Detection of Extended Spectrum Beta-Lactamases from a Pet Blue-Fronted-Amazon Parrot (*Amazona aestiva*)

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ABSTRACT

**Background:** The indiscriminate use of antibiotics both in humans and in animals, has contributed to the development of bacterial resistance. One of the key mechanisms in the resistance of enterobacteria to antibiotics is the production of extended-spectrum β-lactamases (ESBLs), which reduce the therapeutic options available. Several studies have been performed in different animal species in order to isolate and identify multidrug-resistant strains and determine their antibiotic sensitivity profile. The purpose of this study was to detect ESBL-producing enterobacteria in isolates from a pet blue-fronted amazon parrot.

**Case:** A 16-year-old pet blue-fronted-amazon parrot (*Amazona aestiva*) weighing 0.445 kg, from the city of Presidente Castelo Branco, Paraná, Brazil, was admitted into a specialized veterinary clinic in the city of Maringá. This parrot was presented with agitation and aggressive behavior. It was fed with sunflower seeds, and its cage was small and unsanitary especially in the feeding and drinking areas, which were heavily contaminated with feces. The parrot had been diagnosed with acute sinusitis approximately one year prior, with a history of treatment with tylosin, thuya (a herbal remedy popularly used for fowl pox), and a mineral-vitamin complex. The clinical symptoms included dyspnea and bilateral increase of facial swelling, with a mass in the peri-nasal region. A membranous red tissue not related to the nictitating membrane was observed in the lower left eyelid. To obtain a better clinical evaluation, the bird was anesthetized with an intramuscular injection of dextroketamine 50 mg/kg (KetaminS+™). Subsequently, physical examination, head radiographic examinations, and an attempt at sinus draining were performed. The radiographic examinations showed a decreased amount of air in the nasal sinuses. For treatment, oral itraconazole and vitamin A were prescribed, and a dietary improvement, prioritizing the offering of fruits and industrialized food for parrots was proposed. Samples were collected from the cloaca and choanae by making rotating movements with compressed sterile swabs in the corresponding locations. All samples were kept in Amies transport media with activated coal and forwarded under refrigeration to the Laboratory of Preventive Veterinary Medicine and Public Health in the Post-Graduation Program in Animal Sciences with Emphasis in Bioactive Products at Universidade Paranaense (UNIPAR).

**Discussion:** The samples were subjected to phenotypic antimicrobial sensitivity tests and phenotypic testing for detecting ESBL-producing strains. *Escherichia coli* was identified and isolated from the cloacal sample. Phenotypic tests for antimicrobial sensitivity, detected resistance to the following antimicrobials: ceftriaxone, ceftiofur, cefotaxime, cefepime, ampicillin, amoxicillin, amoxicillin + clavulanate, and tetracycline. The phenotypic test for detecting ESBL-producing strains was positive. The findings in this study had no relation with the clinical symptoms presented by the parrot. However, in the samples collected, it was possible to detect the presence of ESBL-producing *Escherichia coli*, indicating that this parrot had strains in its cloacal microbiota that were either multidrug-resistant or at a subclinical stage of an infection. This knowledge is important, since the presence of this bacteria in a pet represents an important factor in the dissemination of multidrug-resistant strains into the environment, as well as being a source of contamination for both humans and other animals.

**Keywords:** antibiotics, birds, enterobacteria, gram-negative bacteria, health risk.
INTRODUCTION

The indiscriminate use of antibiotics, both in humans and in animals, has contributed to the development of resistance of bacteria to antibiotics [6,11]. In humans, 30 to 35% of all cases of sepsis, more than 70% of urinary infections, and the most common and relevant intestinal infections are caused by bacteria of the family Enterobacteriaceae. Production of extended spectrum β-lactamases (ESBLs) is one of the main mechanisms by which Enterobacteriaceae develop resistance, thus reducing the available therapeutic options [5].

Several bird species such as the Brazilian blue-fronted-amazon parrot (Amazona aestiva), live with humans. This bird is known for its beautiful and colorful feathers as well as its capacity for learning vocabulary and imitating human speech. It is therefore, very popular as a pet [2].

Even though this species has been quite extensively studied in clinical terms, there are few studies related to bacterial infections, especially regarding ESBL resistance in these parrots. Thus, the purpose of this study was to detect ESBL-producing Enterobacteriaceae in isolates from a pet blue-fronted-amazon parrot.

CASE

A 16-year-old pet blue-fronted-amazon parrot (Amazona aestiva) weighing 0.445 kg, from the city of Presidente Castelo Branco, Paraná, Brazil (Figure 1), was admitted into a specialized veterinary clinic in the city of Maringá. This parrot was presented with agitation and aggressive behavior. It was fed with sunflower seeds, its cage was small and unsanitary, especially in the feeding and drinking areas, which were heavily contaminated with feces. The parrot had been diagnosed with acute sinusitis approximately one year prior, with a history of treatment with tylosin, thuya (a herbal remedy popularly used for fowl pox), and a mineral-vitamin complex.

The clinical symptoms included dyspnea and bilateral increase of facial swelling, with emphasis on the presence of a mass in the perinasal region. A membranous red tissue not related to the nictitating membrane was observed in the lower left eyelid (Figure 1). In order to obtain a better clinical evaluation, the bird was anesthetized with an intramuscular injection of dextroketamine 50 mg/kg (KetaminS+™). Subsequently, physical examination, head radiographic examinations, and an attempt at sinus draining were performed. The radiographic examinations showed a decreased amount of air in the nasal sinuses.

For treatment, oral itraconazole and vitamin A were prescribed, and a dietary improvement prioritizing the offering of fruits and industrialized food for parrots was proposed.

Samples were collected from the cloaca and choanae by making rotating movements with compressed sterile swabs in the corresponding locations. All samples were kept in Amies transport medium with activated coal and forwarded under refrigeration to the Laboratory of Preventive Veterinary Medicine and Public Health in the Post-Graduation Program in Animal Sciences with Emphasis in Bioactive Products at Universidade Paranaense (UNIPAR).

At the laboratory, the swabs containing the samples were placed in tubes containing 3.0 mL of Brain Heart Infusion (BHI) medium, and incubated at 37°C for 24 h. After incubation, the cultures obtained were sown by striation in plates containing MacConkey agar with 10 µ/mL cefotaxime, and MacConkey agar with 50 µg/mL nalidixic acid. These plates were incubated at 37°C for 24 h in order to isolate the colonies resistant to cephalosporines and fluoroquinolones. The isolated colonies were preserved in BHI medium and then were stored in 80% glycerol at -20°C for preservation [8].

The biochemical identification of the bacteria belonging to the Enterobacteriaceae family was made using the “Kit para Enterobactérias” Enterobacteria Kit®, according to the manufacturer’s instructions [8].

In order to determine the bacterial resistance profile, the agar diffusion disk method was used, according to the recommendations by the Clinical and Laboratory Standards Institute [3]. The tested discs were the following: gentamicin, ciprofloxacin, ceftazidime, sulfazotrim, amikacin, aztreonam, chloramphenicol, ampicillin, tobramycin, cefotaxin, ceftriaxone, cefepime, cefotaxime, tetracycline, amoxicillin, amoxicillin + clavulanate, imipenem, meropenem, ertapenem, norfloxacin, nalidixic acid, enrofloxacin, and ceftiouf.

The double disc synergy test was performed, with the discs containing cefotaxime, ceftazidime, ceftriaxone, and aztreonam distributed at a 20-mm distance from another disc containing amoxicillin + clavulanate (20/10 µg). Any increase or distortion in the inhibition zone of any of the antibiotics towards the amoxicillin + clavulanate disc was considered suggestive of ESBL production [1].
Bacterial growth in BHI was observed in all samples collected. However, when transferring them to MacConkey agar with ceftaxime, only the cloaca sample presented bacterial growth, allowing for isolation and identification of *Escherichia coli*.

In the phenotypic tests for the sensitivity to antimicrobials, resistance to the following antimicrobials was observed: ceftriaxone, ceftiofur, ceftaxime, cefepime, ampicillin, amoxicillin, amoxicillin + clavulanate, and tetracycline (Figure 2). The phenotypic test for detecting ESBL-producing strains was positive (Figure 3).

**DISCUSSION**

The presence of multidrug-resistant bacteria in animals is not always related to the incorrect use of antibiotics or to mutation events, due to selective pressures that make bacteria resistant. In many cases, animals may have acquired multidrug-resistant strains from the environment itself [11]. Given that the strain in this study was isolated from a pet bird, it cannot be ruled out that the bird may have acquired the strain directly from its care givers, because different studies have isolated multidrug-resistant *E. coli* strains in healthy humans [10].

Despite these assumptions, the relevance of this study lies in the isolation of an ESBL-producing strain from a pet, which can be a source of contamination and dissemination of such bacteria. ESBL-producing strains are particularly important since they usually present cross-resistance to quinolones and trimethoprim + sulfamethoxazole [7]. Such multidrug-resistant profiles reduce the therapeutic options, making it difficult to treat infections in both people and animals. The early identification of ESBL production allows for a greater control of the dissemination of the bacteria to the community and the environment, as well speeds up appropriate treatment [7].

Regarding the isolation of resistant bacteria in birds, Horn et al. [4], isolated 61 bacterial strains from 42 cloacal and 9 necropsy swabs from canaries.
(Serinus canarius) in the city of Fortaleza, in the State of Ceará, Brazil. The species found were Escherichia coli, Klebsiella spp., Enterobacter spp., Pantoea agglomerans, Serratia spp., and Proteus mirabilis, with E. coli being the most prevalent one, with a total of 25 isolations. Regarding sensitivity to antimicrobials, the strains presented higher resistance to sulfonamides (55.70%), ampicillin (54.10%), and tetracycline (39.30%). Additionally, the total number of strains resistant to multiple drugs was 55.70%, concluding that the birds analyzed, as well as being colonized by enterobacteria, also presented strains with a high level of resistance to some antimicrobial classes.

Similar results were obtained by Santos et al. [9], who identified a greater prevalence of E. coli (70.50%), Staphylococcus aureus (49.00%), and Streptococcus spp. (25.40%) when analyzing samples from cloacal swabs of 51 captive cracidae birds from 10 different species in the State of Rio Grande do Sul, Brazil. With respect to sensitivity, all 93 isolated strains were sensitive to only imipenem, and it was observed that the birds studied presented multidrug-resistant bacteria from several genera and species in their cloacal microbiota, which presented different resistance mechanisms.

The findings in this study had no relation with the clinical symptoms presented by the parrot. However, in the samples collected, it was possible to detect the presence of ESBL-producing Escherichia coli, indicating that this parrot had strains in its cloacal microbiota that were either multidrug-resistant or at a subclinical stage of an infection. Such knowledge is important, since the presence of this bacteria in a pet represents an important factor in the dissemination of multidrug-resistant strains into the environment, as well as being a source of contamination for both humans and other animals, thus being considered a unique health risk.

MANUFACTURERS

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REFERENCES

