THE INCIDENCE OF SALMONELLA BACTERIA IN MEAT AND
MEAT PRODUCTS
DURING THE PERIOD 2009 - 2011 IN DOLJ COUNTY

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ABSTRACT

During the period 2009-2011, 3971 meat samples and meat preparations collected both from agri-processing units and from alimentary cold stores and food marketing network were analysed within the Sanitary-Veterinary Laboratory of Dolj County, thus seeking to isolate and identify the serotypes of Salmonella spp. incriminated in triggering food poisoning.

Of the total number of samples examined, six samples were positive representing 0.15 percent. The positive samples were composed of pork minced meat (3), a mixture of prepared beef and pork meat (cattle-swine) (1), refrigerated minced pork meat rolls (1) and turkey wings (1).

All strains of Salmonella spp. isolated in the Sanitary-Veterinary Laboratory of Dolj County were submitted for serotyping of relevant isolates of Salmonella spp in the reference laboratories within the Institute for Diagnosis and Animal Health (IDSA) and the Institute of Hygiene and Public Veterinary Health (IISPV). Four strains belong to the Salmonella Typhimurium serotype, one strain belongs to Salmonella Goldcoast and one strain belongs to Salmonella Hadar. The dominant serovariant identified among tested isolates is Salmonella Typhimurium (66.33 %).

Of the four serovariants of isolated Salmonella Typhimurium, three serovariants presented the classical antigenic structure and one serovariant exhibited a different antigenic structure, lacking the 1,2 factors – initially denominated as Salmonella Typhimurium-like; it has been recently denominated as Monophasic Salmonella Typhimurium.

Key words: Salmonella spp., antigenic structure, food poisoning.

INTRODUCTION

The Salmonella genus is a member of the family Enterobacteriaceae comprising over 2400 serotypes; and bacteria contained within the genus Salmonella have a worldwide proliferation therefore affecting the entire fauna, including the human being (4). Due to its high pathogenicity for human beings through contamination of animal products especially, it is of specific interest for food microbiology (2) Primary Salmonella infections determine high economic damages namely by seizing the products and by-
products derived from animals slaughtered (5). The spreading of Salmonella infections to human beings is increasing subsequent to handling of food products contaminated through raw materials or during their preparation, manipulation, storage and commercialization. The storage inside cold stores prevents the multiplication of Salmonella strains but this does not destroy them. In most cases disorders in humans are a consequence of the consumption of potentially contaminated food resulted from the meat of animals slaughtered upon requirement which were carrying and spreading germs of the genus Salmonella, or the consumption of animal products which were sterilized insufficiently or preserved inadequately. In addition, it must not be forgotten that food products could be contaminated by people carrying Salmonella or infected with Salmonella during the handling of food products as raw materials or finished products. The key feature of food products contaminated with Salmonella is the fact that no organoleptic defects emerges which might draw the attention of the potential presence of germs, which means that the appearance, colour, consistency, smell and taste of food products will remain unchanged (1, 3). The HACCP plan should include control measures for prevention, destruction or elimination of this bacteria and in order to avoid recontamination (1).

**MATERIAL AND METHODS**

Meat samples and meat preparations and products collected from slaughterhouses, processing units, cold stores and food marketing network have been analyzed. The detection and identification of bacteria of the genus Salmonella was carried out according to the reference method SR EN ISO 6579/2003/AC/2009, but alternative methods (Vidas and Vitek methods) were also used; these methods offers the additional advantage of getting the result within a shorter period of time. However, in the event of a positive result obtained through these alternative methods confirmation by means of classical methods should also be carried out. According to the reference method the detection of bacteria of the genus Salmonella requires four successive stages, namely: pre-enriching in non-selective liquid media; isolation and identification; confirmation.
The detection of Salmonella represents the determination of the presence or absence of Salmonella in a certain mass or product bulk when tests are carried out according to the classical method. The pre-enriching is performed by means of buffered peptone water (225ml) which has been pre-warmed to room temperature and has been inoculated to the sample to be analysed (25g), then it is incubated at 37°C ±1°C for a period of 18h±2h. The following have been used as liquid selective media: Rappaport media - Vassiliadis cu soia (bulion RVS) și bulionul Muller - Kauffmann tetrationat/novobiocină (bulion MKTTn). These media are incubated at specific temperatures according to the working standard procedure, for a period of 24h±3h. For the purpose of isolation and identification two solid growth media are inoculated: agar with xylose - lysine - deoxycholate (XLD) – the first selective medium and the second selective medium – any other solid growth medium complementary with the XLD agar and appropriate especially for the isolation of lactose positive Salmonella and the genus Salmonella Typhi and Salmonella Paratyphi (Istrati Meitert - IM or Edel and Kampelmaker - EK); the selective growth medium Edel and Kampelmaker is currently used in the Sanitary-Veterinary Laboratory of Dolj County. The XLD agar is being incubated at 37°C±1°C and it is examined after a period of 24h±3h. The second selective agar is being incubated according to the manufacturer's regulations. The typical colonies of Salmonella grown on XLD agar have a black spot in the middle and a transparent bright reddish area due to the change of the indicator colour. The confirmation of the supposed Salmonella colonies is carried out by means of biochemical or serological tests using identification kits. The recognition of Salmonella colonies is mainly an issue related to experience and their appearance may vary anyway, not only from serovar to serovar but also from a certain selective growth media batch to another selective growth media batch used.

RESULTS AND DISCUSSIONS

Throughout the survey period 3971 samples on different matrices were analyzed within Dolj Sanitary Veterinary and Food Safety Laboratory in order to detect bacteria of the genus Salmonella in meat and meat preparations and products.
### Samples collected on different matrices during the period examined

<table>
<thead>
<tr>
<th>MATRIX</th>
<th>Samples examined 2009</th>
<th>Samples examined 2010</th>
<th>Samples examined 2011</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovine, sheep, caprine carcases</td>
<td>34</td>
<td>47</td>
<td>30</td>
</tr>
<tr>
<td>Pigs carcases</td>
<td>41</td>
<td>42</td>
<td>40</td>
</tr>
<tr>
<td>Poultry carcases</td>
<td>38</td>
<td>69</td>
<td>21</td>
</tr>
<tr>
<td>Fresh meat and comestible by-products</td>
<td>293</td>
<td>286</td>
<td>38</td>
</tr>
<tr>
<td>Minced meat and prepared meat derived from poultry, meant to be cooked</td>
<td>25</td>
<td>33</td>
<td>27</td>
</tr>
<tr>
<td>Minced meat and prepared meat derived from species other than poultry, meant to be eaten after being cooked</td>
<td>389</td>
<td>563</td>
<td>330</td>
</tr>
<tr>
<td>Processed chicken meat meant to be prepared (cooked)</td>
<td>56</td>
<td>40</td>
<td>35</td>
</tr>
<tr>
<td>Heat treated processed meat</td>
<td>500</td>
<td>434</td>
<td>560</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>1376</strong></td>
<td><strong>1514</strong></td>
<td><strong>1081</strong></td>
</tr>
</tbody>
</table>

Subsequent to the carrying out of analyses and data processing it results the fact that in 2009 two samples were found positive out of a total number of 1376 samples representing a 0.15 percentage of meat and meat preparations’ samples. In 2010, two samples were found positive out of a total number of 1514 samples representing a 0.14 percentage; in 2011, two samples were found positive out of a total number of 1081 samples examined representing a 0.19 percentage.

### Positive cases and percentage of positive samples reported to the total examined

<table>
<thead>
<tr>
<th>Period Samples</th>
<th>2009</th>
<th>2010</th>
<th>2011</th>
</tr>
</thead>
<tbody>
<tr>
<td>Examined</td>
<td>1376</td>
<td>1514</td>
<td>1081</td>
</tr>
<tr>
<td>Positive</td>
<td>2</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>% positive</td>
<td>0.15</td>
<td>0.14</td>
<td>0.19</td>
</tr>
</tbody>
</table>
The six positive confirmed samples derive from different matrices, namely: three samples are pork minced meat, one sample was collected from a mixture of prepared beef and pork meat (cattle-swine), one sample was collected from refrigerated minced pork meat rolls and one sample was collected from turkey wings.

It should be specified that Salmonella spp. strains, dated 2009, were isolated in the following assortments: mixture of prepared beef and pork meat (crude frozen product), sample collected from a cold store; the pork minced meat was taken from a supermarket. Salmonella group OC was isolated from the pork minced meat sample according to the reference method SR EN ISO 6579/AC/2006. This strain was confirmed at IDSA and IISPV in Bucharest as being a Salmonella Goldcoast strain. Subsequent to the carrying out of the product traceability it has been determined that the manufacturer that had supplied the pork minced meat to the supermarket had purchased the raw material from a swine farm endowed with its own slaughterhouses which had a past history with this type of Salmonella (S. Goldcoast) detected in its own head.

All strains of Salmonella spp. isolated in LSVSA Craiova were submitted for serotyping in the reference laboratories within IDSA and IISPV.

<table>
<thead>
<tr>
<th>Period</th>
<th>Matrix</th>
<th>2009</th>
<th>2010</th>
<th>2011</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Samples examined</td>
<td>Positive samples</td>
<td>Samples examined</td>
<td>Positive samples</td>
</tr>
<tr>
<td></td>
<td>Fresh meat</td>
<td>293</td>
<td>-</td>
<td>286</td>
</tr>
<tr>
<td></td>
<td>Minced meat and prepared meat</td>
<td>389</td>
<td>2</td>
<td>563</td>
</tr>
</tbody>
</table>

Out of the six isolated strains, four strains belong to the Salmonella Typhimurium serotype, one strain belongs to Salmonella Goldcoast and one strain belongs to Salmonella Hadar.

The dominant serovariant identified among tested isolates is Salmonella Typhimurium representing 66.33 percentage of the total number of isolated germs of Salmonella species.

Among the serovariants of isolated Salmonella Typhimurium, three serovariants presented the classical antigenic structure and one serovariant
exhibited a different antigenic structure, lacking the 1,2 factors – initially
denominated as Salmonella Typhimurium-like; it has been recently
denominated as Monophasic Salmonella Typhimurium.

![Chart 1](image)

Chart 1 - the percentage of the four genii isolated from the total number of samples examined

**CONCLUSIONS**

Bacteria of the genus Salmonella identified, isolated and confirmed
throughout the period examined were six; they did not exceed the said 1
percentage of the samples examined each year.
Subsequent to the carrying out of this study, it can be concluded that the
majority serovariants isolated were Salmonella typhimurium.
The highest percentage of germs of the genus Salmonella was detected in
meat half-cooked products: minced meat, prepared meat and minced meat
rolls paste.
The isolation of germs of the genus Salmonella spp. in meat and meat
products is mainly the consequence of intense processing and handling
labours carried out by people as the highest percentage was isolated from
sorts enduring multiple operations.
The epidemiologic surveys performed led to the conclusion that
contamination with Salmonella inside the slaughterhouses cannot be
neglected when slaughtering animals carrying and spreading bacteria of the
genus Salmonella and in cases when principles regarding products handling, instruments sterilisation are disregarded as well as in cases of non-compliance with the principle of the two knives.

REFERENCES