Antiherpetic Drugs in Equine Medicine



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KEYWORDS

- Equine Herpesvirus EHV-1 EHV-5 Antiviral Nucleoside analogs Acyclovir
- Valacyclovir

KEY POINTS

- Equine herpesvirus (EHV)-1 differentially affects different classes of horses but can be particularly devastating to neonatal foals, pregnant mares, and adult performance horses.
- Recent high-profile outbreaks of EHV myeloencephalopathy (EHM) have had an impact on the equine industry and stimulated interest in antiherpetic interventions.
- Several antiherpetic drugs that are active against EHV-1 in the laboratory have been investigated both clinically and experimentally in horses and foals.
- The recent association between equine pulmonary multinodular fibrosis (EMPF) and EHV-5 has resulted in the empiric use of antiherpetic drugs for this condition.
- Little is currently known about resistance patterns of EHV-1 or EHV-5 for antiherpetic drugs in horses.

INTRODUCTION

Herpesviruses comprise a large, ancient family of viruses that infect most if not all vertebrates and even lower organisms.¹ The herpesviruses are specialists that have coevolved with their host species over many years, evading multiple steps of immunity.² Perhaps as a consequence of this immune evasion, current equine vaccines can decrease the replication and clinical signs associated with several herpesvirus infections but cannot completely prevent infection.^{3–6} Currently, 9 herpesviruses have been described from equids and are appropriately named EHV-1 through EHV-9.¹ These 9 herpesviruses belong to 2 separate subfamilies, either the Alphaherpesvirinae (EHV-1, EHV-3, EHV-4, EHV-6, EHV-8, and EHV-9) or the Gammaherpesvirinae (EHV-2, EHV-5, and EHV-7). Several of these viruses, such as EHV-1, EHV-2, EHV-3, EHV-4, and EHV-5, are associated with clinical disease in horses. Whereas in vitro antiherpetic drug susceptibility testing (Table 1) has been performed with EHV-1, EHV-3,

The author has nothing to disclose.

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Virus EHV-1	Drug Acyclovir	Virus Strain Rac-H, H-45 Kentucky D Quai Hais 94P247, 97P70, and 99P96; 97P82, 99P136,	Cell Type PK13 PRK R13	Half-maximal Inhibitory Concentration (μg/mL) 0.45 7 2.6	Reference Rollinson & White, 85 1983 De Clerq et al, 109 1986
		Rac-H, H-45 Kentucky D Quai Hais 94P247, 97P70, and	PK13 PRK R13	0.45 7	Rollinson & White, 85 1983
EHV-I	Acyclovir	Kentucky D Quai Hais 94P247, 97P70, and	PRK R13	7	·
		Quai Hais 94P247, 97P70, and	R13	=	De Clerq et al, 1986
		94P247, 97P70, and			
					Boyd et al, ⁶⁰ 1987
		and 03P37	EEL	1.7–3	Garre et al, ⁸⁴ 2007
		89c25	RK13	2.3-3.1	Azab et al, ⁸² 2010
		T953 (Findlay OH 2003)	ELF	11.4 ± 1.5	(Maxwell LK, Bentz BG, Gilliam LL, et al. Efficacy of the early administration of valacyclovir for the therapy of neuropathogenic EHV-1 in horses. Submitted for publication.)
		T953 (Findlay OH 2003)	PBMC	0.8	(Maxwell LK, Bentz BG, Gilliam LL, et al. Efficacy of the early administration of valacyclovir for the therapy of neuropathogenic EHV-1 in horses. Submitted for publication.)
	Penciclovir	Quai Hais	R13	1.6	Boyd et al, ⁶⁰ 1987
		AB4	RK13	1.3-1.9	de la Fuente et al, ⁷⁶ 1992
		T953 (Findlay OH 2003)	ELF	4.8 ± 0.7	(Maxwell LK, Bentz BG, Gilliam LL, et al. Efficacy of the early administration of valacyclovir for the therapy of neuropathogenic EHV-1 in horses. Submitted for publication.)
	Ganciclovir	Rac-H, H-45	PK13	0.02	Rollinson and White, 85 1983
		Kentucky D	E Derm	0.03	Smith et al, 110 1983
		Rac-H, H-45	RK13	0.02-0.1	Rollinson, 111 1987
		94P247, 97P70, and 99P96; 97P82, 99P136, and 03P37	EEL	0.1–4	Garre et al, ⁸⁴ 2007
		89c25	RK13	0.1-0.7	Azab et al, ⁸² 2010

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