

The Current State of Oncolytic Herpes Simplex Virus for Glioblastoma Treatment

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Abstract: Glioblastoma (GBM) is a lethal primary malignant brain tumor with no current effective treatments. The recent emergence of immuno-virotherapy and FDA approval of T-VEC have generated a great expectation towards oncolytic herpes simplex viruses (oHSVs) as a promising treatment option for GBM. Since the generation and testing of the first genetically engineered oHSV in glioma in the early 1990s, oHSV-based therapies have shown a long way of great progress in terms of anti-GBM efficacy and safety, both preclinically and clinically. Here, we revisit the literature to understand the recent advancement of oHSV in the treatment of GBM. In addition, we discuss current obstacles to oHSV-based therapies and possible strategies to overcome these pitfalls.

Keywords: glioblastoma, oncolytic herpes simplex virus, armed oHSV, re-targeted oHSV, combination therapy

Introduction

Oncolytic viruses (OVs)^{1,2} such as oncolytic herpes simplex virus (oHSV) are an emerging treatment strategy for glioblastoma (GBM).^{3–10} GBM is the most common primary malignant brain tumor that is almost always lethal and has no effective treatments.^{11,12} OVs selectively replicate in and kill cancer cells and induce immunogenic cell death while stimulating anti-tumor immunity (ie, in situ vaccine effect).^{13,14} Among all OVs, oHSV is the furthest along in the clinic, with an oHSV talimogene laherparepvec (T-VEC) approved for use in advanced melanoma.¹⁵ The highly stable genome, potent cytolytic capability, convenience in genome engineering, and availability of effective anti-herpetic drugs to treat adverse reactions make oHSV an attractive class of anti-cancer therapeutic.¹⁶ The first genetically modified oHSV for glioma treatment was reported in 1991.¹⁷ Since then, many oHSVs such as unarmed – genetically modified oHSVs but not armed with transgene, re-targeted – genetically altered for tumor receptor-specific viral entry, and armed – engineered to express therapeutic transgene variants, etc., have been developed and tested in GBM preclinically, either alone or in combination with other anti-cancer agents (Table 1–3, Figure 1–3). Some of them are currently under clinical trial evaluation and early oHSV trials confirmed their safety and signs of efficacy in GBM patients (Table 4, figure 1). Despite significant therapeutic advancement, various challenges to oHSV therapy still currently exist (such as suboptimal viral delivery, inefficient viral entry to cancer cells, limited oHSV replication/spread in the tumor microenvironment (TME), host's anti-viral response, etc.)

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Table I List of Unarmed oHSVs for GBM Treatment

oHSV	Strain	Genomic Modification	Tumor Model	PFU (Doses)	Results and References
dlsptk	F	-TK	U87	Subcutaneous: 5x10 ⁶ (1 st dose) and 1x10 ⁷ (2 nd dose)	Virus treated tumors were smaller than control tumors at day 28. ¹⁷
				Intracerebral: 1x10 ⁵ (1 dose)	Treatment with dlsptk significantly extended survival compared to mock group. ¹⁷
hrR3	F	ΔICP6, +LacZ	U87	5x10 ⁶ (1 dose)	The mean tumor growth rate was significantly inhibited by hrR3 treatment compared to control treatment. ²¹⁰
HSV1716	17+	-/-γ34.5	NT2	-	HSV1716 treated mice survived significantly longer than saline-treated group (25 weeks vs 9 weeks). ²⁴
			Toxicity study in naïve mice	LD ₅₀ 7x10 ⁶ (1 dose)	Non-neurovirulent in mice following intracerebral injection. ²¹
HSV3616	F	-/-γ34.5	U87	2x10 ⁷ (1 dose)	R3616 treatment led to tumor eradication by day 60 in 3/21 animals versus all mice in control group treated with buffer solution died by day 30. ²⁵
G207	F	-/-γ34.5, ΔICP6, +LacZ	U87	5x10 ⁷ (1 dose)	Mean tumor volume was significantly reduced by G207 treatment compared to control treatment. ²⁶
G47Δ	F	-/-γ34.5, ΔICP6, ΔICP47, +LacZ	U87	5x10 ⁶ (2 doses)	Average tumor volume was significantly smaller in G47Δ group compared to G207 and control groups. ³¹
rQNestin34.5v.2	F	-γ34.5, ΔICP6, γ34.5 driven by nestin enhancer/promoter, +EGFP	U87ΔEGFR	3.5x10 ⁷ (1 dose)	rQNestin34.5v.2 significantly extended survival compared to mock group. ³⁵
HSVQ1	F	-γ34.5, ΔICP6, +EGFP			
C130	AD169/ F	-/-γ34.5, HCMV-TRSI	U87	5x10 ⁵ (1 dose)	C130 or C134 significantly extended survival compared to both doses of C101. ³⁹
C134	AD169/ F	-/-γ34.5, HCMV-IRSI			
C101	F	-/-γ34.5, +EGFP			
MG18L	F	-US3, ΔICP6, +LacZ	BT74	2x10 ⁶ (1 dose)	MG18L significantly extended survival compared to mock group. ¹⁸¹

Abbreviations: PFU, plaque forming unit; (-), single deletion; (-/-), diploid deletion; Δ, mutation/inactivation; (+), insertion.

that compromise oHSV-mediated anti-tumor efficacy (Figure 4).^{4,18} With the advance in biotechnology and immuno-oncology, many multimodal strategies are budding out for GBM to overcome existing challenges (Figure 4).^{4,18} In this review, we will first highlight the recent preclinical and clinical studies

conducted in GBM using oHSVs (unarmed, re-targeted, or armed), either as monotherapy or in combination. We will then provide a short overview of the current pitfalls of oHSV-based therapies and outline potential strategies to overcome the existing treatment hurdles.

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Table 2 List of Re-Targeted oHSVs for GBM Treatment

oHSV	Strain	Modifications	GBM Model	Unit (Doses)	Results and References
R-LM113	F	Nectin 1/HVEM-de-targeted, hHER2-re-targeted	mGBM-HER2	3x10 ⁵ pfu (1 dose)	R-LM113 significantly extended median survival compared to UV-inactivated R-LM113 group in both models. ^{46,211}
			BALB/c-HGG-HER2	5.5x10 ⁵ pfu (1 dose)	
R-115	F	Nectin 1/HVEM de-targeted, hHER2-re-targeted, +mIL-12	mHGG ^{p_{def}} -hHER2	2x10 ⁶ pfu (1 dose)	R-115 treatment led to tumor eradication in 30% of animals and these long-term survivors resisted tumor re-challenge. The efficacy of R-115 was associated with production of antibodies against mHGG ^{p_{def}} -hHER2 cells in 83% treated animals compared to 40% in R-LM113 (IL-12 unarmed)-treated or 0% in mock-treated mice. ⁴⁷
KNE	KOS	EGFR-re-targeted	GBM30	8.6x10 ⁸ gc (1 dose)	All PBS-treated mice died within 35 days (median survival 23 days) of tumor cell implantation. In contrast, 63–73% of mice treated with KNE survived for 90 days. ⁴²
KGE-4:T124	KOS	EGFR-re-targeted, +4 tandem miR-124 target sites in ICP4	GBM30	1.8x10 ⁸ gc (1 dose)	The median survival of PBS-treated group was 21.5 days, which was significantly shorter than the median survival (85.5 days) of KGE-4:T124-treated group. The median survival of KGE-4:T124 was no different than control KGE virus. ⁴⁹
KMMP9	KOS	EGFR/EGFRvIII-re-targeted, +4 tandem miR-124 target sites in ICP4, +MMP9	GBM30	1x10 ⁴ pfu (1 dose)	KMMP9 treatment resulted in 70% long-term survivors compared to 0% in KGW (MMP9-unarmed)-treated mice. ⁵¹

Abbreviation: PFU, plaque-forming unit; gc, genome copy; (+), insertion.

Unarmed oHSVs for GBM

The HSV genome comprises several essential and non-essential genes that can be modified to ensure selective replication in cancer cells while sparing normal cells.¹⁶ The frequently mutated gene to engineer HSV is γ 34.5 – a pivotal determinant for viral neuropathogenicity. When HSV infects normal cells, protein kinase R (PKR) phosphorylates eukaryotic translation initiation factor- α (eIF2 α) and prevents viral protein synthesis. HSV with intact γ 34.5 complexes with protein phosphatase 1 alpha (PP1 α) to dephosphorylate eIF2 α and overcomes anti-viral responses in healthy cells.¹⁹ Deletion of γ 34.5 in HSV results in no eIF2 α dephosphorylation, no viral protein translation, and thus, no viral replication in healthy cells.²⁰ On the other hand, γ 34.5-deleted oHSV freely replicates in cancer cells due to the defective anti-viral PKR-eIF2 α pathway. HSV1716 and R3616 are the first generation oHSVs with γ 34.5 diploid deletion.^{21,22} Both were safer than their wild-type parent viruses^{21–23} and demonstrated superior anti-tumor efficacy than the control

treatment.^{24,25} Later, the second generation of oHSV (eg, G207) was engineered, with diploid deletion of γ 34.5 combined with inactivation of ICP6 – a larger subunit of ribonucleotide reductase (RR).²⁶ G207 effectively lysed human glioma cells, controlled tumor growth, and prolonged survival of mice bearing orthotopic and subcutaneous brain tumor xenografts.²⁶ Although γ 34.5 deletion improves safety, loss of γ 34.5 genes severely compromised virus replication in GBM stem cells (GSCs),²⁷ which are oftentimes associated with resistance to standard therapies and immunosuppression.^{28–30} To counter this issue, the next generation of G207-derivative virus, G47 Δ , was created with the deletion of an additional gene, ICP47.³¹ ICP47 deletion complements the loss of γ 34.5 and restores oHSV replication in GSCs.³² G47 Δ was safe and significantly more efficacious than its parent G207 in controlling tumor growth in vivo in both immunodeficient and immunocompetent brain tumor models.³¹ Intratumoral G47 Δ -Empty (G47 Δ -E, a derivative of G47 Δ without transgene expression) treatment resulted in

Table 3 List of Armed oHSVs for GBM Treatment

Armed Transgene	oHSV	Strain	Genomic Modification	GBM Model	PFU (Doses)	Results and References
Cytokines	G47Δ-mIL12	F	ΔICP6, -/-γ34.5, ΔICP47, +LacZ, +mIL-12	005 GSC	5×10 ⁵ (2 doses)	G47Δ-mIL12 significantly extended survival with 10% mice surviving long term compared to 0% in control virus (G47Δ-Empty) group. ³³
	M002	F	-/-γ34.5, +mIL-12	Patient's GBM tumors	1×10 ⁷ (1 dose)	- Pediatric GBM samples with high nectin-1 expression: M002 treatment led to tumor eradication in 8/10 animals compared to 3/10 animals in the G207 group. ⁵⁸ - Pediatric GBM samples with low nectin-1 expression and adult GBM samples: M002 treatment showed trend of improved median survival versus G207 treatment. ⁵⁸
	oHSV-Fit3-L	F	ΔICP6, -/-γ34.5, ΔICP47, +LacZ, +Fit3-L	CT-2A	2×10 ⁶ (1 dose)	Treatment with G47Δ-Fit3L led to 40% long-term survivors compared to 0% in G47Δ-Empty treatment. ⁶⁰
	oHSV-TRAIL	F	ΔICP6, -/-γ34.5, ΔICP47, +LacZ, +TRAIL	LN229-FmC GBM	2×10 ⁶ (2 doses)	The median survival of mice treated with oHSV-TRAIL was 69 days, which was significantly longer than PBS group (50.5 days) or unarmed oHSV group (49 days). ⁶¹
					GSC23-FmC - 2×10 ⁶ (2 doses) and GSC31 2×10 ⁶ (1 dose)	- GSC23-FmC: The median survival of mock group (44 days) was significantly lower than oHSV-TRAIL (55 days). ⁶² - GSC31: oHSV-TRAIL significantly extended median survival (81 days) with 40% long-term survivors compared to 0% (median survival - 26 days) in the mock mock group. ⁶²
					GSC23-FmC and GSC31	

Angiogenic inhibitors	G47Δ-mAngio	F	ΔICP6, -/-γ34.5, ΔICP47, +LacZ, +mAngio	U87	5 × 10 ⁶ (2 doses)	- Subcutaneous U87 model: G47Δ-mAngio delayed tumor growth by 4 days compared to G47Δ-E. ⁶⁷ - Intracranial U87 model: Treatment with G47Δ-mAngio resulted in 40% of mice surviving over 150 days and significantly prolonged animal survival compared to G47Δ-E. ⁶⁷
						The median survival of G47Δ-mAngio was 66 days, which was significantly longer than G47Δ-Empty-treated group (53 days). ⁵⁷
				MGG4	1 × 10 ⁶ (1 dose)	The median survival of G47Δ-mAngio was 113 days, which was significantly longer than G47Δ-Empty group (98 days). ⁵⁷
	VAE	F	ΔICP6, -/-γ34.5, +endo-angio	Human stable GSC lines	Subcutaneous - 1 × 10 ⁷ (1 dose) and intracranial 2 × 10 ⁶ (1 dose)	Mice treated with VAE showed significant median survival extension compared with mice treated with control r-HSV (ΔICP6, -/-γ34.5) or Endostar. ⁶⁵
	RAMBO	F	ΔICP6, -/-γ34.5, +Vstat120	U87ΔEGFR	1 × 10 ⁵ (1 dose)	The median survival of RAMBO was 28 days, which was statistically longer PBS-treated group (17 days). ⁶⁶
				MGG23	1 × 10 ⁵ (1 dose)	The median survival of RAMBO was 65 days, which was significantly longer than mice treated with PBS. ⁶⁶
				U87ΔEGFR	0.5 × 10 ⁵ (1 dose)	The median survival of RAMBO was 29 days, which was statistically longer than 19 days in PBS-treated group. ¹⁴⁷
GADD34	NG34	F	ΔICP6, -/-γ34.5, +GADD34 driven by nestin enhancer/promoter	U87ΔEGFR-RliFluc and G35 and GL26IN4	2 × 10 ⁵ (1 dose)	NG34 was shown to produce therapeutic efficacy as potent as rQnestin34.5 but with better tolerability. ⁷²
Anti-PD-1	NG34scFvPD-I	F	ΔICP6, -/-γ34.5, +GADD34 driven by nestin enhancer/promoter, +scFvPD-I	GL26IN4	1.5 × 10 ⁶ (1 dose)	NG34scFvPD-I treatment doubled the median survival time compared to non-anti-PD-1 expressing NG34 virus (69 vs 36.5 days). ⁷⁹
				CT-2A	1.5 × 10 ⁶ (1 dose)	Neither NG34scFvPD-I nor NG34 led to a significant increase in median survival compared to mock group. However, 2/12 mice treated with NG34scFvPD-I survived for 120 days. ⁷⁹
CDH-I	Ov-CDH-I	F	ΔICP6, -/-γ34.5, +CDH-I	GBM30, GL26IN4 and U87ΔEGFR	2 × 10 ⁵ (1 dose)	Ov-CDH-I significantly prolonged median survival compared to unarmored Ov-Q1 in all three models. ⁸⁴

(Continued)

Table 3 (Continued).

Armed Transgene	oHSV	Strain	Genomic Modification	GBM Model	PFU (Doses)	Results and References
Chondroitinase ABC	OV-Chase	F	Δ ICP6, $-/\gamma$ 34.5, +Chase	Gli36 Δ EGFR-H2B-RFP and U87 Δ EGFR	Subcutaneous -10^6 v.p. (6-7 doses), intracranial -3×10^5 v.p. (1 dose)	In all models, treatment with OV-Chase significantly controlled tumor growth, resulted in extended survival compared to mock and unarmed oHSV. ⁸⁵
	OV-ChaseM	F	Δ ICP6, $-/\gamma$ 34.5, +ChaseM	GBM30	3×10^5 v.p. (1 dose)	The median survival of OV-ChaseM treated group was 33 days versus 22 days in group receiving PBS. ⁸⁸
PTEN α	HSV-P10	F	Δ ICP6, $-/\gamma$ 34.5, +PTEN α , +eGFP	U87 Δ EGFR	1×10^5 (1 dose)	HSV-P10 significantly prolonged median survival, resulted in and 30% long-term survivors compared to 10% long-term survivors in unarmed HSVQ treatment group. ⁹²
ULBP3	oHSV ^{ULBP3}	F	Δ ICP4, +miR-124, +ULBP3	XFM-Luc: PDGF;Cre	1×10^6 (1 dose)	OHSV ^{ULBP3} inhibited tumor growth and prolonged median survival (18 days) compared to unarmed oHSV (8 days). ⁹⁵
	oHSV ^{ULBP3-MMP9}	F	Δ ICP4, +miR-124, +ULBP3, +MMP9	XFM-Luc: PDGF;Cre	1×10^6 (1 dose)	OHSV ^{ULBP3} showed statistically significant prolongation of median survival compared to OHSV ^{ULBP3-MMP9} . ⁹⁶
Pro-drug activating genes	ONCR-1	F	Δ ICP4, $-/\gamma$ 34.5, +miR-124, +ULBP3, +MMP9	U251	3×10^5 - 3×10^6 (1 dose)	ONCR-1 inhibited tumor growth and prolonged survival compared to mock group. ⁹⁸
	MGH2	F	Δ ICP6, $-/\gamma$ 34.5, -UL39, +CYP2B1, +shICE	Gli36 Δ EGF	1.4×10^6	MGH2 did not prolong median survival compared to mock group. ⁹⁹

Abbreviations: PFU, plaque forming unit; (-), single deletion; (-/-), diploid deletion; Δ , mutation/inactivation; (+), insertion.

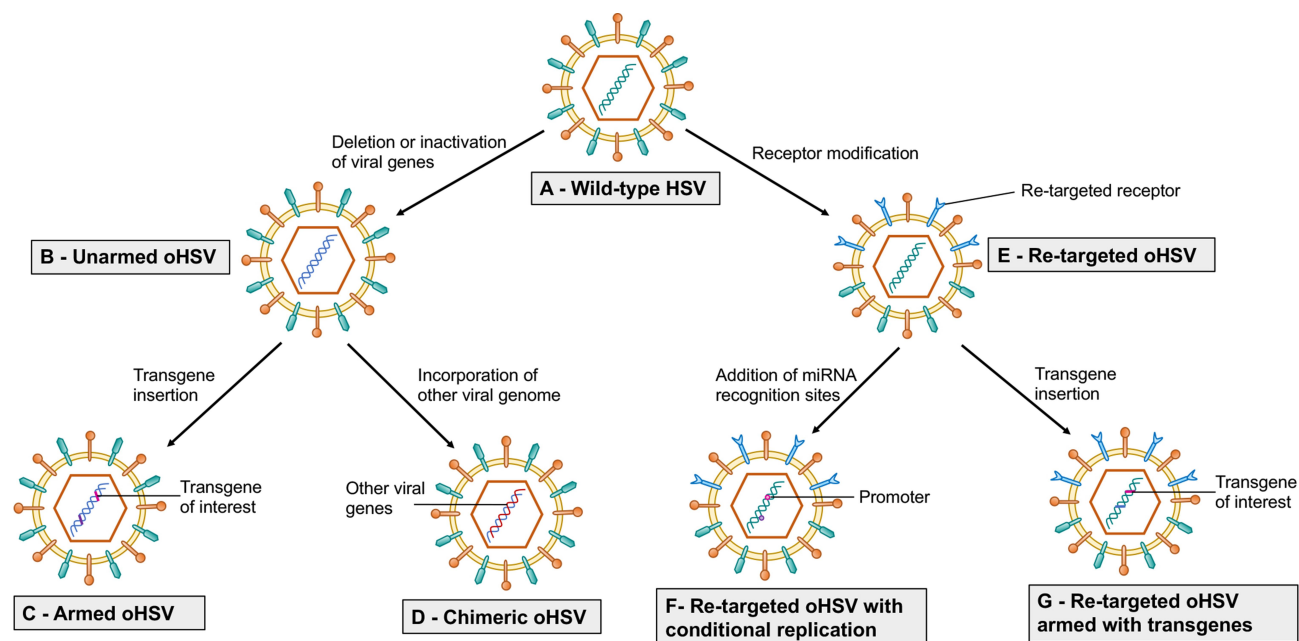


Figure 1 Strategy to enhance efficacy of oHSV as monotherapy. This figure represents different strategies to enhance the anti-GBM efficacy of oHSV. (A–B) Wild-type HSV can be genetically modified to remove or inactivate various genes to create unarmed oHSV. This is to ensure virus replication selectively occurs in cancer cells, thus enhancing the safety of oHSV (eg, *dlspk*, *hrR3*, *I1716*, *3616*, *G207*, *G47Δ*, *HSVQ1*, *rQNestin34.5*, *MG18L*). (C) To further enhance oHSV efficacy, different transgene variants are incorporated into oHSV genome to generate armed oHSV. Transgene expression increases oHSV anti-cancer efficacy through enhancement of viral spread (eg, *OV-CDH1*, *OV-ChaseM*) or activation of the host immune response (eg, *G47Δ-mIL12*, *αMPD-1 scFv*) or other means (See Table 3). (D) In addition, insertion of transgene that has similar function to oHSV genes but originating from other viruses is an alternative approach to enhance oHSV replication/oncolysis. oHSVs that are engineered by this method are defined as chimeric oHSV (eg, C130, C134). (E) Wild-type HSV can also be modified to enhance cancer-specificity, based on cellular surface receptor expression profile. They are known as re-targeted oHSVs, which selectively interact with various receptors that exclusively overexpressed in GBM cells but not in normal neurons/or glial cells (eg, *R-LM113*, *KNE*). This approach allows maximal oncolysis because HSV genome remains intact. (F) To enhance safety and avoid off-target effects, re-targeted oHSVs (E) can be further genetically modified to include miRNA recognition binding sites (such as *miR-124* in *KGE-4:T124*) whose expression are distinct in GBM cells compared to healthy neurons/glial cells. These recognition-binding sites are incorporated into the untranslated region (UTR) of important genes responsible for viral pathogenicity. Based on the difference between GBM cells and healthy neurons/glial cells, these recognition binding sites can be occupied and the sequential genes are not translated in normal cells, whereas the recognition binding sites are free and viral translation takes place in GBM cells. In addition, re-targeted oHSVs (E) can be armed with different transgene of interest (G) to further enhance anti-tumor efficacy (eg, *R-I15*, *KMMP9*).

a significant but modest extension of survival of mice bearing orthotopic immunocompetent 005 GSC-derived tumors, which was associated with a significant reduction of regulatory T cells (vs mock) and T cell-dependent anti-tumor efficacy.³³

Another strategy being utilized to counteract attenuated oHSV replication is restoring one copy of γ 34.5 under the transcriptional control of the nestin promoter/enhancer element. *rQNestin34.5*³⁴ and *rQNestin34.5v.2*³⁵ were engineered using this strategy. Nestin-driven γ 34.5 expression exclusively takes place in GBM cells, which have an overwhelmingly higher level of nestin expression compared to healthy cells.^{34,35} *rQNestin34.5* injection in mouse brain was safe and significantly prolonged survival of mice bearing orthotopic U87ΔEGFR tumors compared to control.³⁵ Cassady et al described a different approach to improve oHSV replication and generated chimeric oHSVs that express the human cytomegalovirus (HCMV) genes, *TRS1* (designated C130) or *IRS1* (designated C134).³⁶

TRS1 and *IRS1* are PKR evasion genes that evade the PKR-eIF2 α translational shutoff pathway,³⁷ and therefore, transfer of these HCMV genes to γ 34.5-deleted oHSV resulted in virus replication/production and killing of GBM cells as efficient as wild-type HSV.³⁸ C130 and C134 chimeric oHSVs demonstrated superior anti-glioma efficacy than C101 (a γ 34.5-deleted oHSV without *TRS1* or *IRS1* expression) in an immunodeficient U87 glioma model.³⁹ C134 treatment also elicited a durable and circulating anti-tumor memory response in an immunocompetent neuro2A brain tumor model.⁴⁰ Intracerebral injections of C134 did not produce toxicity in HSV-susceptible CBA/J mouse and non-human primates.⁴¹

Overall, unarmed or chimeric oHSVs are safe but their anti-GBM efficacy is modest at best (see Table 1), and in view of this observation, extensive research efforts have been made (which we have discussed below) to develop novel oHSVs and oHSV-based treatment strategies (such as receptor re-targeting, arming with therapeutic

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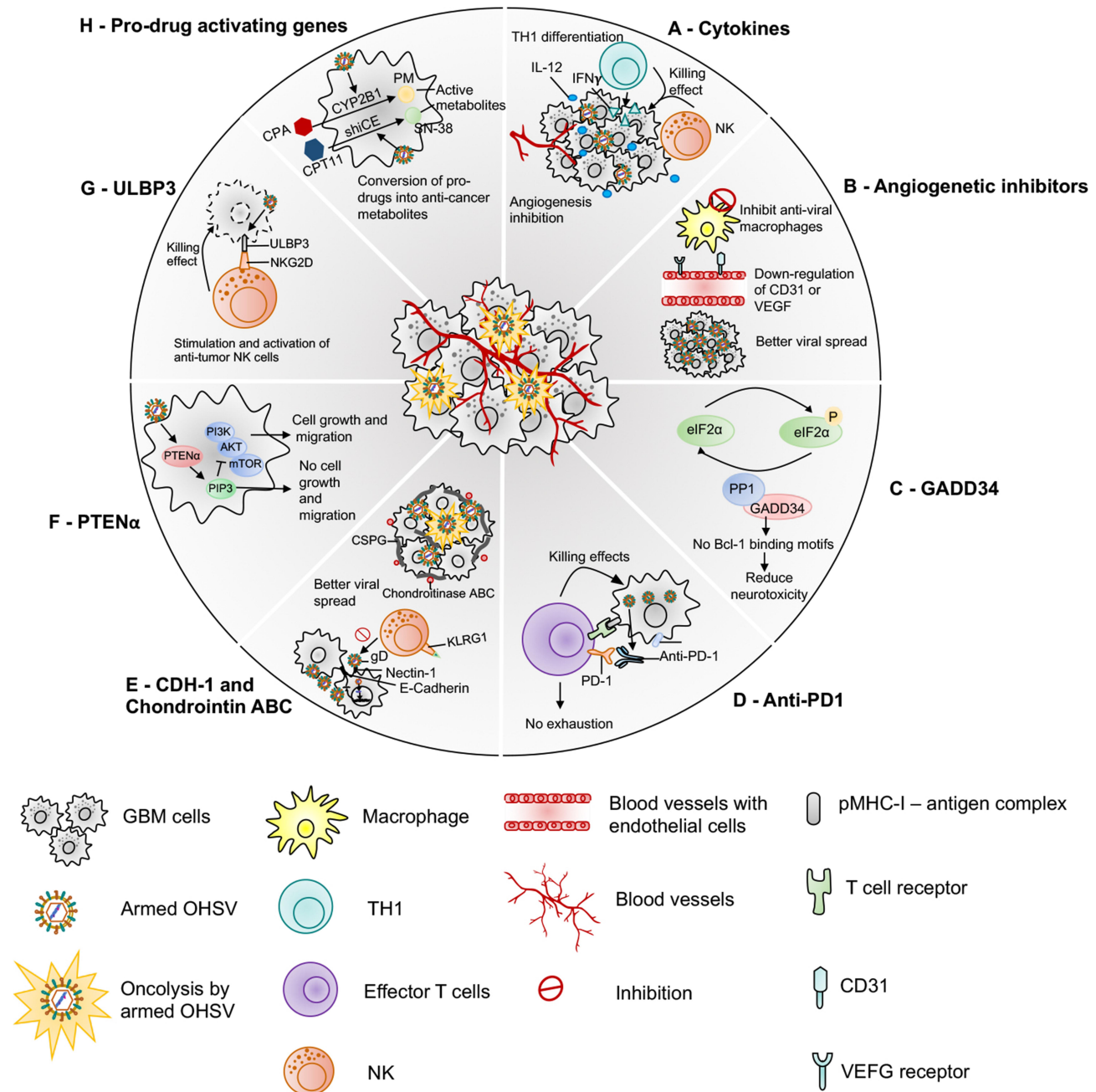


Figure 2 Armed oHSVs for GBM treatment. This figure represents different transgene variants that are inserted in the oHSV genome to enhance the anti-tumor efficacy of oHSV. (A) oHSV armed with cytokine such as IL-12, which induces Th1 differentiation, stimulates growth and cytotoxicity of NKs, increases IFN- γ production, and inhibits angiogenesis. (B) oHSV armed with angiogenic inhibitors decrease tumor vascularity (CD31⁺ vessels) or VEGF, inhibit anti-viral macrophages, and increase viral spread. (C) oHSV can be engineered to replace one copy of γ 34.5 with GADD34. GADD34 binds to PP1 and promotes eIF-2 α dephosphorylation - a function that corresponds to γ 34.5 in HSV. Since GADD34 does not possess beclin-1-binding motifs of γ 34.5, it does not produce neurotoxicity as wild-type HSV containing both copies of γ 34.5. (D) Localized expression of anti-PD-1 by an oHSV inhibits PD-1/PD-L1 engagement, prevents T cell exhaustion and unleashes anti-tumor immunity. (E) oHSV can be armed with CDH-1 gene, which encodes for E-cadherin – an adhesion molecule and a ligand for an inhibitory receptor expressed on NK cells (KLRG1). E-cadherin can cooperate with nectin-1 and promote cell-to-cell adherent junctions and enhance cell-to-cell oHSV spread. Another strategy to increase viral spread is to create an oHSV that expresses chondroitinase ABC, which removes side chain of CSPG, prevents extracellular space tortuosity, and thus, facilitates viral spread. (F) PTEN α expressed by an oHSV metabolizes PIP₃ prevents activation of the PI3K/AKT/mTOR signaling pathway and inhibits tumor growth and migration. (G) oHSV armed with ULBP3 enhances anti-tumor immunity. ULBP3 is a ligand for NKG2D. ULBP3-NKG2D interaction augments the anti-tumor activity of NK cells. (H) oHSV can be armed with pro-drug activating genes such as CYP2B1 (which converts CPA to PM) and shiCE (which converts CPT11 to SN-38) to enhance the conversion of these pro-drugs into their anti-cancer active metabolites.

Abbreviations: IL-12, interleukin-12; NKs, natural killer cells; IFN- γ , interferon-gamma; VEGF, vascular endothelial growth factor; GADD34, growth arrest and DNA damage-inducible protein 34; Bcl-1, beclin-1; PD-1, programmed death-1; CDH-1, cadherin-1; KLRG1, killer cell lectin-like receptor G1; CSPG, chondroitin sulfate proteoglycan; PTEN α , phosphatase and tensin homolog deleted on chromosome 10 alpha; PIP₃, phosphatidylinositol (3,4,5)-trisphosphate; PI3K/AKT/mTOR, phosphoinositide 3-kinase/AKT/mammalian target of rapamycin; ULBP3, ULI6-binding protein 3; NKG2D, natural killer group 2; member D; CPA, cyclophosphamide; CPT11, irinotecan; CYP2B1, CPA-activating cytochrome P4502B1; shiCE, CPT11-activating secreted human intestinal carboxylesterase; PM, phosphoramidate mustard.

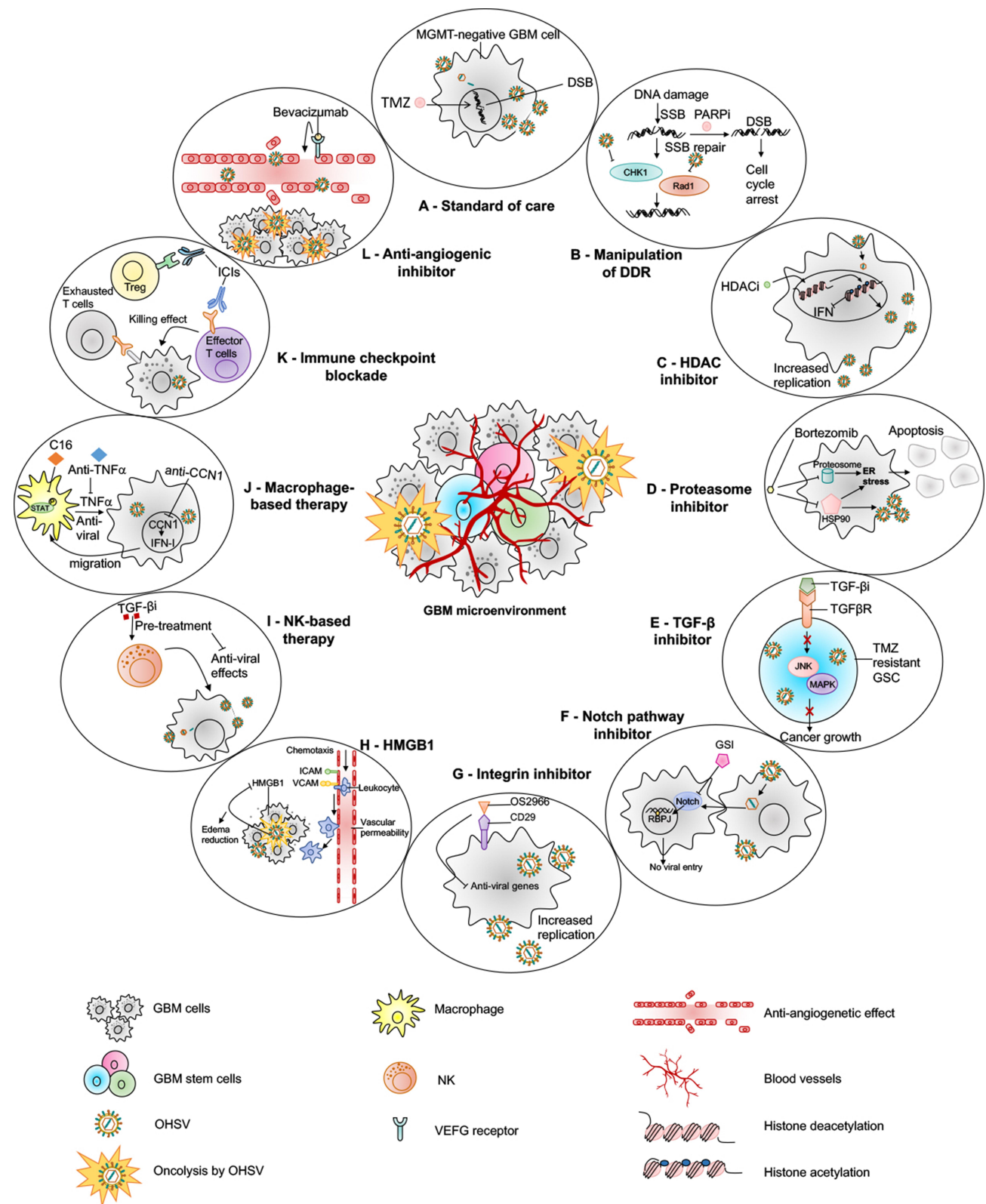


Figure 3 oHSV-based combination therapies for GBM. This figure represents strategies to combine oHSV with different anti-cancer therapies for GBM treatment. (A) oHSV can be synergistically combined with TMZ, a DNA-alkylating agent and an immunomodulator, to induce DNA damage response in MGMT-negative GBM cells. (B) oHSV replication promotes degradation of Rad51 and Chk1 whose functions are important for SSB repair mechanism. Inhibition of PARP in DNA-damaged cells facilitates the conversion of SSB to DSB. The combination of oHSV and PARPi synergistically induces cell cycle arrest in GBM. (C) HDAC is an enzyme that controls cancer cell survival/progression and upregulates IFN genes. Treatment with HDAC inhibitors prior to oHSV infection inhibits induction of anti-viral IFN genes, resulting in increased transcription of viral genes and improved virus replication. (D) Bortezomib is a proteasome inhibitor; which induces unfolded protein response (UPR), characterized by induction of heat-shock proteins (HSP) 40, 70 and 90, and ER stress in cancer

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transgenes, combination therapies, etc.) to improve oHSV's cancer selectivity and anti-GBM efficacy, without compromising safety.

Re-Targeted oHSVs for GBM

oHSVs can be engineered to specifically infect cancer cells based on their receptor expression profile.⁴² Epidermal growth factor receptor (EGFR) is overexpressed in ~60% of primary GBMs.⁴³ To specifically target EGFR⁺ GBM, an EGFR-retargeted gB:NT recombinant virus (designated KNE) was generated by replacing 2–24 amino acid residues of HSV glycoprotein D (gD, a natural oHSV receptor) with a single-chain fragment variant (scFv) antibody directed against EGFR.⁴² KNE significantly controlled tumor growth and prolonged survival of mice in an orthotopic GBM xenograft model.⁴² Like EGFR, human epidermal growth factor receptor 2 (HER2) is also expressed by ~80% of primary GBMs, but not by neurons or glial cells.^{44,45} Therefore, prior to KNE, an oHSV-1 re-targeted to HER2 receptor (designated R-LM113) was developed.⁴⁶ R-LM113 was significantly efficacious against HER2 overexpressing murine GBM xenografts, as demonstrated by 21-day extension of median survival compared to the control group.⁴⁶ R-115 is another HER2 re-targeted oHSV-1, which is armed with murine IL-12.⁴⁷ R-115 treatment (vs control) also significantly prolonged the median survival of mice bearing orthotopic poorly immunogenic high-grade gliomas. Most importantly, R-115 treatment led to the eradication of established tumors in about 30% of animals and they were protected lethal tumor re-challenge, indicating a cancer-specific immunological memory

response.⁴⁷ This promising anti-tumor efficacy of R-115 monotherapy was associated with the production of antibodies against implanted tumor cells in 83% treated animals (compared to 40% in IL-12 unarmed R-LM113 group or 0% in mock group) and increased infiltration of CD4⁺ and CD8⁺ T cells in the tumor mass.⁴⁷ Despite re-targeted oHSVs elicited promising oncolytic activity and cancer selectivity, most cancer cell surface markers are shared with normal cells. Therefore, to further enhance cancer specificity, an additional mechanism was explored to better localize re-targeted oHSV replication within GBM cells. GBM has a distinguished microRNA (miRNA) expression profile compared to the normal brain.⁴⁸ miRNA-124 (miR-124) is specifically expressed in neurons but not in GBM cells.⁴⁸ To utilize this distinct miR-124 feature, FGE-4:T124, the next generation of EGFR re-targeted KNE virus, was engineered, which contains the insertion of four copies of miR-124 recognition site (T124+) into the 3'UTR of the viral ICP4 gene.⁴⁹ ICP4 is critically required for transcription of early/late proteins and viral DNA synthesis.⁵⁰ The normal healthy cells have a high expression of miR-124, which binds to miR-124 recognition sites and inhibits T124+ virus replication. In contrast, the T124+ virus replicated robustly in glioma cells, and lentiviral expression of miR-124 in tumor cells selectively blocked its replication, confirming tumor-specificity of T124+.⁴⁹ Re-targeted oHSVs can be armed with various transgene variants to further enhance the antitumor efficacy. For instance, KMMP9 is an EGFR/EGFRvIII re-targeted oHSV that has an insertion of repeated miR-124 recognition site into the viral ICP4 gene and armed with matrix metalloproteinase 9 in the UL3-UL4

←
Figure 3 (Continued).

cells. Bortezomib-induced ER stress and UPR significantly enhance oHSV replication and synergistic killing of GBM cells. (E) TGF- β plays a critical role in GBM pathogenesis and in maintaining the stemness of GSCs. In TMZ-resistant GSC model, the combination of oHSV and TGF- β synergistically kills TMZ-resistant recurrent GSCs, increases oHSV replication and induces JNK-MAPK signaling blockade, and eventually inhibits tumor progression. (F) oHSV infection of tumor cells leads to activation of notch signaling in adjacent non-infected tumor cells. Notch signaling pathway plays a critical role in cell-cell interaction and viral spread. Inhibition of notch signaling pathway such as GSI results in increased killing of GBM cells after oHSV therapy. (G) ITGB1, also referred as CD29, plays a critical role in tumor cell proliferation and progression. OS2966 (a humanized CD29 blocking antibody) blocks CD29, reduces the expression of anti-viral genes (IFN α , IFN β , Stat1, OAS1, OAS2, IRF3, IRF9, and PKR), suppresses oHSV-induced macrophage activation, resulting in enhanced oHSV replication and oncolysis. (H) GBM cells that are infected by oHSV upregulate HMGB1. HMGB1 causes upregulation of ICAM and VCAM, increases vascular permeability and PMBC infiltration to the tumor, leading to edema that might cause CNS injuries. Thus, combination of anti-HMGB1 and oHSV increases survival by reducing brain injuries. (I) Recruitment of NKs after oHSV administration can limit oHSV replication and oHSV-mediated anti-tumor efficacy. Transient inhibition of anti-viral effects of NKs by TGF- β inhibitors enhances viral replication and viral yield. (J) Similarly, transient blockade of TNF α , produced from anti-viral macrophages, by TNF α blocking antibodies or inhibition of STAT1/3 phosphorylation by C16 enhances oHSV replication. In addition, virus-infected cells upregulate CCN1, which in turn activates an intracellular type I IFN response and increases infiltration of macrophages to the site of infection. Treatment with anti-CCN1 reduces virus clearance by macrophages, resulting in better anti-tumor efficacy. (K) Administration of immune checkpoint blockade such as anti-PD-1, anti-PD-L1, anti-CTLA4 prevents T cell exhaustion and enhances oHSV-mediated-anti-tumor immunity. (L) Bevacizumab binds to VEGF, reduces tumor vascularization, decreases vascular permeability, and inhibits tumor growth; however, it also induces tumor cell invasion. An oHSV expressing anti-angiogenic vasculostatin that contains an integrin-antagonizing RGD (Arg-Gly-Asp) motif significantly inhibits glioma cell migration/invasion following bevacizumab treatment, leading to a significant extension of survival compared to bevacizumab monotherapy. **Abbreviations:** TMZ, temozolomide; MGMT, O-6-methylguanine-DNA methyltransferase; DSB, double-strand break; SSB, single-strand break; PARP, poly(ADP-ribose) polymerase; DDR, DNA damage response; HDAC, histone deacetylase; IFN, interferon; UPR, unfolded protein response; ER, endoplasmic reticulum; TNF- α , tumor necrosis factor-alpha; TGF- β , transforming growth factor-beta; GSC, glioma stem cells; RBPJ, recombination signal binding protein for immunoglobulin kappa J region; JNK, c-Jun NH2-terminal kinases; ICAM, intercellular adhesion molecule; VCAM, vascular cell adhesion molecule; MAPK, mitogen-activated protein kinase; GSI, gamma-secretase inhibitor; ITGB1, integrin β 1; OAS1, 2'-5'-Oligoadenylate Synthetase; IRF, interferon regulatory factor; PKR, protein kinase R; HMGB1, high mobility group box 1; CNS, central nervous system; STAT, signal transducer and activator of transcription; CCN1, cellular communication network factor 1; PD-1, programmed death-1; PD-L1, programmed death ligand-1; CTLA4, cytotoxic T-lymphocyte-associated protein 4.

Table 4 Clinical Studies with oHSVs in GBM

oHSV	Combination	Condition	N	Phase	Status	Country	References
HSV1716	N/A	Recurrent malignant glioma	9	N/A	Completed	UK	[187]
	N/A	High-grade glioma	12	N/A	Completed	UK	[188]
	N/A	High-grade glioma	12	N/A	Completed	UK	[186]
	Dexamethasone	Refractory or recurrent high-grade glioma	2	I	Terminated	US	NCT02031965
G207	N/A	Recurrent malignant glioma	21	I	Completed	US	[190]
	N/A	Recurrent Brain Cancer	65	Ib/2	Completed	US	NCT00028158
	N/A	Recurrent GBM	6	Ib	Completed	US	[191]
	Radiation	Recurrent malignant glioma	9	I	Completed	US	[101]
	Radiation	Progressive or recurrent supratentorial brain tumors	12	I	Active, not recruiting	US	[194] NCT02457845
	Radiation	Recurrent or refractory cerebellar brain tumors	15	I	Recruiting	US	[196] NCT03911388
	Radiation	Recurrent high-grade glioma	30	2	Not yet recruiting	US	NCT04482933
G47Δ	N/A	Progressive GBM	21	I/2	Completed recruiting	Japan	UMIN000002661
	N/A	Residual or recurrent GBM	30	2	Completed recruiting	Japan	UMIN000015995
M032	N/A	Recurrent/progressive GBM	36	I	Recruiting	US	NCT02062827
rQNestin34.5v.2	Cyclophosphamide	Recurrent malignant glioma	108	I	Recruiting	US	NCT03152318
C134	N/A	Recurrent GBM	24	I	Active, not recruiting	US	NCT03657576

Abbreviations: N, number of participants; N/A, not applicable.

intergenic region.⁵¹ At a low viral dose, KMMP9 exhibited significantly enhanced anti-tumor efficacy with 70% long-term survivors compared to 0% with re-targeted unarmed KGW virus in a GBM xenograft model.⁵¹ The preclinical efficacy of several re-targeted oHSVs in GBM is listed in Table 2. Overall, the re-targeting strategy not only confers tumor specificity/safety but also ensures enhanced oncolytic activity with promising anti-GBM efficacy.

Armed oHSVs for GBM

While OV's possess an inherent ability to selectively replicate and lyse cancer cells, the success of oncolytic virotherapy requires the inevitable contribution of the host immune system.⁵² Therefore, the focus of OV therapies has recently shifted from their direct oncolytic effect to secondary effects (such as immunomodulation and anti-angiogenesis), which

can be better achieved by localized expression of different immunomodulatory or anti-cancer transgenes. The unique ability of OV's to selectively replicate in cancer cells makes them a good candidate to safely deliver anti-cancer agents locally in the TME and reducing the possible systemic side effects. For example, systemic delivery of interleukin 12 (IL-12) as an immunotherapeutic agent induces toxicity,⁵³ but local IL-12 expression by an OV in the TME⁵⁴ or in the cerebellum⁵⁵ was found safe and showed better anti-tumor response compared to parental unarmed OV's.⁵⁴ Research in armed oHSVs and their application in GBM have bloomed in recent years, as indicated by the development of a significant number of oHSVs expressing various anti-cancer agents (Table 3, Figure 2). In this section, we will provide an overview of the pre-clinical therapeutic efficacy of recently developed armed oHSVs for GBM treatment.

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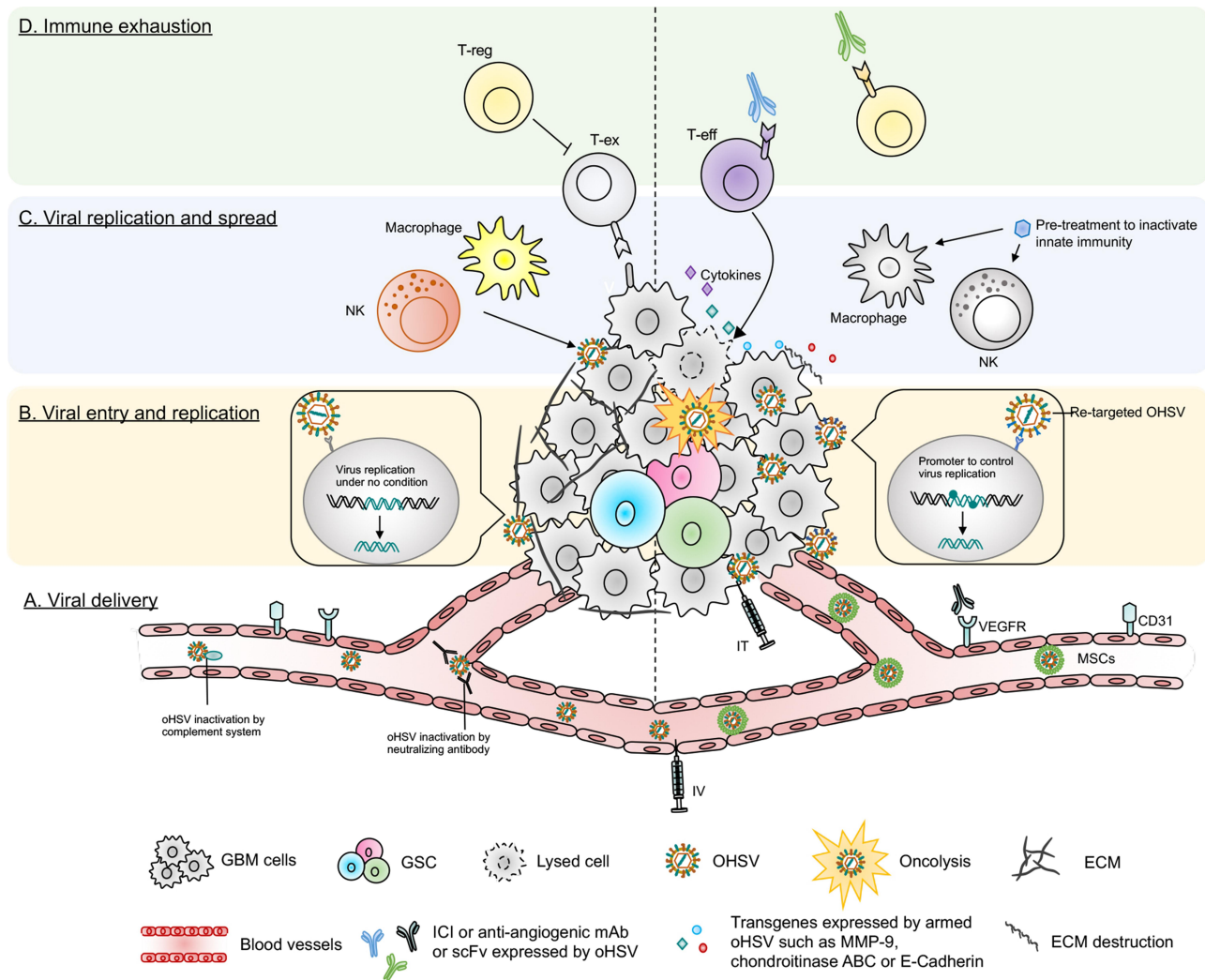


Figure 4 Barriers of current oHSV therapy in GBM and their potential solutions. This figure represents various obstacles of oHSV therapy in GBM (left side) and proposed solutions to overcome these hurdles (right side). (A) Viral delivery – oHSV can be delivered systemically (IV) or locally (IT). IT injection can cause backflow of virus solution to the needle or catheter, leading to insufficient viral dose. IV can be another alternative route to deliver oHSV. Systemic administration of naked OVs can be inactivated by host serum/complement and other immune factors. Systemically delivered oHSVs can be protected from serum/immune factors if delivered by carrier cells, such as MSCs. (B) Viral entry and replication – The higher the expression of HSV’s entry surface receptors such as nectin-1, the higher the entry of oHSVs (such as G207, M032) to cancer cells. Genomic modification attenuates virus replication (such as G207) in tumor cells, especially in GSCs. In contrast, receptor re-targeting can produce off-target effects in healthy cells that also express the same receptors. This issue can be overcome by incorporating miRNA recognition binding sites (such as miR-124 in KGE-4:T124) whose expression are distinct in GBM cells compared to healthy neurons/glia cells. (C) Viral replication and spread – Condensed ECM can limit viral spread in the TME. oHSVs can be engineered to express E-cadherin, chondroitinase ABC, and/or MMP-9 to destroy the ECM components and facilitate viral spread. In addition, anti-viral innate immunity can also limit viral replication/spread in the TME, as characterized by recruitment of macrophage or NKs that clear virus or virus-infected cells. Transient inhibition of anti-viral innate immune response by pre-treatment with different therapies such as TNF- α or TGF- β inhibitors can help to increase oHSV replication/spread. (D) Immune exhaustion – Successful viral delivery, entry, replication and spread will eventually activate the host’s adaptive anti-tumor immunity, leading to recruitment of T cells to the TME. The recruited T cells oftentimes fail to produce anti-cancer effects due to the expression of T cell exhaustion markers. The presence of Tregs and angiogenesis also contribute to GBM immunosuppression. oHSVs can be combined with systemic ICI or systemic anti-angiogenic mAb, or oHSVs can be engineered to locally express scFv of ICI or anti-angiogenic mAb that should overcome GBM immunosuppression. In addition, oHSVs can be armed with different transgene variants (such as cytokines, co-stimulatory ligands) and testing them in combination with ICIs or other immunotherapies such as anti-cancer vaccines (eg. DCs loaded with tumor-associated antigens) to improve anti-tumor efficacy.

Abbreviations: IV, intravenous; IT, intra-tumoral; MSCs, mesenchymal stem cells; TME, tumor microenvironment; ECM, extracellular matrix; MMP, matrix metalloproteinase; NK, natural killer cells; TNF- α , tumor necrosis factor-alpha; TGF- β , transforming growth factor-beta; scFv, single-chain fragment variant; ICI, immune checkpoint inhibitor; mAb, monoclonal antibody; VEGF, vascular endothelial growth factor; T-reg, regulatory T cell; T-ex, exhausted T cell; T-eff, effector T cell.

Arming with Cytokines

Among many cytokines explored to enhance the anti-tumor efficacy of OVs, IL-12 is the most commonly used cytokine.^{54,56} IL-12 regulates innate and adaptive immunity, induces Th1 differentiation, stimulates growth and

cytotoxicity of natural killer cells (NKs), increases IFN- γ production, and inhibits angiogenesis.^{33,56} Cheema et al constructed G47 Δ -mIL12 (an oHSV expressing murine IL-12), which has similar genomic modifications as G47 Δ ³³ (Table 3), with the additional insertion of murine IL-12

cDNA in the ICP6 region.³³ G47Δ-mIL12 treatment significantly inhibited GBM angiogenesis and extended survival (vs unarmed oHSV) in orthotopic U87 and human GSC-derived GBM xenograft models.⁵⁷ Although G47Δ-mIL12 was efficacious in xenograft models, the contribution of the immune system to the efficacy of oHSV or oHSV product (ie, IL-12) could not be evaluated in those models. In immunocompetent orthotopic mouse 005 GSC-derived GBM model, intratumoral G47Δ-mIL12 treatment led to a significant extension of median survival with 10% mice surviving long term compared to treatment with unarmed G47Δ-Empty (G47Δ-E).³³ The promising G47Δ-mIL12's efficacy was associated with a significant reduction of tumor cells (GFP⁺ 005 GSCs) and CD4⁺FoxP3⁺ regulatory T cells (Tregs), and inhibition of CD31⁺ tumor vascularity, and the efficacy was dependent on T cells but not NK cells.³³ Another IL-12 expressing oHSV, M002, showed a trend of improved median survival in three patient-derived pediatric or adult xenograft GBM models compared to unarmed G207.⁵⁸ Like 005 model, M002 was extremely efficient in an intracranial syngeneic 4C8 murine glioma model, since M002 treatment resulted in 80% long-term survivors.⁵⁹ Like IL-12, other cytokines such as FMS-like tyrosine kinase 3 ligand (Flt3L) or tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) were inserted into the backbone of G47Δ and generated G47Δ-Flt3-L⁶⁰ and oHSV-TRAIL viruses,⁶¹ respectively. Both G47Δ-Flt3-L and oHSV-TRAIL were efficacious in GBM^{60–62} (Table 3). Importantly, recently, Jahan et al demonstrated that oHSV-TRAIL can overcome TMZ-resistance in GBM.⁶² For instance, intracranial oHSV-TRAIL treatment of mice bearing TMZ-insensitive, recurrent human GSC-derived orthotopic tumors led to potent inhibition of tumor growth (examined by *in vivo* imaging) and extension of survival with a 40% cure rate. The treatment efficacy of oHSV-TRAIL in chemo-resistant tumors was associated with robust induction of tumor apoptosis.⁶² TRAIL has a short half-life and produces off-target toxicity⁶³ and the clinical trials of TRAIL receptor agonist have been discontinued in the clinic due to insufficient evidence of anti-cancer efficacy.⁶⁴ Local expression of TRAIL by an oHSV offers a promising strategy to avoid systemic off-target toxicity.

Arming with Angiogenic Inhibitors

Preclinical studies with oHSV expressing various angiogenic inhibitors consistently exhibited improved survival outcomes compared to mock or unarmed viruses.^{57,65–67}

G47Δ-mAngio, an oHSV armed with murine angiostatin (mAngio), controlled tumor burden and significantly extended median survival compared to G47Δ-E in two immunodeficient intracranial GBM models.^{57,67} A closer look at the TME revealed that G47Δ-mAngio treatment resulted in significantly increased virus spread (LacZ expression), decreased tumor vascularity (CD31⁺ vessels) and anti-viral macrophage density, and reduced expression of vascular endothelial growth factor (VEGF) compared to G47Δ-E.⁵⁷

Zhang et al constructed an oHSV expressing endostatin-angiostatin (endo-angio) fusion gene (designated VAE) by deleting both copies of γ 34.5, inactivating ICP6, and inserting endo-angio fusion gene into ICP6 region.⁶⁵ VAE demonstrated a significant improvement in median survival in two (subcutaneous and intracranial) GSC-derived xenograft models compared to γ 34.5/ICP6-deleted r-HSV-1 and recombinant human endostatin (Endostar).⁶⁵ The better response in VAE-treated group was associated with a reduction in mean micro-vessel density (MVD) compared to r-HSV-1 or Endostar.⁶⁵ Tomita et al recently described antitumor efficacy of RAMBO – an oHSV (γ 34.5/ICP6 deleted) expressing vasculostatin (Vstat120) – in athymic mice bearing intracerebral U87ΔEGFR tumors.⁶⁶ The median survival of PBS-treated mice was significantly shorter (17 days) than mice treated with RAMBO (28 days).⁶⁶ Mechanistically, Vstat120 expression inhibited TNF- α production by blocking brain-specific angiogenesis inhibitor 1 (BAI1)-mediated macrophage response to viral infection, resulting in an increased viral spread.⁶⁸

Arming with Growth Arrest and DNA Damage-Inducible Protein 34 (GADD34)

GADD34 or protein phosphatase 1 regulatory subunit 15A (PPP1R15A) is a member of a group of genes whose transcript levels are increased by DNA damage, growth factor deprivation, and other forms of cell stress.⁶⁹ GADD34 binds to PPI and promotes eIF-2 α dephosphorylation,⁷⁰ a function that corresponds to γ 34.5 in HSV.⁷¹ Interestingly, while GADD34 shares sequence homology with the C terminus of γ 34.5,⁷⁰ it does not possess the beclin-1-binding motifs of γ 34.5 – a feature that is responsible for neurotoxicity by HSV.⁷² Therefore, the replacement of γ 34.5 by GADD34 is thought to reduce neurotoxicity. Based on this principle, NG34, a novel oHSV (γ 34.5/ICP6-deleted) expressing GADD34, was

created.⁷² Noticeably, in NG34, GADD34 expression is transcriptionally controlled by a nestin promoter/enhancer; thus, its expression occurs selectively in GBM.⁷² NG34 demonstrated a similar replication/propagation pattern in GBM cells as the parental unarmed rQNestin34.5 virus.⁷² In both human xenograft and immunocompetent GBM models, NG34 produced therapeutic efficacy, which was as potent as rQnestin34.5 but with better tolerability.⁷²

Arming with Programmed Death-1 (PD-1) Inhibitor

Immune checkpoint co-inhibitory molecule PD-1 and its ligand PD-L1 play a critical role in regulating immune responses and suppressing anti-tumor immunity. Anti-PD-1 inhibits PD-1/PD-L1 interaction, prevents T cell exhaustion, and unleashes anti-tumor immunity.^{73,74} Despite being extensively studied, late-phase clinical trials with anti-PD-1 in GBM did not signify its therapeutic benefits.^{75,76} The inability of anti-PD-1 to cross the blood-brain barrier (BBB) and highly immunosuppressive TME with a low number of tumor-infiltrating lymphocytes (TILs) are probably the major contributors to treatment failure.⁷⁷ Local oHSV expression of anti-PD-1 may overcome these obstacles. OVH- α MPD-1, an oHSV encoding a single-chain fragment variable antibody against PD-1 (α MPD-1 scFv), was generated that demonstrated a promising anti-tumor efficacy (ie, enhanced antigen presentation, T cell-mediated cytotoxicity, and reduction of tumor burden) in a liver cancer model, but its efficacy has not been tested yet in GBM.⁷⁸ Recently, Passaro et al constructed another oHSV by inserting an scFv against PD-1 (designated as NG34scFvPD-1) in the previously mentioned NG34 backbone.⁷⁹ In a syngeneic GL261N4 immunocompetent GBM mouse model, in situ expression of a PD-1 blockade by NG34scFvPD-1 produced a durable response and long-term memory protection, but surprisingly, these anti-tumor effects were statistically similar to that of parental NG34 virus.⁷⁹

Arming with Cadherin-1 (CDH-1)

The limited virus spread in the TME due to rapid innate immune clearance of the virus is considered as one of the major obstacles of oHSV virotherapy.⁸⁰ The interaction between viral glycoprotein D (gD) and nectin-1 receptor is an important step for HSV-1 entry.⁸¹ CDH-1 gene encodes for E-cadherin – an adhesion molecule and

a ligand for killer cell lectin-like receptor G1 (KLRG1), which is an inhibitory receptor expressed on NK cells.⁸² E-cadherin can cooperate with nectin-1 and promote cell-to-cell adherent junctions,⁸³ and thus, its overexpression can enhance cell-to-cell oHSV spread. Xu and colleagues engineered an oHSV to express CDH-1 (named OV-CDH1).⁸⁴ In vitro, OV-CDH1 infection resulted in ectopic expression of E-cadherin that enhanced viral spread/entry by promoting cell-to-cell infection and reduced viral clearance by protecting infected GBM cells from KLRG1⁺ NK cell killing.⁸⁴ In vivo, OV-CDH1 treatment significantly prolonged median survival compared to unarmed virus (OV-Q1) in two xenografts and one immunocompetent GBM models.⁸⁴ The better outcome in OV-CDH1-treated group in the immunocompetent GBM model was associated with an improved viral spread rather than inhibition of the activity of NK cells.⁸⁴

Arming with Chondroitinase ABC

Chondroitinase ABC is a bacteria-derived enzyme that cleaves and removes the side chain of chondroitin sulfate proteoglycans (CSPG).⁸⁵ CSPG is upregulated in 60–65% GBM cases,⁸⁶ tortois the extracellular space, and limits OV spread.^{85,87} In a mathematical model, chondroitinase ABC enhanced OV spread and anti-tumor efficacy.⁸⁷ An oHSV expressing chondroitinase ABC (designated OV-Chase) exhibited widespread three-dimensional glioma cultures compared to the control virus (rHSVQ). In vivo, OV-Chase treatment (vs rHSVQ) significantly extended median survival in subcutaneous and intracranial GBM models.⁸⁵ The humanized version of OV-Chase (designated OV-ChaseM) also enhanced median survival of athymic mice bearing GBM xenograft (33 days vs 22 days in the mock group).⁸⁸

Arming with Phosphatase and Tensin Homolog Deleted on Chromosome 10 Alpha (PTEN α)

PTEN is a tumor suppressor protein phosphatase that metabolizes phosphatidylinositol (3,4,5)-trisphosphate (PIP₃), which directly opposes activation of the PI3K/AKT/mTOR signaling pathway.⁸⁹ Loss or mutation of PTEN, commonly occurred in GBM,^{89,90} activates the PI3K/AKT pathway and promotes cell growth/survival and migration.⁹¹ Local expression of PTEN α , an isoform of PTEN, by an oHSV (termed HSV-P10) significantly prolonged median survival with 30% long-term survivors

in a U87ΔEGFR xenograft model compared to 10% in the unarmed oHSV (HSVQ) group.⁹²

Arming with ULI 6-Binding Protein 3 (ULBP3)

Human ULBP3, a ligand for NKG2D (natural killer group 2, member D), is overexpressed on certain tumor cells,⁹³ including brain tumors.⁹⁴ ULBP3-NKG2D interaction augments the anti-tumor activity of NK cells.⁹³ To exploit this anti-tumor mechanism, an oHSV armed with human ULBP3 gene (oHSV^{ULBP3}) was created and tested in a genetically engineered IDH wild-type GBM mouse model (XFM-Luc: PDGF, Cre).⁹⁵ Intra-tumoral oHSV^{ULBP3} treatment inhibited tumor growth and prolonged survival, whereas oHSV lacking ULBP3 did not.⁹⁵ The survival efficacy was associated with an increased accumulation of activated macrophages and CD8⁺ T cells in the TME. The same group later reported an increased tumoral expression of VEGF and matrix metalloproteinase 9 (MMP-9) by oHSV^{ULBP3} treatment. A vast portion of virally enhanced VEGF remained trapped in the extracellular matrix⁹⁶ and cannot fully exert its effects until trapped VEGF is cleaved by MMP-9 and subsequently released.⁹⁷ To determine the role of MMP-9 on VEGF signaling and its subsequent effect on oHSV^{ULBP3} efficacy, Wirsching et al engineered an oHSV^{ULBP3-MMP9} to co-express ULBP3 and MMP-9.⁹⁶ Although oHSV^{ULBP3} significantly prolonged survival of mice bearing XFM-Luc: PDGF, Cre GBM tumors compared to mock, MMP-9 expression, ie, oHSV^{ULBP3-MMP9} treatment nearly abolished the survival benefit of oHSV^{ULBP3} to a mock level.⁹⁶ The use of a VEGF-neutralizing antibody (B20) rescued anti-tumor effects of oHSV^{ULBP3-MMP9}, confirming the role of VEGF in abrogating treatment efficacy.⁹⁶ Another oHSV (designated ONCR-1) was engineered by deleting one copy of γ 34.5, inserting miR-124 binding sites into the ICP4 locus and arming with both ULBP3 and MMP-9.⁹⁸ Although ONCR-1 controlled tumor growth and extended survival in mice bearing subcutaneous or orthotopic human U251 GBM tumors, it was not determined whether the absence of MMP-9 expression would have facilitated a better spread/efficacy of a single transgene (ULBP3)-armed virus.⁹⁸

Arming with Prodrug-Activating Genes

Oncolytic effects of an oHSV can be enhanced by arming with pro-drug activating genes.⁹⁹ MGH2 is an oHSV armed with two prodrug-activating genes, cyclophosphamide (CPA)-activating cytochrome P4502B1 (CYP2B1)

and CPT11-activating secreted human intestinal carboxylesterase (shiCE).⁹⁹ CYP2B1 converts cyclophosphamide (CPA) into active anti-cancer DNA-alkylating metabolite, phosphoramidate mustard (PM), whereas shiCE converts irinotecan (CPT11) into active topoisomerase I inhibitor, SN-38. In athymic mice harboring human GBM tumor (Gli36ΔEGFR), MGH2 was not effective alone (vs PBS), but in combination with CPA and CPT11, MGH2 displayed a significantly robust anti-tumor efficacy in comparison to controls.⁹⁹ MGH2 was later modified to exclude a green fluorescent protein (GFP) expression cassette from its genome, and named MGH2.1, for its assessment in clinical trials.¹⁰⁰ The intracerebral administration of MGH2.1 was safe and did not affect the endogenous metabolism of CPA and CPT11.¹⁰⁰

The aforementioned reported studies illustrate that oHSVs armed with therapeutic transgene(s) in general produce superior anti-tumor efficacy than unarmed oHSVs (Table 3). The therapeutic efficacy of armed oHSVs varied from model to model, and none of the armed oHSVs led to complete tumor eradication in 100% of animals tested compared to unarmed oHSVs or control treatments (Table 3), indicating further research is needed to improve the efficacy of armed oHSVs.

Carrier Cell-Based oHSV Treatment for GBM

Intra-tumoral injection is the primary mode of viral delivery.^{101–104} Intra-tumoral injection can cause backflow of virus solution to the needle or catheter, leading to significant loss of infused solution, and thus, patients may not receive the required virus dose.^{104,105} Although this hurdle can be overcome by multiple injections into several sites, each injection increases the risk of intracranial bleeding and influx of cerebrospinal fluid into the resection cavity that could further rinse out the injected virus.^{5,104} Intravenous administration is another important viral delivery method; however, systemic (even local) administration of naked OVs can be inactivated by host serum/complement and other immune factors,¹⁴ and thus, alternative approaches were explored. In 2008, Sonabend et al reported the first study utilizing mesenchymal stem cells (MSCs) as a carrier to deliver OV for GBM treatment.¹⁰⁶ In 2014, Duebgen et al loaded human MSCs with oHSVs (MSC-oHSVs) and demonstrated that MSC-oHSVs were effective in producing oHSV progeny and anti-tumor efficacy.¹⁰⁷ Intracranial injection of

synthetic extracellular matrix-encapsulated MSC-oHSVs resulted in a significant extension of median survival compared to naked oHSV in a preclinical model of GBM resection.¹⁰⁷ Similarly, MSC-oHSV-TRAIL (MSC loaded with oHSV-TRAIL)⁶² effectively induced apoptosis-mediated killing of oHSV- and TRAIL-resistant GBM cells and prolonged median survival of mice bearing orthotopic tumors.¹⁰⁷ Although MSC-oHSV and MSC-oHSV-TRAIL were efficacious in immunodeficient GBM models, it is not clear whether they will be similarly effective against syngeneic GBM tumors.^{108,109} Recently, few more research groups utilized a similar carrier cell-based treatment approach for GBM.^{110–112} These studies provide an excellent platform to further investigate oHSV-loaded carrier cells in GBM.

oHSV-Based Combination Therapies for GBM

As discussed above, anti-tumor efficacy of unarmed, even armed viruses is limited. Thus, the focus of oHSV research has been shifted from monotherapy to combination therapy. Multimodal approaches were implemented to combine oHSV with chemotherapy, antiangiogenic agents, etc. (Figure 3). Here, we have covered different combinatorial treatment strategies that were developed in recent years.

Combination with Standard Chemotherapy

Temozolomide (TMZ), an oral DNA-alkylating agent and immunomodulator,¹¹³ is the first-line chemotherapy in GBM; however, at least 50% of GBM patients do not respond to TMZ.¹¹⁴ Expression of O-6-methylguanine-DNA methyltransferase (MGMT) is primarily associated with TMZ resistance, in addition to mutations in DNA mismatch repair proteins.^{114,115} OV-ChaseM sensitized glioma cells to TMZ and enhanced survival.⁸⁸ It was not clear from this study whether OV-ChaseM can overcome MGMT-mediated TMZ resistance. TMZ was also shown to synergize with G47Δ in killing human GSCs in vitro and enhancing survival in human GSC-derived orthotopic tumors in vivo.¹¹⁶ TMZ-oHSV interaction can be immunologically antagonistic since TMZ abrogated the efficacy of G47Δ-mIL12 in an immunocompetent orthotopic 005 GSC-derived GBM model.¹¹⁷ The antagonistic interaction was probably associated with MGMT expression and immune inhibitory effects of TMZ.¹¹⁷ The addition of an MGMT inhibitor, O6-BG, to TMZ did not overcome the

negative effects of TMZ.¹¹⁷ Synergy in the human GSC model and antagonistic effects in the mouse model can be explained by the fact that (i) human cancer cells are typically more permissive to oHSV replication than mouse cancer cells;³¹ (ii) oHSV impairs DNA damage response in human GSCs,¹¹⁶ but it is not known whether this occurs in mouse GSCs; and (iii) the schedule of TMZ treatment can also affect oHSV efficacy, since OV treatment prior to TMZ significantly extended survival in GL261 GBM model,¹¹⁸ whereas antagonism observed when G47Δ-mIL12 and TMZ were concurrently applied in the 005 model.¹¹⁷

Inhibition of Glioma Invasion and Angiogenesis

GBM is a highly vascularized and invasive tumor.^{119,120} Anti-angiogenic therapy is somewhat effective in GBM^{121–123} and synergizes with oHSV.^{57,67,124} Bevacizumab, an anti-VEGF monoclonal antibody, inhibits GBM growth by reducing tumor vascularity;^{122,123} however, bevacizumab can cause tumor invasion and resistance via several mechanisms including an integrin pathway-mediated invasion.^{125,126} To overcome bevacizumab-induced/integrin pathway-mediated invasion,⁶⁶ Tomita et al utilized RAMBO, an oHSV expressing anti-angiogenic vasculostatin (Vstat120) that contains an integrin-antagonizing RGD (Arg-Gly-Asp) motif,¹²⁷ which significantly inhibited glioma cell migration/invasion following bevacizumab treatment in vitro. In vivo, the combination (RAMBO + Bevacizumab) significantly reduced the depth of tumor invasion and extended survival of mice bearing orthotopic GBM xenograft compared to bevacizumab monotherapy.⁶⁶ Mechanistically, RAMBO prevented bevacizumab-induced cysteine-rich protein 61 (CCN1) expression and AKT phosphorylation and eventually suppressed glioma invasion.⁶⁶ Like RAMBO, G47Δ-mAngio⁶⁷ was also effective in reducing bevacizumab-induced tumor invasion in vivo and extending survival in a U87 glioma model.⁶⁷ The combination efficacy was associated with inhibition of tumoral VEGF and expression of invasion markers such as matrix metalloproteinases-2 (MMP2), MMP9, and collagen.⁶⁷ Like the G47Δ-mAngio virus, G47Δ-mIL12 is also highly anti-angiogenic³³ due to the presence of anti-angiogenic cytokine IL-12. G47Δ-mIL12 in combination with a VEGFR tyrosine kinase inhibitor axitinib significantly prolonged survival of mice bearing mouse or human GSC-derived

GBM tumors compared to monotherapies via multifaceted mechanisms: direct oncolysis, increased macrophage recruitment, reduction of CD34⁺ tumor vascularity, and inhibition of PDGFR/ERK pathway in an immunodeficient model, and T cell-dependent efficacy in an immunocompetent model.¹²⁴ No additional benefit was observed when anti-CTLA4 was added to the combination therapy (G47Δ-mIL12+axitinib).¹²⁴

Manipulation of DNA Damage Response (DDR)

oHSV G47Δ induces DDR and synergizes with DNA-damaging agents such as TMZ.¹¹⁶ Poly (ADP-ribose) polymerase (PARP) plays a critical role in DNA repair. Ning et al¹²⁸ demonstrated that PARP activity is uniformly inhibited by PARPi (such as Olaparib) in both PARPi-sensitive and PARPi-resistant GSCs. Importantly, oHSV synergistically interacted with PARPi and induced cytotoxic effects against GSCs, irrespective of their sensitivity to PARPi, through proteasomal degradation of DDR proteins, Rad51 and Chk1. The combination (Olaparib + oHSV-MG18L) significantly extended the survival of mice bearing PARPi-sensitive or PARPi-resistant GSC-derived tumors compared to monotherapies. This combination strategy is unique and should also be effective to other solid tumors, irrespective of their sensitivity to PARPi.

Inhibition of Histone Deacetylase (HDAC)

Otsuki et al¹²⁹ demonstrated that treatment of glioma cells with valproic acid (VPA), a commonly used antiepileptic agent that inhibits HDAC (an enzyme that controls cancer cell survival, progression, and differentiation¹³⁰), prior to oHSV infection inhibited induction of anti-viral interferon genes, resulting in increased transcription of viral genes and improved virus replication in vitro. VPA pre-treatment also increased oHSV propagation and oHSV-mediated anti-tumor efficacy in vivo in GBM xenograft model.¹²⁹ VPA can also enhance oHSV efficacy through suppression of anti-viral innate immunity.¹³¹ For instance, VPA administration prior to oHSV therapy reduced NK cell activity against oHSV-infected GBM cells through inhibition of STAT5/T-bet signaling and IFN-γ production.¹³¹ Like VPA, tubacin (an HDAC6-specific inhibitor) also increased rQNestin34.5 replication in GSCs by counteracting antiviral effects of type I interferons and by altering

the post-entry trafficking of oHSV toward nuclei rather than to the lysosome,¹³² and improved survival in an orthotopic GSC-derived xenograft model.¹³²

Inhibition of Proteasome

Proteasomes are cellular protein complexes that degrade damaged proteins through proteolysis. Proteasome inhibition can result in the induction of unfolded protein response (UPR) and endoplasmic reticulum (ER) stress, and thus, it offers a way to target GBM.^{133,134} Bortezomib (a proteasome inhibitor) induces ER stress and UPR, which contributed to significantly enhanced oHSV (34.5ENVE) replication and synergistic killing of glioma cells.¹³⁵ HSP90 was found to be the most critical UPR for increased oHSV replication. In an intracranial human GBM model, the combination therapy dramatically enhanced survival compared to monotherapies.¹³⁵ In a follow-up study,¹³⁶ the same group demonstrated that bortezomib pre-treatment of tumor cells enhanced oHSV-induced necroptotic cell death through induction of RIPK1 and mitochondrial reactive oxygen species (ROS). This coincided with an increased secretion of IL1α from combination-treated tumor cells.¹³⁶ To confirm whether IL-1α sensitizes tumor cells to natural killer (NK) cells,^{137,138} a co-culture study (treated brain tumor cells and naïve NK cells) was conducted and revealed a significantly increased secretion/expression of killer cytokines (IFN-γ and TNF-α) and apoptotic markers (cleaved caspase 3, 8 and cleaved PARP) in the combination group than single treatment groups. In vitro findings were also reproduced in vivo and it was reported that the addition of an NK cell-based adjuvant therapy to the combination further significantly enhanced the efficacy.¹³⁶

Blockade of TGF-β Signaling

Transforming growth factor-beta (TGF-β) plays a critical role in GBM pathogenesis and maintaining GSC stemness.¹³⁹ In a TMZ-resistant GSC-derived recurrent GBM model, the combination of oHSV (MG18L) and TGF-β receptor kinase inhibitor (TGF-βi) synergistically killed GSCs in vitro and produced 60% long-term survivors in vivo (vs 0% with monotherapies).¹⁴⁰ In clinical setting, galunisertib (a TGF-βi) in combination with TMZ/radiotherapy was not effective,¹⁴¹ therefore, further pre-clinical research is needed to understand/optimize the therapeutic potential of TGF-βi in the context of combination therapies.

Inhibition of Notch Signaling

Notch signaling pathway is important for cell–cell interactions, tumor progression, GSC maintenance, and treatment resistance.¹⁴² oHSV infection of tumor cells activated notch in adjacent tumor cells, and miR-H16 was found to be responsible for notch induction through downregulation of FIH1 (a direct target of miR-H16).¹⁴³ Notch inhibition, prior to oHSV treatment, by a gamma-secretase inhibitor (GSI) (RO4929097) resulted in increased killing of glioma cells and improved survival compared to monotherapies.¹⁴³ Several GSIs were evaluated in the clinic, and hence, oHSV + GSI combination has translational significance.

Inhibition of Integrin Signaling

Integrin $\beta 1$ (ITGB1), also referred to as CD29, plays an important role in tumor progression¹⁴⁴ and resistance to bevacizumab.¹⁴⁵ Blockade of ITGB1 with OS2966 (a humanized CD29 blocking antibody) overturned bevacizumab-induced resistance.¹⁴⁵ Although OS2966 enhanced oHSV replication and GBM cell cytotoxicity in vitro,¹⁴⁶ systemic OS2966 delivery did not enhance the anti-tumor efficacy of oHSV (rHSVQ1-IE4/5-Luciferase) in vivo. Insufficient delivery of OS2966 to the brain may have contributed to the failure to obtain a combination effect. To overcome this issue, OS2966 was directly injected intracranially into GBM xenograft 2 days prior to oHSV treatment, and this time, the combination significantly extended survival compared to monotherapies.¹⁴⁶ In contrast to OS2966, systemic delivery of cilengitide (another integrin inhibitor) significantly enhanced the anti-glioma efficacy of RAMBO compared to single treatments.¹⁴⁷ Cilengitide has been evaluated in GBM patients¹⁴⁸ and could be a better choice than OS2966 for combination studies.

Inhibition of NK Cell-Related Anti-Viral Response

Intra-tumoral oHSV treatment recruits NK cells within hours of infection,¹⁴ which substantially limits oHSV efficacy.¹⁴⁹ Han et al¹⁵⁰ demonstrated that TGF- $\beta 1$ (an immunosuppressive cytokine) pre-treatment impaired NK's anti-viral activity, leading to significantly increased virus yield from infected glioma cells in a co-culture study. In an orthotopic U87 Δ EGFR model, systemic delivery of TGF- $\beta 1$ prior to intracranial oHSV treatment significantly increased oHSV titers compared oHSV monotherapy, and this TGF- $\beta 1$ -mediated oHSV replication was dramatically decreased following treatment with 1D11 (a TGF- β

neutralizing antibody). In a syngeneic 4C8 GBM model, a single dose of TGF- β prior to oHSV therapy inhibited recruitment/activation of NK cells (and macrophages), leading to efficient tumor control and extension of survival.¹⁵⁰ NK depletion also enhanced oHSV efficacy, which was similar to the observed efficacy with TGF- $\beta 1$ + oHSV combination.¹⁵⁰ This study did not test whether transient inhibition of innate immunity has had any detrimental impact on adaptive immunity.

Inhibition of Macrophage-Mediated Anti-Viral Immunity

Infiltrating macrophages/resident microglia following HSV infection^{151,152} induce an early anti-viral response,¹⁵³ including the production of TNF- α ,¹⁵⁴ which negatively influences virus replication in glioma cells and oHSV (rQNestin34.5)-induced efficacy in an intracerebral U87 Δ EGFR model.¹⁵¹ Transient TNF α blockade abrogated the negative effects of TNF- α , and thereby, increased oHSV replication and anti-GBM efficacy in a xenograft model.¹⁵¹ Pointer et al¹⁵⁵ and Yoo et al¹⁵⁶ also reported potentiation of oHSV virotherapy in GBM by TNF- α inhibition using a TNF- α blocking antibody or a MEK kinase inhibitor (trametinib), respectively. These studies suggest that FDA-approved TNF- α inhibitors¹⁵⁷ can be utilized to improve oHSV efficacy.

Cysteine-rich 61 (CCN1), an extracellular matrix protein, is upregulated in GBM cells in response to oHSV infection.¹⁵⁸ Virally enhanced CCN1 activates type I IFN response¹⁵⁹ and increases macrophage migration/activation that contributes to viral clearance.¹⁶⁰ Anti-CCN1 neutralizing antibody nullified the anti-viral effects of CCN1 and improved oHSV efficacy in vivo, compared to control.¹⁶⁰ Besides CCN1 activation, recently, Delwar et al reported limited oHSV replication in macrophages/microglia due to STAT1/3 phosphorylation, which eventually formed a non-permissive barrier and prevented oHSV dissemination in the glioma TME.¹⁶¹ STAT1/3 inhibition by C16, a derivative of oxindole/imidazole, rescued oHSV-1 replication, both in vitro and in vivo.¹⁶¹

Immune Checkpoint Inhibition

GBM is a highly immunosuppressive tumor and does not respond to immune checkpoint inhibitor (ICI).^{75,162} Immunosuppression in GBM is governed by various mechanisms¹⁶³ including, but not limited to, T-cell expression of co-inhibitory molecules such as PD-1, PD-L1, CTLA-

4, etc.^{164–168} Inhibition of PD-1/PD-L1 and/or CTLA-4/B-7 interaction can be a critical turning point to improve oHSV efficacy.^{152,169,170} In ICI-resistant GSC-like GBM models (005 and CT-2A),^{152,163,171–174} intra-tumoral IL-12 expression by G47Δ-mIL12 dramatically sensitized tumors to ICIs (anti-PD-1 + anti-CTLA-4), leading to 50–89% long-term survivors and memory protection.¹⁵² This remarkable therapeutic efficacy was associated with macrophage polarization and total dependence on CD4⁺ T cells.¹⁵² ICI therapy also demonstrated synergy with oHSV^{ULBP3,95} as evidenced by inhibition of contralateral tumor growth and extension of survival of mice bearing bilateral ICI-resistant XFM-Luc:PDGF, Cre IDH wild-type GBM tumors.⁹⁵

Inhibition of the Mammalian Target of Rapamycin (mTOR)

mTOR is a key mediator of EGFR/PI3K/Akt signaling that regulates cell growth, proliferation, and survival via two functionally and structurally distinct multiprotein complexes known as mTORC1 and mTORC2.^{175,176} GBM patients treated with rapamycin, a solely allosteric inhibitor of mTORC1, exhibited a high rate of resistance and worse progression due to mTORC1-mediated feedback and mTORC2 activation.¹⁷⁵ Interestingly, rapamycin was previously shown to enhance oHSV production and spread in several difficult-to-treat human tumors.¹⁷⁷ Recently, a new generation of mTOR inhibitor, ATP-competitive active-site mTORC1 and 2 inhibitors (asTORi) enhanced ICP0-deleted HSV1 infection specifically in transformed cells (such as human GBM cells) via inhibition of cellular type-I IFN responses, while reducing infection in non-transformed cells.¹⁷⁸ The promising outcome of this combination study warrants further pre-clinical research in vivo in GBM models and clinical translation in GBM patients.

Other Combination Therapies

In addition to the above-described combination studies, many other oHSV-based combination therapies were described in the past.¹⁷⁹ For example, combining oHSV with ionizing radiation,¹⁸⁰ PI3K/Akt pathway inhibitors,¹⁸¹ etoposide¹⁸² or HMGB1 blockade¹⁸³ significantly enhances anti-tumor efficacy in preclinical GBM models.

Clinical Evaluation of oHSVs in GBM and Other Brain Malignancies

Six oHSVs (1716, G207, G47Δ, M032, C134, and rQNestin34.5v.2) have been or are being tested in patients

with brain tumors (Table 4).^{5,10,184,185} Early trials with HSV1716 demonstrated no evidence of HSV-mediated adverse events or toxicities.^{186–188} In the very first study, nine recurrent high-grade glioma (HGG) patients received a stereotactic injection of HSV1716, with doses escalating from 10³–10⁵ plaque-forming unit (pfu) in 1 mL.¹⁸⁷ Four patients survived longer than 14 months following treatment.¹⁸⁷ No evidence of shedding or reactivation of HSV was observed.¹⁸⁷ In the second study, 12 HGG patients were intratumorally injected with 10⁵ pfu of HSV1716.¹⁸⁸ Tumors were resected 4–9 days after treatment. Infectious HSV was recovered from injected sites in two patients. The viral DNA was detected in 10 patients at the injected site and in 4 patients at the distal tumor site.¹⁸⁸ An immune response (ie, changes in the levels of IgG and IgM) to HSV1716 was observed in two patients. Overall, this study confirmed the safety and replication of HSV in HGG tumors. In the third study, 12 recurrent or newly diagnosed HGG patients first underwent maximal surgical resection and then received HSV1716 (10⁵ pfu) into 8 to 10 sites in the cavity wall.¹⁸⁶ Three patients were clinically stable for 15–22 months following virotherapy. Like the second trial, two of three seronegative patients were seroconverted. Imaging demonstrated a reduction of the residual tumor over a period of 22 months.¹⁸⁶ A Phase I trial using HSV1716 in pediatric patients with surgically removable refractory or recurrent HGG was terminated in 2016 and no results of this trial were posted thus far (NCT02031965).

The safety and efficacy of G207 were reported preclinically.¹⁸⁹ Four Phase I/Ib clinical studies involving G207, either alone or in combination with radiation, were reported in patients with malignant glioma or recurrent GBM.^{101,184,190,191} In the first study, which was the first oHSV clinical trial in the US, 21 recurrent HGG patients received an intra-tumoral injection of G207 up to 3 × 10⁹ pfu at five sites.¹⁹⁰ G207 was safe with no virus shedding observed in saliva or conjunctival secretions. Four patients survived 7–19 months following virus inoculation.¹⁹⁰ Next, a Phase Ib trial was initiated in six recurrent GBM patients who received two doses of G207 totaling 1.15 × 10⁹ pfu, with 13% of this dose injected stereotactically into the tumor, followed by *en bloc* tumor resection and administration of the remaining virus dose at multiple sites into the resected cavity wall. Replication and antitumor activity (ie radiographic and neuropathologic response) of G207 were reported.¹⁹¹ Markert and Colleague published a case report demonstrating that G207 treatment (1 × 10⁷

pfu in 120 μ L volume stereotactically injected into the enhancing region of the resected tumor) with minimal adjunctive chemotherapy in a recurrent GBM patient led to 6 years of disease-free survival, indicating a long-term response.¹⁹² G207 was also tested in combination with irradiation in a Phase I clinical trial in recurrent GBM.¹⁰¹ Nine progressive recurrent malignant glioma patients (seven of them bearing GBMs) were stereotactically/intra-tumorally treated with G207 (1×10^9 pfu) 24 hours prior to single focal 5 Gy radiation. The combination therapy was well tolerated and no patients developed HSV-related encephalitis. Three instances of marked radiographic response to treatment were observed.¹⁰¹ Based on the safety/efficacy of three published G207 studies^{101,190,191} and the high sensitivity of pediatric brain tumors to killing by G207,¹⁹³ the same group designed a clinical trial protocol to evaluate G207, either alone or in combination with radiation, in six pediatric brain tumor patients (five GBM and one anaplastic astrocytoma).¹⁹⁴ Patients were safely implanted with 3–4 intra-tumoral silastic catheters for delivery of G207.¹⁹⁵ Recently, the same group has also designed a first-in-human immunotherapy Phase I clinical trial to treat pediatric recurrent cerebellar malignancies and a Phase II trial to treat pediatric recurrent HGG with G207 combined with single low-dose radiation (5 Gy). No results are available from these studies yet¹⁹⁶ (NCT04482933).

In comparison to G207, G47 Δ (which is constructed from G207 by adding a deletion of ICP47 gene) is a relatively newer version of unarmed oHSV. G47 Δ was found safe in Phase I–IIa clinical trial in 21 patients with recurrent GBM (UMIN000002661).¹⁹⁷ In a follow-up Phase II clinical trial in patients with residual or recurrent GBM (UMIN000015995), G47 Δ demonstrated anti-tumor efficacy. In this study, G47 Δ (1×10^9 pfu as used for G207 in)¹⁰¹ was stereotactically injected into the tumor twice within 2 weeks and every 4 weeks thereafter for a maximum of 6 times. The treatment was well tolerated.¹⁹⁷

Like the G47 Δ virus, the safety and efficacy of C134 and rQNestin34.5v.2 are also being tested in recurrent GBM patients (NCT03657576 and NCT03152318).^{5,6} Since M002 (an oHSV expressing murine IL-12) was efficacious in mouse GBM models,^{198–200} the same group generated M032 virus (the clinical version of M002 but expressing human IL-12) and demonstrated its safety after intracerebral administration to HSV-sensitive non-human primates.⁵⁵ M032 is currently in a clinical trial in patients with recurrent or progressive GBM.²⁰¹ The

promising safety profile and preliminary signs of efficacy exhibited by several oHSVs in patients warrant a further clinical evaluation in a larger patient cohort.

Current Challenges and Potential Remedies

FDA approval of T-VEC in 2015 and its promising efficacy in non-GBM cancers have fueled the field of oncolytic virotherapy. Since then, oHSV therapy for GBM has evolved rapidly preclinically. Currently, several oHSVs are under clinical trial evaluation and early trial results demonstrated safety and signs of efficacy in GBM patients (Table 4). Despite significant advancement, there are various treatment challenges to oHSV exist, which include but are not limited to: suboptimal viral delivery, insufficient viral entry/replication, limited oHSV spread, and host anti-viral immune response (Figure 4).^{4,14} As discussed above, although direct oHSV injection into the tumor is the most commonly used viral delivery method, it is not ideal for tumors that are located in an eloquent area of the brain or when the tumor is metastatic. In addition, the risks associated with neurosurgical procedures make repeated virus administration difficult. Systemic intravascular delivery can be useful in these scenarios, but systemic injection has also a significant number of drawbacks, such as (i) loss of virus through liver metabolism; (ii) potential risk of off-target distribution/replication; (iii) serum/complement factors can neutralize oHSVs; (iv) BBB can limit viral entry to the tumor, etc. To overcome such limitations, other approaches were explored, such as intraventricular injection^{24,190,202} and carrier cell delivery.^{106,107,110–112} GBM has aberrant vasculature and its outgrowth can destroy BBB.²⁰³ Intraventricular method may provide viral access to the tumor without the need for repeated invasive surgical procedures. However, this approach requires further preclinical validation to assess its risk and effectiveness. Delivery of oHSV by MSCs could be another potential treatment approach, which avoids serum inhibitory factors and demonstrated promising efficacy preclinically. Carrier cell-based approach is under at developmental stage and needs to be extensively tested in immunocompetent/immunodeficient GBM models to determine its utility for patients.

Following viral delivery, the next important step for effective treatment response is efficient viral entry/replication. HSV uses heparan sulfate proteoglycans, herpes virus entry mediators, and nectin-1 for its entry into host cells,²⁰⁴ and thus, their expression on tumor cells may

predict the treatment efficacy. Recently, Friedman et al demonstrated that sensitivity of pediatric brain tumor cells to oHSVs (G207, M032) was correlated with high nectin-1 expression,⁵⁸ indicating receptor expression profile is critical for optimal efficacy. Based on the abundance of tumor cell surface receptors (such as nectin-1, EGFR and HER2), re-targeted oHSVs can be used to enhance viral entry to cancer cells.^{42,46,47,49,51} Although re-targeting strategy looks promising, new oHSVs with advanced combinations of targets will be needed to address intra- or inter-tumoral heterogeneity. In addition, re-targeted oHSVs might produce off-target effects on healthy cells, since the same receptors (EGFR and HER2) can be shared with non-cancerous cells.^{205,206} To overcome this issue, re-targeted oHSVs can be further modified to allow virus replication selectively in GBM cells that have distinguished miRNA expression profile compared to normal/healthy brain cells.^{49,51}

Another important issue that needs to be addressed before utilizing the full potential of oHSVs is maximizing oHSV replication/spread. The anti-viral innate immunity and condensed extracellular matrix play important role in limiting viral replication/spread. Various approaches have been employed to address these issues such as the development of HCMV/HSV – a chimeric oHSV that replicates in a similar fashion as wild-type virus,^{36,38–41} or generation of armed oHSVs that express E-cadherin,⁸⁴ chondroitinase ABC,⁸⁸ or MMP-9⁹⁶ to enhance cell-cell oHSV spread, or transient inhibition of anti-viral innate immunity prior to oHSV infection to improve virus replication and spread.^{150,151,155,156,160,161} It is still not completely clear how the host's innate resistance mechanisms/pathways (eg IFN signaling pathways and cGAS/STING pathway) limit oHSV replication/spread in cancer cells/TME or whether inhibition of innate (macrophage- or NK-related) immune pathways^{150,151,155,156,160,161} will somewhat compromise adaptive anti-tumor immunity. Further understanding of the host's anti-viral signaling pathways will help to design new combination strategies for GBM.

Successful viral delivery, entry, replication and spread should eventually activate the host's adaptive anti-tumor immunity, leading to recruitment of T cells to the TME. However, recruited T cells are often counter-balanced by the expression of the exhaustion markers such as PD-1. Anti-PD-1 monotherapy failed to reach intended treatment goals in GBM (Phase III CheckMate-143),^{75,207} illustrating that poor tumor immunogenicity and immunosuppression led to treatment-insensitive lesions.^{207,208} In advanced melanoma

patients, T-VEC promoted intratumoral T cell infiltration and sensitized tumors to anti-PD-1.¹⁷⁰ A combination of ICIs and oHSV was shown to eradicate the intracranial tumor in a murine GSC model,¹⁵² indicating the potential promise of this combination approach for poorly immunogenic GBMs. Even though T-VEC + ICI combination trials are currently running in non-GBM cancers, more research is needed to optimize new viral vectors and design more rationale combination clinical trials for GBM. This may include the generation of new oHSVs expressing immune modulators (eg, IL-2, CD40L, OX40L and 4–1BBL) and testing them in combination with ICIs or other anti-cancer therapies. Viral oncolysis exposes tumor antigens to dendritic cells (DCs); thus, new oHSVs can also be combined with tumor antigen-loaded DC-based vaccine to provoke a tumor-specific T-cell response.²⁰⁹

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The authors report no conflicts of interest in this work.

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