Review

ELAVL1 Role in Cell Fusion and Tunneling Membrane Nanotube Formations with Implication to Treat Glioma Heterogeneity

Natalia Filippova * and Louis B. Nabors *

Department of Neurology, School of Medicine, University of Alabama at Birmingham, Birmingham, AL 35294, USA

* Correspondence: as1999@uab.edu (N.F.); bnabors@uab.edu (L.B.N.); Tel.: +1-205-934-2544 (N.F.); +1-205-934-1432 (L.B.N.)

Received: 24 September 2020; Accepted: 18 October 2020; Published: 21 October 2020

Simple Summary: Despite the numerous novel pharmacological and immunological approaches for multimodal glioma treatments that have been proposed in recent years, glioma phenotypic and genotypic spatial profiles in the course of treatments remain heterogeneous and, therefore, represent the biggest challenge for patient outcome. The elimination of glioma heterogeneity is an established chemotherapeutic goal. The role of the mRNA-binding protein of ELAV-family HuR in homotypic and heterotypic cell fusions via permanent intercellular membrane fusions and temporal intercellular tunneling nanotube formations in the glioma microenvironment leading to glioma heterogeneity will be discussed in our review with implications of HuR inhibitors in the prevention of these processes.

Abstract: Homotypic and heterotypic cell fusions via permanent membrane fusions and temporal tunneling nanotube formations in the glioma microenvironment were recently documented in vitro and in vivo and mediate glioma survival, plasticity, and recurrence. Chronic inflammation, a hypoxic environment, aberrant mitochondrial function, and ER stress due to unfolded protein accumulation upregulate cell fusion events, which leads to tumor heterogeneity and represents an adaptive mechanism to promote tumor cell survival and plasticity in cytotoxic, nutrient-deprived, mechanically stressed, and inflammatory microenvironments. Cell fusion is a multistep process, which consists of the activation of the cellular stress response, autophagy formation, rearrangement of cytoskeletal architecture in the areas of cell-to-cell contacts, and the expression of proinflammatory cytokines and fusogenic proteins. The mRNA-binding protein of ELAV-family HuR is a critical node, which orchestrates the stress response, autophagy formation, cytoskeletal architecture, and the expression of proinflammatory cytokines and fusogenic proteins. HuR is overexpressed in gliomas and is associated with poor prognosis and treatment resistance. Our review provides a link between the HuR role in the regulation of cell fusion and tunneling nanotube formations in the glioma microenvironment and the potential suppression of these processes by different classes of HuR inhibitors.

Keywords: cell-to-cell communication; tumor microenvironment; glioma; HuR; heterogeneity; tunneling membrane nanotubes; cell fusion; inhibitors

1. Introduction

Glioma is the most devastating and incurable disease with a relative risk of 3–4 timers higher in the elderly population than young adults. It is characterized by high tissue heterogeneity and undergoes a fast transformation from low-grade (I–II) to high-grade (III–IV) malignancy [1–3]. The transition to a high-grade tumor occurs in approximately 95% of low-grade gliomas; less than 3% of these patients remain alive after five years post-diagnosis [3,4].
Comprehensive genomic profiling of low- and high-grade gliomas confirmed an increase in glioma heterogeneity with tumor grade and treatment resistance. Glioma heterogeneity represents the biggest challenge for glioma treatment, which depends on tumor grade and is customized to individual patients. Surgery, radiotherapy, and chemotherapy with temozolomide (TMZ) represent the first line of treatments for glioma [5,6]. More modern treatments consist of treatment with the genetically engineered autologous or allogeneic T cells expressing chimeric antigen receptors (CARs) directed against specific antigens, the combinations of Optune treatment (an application of the electric field, which induces apoptosis in rapidly dividing tumor cells) and TMZ, the combination of the ketogenic diet with glutamine antagonist 6-diazo-5-oxo-l-norleucine, which induces energy depletion and destroys the mesenchymal type of glioma stem cells [7–9]. However, due to tumor heterogeneity, genomic instability, and transforming ability, the current tumor treatments are efficient only in terms of the prolonging time of relapse; therefore, gliomas remain incurable [1–3,5].

The needs in new glioma treatment modalities, which eliminate the development of tumor heterogeneity and transforming ability, are evident. Intercellular gene transfers between glioma and normal host cells in the glioma microenvironment via permanent cell fusion and temporal tunneling tube formations have been recently discovered and may lead to a de novo glioma genotype and phenotype inside of the progressing tumor. Our review overviews mechanisms and cell-signaling pathways, which lead to cell fusion and tunneling membrane tube formations in the glioma microenvironment and discusses pharmacological approaches of the interception of these processes.

2. Historical Background

Cell fusions via permanent intercellular membrane fusions were discovered more than a century ago under normal physiological conditions such as tissue and organ development, cell fertilization, tissue repair and regeneration, and angiogenesis [10,11]. Only select types of mammalian cells (to include bone marrow-derived mesenchymal stem cells, bone marrow-derived endothelial progenitor cells recruited to the new blood vessels, epithelial cells of the placenta, fiber cells in the eye, macrophages at sites of chronic inflammation, skeletal muscle cells, and gametes) undergo permanent intercellular membrane fusion under normal physiological conditions [12]. The temporal cell fusions via tunneling membrane nanotube formations between mammalian cells were first discovered by Rustom and colleagues in 2004 [13] and occur between different cell types (to include monocytes, lymphocytes, neurons, astrocytes, cardiomyocytes, endothelial cells, fibroblasts, and mesenchymal stem cells under diverse stress conditions) [13–19]. Tunneling membrane nanotubes vary in length from 5 to 100 µm, and in width from 50 nm to up to 1 µm. Depending on the interconnected cell types and the tunneling nanotube compositions and sizes, the following terms have been utilized to reference the tunneling membrane tube structures: tunneling membrane nanotubes, cytoplasmic intercellular bridges, intercellular nanotubular highways, membrane microtubes [20]. The tunneling membrane nanotubes are not adherent to the substrate and have a lifetime from a couple of minutes to several hours. Under normal physiological conditions, bidirectional intercellular transfers of cellular materials via permanent cell fusion and temporal tunneling nanotube formations lead to tissue development and repair [17–19]. However, under pathological conditions such as viral infection, chronic inflammation, diabetes, and cancer, the intercellular transfers of biological materials (including chromosomal and extrachromosomal genetic materials, cellular organelles, viruses) via permanent and temporal cell fusions lead to de novo cell transformations and tissue heterogeneity, and thus to the development of drug resistance and disease progression [20–26].

3. Expression of Fusogens, Fusogen Receptors, and Tunneling Membrane Nanotube-Related Transcripts in Gliomas

Both intercellular tunneling nanotubes and permanent intercellular membrane fusions are reported in gliomas and represent diverse multistep processes, which require an activation of the cellular stress response, the rearrangement of the actin-dependent cytoskeleton, the expression of the fusogenic
The type 2 tunneling nanotube transcripts (GJA1, GAP43) [40,41], which overlap with gap junction proteins, and phosphatidylserine enrichment on the membrane surfaces [24–35]. We utilized the viruses; (ii) the fusogen transcripts encoding endogenous retroviral envelope proteins (ERVW-1, ERVK3-1, ERV3-1, ERVMER34-1, ERVV-1, ERVFDRD-1); (iii) the fusogen transcripts encoding proteins essential for sexual reproduction and gamete fusions (transcripts of IZUMO and IZUMOR families, GLIPR1L1, CD9); (iv) the muscle-specific fusogen transcripts (myomaker and myomixer) at low levels; (v) the transcripts of fusogens involved in intercellular and extracellular vesicle-specific transfers (SNARE-family transcripts, transcripts of small dynamin-like GTPases including atlastins, mitofusins, dynamins). The type 1 tunneling membrane nanotube-related transcripts (TNFAIP2, S100A4, ERp29) [36–39] were significantly overexpressed in gliomas compared to normal brain. The type 2 tunneling nanotube transcripts (GJA1, GAP43) [40,41], which overlap with gap junction formations, exhibited a diverse expression in gliomas (GJA1 transcript was significantly overexpressed; however, GAP43 expression was significantly decreased compared to normal brain samples). It will be important to mention that the higher tumor grades correlated with the significant increase in the TNFAIP2 and S100A4 transcript expression. The Ttyh1 transcript, a biomarker of the recently identified the Ttyh1-dependent subset of the tunneling microtubes with widths up to 2.5 μm [32,42], exhibited significant overexpression in gliomas compared to normal brain. However, it was reported that Ttyh1 is downregulated in 1p/19q-co-deleted versus 1p/19q intact human gliomas by utilizing RNA-Seq gene-expression analysis [32]. Because of deficient Ttyh1-dependent tunneling microtube formation and function, 1p/19q-co-deleted oligodendroglomas have been reported as invasion-deficient [43], suggesting the diverse roles of tunneling nanotube and tunneling microtubes in gliomas with different genetic backgrounds [32,43]. Figure 1 illustrates the expression of fusogens, fusogen receptors, and the tunneling membrane nanotube type 1- and type 2-related transcripts in gliomas normalized to the expression of the corresponding transcripts in the normal brain and the corresponding expression of the house-keeping gene ACTB (actin).

**Figure 1.** Expression of fusogens and the tunneling membrane nanotube-related transcripts in gliomas compared to normal brain. Graphs illustrate normalized expression of the fusogens and their receptors’ transcripts (left) and the tunneling membrane nanotube-related transcripts (right) in gliomas compared to normal brain. The following studies from the R2: platform have been utilized for the analysis: Rembrandt, Madhavan, Mas 5.0-U132p2 study for gliomas and Harris, Mas 5.0-U132p2 study for normal brain. Data are shown as mean ± SD. Detailed methods associated with transcriptic profiling were provided in the R: platform (Rembrandt, Madhavan, gliomas and Harris, normal brain studies).
The publicly available transcriptomic data (RNA-seq) of five patient-derived glioma xenograft (PDGx) neurosphere cell lines of classic, proneural, and mesenchymal subtypes presented in the Gene Expression Omnibus (GEO) repository database (GSE158271) were utilized for the analysis of the expression of the fusogens and the tunneling nanotube-related transcripts in gliomas of different subtypes. The analyses of the transcriptome signatures of five PDGx neurosphere cell lines confirmed the expression of a variety of fusogen and tunneling membrane nanotube-related transcripts in all glioma subtypes. Tables 1 and 2 provide a summary of the normalized to the house-keeping gene ACTB transcript expression in the different glioma subtypes.

Table 1. Expression of the fusogens and the fusogen receptor transcripts in PDGx neurospheres of different subtypes: proneural (PDGx1, PDGx2), classic (PDGx3, PDGx4), and mesenchymal (PDGx5).

| Endogenous Retroviral Fusogens | PDGx1 | PDGx2 | PDGx3 | PDGx4 | PDGx5 |
|-------------------------------|-------|-------|-------|-------|-------|
| ERVK13-1                      | +++   | +++   | +++   | +++   | +++   |
| ERV3-1                        | ++    | +++   | +++   | +++   | ++    |
| ERVMER34-1                    | +     | +     | +     | +     | none  |
| EBI3                          | none  | none  | +     | none  | ++    |
| ERVW-1                        | +     | none  | none  | none  | none  |
| ERVV-1                        | ++    | ++    | ++    | ++    | ++    |
| ERVV-2                        | none  | none  | none  | none  | none  |
| Fusogen Receptors             |       |       |       |       |       |
| CTNNB1                        | +++   | +++   | +++   | +++   | +++   |
| ICAM1                         | none  | none  | none  | none  | none  |
| ITGA4                         | none  | ++    | ++    | ++    | ++    |
| ITGB4                         | +     | none  | none  | none  | none  |
| VCAM1                         | none  | none  | ++    | ++    | ++    |
| Germ Cell Fusogens            |       |       |       |       |       |
| CD9                           | +++   | +++   | +++   | +++   | ++    |
| IZUMO1                        | +     | +     | +     | +     | none  |
| IZUMOR1                       | +     | +     | +     | +     | none  |
| IZUMO3                        | none  | none  | none  | none  | none  |
| IZUMO4                        | ++    | ++    | ++    | ++    | +     |
| SPA17                         | ++    | ++    | ++    | ++    | +     |
| SPAG8                         | +     | +     | +     | +     | +     |
| Myoblast Fusion Factors       |       |       |       |       |       |
| MYMK                          | none  | none  | none  | none  | none  |
| MYMX                          | none  | none  | +     | none  | none  |

Table 2. Expression of the tunneling nanotube-related transcripts in PDGx neurospheres of different subtypes: proneural (PDGx1, PDGx2), classic (PDGx3, PDGx4), and mesenchymal (PDGx5).

| TNT Type i and Type ii Transcripts | PDGx1 | PDGx2 | PDGx3 | PDGx4 | PDGx5 |
|-----------------------------------|-------|-------|-------|-------|-------|
| TNFAIP2                           | +++   | ++    | ++    | +++   | ++    |
| RALA                              | +++   | +++   | +++   | +++   | ++    |
| EXOC6                             | ++    | ++    | +++   | +++   | ++    |
| SI100A4                           | +++   | +     | ++    | ++    | +     |
| GAP43                             | +++   | +++   | +++   | +++   | none  |
| GJA1                              | +++   | +++   | +++   | +++   | +     |
Detailed methods associated with transcriptomic profiling were provided in the Gene Expression Omnibus repository database (GSE158271). The expression levels were defined as (i) strong (+++) if the expression was in the 70–100% range of the maximum expression of the corresponding transcripts; (ii) medium (+++) if the expression was in the 30–70% range of the maximum expression of the corresponding transcripts; (iii) weak (+) if the expression was in the 1–30% range of the maximum expression of corresponding transcripts.

The clinical outcome data of patients with gliomas harboring low or high expressions of different fusogens and the tunneling membrane nanotube-related transcripts were obtained from Rembrandt, Madhavan-550 MAS.5.0-U132p2 study, R2: platform. The analyses of the overall survival of patients with gliomas harboring a low or high expression of the TNF Alpha Induced Protein 2 (TNFAIP2) transcript indicates that high expression levels of TNFAIP2 were associated with poor prognosis ($p = 1.2 \times 10^{-4}$). Moreover, upregulations of the TNFAIP2-stabilizing chaperone ERp29 [38], the TNFAIP2-associated small GTPase RALA, and S100A4 protein, which is responsible for the directional intercellular mitochondria transfer through the tunneling membrane nanotubes [39], significantly worsened patient outcome ($p = 2.1 \times 10^{-9}$, $p = 3.3 \times 10^{-4}$, and $p = 7.4 \times 10^{-12}$, respectively). High levels of ERWV-1, ERVK3-1, ERVK13-1, and CD9 fusogens were associated with overall poor survival as well ($p = 1.5 \times 10^{-4}$, $p = 1.6 \times 10^{-3}$, $p = 0.04$, and $p = 2.3 \times 10^{-6}$, respectively). Figure 2 illustrates the Kaplan–Meier overall survival curves for patients with gliomas harboring low or high expression levels of the above transcripts with a significant influence on overall patient survival.

**Kaplan-Meier overall survival curves**

![Figure 2](image_url)

**Figure 2.** Expression of the tunneling membrane nanotube-related transcripts (TNFAIP2, S100A4, ERp29, RALA) and the fusogen transcripts (ERWV-1, ERVK3-1, ERVK13-1, CD9) significantly decreased overall survival of glioma patients. Graphs illustrate the Kaplan–Meier overall survival curves for patients with gliomas harboring low (red) or high (blue) expression of the corresponding transcripts (Rembrandt, Madhavan, Mas 5.0-U132p2 study from R2: platform, median cut-off modus). For all illustrated transcripts, the differences were significant, $p < 0.05$.

Thus, our analysis confirms the enrichment of fusogens and the tunneling membrane nanotube-related transcripts in gliomas and their negative impact on overall patient survival.
4. HuR-Dependent Cell-Signaling Pathways of Cell Fusion and Tunneling Nanotube Formations Leading to Glioma Heterogeneity

Hypoxia, mechanical stress, chronic inflammation, cytotoxic stress, and oncometabolites associated with free radical formations are reported to potentiate intercellular membrane fusion events, and these conditions are often associated with the glioma microenvironment [14,20–23,27,35,44–47]. The mRNA-binding protein of ELAV-family HuR is a valuable biomarker of brain tumor progression [48–51] and is involved in the regulation of the key cell-signaling pathways responsible for the inflammatory glioma microenvironment, the hypoxia-related stress response, the transitions of classic and proneural glioma subtypes to the mesenchymal subtype, the metabolic stress, and the reactive oxygen species (ROS) generation associated with D-2HG oncometabolite production in low-grade gliomas harboring single alleles with IDH1-R132H/C/S mutations [52–55]. HuR exhibits strong overexpression in gliomas and shuttles from the nucleus to the cytoplasm to stabilize and promote the transfer and translation of mRNA transcripts enriched with adenine/uridine motifs in 3'UTR [53–56]. Figure 3 summarizes the key transcripts, which positively correlate with the formation of tunneling membrane nanotubes and cell-to-cell fusion and are reported as being directly upregulated by HuR in gliomas.
The proinflammatory cytokines, including IL-1, IL-6, and TNF-alpha, increase the probabilities of lipid protrusions, essential for the cell-to-cell fusion, through the regulation of arachidonic acid (AA) and sphingolipid metabolisms [57,58]. Additionally, the cytosolic phospholipase A2-alpha (cPLA_2α), which is known as the key enzyme that catalyzes the membrane glycerophospholipids at the sn-2 position to form AA, is the direct HuR-mRNA target upregulated in gliomas [58–62]. COX-2 (an inducible form of the cyclooxygenase enzyme that catalyzes the first step in the synthesis of prostanoids) is overexpressed in gliomas and is an established HuR-mRNA target. COX-2, in combination with PGE2, may influence ROS generation and controls the cellular redox state; therefore, it impacts cell fusion and tunneling membrane nanotube formation [63,64]. Direct COX-2 mRNA stabilization by HuR leading to an increase in COX-2 expression has been demonstrated in breast carcinoma [65]; also, the constitutive overexpression of COX-2 in ovarian and colon cancers is the result of HuR overexpression [63,66]. The positive interplay between HuR and COX-2 has been reported in gliomas as well [67].

Based on the combination of histological, ultrastructural, and genetic evidences, several models have emerged and described fusion steps, the diversity of chemokines, and a variety of cytoskeleton–lipid interactions leading to the formations of cell-to-cell extended contact zones, invasive protrusions, and tunneling membrane nanotubes [10,68]. However, all models implicate the rearrangement of the actin-dependent cytoskeleton to match fusion machineries in the fusing cells. In most cases in mammalian cells, the cytoskeleton–lipid interactions during membrane fusion culminate with the road-like structures filled with bundles of parallel F-actin filaments, which allow the intercellular material transfer. Several lines of evidence suggest that the formation of these cytoskeletal intercellular roads is cytochalasin B-sensitive and employs several HuR-dependent chemokines, matrix metallopeptidase, GTPases, and cytoskeletal components (to include CXCL1, CXCL3, CXCL5, MMP9, ACTB, VIM, Ral11, RaLa, RalBP1, Exo84, Sec5) [31,34,36,69–74]. It will be worth mentioning that some forms of membrane fusion (such as in the hypoxia condition) require the autophagy-dependent degradation of macromolecular membrane complexes. HuR promotes autophagosome formation by regulating the expression of several essential autophagy-related gene products (to include ATG5, ATG7, ATG12, ATG16L1) [75,76]. Hence, we conclude that the mRNA-binding protein HuR may serve as a potential positive regulator of the intercellular membrane fusions on the molecular level in gliomas.

Glioma-to-stromal cell communication via intercellular gene transfer leads to gene reprogramming in fused cells and the creation of de novo tumorigenicity and plasticity of glioma stem cells. The following glioma/microenvironmental transfers of biological materials via permanent and temporal cell fusions have been identified: (i) the bidirectional acquisition of the whole genome between glioma stem cells and bone marrow mesenchymal cells, between glioma stem cells and endothelial cells or pericytes, between glioma stem cells and monocytes (macrophages and microglia), between glioma cells and neural stem cells, and between glioma stem cells and macrophages fused with T cells [28,29,77–83]; (ii) the mitochondria and the cargo vesicle transfers through the tunneling membrane nanotubes between glioma cells and reactive astrocytes, between glioma and neuronal cells, between endothelial cells and pericytes in the glioma microenvironment, between glioma cells and macrophages, between glioma cells by themselves [22,23,30–34,84]; (iii) viruses and viral genome transfers between glioma cells by themselves, and between T cells and macrophages fused with glioma cells [32,33,85]. It is worth mentioning that proinflammatory monocytes and macrophages infiltrate the glioma microenvironment in a HuR-dependent manner and positively contribute to cell fusion and tunneling nanotube formations, and therefore promote glioma plasticity, tissue heterogeneity, and angiogenesis [86,87]. Moreover, growth factor- and integrin-mediated HuR expression and nuclear/cytoplasmic translocation have been reported in different cell types, which commonly reside in the glioma microenvironment [88–90]. Hence, we suggest that HuR overexpression stimulates the proinflammatory cellular composition of the glioma microenvironment, which provides positive feedback for HuR overexpression and nuclear/cytoplasmic shuttling in glioma cells and is associated with cell fusion and tunneling nanotube formations, which favor the development of glioma heterogeneity and treatment resistance.
5. Potential Pharmacological Modulators of Cell Fusion

Currently, several classes of inhibitors of cell fusion are under development worldwide: (i) the entry inhibitors, also known as the antiretroviral drugs, which inhibit fusogenic protein function; (ii) the F-actin depolymerizing agents (such as cytochalasin B), which inhibit actin-dependent cytoskeletal rearrangements essential for formations of the tunneling membrane nanotubes and the intercellular transfer; (iii) the modulators of the membrane fluidity; (iv) the inhibitors of autophagy; (v) the inhibitors of epithelial–mesenchymal and proneural–mesenchymal transitions in gliomas. The FDA-approved group of the entry inhibitors related to antiviral drugs includes 46 compounds, which represent the fusion-inhibitors, the CCR5 antagonists, and the post-attachment inhibitors. Although the actin-modifying agents exhibited strong potency in the inhibition of cell fusion in vitro, the actin-polymerization or depolymerization inhibitors are not the viable therapeutics due to their overall cytotoxicity. There are only two FDA-approved cytoskeletal modulators: (i) HALAVEN (eribulin mesylate), which has a broad spectrum of antitumor effects and is mostly utilized for the treatment of inoperable liposarcoma and metastatic breast cancer [91]; (ii) methyl-β-cyclodextrin (MCD) and its derivatives, which, as it was shown recently, induce actin depolymerization [92]. Interferon, Resveratrol, Miltefosine, Perifosine, Filipin, MCD, Emodin, monounsaturated and polyunsaturated fatty acids (MUFAs and PUFAs, respectively), diets with linoleic acid, oleic acid, marine fish oils are believed to be the perturbators of the cell membrane fluidity and are utilized for cancer treatment or prevention [93]. Chloroquine or hydroxychloroquine (HCQ) is an FDA-approved inhibitor of autophagy; ROC-325, Lys05, and DQ661 are the newest most potent inhibitors of autophagy from preclinical trials [94,95]. Note that HALAVEN, the actin-modifying agent, prevents glioma transitions to the mesenchymal subtype induced by the hypoxic condition. Pirfenidone (an inhibitor of the TGFβ-activated pathways), Quetiapine (an inhibitor of the NF-kB cell-signaling pathways), Rifampin (an inhibitor of the Wnt/beta-catenin cell-signaling pathway), Naproxen (an inhibitor of the non-canonical Wnt signaling pathways), Itraconazole (an inhibitor of the Hh cell-signaling pathways), Metformin (an inhibitor of the mitochondrial respiratory chain complex one and an activator of AMPK kinase) are among the FDA-approved potential inhibitors of transition to the mesenchymal subtype [96]. Most of the above drugs are domain-, gene-, or pathway-specific and have to be utilized in combination with other drugs.

Cell fusion and tunneling nanotube formations are multistep processes, which require diverse sets of genes at different stages; therefore, many inhibitors of cell fusion are stage-dependent. For example, in the PC12 cells, cytochalasin B (the actin depolymerization drug) strongly interfered with the formation of the tunneling nanotubes, but hardly affected their numbers and stability after formation [31]. Wiskott–Aldrich syndrome protein (WASP)-deficient macrophages were able to resemble tunneling nanotubes structures; however, the material transfer in these structures was completely abolished [97]. Small GTPases, such as Ras, Cdc42, Rac1, and RhoA, have been reported to be involved in actin remodeling and tunneling nanotube formations in different cell types. The gene-specific knockdown or inhibition of the individual GTP-binding transcripts usually led to a decrease in tunneling nanotube formation by 20–40%, and rarely by 75–90% compared to the control [36,37,97–99], suggesting that multiple gene-sets might backup mechanisms of tunneling tube formation in the same cell. The steps of nanotube biogenesis, consisting of the tube formation, elongation, and degradation/disassembly, are affected differently, sometimes in opposite directions, by the same gene-sets, adding additional complexity to the development of the nanotube modulators [97]. In this regard, HuR inhibitors could be valuable tools in the modulation of cell fusion and tunneling nanotube formations, illuminating a way to suppress (i) the inflammatory and hypoxic microenvironment (the leading cause of the cell fusion in the gliomas and different types of cancer), (ii) the expression of the multiple sets of the fusogenic proteins and the tunneling nanotube-related transcripts, (iii) the transcripts essential for cytoskeleton remodeling and membrane protrusion formations, (iv) the transcripts and cell-signaling pathways involved in autophagy formation [100–111].
We propose that the HuR inhibitors may produce interference with cell fusion and tunneling membrane nanotube formations due to the fact that HuR is the central node in the regulation of inflammation, stress response, lipid metabolism, autophagy formation, and the actin-related cytoskeletal transcripts essential for cell fusion. To date, the following key compounds/scaffolds have been discovered as the inhibitors of HuR function: MS-444 as a blocker of HuR dimerization and nuclear/cytoplasmic shuttling [100]; CMLD1, CMLD2, quercetin, azaphilone derivatives, DHTS, NSC#84126, and mitoxantrone as the modulators of the HuR/mRNA interaction [101–107]; pyrvinium pamoate, okicenone, trichostatin, 5-aza-2’-deoxycytidine (AZA) as the inhibitors of HuR shuttling [100,108]. The pyrvinium pamoate, an antihelminthic drug, is FDA-approved and has been recently repurposed for the blocking of HuR nuclear/cytoplasmic shuttling [109]. Mitoxantrone is classified as an antitumor antibiotic and as an inhibitor of HuR/Cox2-mRNA interaction. Novantrone (mitoxantrone hydrochloride) injections for reducing neurologic disability and/or the frequency of clinical relapses in patients with secondary (chronic) progressive, progressive relapsing, or worsening relapsing–remitting multiple sclerosis have been approved by the FDA since October 2000 [110]. Several randomized multicenter trials have been initiated with Novantrone as anti-inflammatory drug for treatment of prostate, ovarian, breast, hematologic cancers, and solid or central nervous system (CNS) tumors overexpressing EGFR.

Inflammation has been recognized as one of the risk factors for oncogenic transformations in cancer [112,113]. Inflammatory mediators, such as prostaglandins (PGs), thromboxanes, and leukotrienes, are implicated in the inflammatory processes in the tumor microenvironment; PG production is tightly regulated by the COX-2 enzyme and favors tunneling tube formations. COX-2 selective non-steroid anti-inflammatory drugs (NSAIDs) and non-selective NSAIDs are considered as anti-inflammatory and antitumor chemotherapeutics [114]. Non-selective COX inhibitors, tolfenamic acid and indomethacin, substantially blocked both TNT formations and the spread of HCMV viruses between fibroblasts [115]. Recently, the anti-inflammatory and the TNT- and COX-1-inhibitory properties of NSAIDs have been confirmed by using (phendione)ZnII(NPR)2(H2O)2 and (phendione)ZnII(MFN)2 compounds on human breast cancer cell lines [116]. Although HuR inhibitors are currently available for preclinical evaluations, the combinations of the NSAIDs and gene-specific targeting approaches remain valuable as suppressors of cell fusion and tunneling nanotube formation.

In our recent work, we developed a strategy to search and optimize the inhibitors of HuR dimerization in glioma cells [54]. HuR dimerization is essential for HuR nuclear/cytoplasmic translocation and the high-affinity binding to target mRNA [100]. HuR dimerization/multimerization is mostly observed in cancer cells and is associated with glioma progression [54]. The combination of medicinal chemistry with high-throughput HuR-specific biochemical and cell-based assays led us to the identification of several new inhibitors of HuR dimerization, such as compound #5 (N-[4-(1H-benzi-midazol-2-yl)phenyl]-2-chloro-5-nitrobenzamide), suitable for future optimization [54]. Figure 4 summarizes the impact of HuR overexpression on patient overall survival and illustrates classes of the available HuR inhibitors.
which is associated with intra- and intercellular microtubule formations and could be suppressed
stress to promote cell motility, survival, plasticity, aberrant proliferation, and angiogenesis. Therefore,
regulate transcriptome splicing to keep cell identity [130]. HuR is dimerized and shuttles to the
In the post-embryonic period, HuR is predominantly nuclear, is observed in spliceosomes, and may
differentiation and intercellular membrane fusion on several occasions during normal development.
Several reports suggest that HuR may be directly involved in the intercellular membrane fusion
during development and under the pathological conditions: (i) HuR-deficient mice are embryonic-lethal
due to defects in placenta development [117]; (ii) HuR is essential for the skeletal muscle myotube
formations during embryogenesis [118,119]; (iii) HuR contributes to the post-natal pathological
angiogenesis via the regulation of pruning of the vascular branches and the endothelial cell self-fusion
during this process [88,120–122]; on the other hand, HuR dimerization may promote atherosclerosis
and may enhance the permeability of the vascular endothelial layer [123]; (iv) HuR is essential for
the formation of germ cell syncytium where cells stay connected to one another by intercellular
bridges [124]; (v) under the hypoxic condition, HuR enhances epithelial-to-mesenchymal transition,
which is associated with intra- and intercellular microtubule formations and could be suppressed
by inhibitors of HuR nuclear/cytoplasmic shuttling [125,126]; (vi) HuR promotes integrity of the
gap-junction and the stability of Cx43 transcripts, which are involved in the type ii intercellular
tunneling nanotube and microtube formations [127–129]. Collectively, HuR protein dimerization and
translocation from the nucleus to the cytoplasm is associated with cell reprogramming toward cell
differentiation and intercellular membrane fusion on several occasions during normal development.
In the post-embryonic period, HuR is predominantly nuclear, is observed in spliceosomes, and may
regulate transcriptome splicing to keep cell identity [130]. HuR is dimerized and shuttles to the
cytoplasm under pathological conditions such as obesity, diabetes, cancer, and in response to cellular
stress to promote cell motility, survival, plasticity, aberrant proliferation, and angiogenesis. Therefore,
we suggest that glioma stem cells with cell-cycle checkpoint abnormalities, genomic instabilities, and renewal abilities hijack HuR function in stressed glioma microenvironments to advance glioma heterogeneity and proliferation. Hence, we predict that the inhibitors of the HuR nuclear/cytoplasmic shuttling and dimerization could be valuable pharmacological tools for the inhibition of glioma cell plasticity and proliferation, and therefore the suppression of glioma progression at different stages of disease development.

6. Conclusions and Perspectives

The elimination of glioma heterogeneity evoked by pharmacological/radio treatments or by environmental stress is an established chemotherapeutic goal. Understanding the role of HuR in the regulation of glioma heterogeneity in the tumor microenvironment is a new direction, which may lead to the discovery of common and environment-specific mechanisms of glioma plasticity. The implication of the HuR protein inhibitors in the treatment of glioma heterogeneity is new and has the following rationales: (i) HuR is overexpressed in gliomas; (ii) HuR orchestrates stress responses and chemoresistance; (iii) HuR upregulates cell-signaling pathways and mRNA transcripts essential for the intercellular membrane fusions and the tunneling membrane tube formations leading to the glioma cell survival and plasticity.

Author Contributions: Conceptualization, N.F.; methodology, N.F. and L.B.N.; data analysis N.F.; writing—original draft preparation, N.F.; writing—review and editing, N.F. and L.B.N.; visualization, N.F. and L.B.N. All authors have read and agreed to the published version of the manuscript.

Funding: This work received no external funding.

Acknowledgments: Data utilized in this review for analysis is available in the R2: Genomics Analysis and Visualization Platform and GEO database (GSE158271).

Conflicts of Interest: The authors declare no conflict of interest.

Abbreviations

| Acronym | Definition |
|---------|------------|
| AA      | Arachidonic Acid |
| CNS     | Central Nervous System |
| CARs    | T cells expressing Chimeric Antigen Receptors |
| cPLA2α  | Cytosolic Phospholipase A2-alpha |
| COX2    | Cytochrome Oxidase Subunit 2 |
| FDA     | Food and Drug Administration |
| GEO     | Gene Expression Omnibus repository database |
| HCQ     | Hydroxychloroquine |
| MCD     | Methyl-β-cyclodextrin |
| Novantrone | mitoxantrone hydrochloride |
| NSAIDs  | Non-Steroid Anti-Inflammatory Drugs |
| Optune  | An application of the electric field, which induces apoptosis in rapidly dividing tumor cells |
| PDGx    | Patient-Derived Glioma Xenografts |
| PGE2    | Prostaglandin E2 |
| PG      | Prostaglandins |
| ROSc    | Reactive Oxygen Species |
| R2:     | Genomic Analysis and Visualization platform R: |
| TMZ     | Temozolomide |
| TNFAIP2 | TNF Alpha Induced Protein 2 |
| TNT     | Tunneling Membrane Nanotubes |
| MT      | Membrane Microtubes |
| WASP    | Wiskott-Aldrich Syndrome Protein |
References

1. Parker, N.R.; Ekhong, P.; Parkinson, J.F.; Howell, V.M.; Wheeler, H.R. Molecular Heterogeneity in Glioblastoma: Potential Clinical Implications. *Front. Oncol.* 2015, 5, 55. [CrossRef] [PubMed]

2. Louis, D.N.; Perry, A.; Reifenberger, G.; Von Deimling, A.; Figarella-Branger, D.; Cavenee, W.K.; Ohgaki, H.; Wiestler, O.D.; Kleihues, P.; Ellison, D.W. The 2016 World Health Organization Classification of Tumors of the Central Nervous System: A summary. *Acta Neuropathol.* 2016, 131, 803–820. [CrossRef]

3. Walid, M.S. Prognostic Factors for Long-Term Survival after Glioblastoma. *Perm. J.* 2008, 12, 45–48. [CrossRef] [PubMed]

4. Mariotto, A.B.; Yabroff, K.R.; Shao, Y.; Feuer, E.J.; Brown, M.L. Projections of the Cost of Cancer Care in the United States: 2010–2020. *J. Natl. Cancer Inst.* 2011, 103, 117–128. [CrossRef]

5. Rusthoven, C.G.; Koshy, M.; Sher, D.J.; Perry, J.R.; Laperriere, N.; Mason, W.P. Radiation plus Temozolomide in Patients with Glioblastoma. *N. Engl. J. Med.* 2017, 376, 2195–2197. [CrossRef] [PubMed]

6. Tamimi, A.F.; Juweid, M. Epidemiology and Outcome of Glioblastoma. *Glioblastoma* 2017, 8, 143–153. [CrossRef]

7. Bagley, S.J.; Desai, A.S.; Linette, G.P.; June, C.H.; O’Rourke, D.M. CAR T-cell therapy for glioblastoma: Recent clinical advances and future challenges. *Neuro-Oncology* 2018, 20, 1429–1438. [CrossRef] [PubMed]

8. Branter, J.; Basu, S.; Smith, S. Tumour treating fields in a combinational therapeutic approach. *Onco target* 2018, 9, 36631–36644. [CrossRef]

9. Mukherjee, P.; Augur, Z.M.; Li, M.; Hill, C.; Greenwood, B.; Domin, M.A.; Kondakci, G.; Narain, N.R.; Kiebish, M.A.; Bronson, R.T.; et al. Therapeutic benefit of combining calorie-restricted ketogenic diet and glutamine targeting in late-stage experimental glioblastoma. *Commun. Biol.* 2019, 2, 1–14. [CrossRef]

10. Hernández, J.M.; Podbilewicz, B. The hallmarks of cell-cell fusion. *Development* 2017, 144, 4481–4495. [CrossRef]

11. Sapir, A.; Avinoam, O.; Podbilewicz, B.; Chernomordik, L.V. Viral and Developmental Cell Fusion Mechanisms: Conservation and Divergence. *Dev. Cell* 2008, 14, 11–21. [CrossRef] [PubMed]

12. Segev, N.; Avinoam, O.; Podbilewicz, B. Fusogens. *Curr. Biol.* 2018, 28, R378–R380. [CrossRef] [PubMed]

13. Rustom, A.; Saffrich, R.; Markovic, I.; Walthier, P.; Gerdes, H.-H. Nanotubular Highways for Intercellular Organelle Transport. *Science* 2004, 303, 1007–1010. [CrossRef] [PubMed]

14. Jash, E.; Prasad, P.; Kumar, N.; Sharma, T.; Goldman, A.; Sehrawat, S. Perspective on nanochannels as cellular mediators in different disease conditions. *Cell Commun. Signal.* 2018, 16, 76. [CrossRef]

15. Sowinski, S.; Jolly, C.; Berninghausen, O.; Purbhoo, M.A.; Chauveau, A.; Köhler, K.; Oddes, S.; Eissmann, P.; Brodsky, F.M.; Hopkins, C.; et al. Membrane nanotubes physically connect T cells over long distances presenting a novel route for HIV-1 transmission. *Nat. Cell Biol.* 2008, 10, 211–219. [CrossRef]

16. Abounit, S.; Bousset, L.; Loria, F.; Zhu, S.; De Chaumont, F.; Pieri, L.; Olivo-Marín, J.; Melki, R.; Zurzolo, C. Tunneling nanotubes spread fibrillar α-synuclein by intercellular trafficking of lysosomes. *EMBO J.* 2016, 35, 2120–2138. [CrossRef]

17. Wang, X.; Yu, X.; Xie, C.; Tan, Z.; Tian, Q.; Zhu, D.; Liu, M.; Guan, Y. Rescue of Brain Function Using Tunneling Nanotubes Between Neural Stem Cells and Brain Microvascular Endothelial Cells. *Mol. Neurobiol.* 2015, 53, 2480–2488. [CrossRef]

18. Ahmad, T.; Mukherjee, S.; Pattnaik, B.; Kumar, M.; Singh, S.; Rehman, R.; Tiwari, B.K.; Jha, K.A.; Barhanpurkar, A.P.; Wani, M.R.; et al. MiR01 regulates intercellular mitochondrial transport & enhances mesenchymal stem cell rescue efficacy. *EMBO J.* 2014, 33, 994–1010. [CrossRef]

19. Torralba, D.; Baixauli, F.; Sánchez-Madrid, F. Mitochondria Know No Boundaries: Mechanisms and Functions of Intercellular Mitochondrial Transfer. *Front. Cell Dev. Biol.* 2016, 4, 107. [CrossRef]

20. Roehlecke, C.; Schmidt, M.H. Tunneling Nanotubes and Tumor Microtubes in Cancer. *Cancers* 2020, 12, 857. [CrossRef]

21. English, K.; Shepherd, A.J.; Nzor, N.-E.; Trinh, R.; Kavelaars, A.; Heijnen, C.J. Astrocytes rescue neuronal health after cisplatin treatment through mitochondrial transfer. *Acta Neuropathol. Commun.* 2020, 8, 36. [CrossRef]

22. Weil, S.; Osswald, M.; Solecki, G.; Grosch, J.; Jung, E.; Lemke, D.; Ratliff, M.; Hänggi, D.; Wick, W.; Winkler, F. Tumor microtubes convey resistance to surgical lesions and chemotherapy in gliomas. *Neuro-Oncology* 2017, 19, 1316–1326. [CrossRef] [PubMed]
23. Pasquier, J.; Guerrouahen, B.S.; Al Thawadi, H.; Ghiabi, P.; Maleki, M.; Abu-Kaoud, N.; Jacob, A.; Mirshahi, M.; Galas, L.; Raffi, S.; et al. Preferential transfer of mitochondria from endothelial to cancer cells through tunneling nanotubes modulates chemoresistance. *J. Transl. Med.* **2013**, *11*, 94. [CrossRef] [PubMed]

24. Shabo, I.; Svanvik, J.; Lindström, A.; Lechertier, T.; Trablo, S.; Hulit, J.; Sparey, T.; Pawelek, J. Roles of cell fusion, hybridization and polyploid cell formation in cancer metastasis. *World J. Clin. Oncol.* **2020**, *11*, 121–135. [CrossRef] [PubMed]

25. Kaur, E.; Rajendra, J.; Jadhav, S.; Shridhar, E.; Goda, J.S.; Moiyadi, A.; Dutt, S. Radiation-induced homotypic cell fusions of innately resistant glioblastoma cells mediate their sustained survival and recurrence. *Carcinogenesis* **2015**, *36*, 685–695. [CrossRef]

26. Mercapide, J.; Rappa, G.; Lorico, A. The intrinsic fusogenicity of glioma cells as a factor of transformation and progression in the tumor microenvironment. *Int. J. Cancer* **2011**, *131*, 334–343. [CrossRef]

27. Lu, X.; Kang, Y. Cell fusion as a hidden force in tumor progression. *Cancer Res.* **2009**, *69*, 8536–8539. [CrossRef] [PubMed]

28. Sun, C.; Zhao, D.; Dai, X.; Chen, J.; Rong, X.; Wang, H.; Wang, A.; Li, M.; Dong, J.; Huang, Q.; et al. Fusion of cancer stem cells and mesenchymal stem cells contributes to glioma neovascularization. *Oncol. Rep.* **2015**, *34*, 2022–2030. [CrossRef] [PubMed]

29. El Hallani, S.; Colin, C.; El Houfi, Y.; Idbaih, A.; Boisselier, B.; Marie, Y.; Ravassard, P.; Labussiere, M.; Mokhtari, K.; Thomas, J.-L.; et al. Tumor and Endothelial Cell Hybrids Participate in Glioblastoma Vasculature. *BioMed Res. Int.* **2014**, *2014*, 1–9. [CrossRef]

30. Zhang, L.; Zhang, Y. Tunneling nanotubes between rat primary astrocytes and C6 glioma cells alter proliferation potential of glioma cells. *Neurosci. Bull.* **2015**, *31*, 371–378. [CrossRef]

31. Bukoreshtliev, N.V.; Wang, X.; Hodneland, E.; Gurke, S.; Barroso, J.F.; Gerdes, H.-H. Selective block of tunneling nanotube (TNT) formation inhibits intercellular organelle transfer between PC12 cells. *FEBS Lett.* **2009**, *583*, 1481–1488. [CrossRef] [PubMed]

32. Osswald, M.; Jung, E.; Sahm, F.; Solecki, G.; Venkataramani, V.; Blaes, J.; Weil, S.; Horstmann, H.; Wiestler, B.; Syed, M.; et al. Brain tumour cells interconnect to a functional and resistant network. *Nat. Cell Biol.* **2015**, *528*, 92–98. [CrossRef] [PubMed]

33. Osswald, M.; Solecki, G.; Wick, W.; Winkler, F. A malignant cellular network in gliomas: Potential clinical implications. *Neuro-Oncology* **2016**, *18*, 479–485. [CrossRef] [PubMed]

34. Errede, M.; Mangieri, D.; Longo, G.; Girolamo, F.; De Trizio, I.; Vimercati, A.; Serio, G.; Frei, K.; Perris, R.; Virgininto, D. Tunneling nanotubes evoke pericyte/endothelial communication during normal and tumoral angiogenesis. *Fluids Barriers CNS* **2018**, *15*, 28. [CrossRef] [PubMed]

35. Liang, D. A Salutary Role of Reactive Oxygen Species in Intercellular Tunnel-Mediated Communication. *Front. Cell Dev. Biol.* **2018**, *6*, 2. [CrossRef]

36. Hase, K.; Kimura, S.; Takatsu, H.; Ohmae, M.; Kawano, S.; Kitamura, H.; Ito, M.; Watarai, H.; Hazelett, D.; Yeaman, C.; Yeaman, C.; et al. M-Sec promotes membrane nanotube formation by interacting with Ral and the exocyst complex. *Nat. Cell Biol.* **2009**, *11*, 1427–1432. [CrossRef]

37. Ohno, H.; Hase, K.; Kimura, S. M-Sec. *Commun. Integr. Biol.* **2010**, *3*, 231–233. [CrossRef]

38. Pergu, R.; Dagar, S.; Kumar, H.; Kumar, R.; Bhattacharya, J.; Malyavarapu, S.V.S. The chaperone ERp29 is required for tunneling nanotube formation by stabilizing MSec. *J. Biol. Chem.* **2019**, *294*, 7177–7193. [CrossRef]

39. Sun, X.; Wang, Y.-P.; Zhang, J.; Tu, J.; Wang, X.-J.; Su, X.-D.; Wang, L.; Zhang, Y. Tunneling-nanotube direction determination in neurons and astrocytes. *Cell Death Dis.* **2012**, *3*, e438. [CrossRef]

40. Gerdes, H.-H.; Rustom, A.; Wang, X. Tunneling nanotubes, an emerging intercellular communication route in development. *Mech. Dev.* **2013**, *130*, 381–387. [CrossRef]

41. Ribeiro-Rodrigues, T.M.; Martins-Marques, T.; Morel, S.; Kwak, B.R.; Girão, H. Role of connexin 43 in different forms of intercellular communication–gap junctions, extracellular vesicles and tunneling nanotubes. *J. Cell Sci.* **2017**, *130*, 3619–3630. [CrossRef] [PubMed]

42. Jung, E.; Osswald, M.; Blaes, J.; Wiestler, B.; Sahm, F.; Schmenger, T.; Solecki, G.; Deumelandt, K.; Kurz, F.T.; Xie, R.; et al. Tweety-Homolog 1 Drives Brain Colonization of Gliomas. *J. Neurosci.* **2017**, *37*, 6837–6850. [CrossRef] [PubMed]

43. Wesseling, P.; Bent, M.V.D.; Perry, A. Oligodendroglioma: Pathology, molecular mechanisms and markers. *Acta Neuropathol.* **2015**, *129*, 809–827. [CrossRef]
44. Desir, S.; Dickson, E.L.; Vogel, R.I.; Thayanithy, V.; Wong, P.; Teoh, D.; Geller, M.A.; Steer, C.J.; Subramanian, S.; Lou, E. Tunneling nanotube formation is stimulated by hypoxia in ovarian cancer cells. Oncotarget 2016, 7, 43150–43161. [CrossRef] [PubMed]

45. Kemerdere, R.; Akgun, M.Y.; Toklu, S.; Alizada, O.; Tanriverdi, T. Preoperative systemic inflammatory markers in low- and high-grade gliomas: A retrospective analysis of 171 patients. Heliyon 2019, 5, e01681. [CrossRef] [PubMed]

46. Galvão, R.P.; Zong, H. Inflammation and Gliomagenesis: Bi-Directional Communication at Early and Late Stages of Tumor Progression. Curr. Pathobiol. Rep. 2013, 1, 19–28. [CrossRef] [PubMed]

47. Katrib, A.; Jeong, H.-H.; Fransen, N.L.; Henzel, K.S.; Miller, J.A. An Inflammatory Landscape for Preoperative Neurologic Deficits in Glioblastoma. Front. Genet. 2019, 10, 488. [CrossRef]

48. Nabors, L.B.; Gillespie, G.Y.; Harkins, L.; King, P.H. HuR, a RNA stability factor, is expressed in malignant brain tumors and binds to adenosine- and uridine-rich elements within the 3′ untranslated regions of cytokine and angiogenic factor mRNAs. Cancer Res. 2001, 61, 2154–2161. [PubMed]

49. Filippova, N.; Yang, X.; Wang, Y.; Gillespie, G.Y.; Langford, C.; King, P.; Wheeler, C.; Nabors, L.B. The RNA-binding protein HuR promotes glioma growth and treatment resistance. Mol. Cancer Res. 2011, 9, 648–659. Available online: https://mcr.aacrjournals.org/content/9/5/648 (accessed on 6 July 2020). [CrossRef]

50. Bolognani, F.; Gallani, A.-I.; Sokol, L.; Baskin, D.S.; Meisner-Kober, N. mRNA stability alterations mediated by HuR are necessary to sustain the fast growth of glioma cells. J. Neurooncol. 2011, 106, 531–542. [CrossRef]

51. Gálbán, S.; Kuwano, Y.; Pullmann, R.; Martindale, J.L.; Kim, H.H.; Lal, A.; Abdelmohsen, K.; Yang, X.; Dang, Y.; Liu, J.O.; et al. RNA-Binding Proteins HuR and PTB Promote the Translation of Hypoxia-Inducible Factor 1α. Mol. Cell. Biol. 2007, 28, 93–107. [CrossRef]

52. Wang, J.; Leavenworth, J.W.; Hjelmeland, A.B.; Patel, N.; Borg, B.; Si, Y.; King, P. Deletion of the RNA regulator HuR in tumor-associated microglia and macrophages stimulates anti-tumor immunity and attenuates glioma growth. Glia 2019, 67, 2424–2439. [CrossRef] [PubMed]

53. Abdelmohsen, K.; Gorospe, M. Posttranscriptional regulation of cancer traits by HuR. Wiley Interdiscip. Rev. RNA 2010, 1, 214–229. [CrossRef] [PubMed]

54. Filippova, N.; Yang, X.; Ananthan, S.; Sorochinsky, A.; Hackney, J.R.; Gentry, Z.; Bae, S.; King, P.; Nabors, L.B. Hu antigen R (HuR) multimerization contributes to glioma disease progression. J. Biol. Chem. 2017, 292, 16999–17010. [CrossRef] [PubMed]

55. Zarei, M.; Lal, S.; Vaziri-Gohar, A.; O’Hayer, K.; Gunda, V.; Singh, P.K.; Brody, J.R.; Winter, J.M. RNA-Binding Protein HuR Regulates Both Mutant and Wild-Type IDH1 in IDH1-Mutated Cancer. Mol. Cancer Res. 2018, 17, 508–520. [CrossRef]

56. Uren, P.J.; Burns, S.C.; Ruan, J.; Singh, K.K.; Smith, A.D.; Penalva, L.O.F. Genomic Analyses of the RNA-binding Protein HuR Antigen R (HuR) Identify a Complex Network of Target Genes and Novel Characteristics of Its Binding Sites. J. Biol. Chem. 2011, 286, 37063–37066. [CrossRef]

57. Astanina, K.; Koch, M.; Jüngst, C.; Zumbusch, A.; Kiener, A.K. Lipid droplets as a novel cargo of tunnelling nanotubes in endothelial cells. Sci. Rep. 2015, 5, 11453. [CrossRef]

58. Nakamura, H.; Murayama, T. The Role of Sphingolipids in Arachidonic Acid Metabolism. J. PharmacoL Sci. 2014, 124, 307–312. [CrossRef]

59. Lin, W.-N.; Lin, C.-C.; Cheng, H.-Y.; Yang, C.-M. Regulation of cyclooxygenase-2 and cytosolic phospholipase A2gene expression by lipopolysaccharide through the RNA-binding protein HuR: Involvement of NADPH oxidase, reactive oxygen species and mitogen-activated protein kinases. Br. J. PharmacoL 2011, 163, 1691–1706. [CrossRef]

60. Liao, W.-L.; Wang, W.-C.; Chang, W.-C.; Tseng, J.T. The RNA-binding Protein HuR Stabilizes Cytosolic Phospholipase A2α mRNA under Interleukin-1β Treatment in Non-small Cell Lung Cancer A549 Cells. J. Biol. Chem. 2011, 286, 35499–35508. [CrossRef] [PubMed]

61. Linkous, A.G.; Yazlovitskaya, E.M.; Hallahan, D.E. Cytosolic Phospholipase A2 and Lysophospholipids in Tumor Angiogenesis. J. NatL. Cancer Inst. 2010, 102, 1398–1412. [CrossRef] [PubMed]

62. Yang, L.; Zhang, H. Expression of Cytosolic Phospholipase A2 Alpha in Glioblastoma Is Associated with Resistance to Chemotherapy. Am. J. Med. Sci. 2018, 356, 391–398. [CrossRef] [PubMed]
63. Dixon, D.A.; Tolley, N.D.; King, P.H.; Nabors, L.B.; McIntyre, T.M.; Zimmerman, G.A.; Prescott, S.M. Altered expression of the mRNA stability factor HuR promotes cyclooxygenase-2 expression in colon cancer cells. *J. Clin. Investig.* 2001, 108, 1657–1665. [CrossRef] [PubMed]

64. Palumbo, P.; Lombardi, F.; Augello, F.R.; Giusti, I.; Dolo, V.; Leocata, P.; Cifone, M.G.; Cinque, B. Biological effects of selective COX-2 inhibitor NS398 on human glioblastoma cell lines. *Cancer Cell Int.* 2020, 20, 1–17. [CrossRef]

65. Sengupta, S.; Jang, B.-C.; Wu, M.-T.; Paik, J.-H.; Furneaux, H.; Hla, T. The RNA-binding Protein HuR Regulates the Expression of Cyclooxygenase-2. *J. Biol. Chem.* 2003, 278, 25227–25233. [CrossRef]

66. Jin, R.; Junutula, J.R.; Matern, H.T.; Ervin, K.E.; Scheller, R.H.; Brunger, A.T. Exo84 and Sec5 are competitive regulatory Sec6/8 effectors to the RaLA GTPase. *EMBO J.* 2005, 24, 2064–2074. [CrossRef]

67. Wang, J.; Hjelmeland, A.B.; Nabors, L.B.; King, P. Anti-cancer effects of the HuR inhibitor, MS-444, in malignant glioma cells. *Cancer Biol. Ther.* 2019, 20, 979–988. [CrossRef]

68. Chernomordik, L.V.; Kozlov, M.M. Mechanics of membrane fusion. *Nat. Struct. Mol. Biol.* 2008, 15, 675–683. [CrossRef]

69. Palanisamy, K.; Tsai, T.-H.; Yu, T.-M.; Sun, K.-T.; Yu, S.-H.; Lin, F.-Y.; Wang, I.-K.; Li, C.-Y. RNA-binding protein, human antigen R regulates hypoxia-induced autophagy by targeting ATG7 mRNA During Seizure-Induced Mmp-9 Expression in Neurons. *Front. Neurosci.* 2018, 12, 224. [CrossRef] [PubMed]

70. Abdelmohsen, K.; Srikanth, S.; Yang, X.; Lal, A.; Kim, H.H.; Kuwano, Y.; Galban, S.; Becker, K.G.; Kamara, D.; De Cabo, R.; et al. Ubiquitin-mediated proteolysis of HuR by heat shock. *EMBO J.* 2009, 28, 1271–1282. [CrossRef] [PubMed]

71. D’Alloia, A.; Berruti, G.; Costa, B.; Schiller, C.; Ambrosini, R.; Pastori, V.; Martegani, E.; Ceriani, M. RalGPS2 is involved in tunneling nanotubes formation in 5637 bladder cancer cells. *Exp. Cell Res.* 2018, 362, 349–361. [CrossRef] [PubMed]

72. Fiftadara, N.H.; Beer, F.; Ono, S.; Ono, S.J. Interaction between activated chemokine receptor 1 and FcrRI at membrane rafts promotes communication and F-actin-rich cytoneme extensions between mast cells. *Int. Immunol.* 2010, 22, 113–128. [CrossRef]

73. Palanisamy, K.; Tsai, T.-H.; Yu, T.-M.; Sun, K.-T.; Yu, S.-H.; Lin, F.-Y.; Wang, I.-K.; Li, C.-Y. RNA-binding protein, human antigen R regulates hypoxia-induced autophagy by targeting ATG7/ATG16L1 expressions and autophagosome formation. *J. Cell. Physiol.* 2018, 234, 7448–7458. [CrossRef]

74. Jin, R.; Kim, C.; Kang, H.; Ahn, S.; Jung, M.; Hong, Y.; Tak, H.; Lee, S.; Kim, W.; Lee, E.K. RNA Binding Protein HuR Promotes Autophagosome Formation by Regulating Expression of Autophagy-Related Proteins 5, 12, and 16 in Human Hepatocellular Carcinoma Cells. *Mol. Cell. Biol.* 2019, 39, e00508-18. [CrossRef]

75. Sun, C.; Dai, X.; Zhao, D.; Wang, H.; Rong, X.; Huang, Q.; Lan, Q. Mesenchymal stem cells promote glioma neovascularization in vivo by fusing with cancer stem cells. *BMC Cancer* 2019, 19, 1240. [CrossRef]

76. Goldenberg, D.M.; Zagzag, D.; Heselmeyer-Haddad, K.M.; Garcia, L.Y.B.; Ried, T.; Loo, M.; Chang, C.-H.; Gold, D.V. Horizontal transmission and retention of malignancy, as well as functional human genes, after spontaneous fusion of human glioblastoma and hamster host cells in vivo. *Int. J. Cancer* 2011, 131, 49–58. [CrossRef]

77. Cao, M.; Chen, L.; Dang, W.-Q.; Zhang, X.-C.; Zhang, X.; Shi, Y.; Yao, X.-H.; Li, Q.; Zhu, J.; Lin, Y.; et al. Hybrids by tumor-associated macrophages × glioblastoma cells entail nuclear reprogramming and glioblastoma invasion. *Cancer Lett.* 2019, 442, 445–452. [CrossRef]

78. Manjunath, Y.; Porciani, D.; Mitchell, J.; Suvilesh, K.N.; Avella, D.M.; Kimchi, E.T.; Staveley-O’Carroll, K.F.; Burke, D.H.; Li, G.; Kaifi, J.T. Tumor-Cell–Macrophage Fusion Cells as Liquid Biomarkers and Tumor Enhancers in Cancer. *Int. J. Mol. Sci.* 2020, 21, 1872. [CrossRef]

79. Bracq, L.; Xie, M.; Lambelé, M.; Vu, L.-T.; Matz, J.; Schmitt, A.; Delon, J.; Zhou, P.; Randriamampita, C.; Bouchet, J.; et al. T Cell-Macrophage Fusion Triggers Multinucleated Giant Cell Formation for HIV-1 Spreading. *J. Virol.* 2017, 91, e01237-17. [CrossRef]
82. Chen, K.A.; Laywell, E.D.; Marshall, G.; Walton, N.; Zheng, T.; Steindler, D.A. Fusion of neural stem cells in culture. *Exp. Neurol.* 2006, 198, 129–135. [CrossRef] [PubMed]

83. Guadagno, E.; Presta, I.; Maisano, D.; Donato, D.; Pirrone, C.K.; Cardillo, G.; Corrado, S.D.; Mignogna, C.; Mancuso, T.; Donato, G.; et al. Role of Macrophages in Brain Tumor Growth and Progression. *Int. J. Mol. Sci.* 2018, 19, 1005. [CrossRef] [PubMed]

84. Civita, P.; Leite, D.M.; Flikington, G.J. Pre-Clinical Drug Testing in 2D and 3D Human In Vitro Models of Glioblastoma Incorporating Non-Neoplastic Astrocytes: Tunneling Nano Tubules and Mitochondrial Transfer Modulates Cell Behavior and Therapeutic Respons. *Int. J. Mol. Sci.* 2019, 20, 6017. [CrossRef] [PubMed]

85. Zaccard, C.R.; Rinaldo, C.R.; Mailliard, R. Linked in: Immunologic membrane nanotube networks. *J. Leukoc. Biol.* 2016, 100, 81–94. [CrossRef] [PubMed]

86. Wei, J.; Chen, P.; Gupta, P.; Ott, M.; Zamler, D.; Kassab, C.; Bhat, K.P.; Curran, M.A.; De Groot, J.F.; Heimberger, A.B. Immune biology of glioma associated macrophages and microglia: Functional and therapeutic implications. *Neuro-Oncology* 2019. [CrossRef]

87. Hanna, S.J.; McCoy-Simandle, K.; Leung, E.; Genna, A.; Condeelis, J.; Cox, D. Tunneling nanotubes, a novel mode of tumor cell–macrophage communication in tumor cell invasion. *J. Cell Sci.* 2019, 132, jcs223231. [CrossRef]

88. Morrison, A.R.; Yarovinsky, T.O.; Young, B.D.; Moraes, F.; Ross, T.D.; Ceneri, N.; Zhang, J.; Zhuang, Z.W.; Sinusas, A.J.; Pardi, R.; et al. Chemokine-coupled β2 integrin-induced macrophage Rac2–Myosin IIA interaction regulates VEGF-A mRNA stability and arteriogenesis. *J. Exp. Med.* 2014, 211, 1957–1968. [CrossRef]

89. Zhang, J.; Modi, Y.; Yarovinsky, T.; Yu, J.; Collinge, M.; Kyriakides, T.; Zhu, Y.; Sessa, W.; Pardi, R.; Bender, J.R. Macrophage β2 Integrin–Mediated, HuR-Dependent Stabilization of Angiogenic Factor–Encoding mRNAs in Inflammatory Angiogenesis. *Am. J. Pathol.* 2012, 180, 1751–1760. [CrossRef]

90. Filippova, N.; Yang, X.; Nabors, L.B. Growth Factor Dependent Regulation of Centrosome Function and Genomic Instability by HuR. *Biomolecules* 2015, 5, 263–281. [CrossRef]

91. Dybdal-Hargreaves, N.; Risinger, A.L.; Mooberger, S.L. Erubulin mesylate: Mechanism of action of a unique microtubule-targeting agent. *Clin. Cancer Res.* 2015, 21, 2445–2452. [CrossRef]

92. Mundhara, N.; Majumder, A.; Panda, D. Methyl-β-cyclodextrin, an actin depolymerizer augments the antiproliferative potential of microtubule-targeting agents. *Sci. Rep.* 2019, 9, 7638. [CrossRef] [PubMed]

93. Zalba, S.; Hagen, T.L.T. Cell membrane modulation as adjuvant in cancer therapy. *Cancer Treat. Rev.* 2017, 52, 48–57. [CrossRef] [PubMed]

94. Kimura, T.; Takabatake, Y.; Takahashi, A.; Isaka, Y. Chloroquine in Cancer Therapy: A Double-Edged Sword of Autophagy. *Cancer Res.* 2013, 73, 3–7. [CrossRef]

95. Jones, T.M.; Carew, J.S.; Nawrocki, S.T. Therapeutic Targeting of Autophagy for Renal Cell Carcinoma Therapy. *Cancers* 2020, 12, 1185. [CrossRef] [PubMed]

96. Kast, R.E.; Skuli, N.; Karpel-Massler, G.; Frosina, G.; Ryken, T.; Halatsch, M.-E. Blocking epithelial-to-mesenchymal transition in glioblastoma with a sextet of repurposed drugs: The EIS regimen. *Oncotarget* 2017, 8, 60727–60749. [CrossRef]

97. Hanna, S.J.; McCoy-Simandle, K.; Miskolci, V.; Guo, P.; Cammer, M.; Hodgson, L.; Cox, D. The Role of Rho-GTPases and actin polymerization during Macrophage Tunneling Nanotube Biogenesis. *Sci. Rep.* 2017, 7, 1–16. [CrossRef]

98. Kim, B.-W.; Lee, J.-S.; Ko, Y.-G. Mycoplasma exploits mammalian tunneling nanotubes for cell-to-cell dissemination. *BMB Rep.* 2019, 52, 490–495. [CrossRef]

99. Delage, E.; Cervantes, D.C.; Penard, E.; Schmitt, C.; Syan, S.; Disanza, A.; Scita, G.; Zurzolo, C. Differential identity of Filopodia and Tunneling Nanotubes revealed by the opposite functions of actin regulatory complexes. *Sci. Rep.* 2016, 6, 39632. [CrossRef]

100. Meisner, N.-C.; Hintersteiner, M.; Mueller, K.; Bauer, R.; Seifert, J.-M.; Naegeli, H.-U.; Ottl, J.; Oberer, L.; Guenat, C.; Moss, S.; et al. Identification and mechanistic characterization of low-molecular-weight inhibitors for HuR. *Nat. Chem. Biol.* 2007, 3, 508–515. [CrossRef]

101. Muralidharan, R.; Mehta, M.; Ahmed, R.; Roy, S.; Xu, L.; Aubé, J.; Chen, A.; Zhao, Y.D.; Herman, T.; Ramesh, R.; et al. HuR-targeted small molecule inhibitor exhibits cytotoxicity towards human lung cancer cells. *Sci. Rep.* 2017, 7, 1–11. [CrossRef] [PubMed]
Cancers 2020, 12, 3069

102. Chae, M.-J.; Sung, H.Y.; Kim, E.-H.; Lee, M.; Kwak, H.; Chae, C.H.; Kim, S.; Park, W.-Y. Chemical inhibitors destabilize HuR binding to the AU-rich element of TNF-α mRNA. Exp. Mol. Med. 2009, 41, 824–831. [CrossRef] [PubMed]

103. Kaur, K.; Wu, X.; Fields, J.K.; Johnson, D.K.; Lan, L.; Pratt, M.; Somova, A.D.; Wang, C.C.C.; Karanicolas, J.; Oakley, B.R.; et al. The functional natural product azaphilone-9 binds to HuR and inhibits HuR-RNA interaction in vitro. PLoS ONE 2017, 12, e0175471. [CrossRef] [PubMed]

104. D’Agostino, V.G.; Lal, P.; Mantelli, B.; Tiedje, C.; Zucal, C.; Thongon, N.; Gaestel, M.; Latorre, E.; Marinelli, L.; Seneci, P.; et al. Dihydroranshinone-I interferes with the RNA-binding activity of HuR affecting its post-transcriptional function. Sci. Rep. 2015, 5, 16478. [CrossRef] [PubMed]

105. Della Volpe, S.; Nasti, R.; Queirolo, M.; Unver, M.Y.; Jumde, V.K.; Dömling, A.; Vasile, F.; Potenza, D.; Ambrosio, F.A.; Costa, G.; et al. Novel Compounds Targeting the RNA-Binding Protein HuR. Structure-Based Design, Synthesis, and Interaction Studies. ACS Med. Chem. Lett. 2019, 10, 615–620. [CrossRef]

106. Kakuguchi, W.; Nomura, T.; Kitamura, T.; Otsuguro, S.; Matsushita, K.; Sakaitani, M.; Maenaka, K.; Tei, K. Suramin, screened from an approved drug library, inhibits HuR functions and attenuates malignant phenotype of oral cancer cells. Cancer Med. 2018, 7, 6269–6280. [CrossRef]

107. Wang, Z.; Bhattacharya, A.; Ivanov, D.N. Identification of Small-Molecule Inhibitors of the HuR-RNA Interaction Using a Fluorescence Polarization Screening Assay Followed by NMR Validation. PLoS ONE 2015, 10, e0138780. [CrossRef]

108. Hostetter, C.; Licata, L.A.; Costantino, C.L.; Witkiewicz, A.; Yeo, C.J.; Brody, J.R.; Keen, J.C. Cytoplasmic accumulation of the RNA binding protein HuR is central to tamoxifen resistance in estrogen receptor positive breast cancer cells. Cancer Biol. Ther. 2008, 7, 1496–1506. [CrossRef]

109. Guo, J.; Lv, J.; Chang, S.; Chen, Z.; Lu, W.; Xu, C.; Liu, M.; Pang, X. Inhibiting cytoplasmic accumulation of HuR synergizes genotoxic agents in urothelial carcinoma of the bladder. Oncotarget 2016, 7, 45249–45262. [CrossRef]

110. Marriott, J.J.; Miyasaki, J.M.; Gronseth, G.; O’Connor, P.W. Evidence Report: The efficacy and safety of mitoxantrone (Novantrone) in the treatment of multiple sclerosis. Neurology 2010, 74, 1463–1470. [CrossRef]

111. Wu, X.; Lan, L.; Wilson, D.M.; Marquez, R.T.; Tsao, W.-C.; Gao, P.; Roy, A.; Turner, B.A.; McDonald, P.; Tunge, J.A.; et al. Identification and Validation of Novel Small Molecule Disruptors of HuR-mRNA Interaction. ACS Chem. Biol. 2015, 10, 1476–1484. [CrossRef] [PubMed]

112. Gandhi, J.; Khera, L.; Gaur, N.; Paul, C.; Kaul, R. Role of Modifier of Inflammation Cyclooxygenase-2 in the phenotype of oral cancer cells. Cancer Med. 2020, 11, 293–298. [CrossRef]

113. Schroer, J.; Shenk, T. Inhibition of cylooxygenase activity blocks cell-to-cell spread of human cytomegalovirus. Proc. Natl. Acad. Sci. USA 2008, 105, 19468–19473. [CrossRef]

114. Deb, J.; Lakshman, T.R.; Ghosh, I.; Jana, S.S.; Paine, T.K. Mechanistic studies of in vitro anti-proliferative and anti-inflammatory activities of the Zn(ii)–NSAID complexes of 1,10-phenanthroline-5,6-dione in MDA-MB-231 cells. Dalton Trans. 2020, 49, 11375–11384. [CrossRef]

115. Hemberger, M.; Kontoyiannis, D.L. The RNA-Binding Protein Elavl1 regulates muscle differentiation genes. Mol. Cell. Biol. 2008, 29, 2762–2776. [CrossRef]

116. Figueroa, A.; Cuadrado, A.; Fan, J.; Atasoy, U.; Muscat, G.E.; Muñoz-Canoves, P.; Gorospe, M.; Emuñoz, A. Role of HuR in Skeletal Myogenesis through Coordinate Regulation of Muscle Differentiation Genes. Mol. Cell. Biol. 2003, 23, 4991–5004. [CrossRef]

117. Dormoy-Raclet, V.; Cammas, A.; Celona, B.; Lian, X.J.; Van Der Giessen, K.; Zivovnjvic, M.; Brunelli, S.; Riuzzi, F.; Sorci, G.; Wilhelm, B.T.; et al. HuR and miR-1192 regulate myogenesis by modulating the translation of HMGBl mRNA. Nat. Commun. 2013, 4, 2388. [CrossRef]

118. Lenard, A.; Daetwylser, S.; Betz, C.; Eliertsdöttir, E.; Belting, H.-G.; Huisken, J.; Affolter, M. Endothelial Cell Self-fusion during Vascular Pruning. PLoS Biol. 2015, 13, e1002126. [CrossRef]
121. Naito, H.; Iba, T.; Takakura, N. Mechanisms of new blood-vessel formation and proliferative heterogeneity of endothelial cells. *Int. Immunol.* 2020, 32, 295–305. [CrossRef] [PubMed]

122. Yang, C.; Kelaini, S.; Caines, R.; Margariti, A. RBPs Play Important Roles in Vascular Endothelial Dysfunction Under Diabetic Conditions. *Front. Physiol.* 2018, 9, 1310. [CrossRef]

123. Bibli, S.-I.; Hu, J.; Sigala, F.; Wittig, I.; Heidler, J.; Zukunft, S.; Tsilimigras, D.I.; Randriamboavonjy, V.; Wittig, J.; Kojonazarov, B.; et al. Cystathionine γ Lyase Sulphhydrates the RNA Binding Protein Human Antigen R to Preserve Endothelial Cell Function and Delay Atherogenesis. *Circulation* 2019, 139, 101–114. [CrossRef] [PubMed]

124. Chi, M.N.; Auriol, J.; Jégou, B.; Kontoyiannis, D.L.; Turner, J.M.; Rooij, D.G.D.; Morello, D. The RNA-binding protein ELAVL1/HuR is essential for mouse spermatogenesis, acting both at meiotic and postmeiotic stages. *Mol. Biol. Cell* 2011, 22, 2875–2885. [CrossRef]

125. Tsai, I.-T.; Kuo, C.-C.; Liou, J.-P.; Chang, J.-Y. Novel microtubule inhibitor MPT0B098 inhibits hypoxia-induced epithelial-to-mesenchymal transition in head and neck squamous cell carcinoma. *J. Biomed. Sci.* 2018, 25, 1–12. [CrossRef]

126. Sun, J.; Gu, X.; Wu, N.; Zhang, P.; Liu, Y.; Jiang, S. Human antigen R enhances the epithelial-mesenchymal transition via regulation of ZEB-1 in the human airway epithelium. *Respir. Res.* 2018, 19, 109. [CrossRef]

127. Ale-Agha, N.; Galban, S.; Sobieroy, C.; Abdelmohsen, K.; Gorospe, M.; Sies, H.; Klotz, L.-O. HuR regulates gap junctional intercellular communication by controlling β-catenin levels and adherens junction integrity. *Hepatology* 2009, 50, 1567–1576. [CrossRef]

128. Salat, C.; Muñoz, M.J.; Sesé, M.; Cajal, S.R.Y.; Aasen, T. Post-transcriptional regulation of connexins. *Biochem. Soc. Trans.* 2015, 43, 465–470. [CrossRef] [PubMed]

129. Rimkute, L.; Jotautis, V.; Marandykina, A.; Sveikatienė, R.; Antanavičiūtė, I.; Skeberdis, V.A. The role of neural connexins in HeLa cell mobility and intercellular communication through tunneling tubes. *BMC Cell Biol.* 2016, 17, 1. [CrossRef]

130. Zhou, Z.; Licklider, L.J.; Gygi, S.P.; Reed, R. Comprehensive proteomic analysis of the human spliceosome. *Nat. Cell Biol.* 2002, 419, 182–185. [CrossRef] [PubMed]

**Publisher’s Note:** MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.

© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).