Dosing	Adequan i.m.		
Schedule For:	polysulfated glycosaminoglycan		

Ē	Administer 1 (500 mg) dose	Every 4 days	For 7 treatments
DOSE 1	DATE		TIME
DOSE 2			
DOSE 3			
DOSE 4			
DOSE 5			
DOSE 3			
DOSE 6		I	
_			
DOSE 7			



Dr.		
Office #		
Cell #		
Follow-up appointment	DATE	TIME
Notes:		

Please see accompanying Full Prescribing Information.



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SINGLE DOSE

Solution 500 mg/5 mL

For Intramuscular Use In Horses



Rev 9/2021

MG #44455

CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION: Each 5 millitiers of Adequan® i.m. contains 500 mg of Polysulfated

DESCRIPTION: Each 5 millifiers of Adequan® i.m. contains 500 mg of Polysulfated Glycosaminoglycan (PSGAG) and Water for Injection q.s. Sodium Hydroxide and/or Hydroxhloric Acid added when necessary to adjust pH. Sodium Chloride may be added to adjust tonicity.

PHARMACOLOGY: Polysullated Glycosaminoglycan is chemically similar to the glycosaminoglycans in articular cartiage matrix. PSAGA is a potent proteolytic enzyme inhibitor and diminishes or reverses the pathologic processes of traumatic or degenerative joint disease which result in a net loss of cartifage matrix components. PSGAG improves joint function by reducing syrroxial fluid protein components. PSGAG improves joint function by reducing syrroxial fluid protein components. PSGAG improves joint function by reducing syrroxial fluid protein components. PSGAG improves joint function by reducing syrroxial fluid protein components. PSGAG improves joint function protein protein components. PSGAG improves joint function protein prote

INDICATIONS: Adequan® i.m. is recommended for the intramuscular treatment of non-infectious degenerative and/or traumatic joint dysfunction and associated lameness of the careal and hock joints in horses.

DOSAGE AND ADMINISTRATION: The recommended dose of Adequan® i.m. in horses is 500 mg every 4 days for 28 days intramuscularly. The injection site must be thoroughly cleansed prior to injection. Do not mix Adequan® i.m. with other drugs or solvents.

CONTRAINDICATIONS: There are no known contraindications to the use of intramuscular Polysulfated Glycosaminoglycan.

WARNINGS: Do not use in horses intended for human consumption. Not for use in humans. Keep this and all medications out of the reach of children.

PRECAUTIONS: The safe use of Adequan® i.m. in horses used for breeding

purposes, during pregnancy, or in lactating marce has not been evaluated.

ANIMAL SAFETY: Toxicity studies were conducted in horses. Doses as high as 2.000 mg were administered intramacularly to 6 horses livite a week for 12 weeks. The properties of the properties

evidence of toxicity.

STORAGE CONDITIONS: Store at 20*-25*C (68*-77*F); (See USP Controlled Room Temperature). Discard unused portion.

Dispose of spent needles in accordance with all federal, state and local environmental laws.

HOW SUPPLIED: Adequan® i.m. solution, 500 mg/5 mL (100 mg/mL) in a 5 mL single dose glass vial.

NDC 10797-995-70 5 mL Single Dose Vials Packaged 7 vials per box
AMERICAN REGENT, INC.
ANIMAL HEALTH INS9501

Approved by FDA under NADA # 140-901

Shirley, NY 11967

(1-888-354-4857)



Solution 100 mg/mL in a 50 mL Preserved Multi-Dose Vial For Intramuscular Use in Horses Only. Not for Intra-Articular Use.



CAUTION: Federal law restricts this drug to use by or on the order of a licensed veterinarian.

DESCRIPTION: Each mL contains Polysulfated Glycosaminoglycan (PSGAG) 100 mg, Benzyl Alcohol

0.9% viv as a preservative, and Water for Injection qs. Sodium Hydroxide and/or Hydroxhloric Acid added when necessary to adjust pH. The solution is clear, coloriess to slightly yellow. PHARMACOLOGY: Polysulfated Glycosaminoglycan is chemically similar to the

glycosaminoglycans in articular cartilage matrix. PSGAG is a potent proteolytic enzyme inhibitor and diminishes or revenues the pathologic processes of traumatic or degenerative joint disease which result in a net loss of cartillage matrix components. PSGAG improves joint function by reducing synovial fluid protein levels and increasing synovial fluid hyalurenic acid concentration in traumatized equine carpal and hock joints.

INDICATIONS: Adequare I.m. Multi-Dose is recommended for the intramuscular treatment of non-infectious degenerative and/or traumatic joint dysfunction and associated lameness of the carpal and hock joints in horses.

DOSAGE AND ADMINISTRATION: Practice aseptic techniques in withdrawing each dose to decrease the possibility of post-injection bacterial infections. Adequately clean and disinfect the stopper prior to entry with a sterile needle and syringe. Use only sterile

needles, and use each needle only once.

The vial stopper may be punctured a maximum of 10 times.

The recommended dose of Adequan® I.m. Multi-Dose in horses is 500 mg every 4 days for 28 days intramuscularly. The injection site must be thoroughly cleansed prior to injection. Do not mix Adequan® I.m. Multi-Dose with other drugs or solvents.

CONTRAINDICATIONS: There are no known contraindications to the use of inframuscular Polysulfated Glycosaminoglycan.

WARNINGS: Do not use in horses intended for human consumption. Not for use in humans. Keep this and all medications out of the reach of children.

PRECAUTIONS: The safe use of Adequan® i.m. Multi-Dose in horses used for breeding purposes, during pregnancy, or in lactating mares has not been evaluated.

SAFETY AND EFFICACY: Safety and efficacy studies utilizing Adequan® i.m. Multi-Dose were not performed. Adequan® i.m. Multi-Dose was approved based on the conclusion that the safety and effectiveness of Adequan® i.m. Multi-Dose will not differ from that demonstrated for the original formulation of Adequan® i.m.

AMMAL SAFETY: Animal Safety studies utilizing Adequant* In. Multi-Doos were not performed. Safety studies were conclusted in horses using the single does formulation. Does as high as 2,500 mg were administrated internanciality to 6 horses believe a week for 12 weeks. This dosage is 5 femile the normalized bodge and 21 mem de ner commercial the impactor regiment with a femile studies. The safety of the sametal had any difficial or liberative yellowed or floatify.

(See USP Controlled Room Temperature). Avoid prolonged exposure to temperatures ≥ 40°C (104°F). Use within 28 days of first puncture and puncture a maximum of 10 times. Dispose of spent needles in accordance with all federal, state and local environmental laws.

neediss in accordance with all rederal, state and local environmental laws.

HOW SUPPLIED: Adequan[®] i.m. Multi-Dose solution, 5,000 mg/50 mL (100 mg/mL) in 50 mL multi-dose glass vials.

Uses graze value.

50 mL Multi-Dose Vials

Packaged 1 vial per box

AMERICAN REGENT, INC.

Mode in U.S.A.

ANNIAL HEALTH.

1N59

Shirley, NY 11967

Rev, 902021

MG #44433

MG #44437

Approved by FDA under NADA # 140-901