) of 16 µg /ml were detected.

According to the norms of the water intended for the watering of poultry, for the lock of the line of watering, 100 % of samples are of unacceptable quality as regards the fecal coliforms, *Escherichia coli*, fecal streptococci and sulfitoréducteurs and entérocoques. While more than 90 % of drinking water samples are of a satisfactory quality according to pH, a nitrite, conductivity, nitrate and Iron.

Simultaneously, the nine discovered risk factors were significantly associated with *Salmonella* contamination. These risk factors highlight the risks of the broiler channel, particularly linked to poor technical and hygiene practices.

On the other hand, a total of 228 samples of turkey meat and giblets were collected randomly from retail outlets in Meknès. Ninety six samples were analysed for the presence of the following bacteria: *Escherichia coli* (*E. coli*), coagulase positive *Staphylococcus* (SA), *Clostridium perfringens* (CP), for total aerobic mesophilic flora (FMAT) and thermo tolerant coliforms (TC). Their rate of compliance with hygiene standards for FMAT, TC, *E. coli*, SA and CP are respectively 64.6, 18.8, 11.4, 53.1 and 64.6%.

The level of contamination from supermarkets was identified as significantly lower ( $p < 0.05$ ) the other sites. According to the microbiological criteria, 80% of samples did not meet the standards for *E. coli* and total coliforms. 64.6 % (n=64) and 53.1% (n = 51) samples are of acceptable quality for CP and SA among which 8,3 % of samples could be linked to a foodborne due to a concentration of coagulase positive *Staphylococcus* coagulase-positive upper in 5 log<sub>10</sub> ufc/g.

After the study of the biochemical and the culture characteristics of the isolated *Salmonella* strains (47/192), followed by serotyping and antibiotic susceptibility testing, the PCR was used to confirm the

identification of the *Salmonella* species and the search for genes encoding the virulence factors. The genotypic comparison within the same serotype is based on the analysis of plasmid content and the pulsed field electrophoresis (PFGE).

Out of 192 samples examined, 24.5% were contaminated with *Salmonella*. The sizes of the plasmids obtained ranged from 1.8 to 128 kb. The highest percentage of resistance was found to the following antimicrobial agents: Bacitracin (97.8%), amoxicillin (61.7%), streptomycin (44.6%), triméthoprime (34%), nalidixic acid colistin (19.1%) and ciprofloxacin (17%), the drug commonly prescribed to treat salmonellosis. On the other hand, 97.8% of isolates were found to be resistant to one or more of the antibiotics tested and 100% of *S. Kentucky* (n=8) are resistant to ciprofloxacin and nalidixic acid.

The tests of sensitivity to antibiotics showed for the first time in Morocco the presence of the serotype highly resistant. It comes to *S. Agona* bla SHV resistant to the C3G carried by a plasmid conjugation: the resistance profile (amoxicillin, chloramphenicol, streptomycin, bacitracin, cefotaxime, ceftazidime, ceftriaxone, colistine) is an ESBL with a minimum inhibitory concentration for ceftriaxone 16 µg/ml.

After transfer by way of conjugation, the plasmid carrying the resistance encoding ESBLs, we identified the bla SHV gene by PCR, then, it is sequenced and identified by bioinformatics methods: it is bla *SHV12* that underwent four mutations compared to wild SHV gene.

All *Salmonella* strains tested were positive for the virulence genes (*spiA*, *sifA*, *spaN*, *sopB*, *sipB*, *iroN*, *orgA*, *sitC* and *prgH*), whereas they are positive for virulence genes *spvC* and *spvB* at respective frequencies of 6.3% and 0%. The results of the molecular epidemiology of the *Salmonella* obtained by analysis of plasmid content and by pulsed field gel electrophoresis (PFGE) showed a genetic diversity within the studied serotype.

In this study, the overall performance of 16S rDNA sequence analysis was excellent: 100% strains possessed a 16S rDNA sequence with ≥ 97% similarity to that of a genus *Salmonella*, in that it separates more or less the different serotypes in clusters or sub clusters and. However, in order to improve this performance, efforts should be made to complete 16S rDNA databases with high-quality sequences and develop electronic tools for sequence comparison and interpretation.

The results of this work show the emergence of *Salmonella* non Typhi resistant to the beta-lactam antibiotics and quinolones isolated from turkey meat. The rationalization of the use of the fluoroquinolones in practice veterinary and medical constitutes an urgency of public health in order to cease selecting resistant mutants and to limit human contamination risk. On the other hand, the presence of virulence genes represents a serious threat to public health.

More studies to track the evolution of virulence factors among *Salmonella* must be encouraged.