Registration of veterinary products manufacturing company

About the Service

Through this service, approved the registration of foreign veterinary products manufacturers through a licensed warehouse

Service Process

Required Documents

- 1. Provide a file that includes the following basic information: -Type and activity of the company. -Number of factories owned by the company and its addresses.
- -The company's relationship with each of these factories and the extent of their legal, technical and commercial responsibility.
- 2. A power of attorney letter from the factory stating that the agency has been granted to the authorized agent in the United Arab Emirates
- 3.A valid certificate of registration of the company in the country of origin, including the date of establishment,
- stating clearly that the veterinary preparations produced by the company are authorized to be sold freely in the country of origin.
- 4. A valid certificate issued by the concerned authorities in the country of the manufacturer proving its commitment to the principles of good practice in pharmaceutical manufacturing (GMP),
- including evidence that the competent authorities have conducted periodic inspections on it.
- 5. In case that the manufacturer site from an origin country that is not included in the list of countries mentioned in the terms and conditions,
- a certificate of registration for the manufacturer site in at least two of these countries must be submitted.
- 6. A list of the preparations produced by the factory. 7. Provide a Site Master File.

Service completion duration

- 20 Days

Service channels

Ministry of Climate change & Environment Website

MOCCAE

Service Code

122-001-025-000