

## 【Abstract】

Progress in the epidemics of Tibetan sheep and yaks in Qinghai Province in recent years.

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To summary what kinds of epidemics are happening on Tibetan sheep and yaks in Qinghai Province. Such as, the diarrhea of sheep and yaks caused by *Escherichia coli* and *Salmonella*; the diarrhea and deaths caused by multiple clostridial diseases; the infectious pleuropneumonia caused by *Mycoplasma* spp. in cattle and sheep; the prevention and treatment of foot rot disease and the purulent necrosis of internal organs caused by *Fusobacterium necrophorum*. The viral diarrhea caused by bovine viral diarrhea virus, rotaviruses and coronaviruses, bovine papillomavirus infections in yaks and cattle; infectious pustular dermatitis virus disease in sheep. Diagnosis and control of blood protozoan diseases in horses, cattle and sheep; the diarrhea caused by coccidia and nematodes in yaks; diagnosis and control of cestode and trematode diseases in yaks, cattle and sheep.

Optimization of single-tube nested PCR for the detection of *Echinococcus* spp.

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*Echinococcosis* is a serious zoonotic life-threatening parasitic disease caused by metacestodes of *Echinococcus* spp., and appropriate sensitive diagnosis and genotyping techniques are required to detect infections and study the genetic characterization of *Echinococcus* spp. isolates. In this study, a single-tube nested PCR (STNPCR) method was developed and evaluated for the detection of *Echinococcus* spp. DNA based on the COI gene. STNPCR was 100 times more sensitive than conventional PCR and showed the same sensitivity to common nested PCR (NPCR); the limit of detection of the developed STNPCR method was estimated to be 10 copies/ $\mu$  L of the recombinant standard plasmids of *Echinococcus* spp. COI gene. The STNPCR method was suitable for epidemiological investigations and characteristic genetic studies of *Echinococcus* spp. tissue samples. The STNPCR method can effectively amplify low concentrations of genomic DNA from calcification samples and cyst residues infected with *Echinococcus* spp. Subsequently, the sequences of positive PCR products were obtained, which were useful for haplotype analysis, genetic diversity, and evolution studies of *Echinococcus* spp., and understanding of *Echinococcus* spp. dissemination and transmission among the hosts.