



Summary of Studies Supporting USDA Product Licensure

Establishment Name	Merial, Inc.
USDA Vet Biologics Establishment Number	298
Product Code	1601.11
True Name	Fowl Laryngotracheitis Vaccine, Modified Live Virus
Tradename(s) / Distributor or Subsidiary (if different from manufacturer)	BIO LARYNGO - No distributor specified Gallivac LT - IFT Corp (Egypt) Gallivac LT - Merial (France) Gallivac LT - No distributor specified LT Blen - Merial Mexico S.A. DE C.V. LT Blen - No distributor specified LT Blen - Saadat International
Date of Compilation Summary	September 21, 2018

Disclaimer: Do not use the following studies to compare one product to another. Slight differences in study design and execution can render the comparisons meaningless.

Study Type	Efficacy
Pertaining to	Infectious laryngotracheitis virus
Study Purpose	Demonstrate efficacy against Fowl Laryngotracheitis
Product Administration	Eyedrop
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	June 30, 1995

Study Type	Efficacy
Pertaining to	Infectious laryngotracheitis virus
Study Purpose	Demonstrate efficacy against Fowl Laryngotracheitis
Product Administration	Eyedrop
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. Study data, however, are no longer available.
USDA Approval Date	January 6, 1976

Study Type	Efficacy
Pertaining to	Infectious laryngotracheitis virus
Study Purpose	Demonstrate efficacy against Fowl Laryngotracheitis
Product Administration	Eyedrop
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. Study data, however, are no longer available.
USDA Approval Date	January 10, 1979

Study Type	Efficacy
Pertaining to	Infectious laryngotracheitis virus
Study Purpose	Demonstrate efficacy against Fowl Laryngotracheitis
Product Administration	Eyedrop
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	April 24, 1997

Study Type	Efficacy
Pertaining to	Infectious laryngotracheitis virus
Study Purpose	Demonstrate efficacy against Fowl Laryngotracheitis
Product Administration	Drinking water
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	April 24, 1997

Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under field conditions
Product Administration	Eyedrop
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. Study data, however, are no longer available.
USDA Approval Date	April 9, 1984

Study Type	Safety
Pertaining to	ALL
Study Purpose	Demonstrate safety under field conditions
Product Administration	Drinking water
Study Animals	Chickens
Challenge Description	
Interval observed after challenge	
Results	Study data were evaluated by USDA-APHIS prior to product licensure and met regulatory standards for acceptance at the time of submission. No data are published because this study was submitted to USDA-APHIS prior to January 1, 2007, and APHIS only requires publication of data submitted after that date.
USDA Approval Date	March 9, 2000