

Toward a noncytotoxic glioblastoma therapy: blocking MCP-1 with the MTZ Regimen

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Abstract: To improve the prognosis of glioblastoma, we developed an adjuvant treatment directed to a neglected aspect of glioblastoma growth, the contribution of nonmalignant monocyte lineage cells (MLCs) (monocyte, macrophage, microglia, dendritic cells) that infiltrated a main tumor mass. These nonmalignant cells contribute to glioblastoma growth and tumor homeostasis. MLCs comprise of approximately 10%–30% of glioblastoma by volume. After integration into the tumor mass, these become polarized toward an M2 immunosuppressive, pro-angiogenic phenotype that promotes continued tumor growth. Glioblastoma cells initiate and promote this process by synthesizing 13 kDa MCP-1 that attracts circulating monocytes to the tumor. Infiltrating monocytes, after polarizing toward an M2 phenotype, synthesize more MCP-1, forming an amplification loop. Three noncytotoxic drugs, an antibiotic – minocycline, an antihypertensive drug – telmisartan, and a bisphosphonate – zoledronic acid, have ancillary attributes of MCP-1 synthesis inhibition and could be re-purposed, singly or in combination, to inhibit or reverse MLC-mediated immunosuppression, angiogenesis, and other growth-enhancing aspects. Minocycline, telmisartan, and zoledronic acid – the MTZ Regimen – have low-toxicity profiles and could be added to standard radiotherapy and temozolomide. Re-purposing older drugs has advantages of established safety and low drug cost. Four core observations support this approach: 1) malignant glioblastoma cells require a reciprocal trophic relationship with nonmalignant macrophages or microglia to thrive; 2) glioblastoma cells secrete MCP-1 to start the cycle, attracting MLCs, which subsequently also secrete MCP-1 perpetuating the recruitment cycle; 3) increasing cytokine levels in the tumor environment generate further immunosuppression and tumor growth; and 4) MTZ regimen may impede MCP-1-driven processes, thereby interfering with glioblastoma growth.

Keywords: cognition-sparing, high-grade glioma, immunosuppression, macrophage, microglia, monocyte

Introduction

Re-purposing of older, already-marketed drugs is not a new concept. With technological advances giving us a greater understanding of molecular pathways in cancer cells, the idea of re-purposing (redirecting, repositioning, etc) a previously approved drug, with a known safety/toxicity profile for lower cost, risk, and time, compared with developing novel pharmaceuticals has appeal.^{1–4}

Our paper represents a contribution to the ongoing efforts in developing treatments for glioblastoma, one of the most treatment-refractory of all human cancers. Challenges to treatment include difficulty in drug delivery across the blood–brain barrier, the wide heterogeneity of glioblastoma cells within individual tumors,^{5–7} and the multiplicity of essential signaling and metabolic pathways within individual tumor cells, the so-called “Nile Tributary Problem”.^{4,8} One area of growing interest is modifying

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tumor-immune system signaling pathways, specifically the cross talk between the tumor cells and the immune cells resident in the tumor microenvironment including monocytes, dendritic cells, macrophages, and microglia, collectively termed monocyte lineage cells (MLCs).

Early in development, the embryonic brain becomes populated by bone marrow-derived MLCs, to finally comprise ~10%–20% of the adult brain, appearing as microglia. Circulating marrow-derived monocytes can also populate an adult brain, particularly during pathological states, including glioblastoma, but also in the setting of infection, trauma, or inflammatory disease such as multiple sclerosis, appearing then in brain as either macrophages or microglia (described in the magisterial review of microglia by Harry⁹). Some of the first micrographs of microglia by Klatzo from 1952 have not been surpassed.¹⁰ Although normal steady-state microglia have been called “quiescent”, they are in fact physiologically active in multiple homeostatic roles essential to brain function by scavenging dead cell remnants, phagocytosis, antigen presentation, synaptic maintenance, neurite pruning, repair and angiogenesis, and extracellular signaling.^{11,12} Microglia are in continuous reciprocal humoral communication with nearby neurons and astrocytes. Importantly for understanding glioblastoma, microglia are normally sessile but become motile on activation.

MLCs, including microglia, have activation states termed M1 or M2, but these might be better viewed as spectral extremes. The M1-phenotype is associated with high levels of cytokine interleukin (IL)-1 β , IL-12, IL-23, TNF- α , nitric oxide synthase, CD40, MHC I, and MHC II expression; these M1 cells are traditionally called pro-inflammatory state/activated microglia. In contrast, M2 cells have low levels of these inflammatory mediators, are profibrotic, synthesize transforming growth factor- β and IL-10, tend to suppress inflammation, and promote neovascularization.^{11,12} In relation to cancer, the M1 phenotype functions more in an immune-boosting/antitumor role, whereas the M2 phenotype is more anti-inflammatory and tumor-supportive.

The critical importance of this tumor-microglia relationship to tumor growth has been observed in a number of recent papers.^{12–17} Furthermore, this ongoing “reciprocal supportive interplay”¹⁷ facilitates more than just enhanced vigor of tumor cell division and enhanced blood supply but potentially also contributes to glioblastoma migratory invasion into the surrounding brain.¹⁸ Hewedi et al recognize “glioblastoma therapies should address this key CD68-positive [ie, monocyte/macrophage] population”.¹⁹ We propose targeting MCP-1, a 13 kDa signaling peptide, synonymous with CC-chemokine ligand 2, which binds to

outer cell membrane receptors CCR2 and/or CCR4 on MLCs and facilitates glioblastoma-microglia interaction.

This paper reviews and extends this concept, fills in details, and draws the following simple and straightforward conclusions:

- The “reactive gliosis”²⁰ seen involved with and surrounding the main glioblastoma tumor mass is not an epiphenomenon but rather is a crucial core element of glioblastoma pathology.
- It is primarily the malignant glioblastoma cell that generates pathological signals, including MCP-1, to systemic monocytes/microglia.
- Otherwise normal but pathologically activated MLCs are important for robust glioblastoma cell growth; these cells are attracted by increased MCP-1 signaling within the tumor mass. MCP-1 is also an important signal for preferred MLC differentiation to the M2 anti-inflammatory/tumor-promoting phenotype.
- These tumor-infiltrating and pathologically functioning MLCs contribute to a feedback loop involving MCP-1 secretion which further perpetuates glioblastoma production of MCP-1.
- MCP-1 represents a critical mechanism of tumor-microglia communication which contributes significantly to tumorigenesis and potential invasion and is a potential therapeutic target.
- Three noncytotoxic drugs, minocycline, telmisartan, and zoledronic acid, the MTZ Regimen, are all approved for use in humans for noncancer conditions, are all lipophilic and highly brain-penetrant, are all capable of targeting the pathologic MCP-1 secretion, and together will provide MCP-1 and consequently MLC-focused treatment.

A well-developed research database supports each of the three MTZ Regimen drugs’ ability to block or reduce MCP-1 signaling and thus MLC trophic functions and pathologic activation. Additionally, each of these drugs is relatively nontoxic in humans, with significant post-marketing data available, and does not appear to have significant drug-drug interactions alone or in combination. We propose this approach either alone or in combination with other anticancer therapies as a novel glioblastoma treatment approach.

Glioblastoma are heavily infiltrated by blood monocytes

Glioblastomas are heavily infiltrated by cells of the immune system, and these cells are pathophysiologically active. In his seminal 1952 paper describing glioblastoma histology, Klatzo observed, “In the majority of tumors studied there was extensive microglial reaction [...] resting forms were

infrequently seen".¹⁰ We have more recently come to understand that these immune cell populations are composed of bone marrow-origin monocytes,^{21,22} monocyte-derived macrophages, and microglia,^{19,23–29} and neutrophils.^{21,30–35} Estimates vary that these bone marrow-derived cells or nonmalignant microglia comprise 10%–30% of the total glioblastoma tumor mass,^{12,23,26,36,37} but there is general agreement that MLCs make up a considerable percent of total mass³⁸ and MLCs have an established role in glioblastoma growth.³⁹ The monocyte-related component arrives both by migration of circulating monocytes/macrophages²⁷ and by mitoses or centripetal migration of brain-resident microglia⁴⁰ and monocytes.³⁸

Reactive gliosis in brain tissue adjacent to a growing glioblastoma²⁰ is not an incidental finding or epiphenomenon. Reactive gliosis is rather an integral feature of glioblastoma and a crucial link in glioblastoma's growth.²⁰ Brain-resident microglia infiltrate and become integrated with malignant glioblastoma cells⁴¹ as do circulating MLCs.

M1 versus M2 phenotypes

Macrophages can be divided into M1 and M2 subtypes, which have contrasting activity. M1 macrophages are traditionally thought of as immunostimulatory with potential antitumor activity, whereas M2 exhibit an immunosuppressive phenotype with tumor-promoting and angiogenic activity.¹⁷ Human glioblastomas are richly infiltrated by MLCs expressing both M1 and M2 markers.^{14,24} Although some studies indicate that these MLCs are more of M2 immunosuppressive, tumor trophic class,²⁶ other studies show that these tissue macrophages are of indeterminate class, with some having attributes of both M1 and M2 phenotypes, whereas others have attributes of neither phenotype.^{24,42}

Furthermore, these M1/M2 phenotypes are not terminally differentiated cells; in glioblastoma cell-conditioned culture medium, M1 macrophages can be induced to transform to an M2 phenotype,⁴³ a finding that was later independently confirmed.²⁸ A differentiation to M2-phenotypic macrophages has also recently been described to be promoted by glioma cell-derived extracellular vesicles.⁴⁴

In another set of experiments, it has been shown that doxycycline (chemically similar, but more hydrophilic and less brain-penetrant compared with minocycline) inhibits M2-type polarization of human and bone marrow-derived mouse macrophages.⁴⁵ Clearly in viral infections of the brain, for example, West Nile virus, circulating monocytes are recruited to brain tissue by virus-mediated increases in MCP-1, and these monocytes appear in brain tissue as microglia.⁴⁶

Disproportionate immunosuppression of glioblastoma

"Gliomas attract brain-resident (microglia) and peripheral macrophages and reprogram these cells into immunosuppressive, proinvasive cells."⁴⁰ Myeloid cell infiltration and the function changes induced in them after contact with or proximity to malignant glioblastoma cells form a central element of the otherwise unexplained immunosuppression common in glioblastoma patients.^{22,32,34,47,48} Of particular note, normal human monocytes exposed in vitro to glioblastoma cells take on an immunosuppressive M2 phenotype, synthesizing IL-10 and transforming growth factor-beta.^{49,50}

The MCP-1 amplification loop

MCP-1 agonism at CCR2 stimulates migration of monocytes/macrophages. Glioblastoma cells themselves synthesize and release MCP-1.^{51,52} While MCP-1 remains important in MLC recruitment and M2 phenotype promotion, there are other molecules which appear to contribute to these processes as well. One molecule that appears to be primarily secreted from the stem cell subpopulation of glioblastoma cells is periostin, which contributes to M2 differentiation.²⁷ Several other studies have demonstrated that malignant glioblastoma cells will also synthesize and secrete granulocyte-macrophage colony-stimulating factor that can also drive transformation from M1 to M2 tumor-infiltrating MLCs.⁴⁰ That said, while it is difficult to compare relative contributions of different molecules having similar effects, it appears that MCP-1 encompasses MLC recruiting, M2 transformation, and generation of a positive feedback loop to the glioblastoma tumor cells, while the others promote only portions of the larger process. Brain-resident microglia have been shown to migrate into a growing glioblastoma tumor; this migration is driven in part by granulocyte-macrophage colony-stimulating factor synthesized by glioblastoma malignant cells themselves,^{40,53–58} and in part by MCP-1 synthesized by other already-resident microglia.

An oddity of human glioblastoma tissue is that it attracts immunosuppressive, regulatory T-cells (Tregs).⁵⁹ Tregs express CCR2 receptors and are tropic along an increasing MCP-1 gradient.⁵⁹ Glioma cell lines D-54, U-87, U-251, and LN-229 synthesize large amounts of MCP-1 that was shown to be chemotactic to Tregs.⁶⁰ Glioblastoma patients with higher circulating plasma MCP-1 titers have a poorer prognosis and faster disease course compared to patients with low plasma titers.⁶¹ Human glioblastoma biopsy tissue overexpresses MCP-1.⁶² Glioblastoma cell lines overexpressing MCP-1 demonstrated enhanced motility in response to MCP-1 exposure.⁶²

Having established the likely importance of MCP-1 in macrophages' promotion of glioblastoma growth, the next section discusses how several currently marketed noncytotoxic drugs can defeat this growth-facilitating aspect.

Resident peritumoral microglia secrete ample MCP-1 which in turn is chemotactic for circulating monocytes, thereby recruiting yet more MLCs. Thus, MCP-1 synthesis by glioblastoma-resident macrophages/microglia forms a positive amplification loop.⁶² Since glioblastoma cells themselves express MCP-1 receptors, MCP-1 mRNA, and MCP-1 protein,^{62–65} they thereby initiate this pathological amplification loop. Wurdinger et al rhetorically wonder “how to break [this] vicious cycle [...] the interactive process between glioma cells and microglia and macrophages [...]?”⁶⁶ The MTZ Regimen now attempts to simply answer that question (Figure 1).

Preclinical experiments pointed to tumor-associated macrophages as a primary mediator of escape from antiangiogenic therapies and prompted a study of the consequences of anti-VEGF monoclonal antibody bevacizumab treatment of recurrent glioblastoma.⁶⁷ They found on autopsy that patients treated with bevacizumab had greatly increased numbers of glioblastoma tissue macrophages compared with those not exposed to bevacizumab.⁶⁷ Of particular note to our intended use of macrophage-directed treatment, Lu-Emerson et al also found that extent of tumor infiltration with macrophages correlated with shorter overall survival.⁶⁷

The drugs

Minocycline

Minocycline is a 457 Da, intermediate-half-life (10–20 hours), lipophilic, tetracycline class antibiotic^{68–70} with excellent blood–brain penetration and brain tissue levels.⁷¹ Minocycline is active against a variety of Gram-positive and Gram-negative bacteria, *Rickettsia*, *Chlamydia*, as well as protozoans *Plasmodia*, *Toxoplasma*, and others.⁷¹ As an aside, significant neuroprotective attributes of minocycline have been the subject of dozens of reports in the last few years, this observation may provide future directions for research.⁷²

In culture, normal microglia and glioma cell lines secreted ample MCP-1 which could be inhibited by 25 μ M minocycline.⁷³ Lipopolysaccharide-stimulated MCP-1 protein production in monocytes was suppressed by minocycline in vitro by inhibition of an I κ B phosphorylation step, keeping NF κ B within cytoplasm.⁷⁴ The related tetracycline class antibiotic doxycycline inhibited M2 polarization in a laser injury model of choroidal neovascularization.⁴⁵

MCP-1 levels are elevated in human Japanese encephalitis-infected tissues from humans and experimentally infected animals.⁷⁵ Correspondingly, high tissue infiltration of monocytes is seen, a process that can be suppressed by minocycline.⁷⁵ Simian immunodeficiency virus-infected macaques had grossly upregulated cerebral MCP-1 and monocyte infiltration that were both partially but significantly mitigated by minocycline.⁷⁶ Ex vivo West Nile Virus

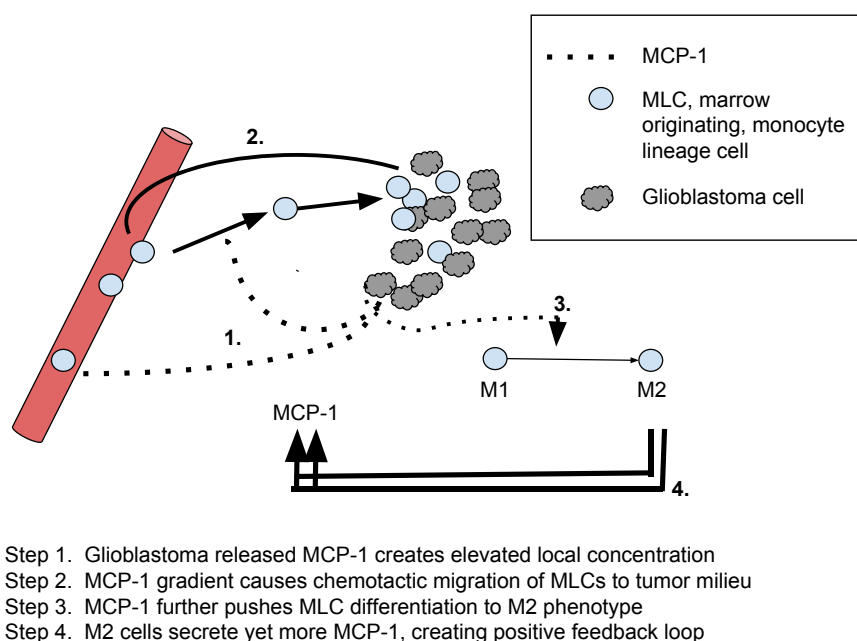


Figure 1 MCP-1 driven, positive feedback loop between glioblastoma and MLCs.
Abbreviation: MLC, monocyte lineage cell.

exposure stimulates microglia to synthesize several cytokines, including MCP-1. This finding was shown to be inhibited by minocycline.⁷⁷ Retinal ischemia–reperfusion injury and cell death was not prevented by minocycline, but the injury-associated centripetal migration of monocytes/macrophages and increase in MCP-1 both were abrogated.⁷⁸

Empirical preclinical studies of minocycline in glioma models have provided evidence of benefit. Growth and core tumor hypoxia of the gliosarcoma line 9L was inhibited by minocycline by unclear means.⁷⁹ Weingart et al found that systemically administered minocycline did not extend survival of 9L gliosarcoma-implanted rats but local delivery after primary resection did,^{80,81} findings confirmed by others in 2014.⁸² Growth of C6 gliomas was inhibited by minocycline.⁸³ An experimental alkylating drug was more effective in killing C6 orthotopic transplants when minocycline was given concomitantly.⁸⁴ Minocycline alone inhibited growth in an orthotopic murine glioma model.⁸⁵ U-87 glioma orthotopic transplants in nude mice grew slower when mice were given intraperitoneal minocycline. GL261 murine glioma cells grew slower in syngeneic mice when intraperitoneal minocycline was given.²⁹

Two clinical studies are in progress, NCT01580969, a Phase II study using minocycline and bevacizumab with radiation for recurrent glioblastoma, and NCT02272270, a Phase I study adding minocycline to standard Stupp Protocol (temozolomide and radiation) for initial post-resection treatment of primary glioblastoma.

Side effects

Skin and other structures can become hyperpigmented after cumulative doses exceed 100 g or duration >1 year.⁸⁶ Autoimmune disturbances such as serum sickness-like disease, drug-induced lupus, and autoimmune hepatitis occur but are rare.⁸⁷ Autoimmune reactions tend not to occur until after >2 years of use.⁸⁸

Telmisartan

Telmisartan is a brain-penetrant, highly lipophilic, 515 Da, intermediate-half-life (~24 hours) angiotensin II receptor blocker marketed to treat high blood pressure.⁸⁹

Telmisartan profoundly downregulated fibronectin mRNA and suppressed MCP-1 over-synthesis in kidneys after experimental unilateral ureter obstruction.⁹⁰ In similar fashion, telmisartan decreased elevated MCP-1 and fibronectin associated with diabetic kidneys.⁹¹

Cerebral MCP-1 increases with age in spontaneously hypertensive stroke-resistant rats well beyond the minimal

increases seen in normotensive rats as they age. Low-dose telmisartan (in doses low enough to not alter hypertension) diminished this progressive increase in cerebral MCP-1 immunohistochemical staining in these spontaneously hypertensive rats with normal aging.^{92,93} In other experiments, telmisartan given in higher doses to normalize these rats' blood pressure lowered cerebral MCP-1 even further.^{92,93} Exposure to homocysteine increased levels of MCP-1 mRNA in, and consequent monocyte binding to, human umbilical vein endothelial cells, an effect blocked by telmisartan and other peroxisome proliferator-activated receptor (PPAR)-gamma agonists.⁹⁴

Hepatic MCP-1 and macrophage infiltration into diet-induced steatohepatitis in susceptible mice was inhibited by telmisartan.⁹⁵ Telmisartan diminished macrophage population, MCP-1 protein, and CCR2 expression in the aorta of spontaneously hypertensive rats.⁹⁶ It also lowered serum MCP-1 level.⁹⁶

Hypertensive humans with increased carotid intima-media thickness had increased serum MCP-1 (thought to contribute to the carotid inflammatory atherosclerotic process) that diminished under telmisartan treatment.⁹⁷ However, although confirming that hypertensive patients had elevated circulating MCP-1, others found no decrease in MCP-1 after successful telmisartan lowering of their blood pressure.⁹⁸ Tumor-infiltrating monocytes/macrophages, as defined by leukocytes that were CD14 and CD68 positive, are known to heavily infiltrate inflammatory breast cancer tissue.⁹⁹ These monocytes/macrophages secreted significant amounts of MCP-1⁹⁹ forming thereby a macrophage amplification circuit that as we indicate here is also operative in glioblastoma. Telmisartan attenuated the elevated myocardial MCP-1 in streptozotocin diabetic rats.¹⁰⁰ Telmisartan decreased MCP-1 mRNA in tumor necrosis factor-alpha-stimulated human umbilical vein endothelial cells.¹⁰¹

Ex vivo lipopolysaccharide-stimulated circulating human monocytes secreted MCP-1 and other inflammation mediators that were decreased by co-incubation with telmisartan.^{102,103} Otherwise normal people with arterial hypertension treated with telmisartan exhibited lowered gene expression of both MCP-1 and PPAR-gamma in circulating monocytes compared to pretreatment levels.¹⁰⁴ In other in vitro experiments, patients with essential hypertension treated for 3 months with telmisartan exhibited reduced circulating monocyte MCP-1 mRNA compared to their pretreatment starting values.¹⁰⁴

Angiotensin II signals via angiotensin 1 receptors. Angiotensin II binding to angiotensin 1 receptors induces rat glomerular endothelia to increase synthesis of MCP-1.¹⁰⁵ This was blocked by telmisartan.¹⁰⁵

Zoledronic acid

Zoledronic acid is a long-half-life, 272 Da bisphosphonate, that although not a low-side effect drug,¹⁰⁶ has seen increasing use as treatment adjuvant in a variety of cancers.¹⁰⁷ Bisphosphonate drugs are commonly known and clinically used for inhibition of osteoclast activity with consequent reduction in bone resorption. Zoledronic acid use in advanced cancer patients has been shown to reduce bone events and hypercalcemia.^{107,108} Zoledronic acid depressed normal osteoblast secretion of MCP-1.¹⁰⁹

In human osteosarcoma, tumor growth directly correlates with and occurs commensurately with osteoclast activation with corresponding osteolysis. Both cell line and human osteosarcomas secrete MCP-1 that forms one of the osteoclast-stimulating paths. Zoledronic acid inhibits both synthesis and secretion of osteosarcoma MCP-1 with consequent diminution of peritumoral osteoclast activity.¹¹⁰ Of particular note to our thinking about glioblastoma, zoledronic acid inhibition of macrophage/osteoclast function reduced growth of the malignant osteosarcoma clone itself.¹¹⁰ Clinical trials of zoledronic acid in breast cancer have prolonged overall survival.¹¹¹

The following four data sets are paradigmatic of our suggested use of zoledronic acid in glioblastoma:

- 1) Rietkotter et al showed that zoledronic acid reduced viability of circulating monocytes and central nervous system microglia¹¹² and microglia exposure to zoledronic acid in ex vivo brain slices also reduced invasion by breast cancer cells without affecting the breast cancer cells themselves.¹¹²
- 2) Indicative of our view of cancer generally as a systemic disease, Hiroshima et al showed that peritoneal macrophages from mice bearing flank-injected pancreatic tumor cells enhanced this cancer's growth when in vitro and in vivo when injected into other tumor-bearing mice.¹¹³
- 3) Ohba et al showed that MCP-1-secreting osteosarcoma cells were blocked from secreting MCP-1 by zoledronic acid.¹¹⁰
- 4) In discussing the correlation between increased tumor-infiltrated macrophages and poorer prognosis, Rogers and Holen presaged our views, stating in 2011, "Tumor cells communicate with the cells of their microenvironment via a series of molecular and cellular interactions to aid their progression to a malignant state and ultimately their metastatic spread". They reviewed data collected prior to 2011 supporting their contention that zoledronic acid exerted an anti-breast cancer effect by defeating MLC's promotion of the malignant clone's growth.¹¹⁴ In this paper we make a similar argument in the setting of glioblastoma.

The MTZ trial

We are planning a pilot clinical trial of the MTZ Regimen in primary glioblastoma, using oral minocycline 100 mg three times daily, oral telmisartan 80 mg twice daily, and intravenous zoledronic acid 4 mg once every 28 days to be given starting postprimary resection, to run concurrent with standard radiation and temozolomide. The primary end points will be tolerability and time to progression, and secondary end point, overall survival.

Discussion

This paper recounts past research data establishing that normal host monocytes are pathologically recruited to and by the glioblastoma tumor. Glioblastomas synthesize and secrete MCP-1 that recruits these monocytes among other effects. Glioblastoma, like other tumors, is not simply a growing autonomous parasitic entity, although it does have elements of this. Glioblastoma is more accurately understood as a malignant clone that secretes cytokines and other hormone-like molecules that signal to normal host cells and organs to support the malignant clone's survival and growth. Glioblastoma is thus integrated into total body physiology.

Given that MCP-1 plays an important role in the destructive inflammatory process of nonalcoholic steatohepatitis,¹¹⁵ we could consider MTZ Regimen or elements of this as part of its treatment.

Psychiatry also is intently interested in microglia–neuron interactions potentially modifiable by minocycline.¹¹⁶ A single case report of minocycline 50 mg po twice daily stopping difficult-to-control seizures in a grade III glioma was interpreted as evidence of its anti-inflammatory effects.¹¹⁷ Going on data and theory that glial activation contributes to pathophysiology of psychotic disorders, three studies, one from 2010 and two from 2014, independently confirmed that minocycline reduced the negative sign/symptom picture that is commonly seen as residue in otherwise well-treated, "schizophrenia".^{118–120} "Schizophrenia" is mentioned in quotes because it is an outmoded term comprising a dozen different pathophysiological entities with prominent psychosis that, as yet, remain unnamed.¹²¹

The malignant MCP-1-centered synergy between the glioblastoma cell and infiltrating microglia or circulating monocytes goes some of the way toward explaining the rarity of metastases outside the central nervous system, even though circulating malignant glioblastoma cells have been easily demonstrated in human disease.¹²²

It is standard military doctrine to destroy a pathway crucial to the enemy along several points along that pathway. Hitting such a pathway at one point only allows too many

chances for the enemy to develop repairs, workarounds, or alternatives that can substitute for or reconstitute the blocked pathway. Destroying multiple points along a crucial pathway significantly impedes an enemy from doing this.

In reviewing the trophic and permissive function of brain microglia for breast cancer metastases Pukrop et al observed “inhibition of proinvasive microglia as a promising anti-metastatic strategy”.¹²³ We provided it here.

MCP-1-triggered hypoxia-related gene transcription and translation via a signal transducer and activator of transcription 3 phosphorylation represents an intermediate step in pluripotent epiblast cells (outer embryonic layer destined to later differentiate into ectoderm and mesoderm layers).¹²⁴ The potential importance of this in glioblastoma and other cancers is that MCP-1 has hypoxia-adapting signaling attributes to cells other than MLCs. In accord with these findings are those of Jung et al who found that minocycline inhibited hypoxia induction factor-1 α transcription and translation.¹²⁵

Baer et al recount details of a reciprocal feedback system between MLCs and tumor-associated new vessel formation, where “physical contact of macrophages with growing blood vessels coordinates vascular fusion of emerging sprouts”.¹²⁶ This feedback (feed-forward) system between MLCs and developing blood vessels is reminiscent of the feed-forward system of Schrufer et al who describe a vessel formation enhancing reciprocal relationship between centripetal-migrating neutrophils and the developing tumor vessel neo-intima that was based on IL-8.^{31,127} We can expect interventions directed against both MLCs and neutrophils each to contribute to suppression of tumor neoangiogenesis.

Radiation of brain tissue during initial glioblastoma treatment often results in memory or intellectual impairments of varying degree.¹²⁸ As a potential added benefit, minocycline reduced memory impairment in radiated rats¹²⁹ and might do so in brain-radiated glioblastoma patients.

As an added benefit of telmisartan, Januel et al showed empirically that those glioblastoma patients coincidentally taking angiotensin-blocking drugs or angiotensin-converting enzyme inhibitors have longer overall survival than those not on these antihypertensive drugs.¹³⁰

Conclusion

Malignant cells co-opt normal cells and physiological systems to serve and enhance growth of the malignant clone/clones. Glioblastoma, like other cancers, can be thought of as an organ system, actively and reciprocally interacting and coordinating with other organ systems of the body, including the immune system. In this case, this collaboration has destructive consequences for the host, yet the elements of

this pathologic relationship incorporate normal physiological systems and communication molecules such as MCP-1.

Current thinking in oncology is not oriented to considering a) the collective effects of drugs or b) effect on and treating contributions of nonmalignant helper cells of the body – in this case MLCs. We tend rather to think of individual effects of the various drugs in a mix. We now propose a paradigm shift to reflect reality more accurately. The reality is that glioblastoma like other cancers has a multiplicity of cross-covering growth paths, most of which must be addressed for successful treatment. Malignant clone-directed treatments must be augmented by simultaneous different treatments directed at halting or limiting the participation of normal host cells and organ systems that subserve and enhance growth of the malignant clone.

Our paper reviewed data showing that growth of glioblastoma cells is connected with and facilitated by surrounding normal microglia and other MLCs, abnormally recruited to and activated by the malignant clone. Our views are in accord with data and views of others. We offer the MTZ Regimen to defeat the role of MLCs that have been abnormally stimulated by MCP-1.

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Author contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in either drafting the article or revising it critically for important intellectual content; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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