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Original Article

A survey of equine anti-Listeria monocytogenes antibodies using Latex Agglutination Test in southeast of Iran

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ABSTRACT

Listeriosis is a zoonotic disease in humans and a wide range of domestic and wild animals and also some birds. The main purpose of the current study is to determination of anti-Listeria monocytogenes sero-prevalence in horses by latex agglutination test in southeast of Iran. A total of 163 serum samples were obtained from apparently healthy horses of equestrian clubs in Kerman and Yazd provinces - Iran. Listeria monocytogenes antibodies were found in 34 out of the 163 sera (20.85%). The latex agglutination test can be considered as an appropriate screening test in the early stages of diagnosis.

ردیاپی پادتن ضد لیستریا مونوسیتوژنز با روش لاتکس اگلوتیناسیون در اسبان جنوب شرق ایران پونا فرامرزپور 1 ، احسان اله سخائی 3 ، مهدی گلچین 3 ، بلال صادقی

ا گروه علوم درمانگاهی دانشکده دامپزشکی دانشگاه شهید باهنر کرمان، کرمان، ایران ۲ گروه علوم درمانگاهی دانشکده دامپزشکی دانشگاه شهید باهنر کرمان، کرمان، ایران ۳ گروه پاتوبیولوژی دانشکده دامپزشکی دانشگاه شهید باهنر کرمان، کرمان، ایران ۳ گروه بهداشت مواد غذایی دانشکده دامپزشکی دانشگاه شهید باهنر کرمان، کرمان، ایران

چحیده

لیستریوز یک بیماری مشترک میان انسان و طیف وسیعی از دامهای اهلی، وحشی و پرندگان می باشد. هدف از انجام مطالعه ی حاضر ردیاپی پادتن ضد لیستریا مونوسیتوژنز با روش لاتکس اگلوتیناسیون در اسبان جنوب شرق ایران می باشد. برای این منظور تعداد ۱۶۳ نمونه سرم از اسب های به ظاهر سالم باشگاه های سوارکاری استان های یزد و کرمان تهیه شد. نتایج مطالعه حاکی از حضور پادتن در ۳۴ نمونه از مجموع ۱۶۳ سرم مورد بررسی (۲۰/۸۵ درصد) می باشد. بر این اساس به نظر می رسد که لاتکس اگلوتیناسیون آزمایش مناسبی برای غربالگری اولیه به منظور تشخیص بیماری می باشد.

واژه های کلیدی: لیستریا مونوسیتوژنز، آزمایش لاتکس اگلوتیناسیون، اسب، ایران

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INTRODUCTION

Listeriosis is one of the food-borne zoonosis diseases around the world caused by Listeria species. This causes considerable morbidity and mortality in humans and animals and the bacterium is considered an important food borne pathogen [1]. Listeria monocytogenes is ubiquitous, aerobic, gram positive and rodshape bacterium, which is capable of causing severe disease in many species as sheep, cattle, goats, horse, humans and chickens [7,19]. There is six species of listeria in this genus that categorized into three groups as genome analyze, that first group contains monocytogenes, L. innocua and L. welshimeri, the second one L. ivanovii, and L. seeligeri, and the third include L. grayi [8]. Listeriosis occurs as a sporadic disease in horses with pathogenic highly intracellular, non-acid resistant bacteria clinically and characterized by meningo-encephalitis. Signs associated with the nervous syndrome include paralysis of the mandibular and pharyngeal muscles, difficulty walking, inappetence, polydipsia, loss of body weight, and collapse. Following the first observation of signs, animals usually die within 3-10 days [9,11]

MATERIALS AND METHODS

Sample collection and processing

A total of 163 serum samples were collected from 93 male and 70 female (clinically healthy) at 9 equestrian clubs in Kerman and Yazd provinces, Iran. The breed composition comprised of Arab and Arab cross horses (50 %) and non-Arab horses (50%) including Darreh-Shouri (5%), Turkmen (12 Thoroughbred (3 %) and mixed breeds (30 %). After collection, the samples were submitted to Microbiology laboratory of School of Shahid Veterinary Medicine, Bahonar University of Kerman, Iran. Sera were stored at -20°C until analysis.

Latex Agglutination Test (LAT)

Latex Agglutination Test Kit (Zist Faravard Pars Co, Rasht, Iran) was used according to the manufacture's protocol. Briefly, 10 µl of serum and 10 µl of antigen coated latex particles were added to on an agglutination card and mixed with a plastic stirrer. The card was rocked from side to side for up to 5 min to provoke the agglutination reaction. Specimens that showed agglutination during this period were recorded as positive, and otherwise negative. Incomplete agglutination recorded as suspected. Specificity and sensitivity assurance was carried out by testing positive and negative controls on daily basis.

RESULTS

According to the Table 1 and 2, antibodies were detected in 34 sera (20.85%) out of 163 samples (21 cases \leq 7 years old and 13 cases >7 years old). Based on the results presented on Table 1 and 2, 21 (15 cases \leq 7 years old and 6

Table 1. Number and frequency (%) of positive, suspected and negative samples among 163 equine serum samples

Gender	Pos	sitive	Su	spect	Neg	Total		
	No*	F (%) **	No	F (%)	No	F (%)	No	
Stallion	21	12.88	22	13.50	50	30.67	93	
Mare	13	7.98	22	13.50	35	21.47	70	
Total	34	20.86	44	27	85	52.14	163	

^{*:} Number of cases **: Frequency (percent)

		Age													Total							
Province	മ	≤ 7 years old									> 7 years old					, Total						
	Gender	Р		N		S		Т	Т Р		N		S		Т	Р		N		S		T
		N _o	ті	No	т	No	п	ā	N _o	т	No	П	No	п	ā	No	т	No	П	No	П	N _o
Yazd	Stallion	7	14.58	31	64.58	10	20.83	48	1	14.28	1	14.28	5	71.42	7	∞	14.54	32	58.19	15	27.27	55
	Mare	ω	8.57	20	57.14	12	34.28	35	1	8.33	4	33.33	7	58.33	12	4	8.52	34	51.06	19	40.42	47
Kerman	Stallion	∞	36.36	11	50	ω	13.63	22	5	31.25	7	43.75	4	25	16	13	34.21	18	47.36	7	18.42	38
	Mare	ω	27.27	6	54.54	2	18.18	11	6	50	ъ	41.66	1	8.33	12	9	39.13	11	47.82	ω	13.04	23
Total		21	18.10	68	58.62	27	23.27	116	13	27.65	17	36.17	17	36.17	47	34	20.85	85	52.14	44	26.99	163

Table 2. The number and frequency of positive, negative and suspected samples (based on the presence of antibodies against Listeria monocytogenes) out of 163 horses in Kerman and Yazd provinces, Iran

cases > 7 years old) male (12.88%) and 13 (6 cases ≤ 7 years old and 7 cases > 7 years old) mares (7.98%) were sero-positive against listeriosis.

DISCUSSION

Our results showed the total seropositivity for all 163 horses examined in this study was 20.85%. Listeria monocytogenes widely exists in the water, soil, plants, feces and feedstuff such as silage, vegetables and moldy forage. This bacteria is considered as one of the most important sources of infection in both domestic and wild animals and also some birds [16]. The prevalence of listeriosis has been mostly reported during the winter which can be due the fact that silage is frequently fed to animals in this season and also the pregnancy of animals predisposes them to this infectious disease [12]. Listeriosis in farm animals often occurs in three main forms: encephalitis, septicemia and abortion [5]. However, the encephalitis form has a lower occurrence rate

in horses [5,7]. Teruya et al., (1977) conducted a study on equine listeriosis in Brazil using tube agglutination method and reported that examined horses 22.7% of 838 seropositive. Solmaz et al., (2002) carried out a similar study in Turkey using the same method and found 176 positive (86.7%) out of 203 horses [15,17]. Guclu et al., (2007) showed 62 cases among 100 tested horses were sero-positive against L. monocytogenes using the Osebold absorption test. Anti L. monocytogenes antibodies were detected at 1:100, 1:200 and 1:400 titers in 29 (46.7%), 31 (50%) and 2 (3.2%) animals, respectively [20]. The results of the present study are similar to the study conducted by Teruya et al., (1977) and Guclu et al., (2007). However, higher prevalence in the study conducted by Solmaz et al., can be due to factors such as climate variation, feedstuff and health management of the animals.

The results of the current study showed more sero-positivity among older males in contrast with previous study showed there is no relationship between age and sex and incidence of listeriosis. Saqib et al., (2015) reported that 23.5% out of 183 equine serum samples were seropositive and number of seropositive males was higher than females [13]. Results of the present study in Yazd and Kerman provinces showed that the higher anti L. monocytogenes antibody titer found in horses in Kerman is possibly due to moderate climate and also the lower temperature and different management and any other factors or any combination(s). The obtained results of the separate studies in each province showed that the sero-positivity in the old males of Kerman follows the general rule which was mentioned earlier. The results of serological studies have shown that the presence of anti L. monocytogenes antibody may be widespread worldwide, but the absence of definitive clinical diagnosis and data makes it impossible to estimate the true incidence of listeriosis in animals and humans. These results have also revealed that the sero-prevalence of positive antibody titer against Listeria monocytogenes in animals varies widely with species tested, geographic location, season, assay type and the criteria used to define positive results [2,3,4,9,18]

It has been reported that the antigenic relationship between various serotypes of L. monocytogenes and a number of gram-positive gram-negative bacteria (such Staphylococcus aureus, Streptococcus faecalis, Arcanobacter pyogenes, Bacillus subtilis and Escherichia coli) may cause false-positive results in serological tests [10,14]. The latex agglutination test can be considered as an appropriate screening test in the early stages of diagnosis, because mentioned method has a favorable sensitivity and specificity. It is also a fast and cheap method which can be performed in all laboratories.

CONCLUSION

The current study detected anti L. monocytogenes antibody in the horses' population of Kerman and Yazd provinces, Iran. Mentioned results showed that the higher antibody titer found in Kerman province is possibly due to moderate climate and also the lower temperature and different management and any other factors or any combination(s).

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ETHICS

All ethical standards have been respected in this study.

CONFLICT OF INTEREST

There is no conflict of interest.

REFERENCES

- [1] Ajay Kumar V, Latha C, Sunil B, Dhanya V. Comparison of different cultural techniques in isolation of Listeria monocytogenes from various samples. J Foodborne Zoonotic Dis. 2014; 2: 10-4.
- [2] Arda M, Müller HE. Listeriosis in animals. Infeks Derg. 1988; 2: 505-521.
- [3] Bhunia AK. Antibodies to Listeria monocytogenes. Crit Rev Microbiol. 1997; 23: 77-107.
- [4] Blenden DC, Kampelmacher EH, Torres-Anjel MJ. Listeriosis. J Am Vet Med Assoc. 1987; 191: 1546-1551.
- [5] Constable P, Hinchcliff KW, Done S, Gruenberg W. Veterinary medicine. Saunders, 11th ed. 2017.
- [6] Goz Y, Babur C, Aydin A, Kilic S. Seroprevalence of toxoplasmosis, brucellosis and listeriosis in horses in Hakkari, eastern region of Turkey. Revue Med Vet. 2007; 158: 534-539.

- [7] Gray ML, Killinger A. Listeria monocytogenes and listeric infections. Bacteriol Rev. 1966; 30(2): 309.
- [8] Hain T, Chatterjee SS, Ghai R, Kuenne CT, Billion A, Steinweg C, et al. Pathogenomics of Listeria spp. Int J of Med Microbiol. 2007; 297(7): 541-57.
- [9] Low JC, Donachie W. A review of Listeria monocytogenes and listeriosis. Vet J. 1997; 153: 9-29.
- [10] Osebold JW, Aalund O. Interpretation of serum agglutinating antibodies to Listeria monocytogenes by immunoglobulin differentiation. J Infect Dis. 1968; 118: 139-148.
- [11] Pirš T, Zdovc I, Gombač M, Švara T, Juntes P, Vengušt M. Listeria monocytogenes septicaemia in a foal. Sci J Vet Fac Uni Ljubljana. 2005; 49.
- [12] Rudol M, Scherer S. High incidence of Listeria monocytogenes in European red smear cheese. IJFM. 2001; 63 (1-2): 91-101.
- [13] Sagib M, Hussain MH, Sajid MS, Mansoor MK, Asi MN, Fadya A, Al-Kitani Zohaib A, Sial AUR, Muhammad G, Ullah I. Sero- epidemiology of equine toxoplasmosis using a latex agglutination test in the three metropolises of Punjab, Pakistan. Trop Biomed. 2015; 32 (2): 276-285.
- Seeliger HPR, Sulzbacher F. Antigenic relationships between Listeria monocytogenes and Staphylococcus aureus. Can J Microbiol. 1956; 2: 220-231.
- [15] Solmaz H, Akkan HA, Tutuncu M, Karaca M, Ekin IH, Kutlu I. Van ve yöresinde atlarda listeriosis in seroprevalansı. Y Y Ü Vet. Fak Derg. 2002; 13: 62-63.
- [16] Tajbakhsh H. General bacteriology, 1st ed. Tehran University Press. 686-687.
- [17] Teruya JM, Santa Rosa CA, Giorgi W, Yanaguita RM. Serological study of listeriosis in domestic animals in Sao PauloBrazil. Int J Zoonoses. 1977; 4: 21-24.
- [18] Vazquez-Boland JA, Kuhn M, Berche P, Chakraborty T, Dominguez-Bernal G, Goebel W, Gonzalez-Zorn B, Wehland J, Kreft J. Listeria and molecular virulence pathogenesis determinants. Clin Microbiol Rev. 2001; 14: 584-640.
- [19] Wesley IV. Listeriosis in animals. Food Sci Tech New York - Marcel Dekker. 1999: 39-74.
- [20] Zeynep Gucl H, Zafer Karaeka K, Babur C, Kilic S. The Seroprevalence of Listeria monocytogenes in sport horses breed in Ankara Province. Turk J Vet Anim Sci. 2007; 31(4): 271-273.