

Mumtaz et al., 2014; Sayarifard et al., 2015). The slight variations of ESR values after amikacin administration can be explained by the low nephrotoxic potential of this aminoglycoside, less than the same for gentamicin and tobramycin (Hottendorf and Gordon, 1980).

Spectinomycin leads to significant increase of ESR, followed by a trend to recovery of the baseline values. The differences were statistically significant, but not highlighted, probably due to the low potential of spectinomycin to cause changes in biochemical and hematological parameters at therapeutic doses (Dinev, 2007). Apramycin brought about the least pronounced and insignificant changes of ESR. We assume the reason for the less distinct changes of goat's ESR in aminocyclitol treatment is due to the lack of nephrotoxic potential of the aminocyclitols (Dowling, 2013). Up to now they are no experimental data suggesting nephrotoxic effects after their application, even after high-dose therapy (Novak et al., 1974). Kanamycin caused decrease of ESR values in the last days of treatment, without restoration - a tendency that is generally contrary to the observed in our study. However, differences in the values of ESR, although statistically significant, are not highly expressed. Lack of research data regarding ESR response to spectinomycin, kanamycin and apramycin administration makes difficult the explanation of the somewhat inconsistent trends observed in our study. We assume that this is due to the use of therapeutic doses of antibiotics, which rarely cause pronounced changes.

In conclusion, according to the experimental results gentamicin and tobramycin cause remarkable changes in goat's ESR, even after treatment with therapeutic doses. This gives grounds to recommend ESR data after treatment with these antibiotics to be cautiously interpreted. Generally, the aminocyclitols cause less pronounced changes in the goat's ESR than the aminoglycosides.

REFERENCES

- Brion, N., Barge, J., Gadenne, J., Dromer, F., Dubois, C., Contro, G., Caron, C., 1984. Gentamicin, netilmicin, dibekacin, and amikacin nephrotoxicity and its relationship to tubular reabsorption in rabbits. *Antimicrobial Agents and Chemotherapy* 25, 168-172.
- Dhar, H., Shah, K., Ghongane, B., Rane, S., 2013. Nephroprotective activity of crocus sativus extract against gentamicin and/or ceftazidime - induced nephrotoxicity in rats. *International Journal of Pharma and Bio Sciences* 4, 864-870.
- Dinev, T., 2007. Comparative investigations on side effects, antimicrobial activity and pharmacokinetics of aminoglycosides and aminocyclitols in goats and goat's isolated microorganisms. PhD Thesis, Trakia University, Stara Zagora, Bulgaria.
- Dinev, T., Kanakov, D., Zapryanova, D., 2005. Investigations on some biochemical and haematological parameters after tobramycin and amikacin treatment in female goats. *Trakia Journal of Sciences* 3, 14-16.
- Dowling, P., 2013. Aminoglycosides and Aminocyclitols. In: Giguere, S. (Ed.), *Antimicrobial Therapy in Veterinary Medicine*, Ames, USA, pp. 233-256.
- Hottendorf, G., Gordon, L., 1980. Comparative low-dose nephrotoxicities of gentamicin, tobramycin, and amikacin. *Antimicrobial Agents and Chemotherapy* 18, 176-181.
- Lacy, M.K., Nicolau, D.P., Nightingale, C.H., Quintiliani, R., 1998. The Pharmacodynamics of Aminoglycosides. *Clinical Infectious Diseases* 27, 127-129.
- Lashev, L., Lasarova, S., 2011. Pharmacokinetics and side-effects of gentamicin in healthy and *Pseudomonas aeruginosa* infected sheep. *Journal of Veterinary Pharmacology and Therapeutics* 24, 237-240.
- Liu, S., Ren, J., Xia, Q., Wu, Y., Han, S., Ren, H., Yan, D., Wang, G., Guo, G., Li, J., 2013. Preliminary case-control study to evaluate diagnostic values of C-reactive protein and erythrocyte sedimentation rate in differentiating active Crohn's disease from intestinal lymphoma, intestinal tuberculosis and Behcet's syndrome. *The American Journal of the Medical Sciences* 346, 467-472.
- Mumtaz, F., Khaliq, T., Rahman, Z., Javed, I., Iftikhar, A., Aslam, M., Ali, A., Ahmad, Z., 2014. Effects of *Rosa damascena* mill flowers, *Cichorium intybus* linn roots and their mixtures on serum electrolytes and hematological parameters against gentamicin induced toxicity in albino rabbits. *Indo American Journal of Pharmaceutical Research* 4, 236-241.
- Novak, E., Gray, J.E., Pfeifer, R.T., 1974. Animal and human tolerance of high-dose intramuscular therapy with spectinomycin. *The Journal of Infectious Diseases* 130, 50-55.
- Rankin, L.I., Luft, F.C., Yum, M.N., Isaacs, L.L., 1980. Comparative nephrotoxicities of dibekacin, amikacin, and gentamicin in a rat model. *Antimicrobial Agents and Chemotherapy* 18, 983-985.
- Sayarifard, A., Javadilarijani, F., Mousavi Movahhed, S., Moghtaderi, M., Javadilarijani, F., 2015. Amikacin-induced Nephrotoxicity in a Child with Idiopathic Nephrotic Syndrome in Iran: A Case Report. *Journal of Pediatric Nephrology* 3, 31-34.
- Schentag, J.J., Plaut, M.E., Cerra, F.B., 1981. Comparative nephrotoxicity of gentamicin and tobramycin: pharmacokinetic and clinical studies in 201 patients. *Antimicrobial Agents and Chemotherapy* 19, 859-866.
- Sumano, H., Gutierrez, L., Velazquez, C., Hayashida, S., 2005. Pharmacokinetics and renal toxicity of three once-a-day doses of amikacin in cows. *Acta Veterinaria Hungarica* 53, 231-240.
- Yazar, E., Elmas, M., Altunok, V., Sivrikaya, A., Oztekin, E., Birdane, Y.O., 2003. Effects of aminoglycoside antibiotics on renal antioxidants, malondialdehyde levels, and some serum biochemical parameters. *The Canadian Journal of Veterinary Research* 67, 239-240.