

License Agreement of Veterinary Drug and Feed Products

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Abstract

Veterinary drugs and feed products are vital for ensuring animal health, productivity, and food safety, but their use requires strict regulatory control. A license agreement serves as a legal authorization issued by a competent authority that permits the manufacture, import, distribution, and sale of veterinary drugs and feed products under defined conditions. This article highlights the purpose of licensing in safeguarding animal and public health, preventing substandard products, and promoting responsible use of veterinary medicines. It discusses the major types of licenses, key components of a license agreement, and the legal and regulatory frameworks governing these products at national and international levels. Overall, license agreements play a crucial role in maintaining product quality, ensuring regulatory compliance, and supporting sustainable livestock production systems.

Keywords: Veterinary drugs; Animal feed products; License agreement; Quality control; Good Manufacturing Practices (GMP); Animal health; Food safety

Introduction

Veterinary drugs and feed products play a crucial role in animal health, food safety, and public health. To ensure their safe manufacture, distribution, and use, governments regulate these products through licensing systems. A license agreement for veterinary drugs and feed products is a legal authorization granted by a competent authority that allows an individual or organization to manufacture, import, distribute, or sell these products under specified conditions.

Meaning of License Agreement

A license agreement is a legally binding document between a regulatory authority and a license holder. It defines the rights, responsibilities, and obligations related to the production and marketing of veterinary drugs and animal feed products. The agreement ensures compliance with established laws, standards, and quality requirements.

Purpose of Licensing

The primary objectives of licensing veterinary drugs and feed products include:

- Ensuring animal health and welfare
- Protecting human health by preventing drug residues in animal-derived foods
- Maintaining product quality, safety, and efficacy
- Preventing the circulation of substandard or counterfeit products
- Regulating the use of antibiotics and growth promoters

Types of Licenses

License agreements may be issued for different activities, such as:

1. **Manufacturing License** – Permits the production of veterinary drugs or feed products.
2. **Import License** – Authorizes the importation of approved products from foreign manufacturers.
3. **Distribution or Wholesale License** – Allows bulk supply to retailers or institutions.
4. **Retail License** – Permits the sale of products to end users.
5. **Feed Additive License** – Regulates products added to animal feed for nutritional or medicinal purposes.

Key Components of a License Agreement

1. **Identification of Parties**- Includes details of the licensing authority and the license holder.
2. **Scope of License**- Specifies the products covered, permitted activities, and geographical area of operation.
3. **Regulatory Compliance**- Requires adherence to Good Manufacturing Practices (GMP), Good Distribution Practices (GDP), and labelling regulations.
4. **Quality Control and Inspection**- Grants authority to conduct inspections, sampling, and audits to ensure compliance.
5. **Validity and Renewal**- Defines the duration of the license and procedures for renewal.
6. **Restrictions and Conditions**- Includes limitations on advertising, prescription-only products, and usage instructions.
7. **Penalties and Cancellation** - Outlines penalties for non-compliance, including suspension or cancellation of the license.

Legal and Regulatory Framework

Veterinary drug and feed product licenses are governed by national laws and

regulations, often aligned with international standards such as those from the World Organization for Animal Health (WOAH), Codex Alimentarius, and relevant pharmaceutical guidelines. Regulatory authorities are responsible for evaluating product safety, efficacy, and quality before granting approval.

Importance of License Agreements

License agreements ensure that veterinary drugs and feed products entering the market are safe, effective, and of consistent quality. They also promote responsible use of veterinary medicines, reduce antimicrobial resistance, and protect consumers from health risks associated with contaminated or improperly used products. Reducing mortality in scavenging systems through better housing, nutrition, and hygiene.

Conclusion

A license agreement for veterinary drug and feed products is an essential regulatory tool that safeguards animal health, public health, and the integrity of the livestock industry. By enforcing legal standards and quality requirements, licensing systems help maintain trust in veterinary products and ensure sustainable animal production practices.

References

- World Organisation for Animal Health (WOAH). (2022). *Terrestrial Animal Health Code*. WOAH, Paris.
- Food and Agriculture Organization of the United Nations (FAO). (2019). *Guide to Good Hygiene Practices for Animal Feed*. FAO Animal Production and Health Guidelines.
- Codex Alimentarius Commission. (2020). *Code of Practice on Good Animal Feeding (CAC/RCP 54-2004)*. FAO/WHO.
- European Medicines Agency (EMA). (2021). *Veterinary medicinal products: Regulation and control*. EMA, European Union.
- World Health Organization (WHO). (2017). *Guidelines on Use of Medically Important Antimicrobials in Food-Producing Animals*. WHO Press.
- United States Food and Drug Administration (FDA). (2022). *Animal Drug User Fee Act and Feed Regulations*. FDA Center for Veterinary Medicine.