

Brain tumors: molecular biology and targeted therapies

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Introduction

Glioblastoma is the most common and most malignant form of primary brain tumors in adults. It frequently develops *de novo* but may also evolve over years from a lower grade astrocytoma [1]. Recent combined modality treatments have led to some improvement in survival. However, most patients will still succumb to their disease (median survival to 15 months and 2-year survival rates of 26%) [2, 3].

Correlative molecular studies and animal models have identified numerous molecular alterations and biological processes involved in initiation and progression of glioma. However, the underlying molecular mechanisms and pathways involved are complex and remain poorly understood. The accumulation of multiple aberrations in regulatory processes enables tumor cells to bypass the effects of many of the available therapies. Molecular alterations underlying such mechanisms comprise aberrations on the genetic level, e.g. point mutations in key genes or respective amplifications and deletions [1], or are a consequence of epigenetic modifications. The latter include aberrant methylation of CpG islands in the regulatory sequence of genes, as well as changes in chromatin structure mediated by mechanisms such as histone acetylation. Hypermethylation of CpG islands in the promoter of genes leads to loss of expression (inactivation), while loss of methylation in normally silenced genes may cause inappropriate expression (e.g. *IGF2* gene), both with tumor promoting effects [4, 5]. Individual molecular tumor profiles are heterogeneous implying that different therapeutic approaches will be necessary for successful personalized treatments.

A first step towards the design of novel therapies is the identification and better understanding of relevant molecular mechanisms driving the aggressive biological behavior that subsequently may be targeted in respective treatment approaches. Hanahan and Weinberg [6] categorized the principle biological requirements for tumor formation as follows: (i) self sufficiency in growth signals, (ii) insensitivity to anti-growth signals, (iii) limitless replicative potential, (iv) sustained angiogenesis, (v) evasion from apoptosis and (vi) tumor infiltration and metastasis. The molecular aberrations conferring these properties in tumors are diverse, although they show some characteristic tumor type and malignancy grade dependent patterns. Molecular tumor profiles may become helpful tools to devise therapeutic strategies aiming for individually tailored treatments.

molecular profiles indicate biological features differentiating glioma subtype and tumor grade

Glioma subtypes exhibit distinct molecular profiles (Figure 1) [1, 7–9]. A hallmark of primary glioblastoma is the amplification of epidermal growth factor receptor gene (*EGFR*) that is often associated with deletion of the *CDKN2A^{P16/Arf}* gene encoding two tumor suppressors, p16 an inhibitor of CDK4 and p14^{ARF} a negative regulator of MDM2. These two aberrations are mutually exclusive with mutations in the *TP53* tumor suppressor gene that represent a hallmark in the evolution of secondary glioblastoma [10, 11]. Primary and secondary glioblastoma, although histologically undistinguishable, are distinct disease entities that occur in different age groups. Childhood glioblastoma, frequently situated in the brain stem, form a third pathogenetically distinct group from their adult counterparts [12, 13].

Oligodendroglioma, characterized by combined loss of heterozygosity on chromosomes 1p and 19q, rarely exhibit mutations in the *TP53* gene or amplification of *EGFR*, but in the majority have an epigenetically inactivated O⁶-methylguanine-DNA methyltransferase (*MGMT*) repair gene that may explain their particular sensitivity to alkylating agent therapy.

The distinct genomic aberrations translate into characteristic differences of global gene expression profiles of glioma subtypes (Figure 2) [14, 15]. Interestingly, differential gene expression profiles distinguishing primary glioblastoma from low grade astrocytoma, and surprisingly also from secondary glioblastoma identified a group of correlated genes related to angiogenesis and hypoxia. This may guide the use of anti-angiogenic treatment strategies [14].

Most glioblastoma (80%) exhibit inactivation of the p14^{ARF}/TP53/MDM2 pathway concurrent with abnormalities in G1-transition control. The latter is a consequence of aberration of either *RB1*, *CDK4*, *CDKN2A* or *CDKN2B^{P15}* genes involved in the same pathway of cell cycle control [16]. Associations of genetic aberrations such as *EGFR* amplification and concurrent deletion of the *CDKN2A^{P16/Arf}* gene locus [7] are suggestive of a cooperative effect. This hypothesis is further supported by *in vivo* and *in vitro* models, and may have relevant consequences for resistance to respective targeted treatments [17, 18].

As molecular data on glioma accumulates, attempts are made to identify predictors of prognosis and response to therapy.

Molecular Hallmarks in Malignant Progression of Gliomas

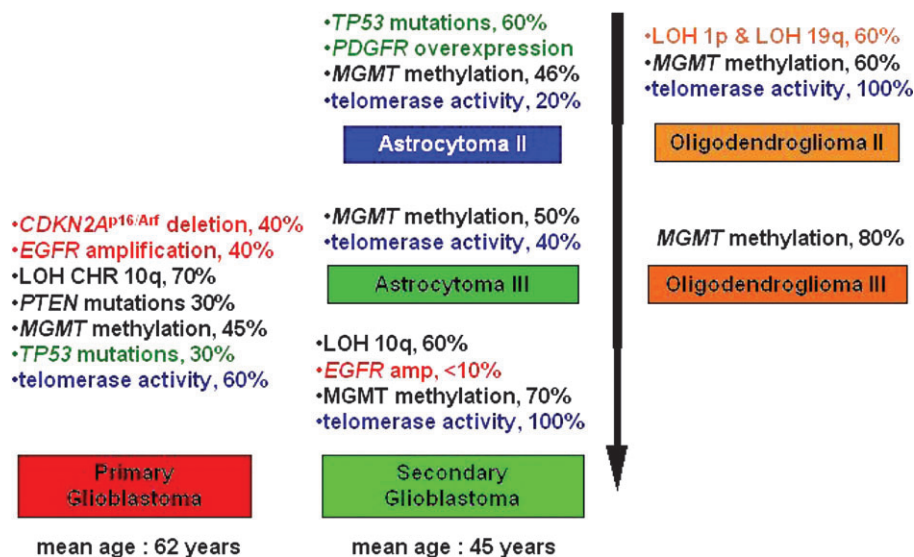


Figure 1. Distinct pathogenetic pathways characterize malignant progression of gliomas. A hallmark of primary glioblastoma is the amplification of the *EGFR* that is often associated with deletion of the *CDKN2A*^{p16/Arf} gene encoding two tumor suppressors. These aberrations are mutually exclusive with mutations in the *TP53* tumor suppressor gene that represent a hallmark in the evolution of secondary glioblastoma. Primary and secondary glioblastoma occur in different age groups. Combined loss of heterozygosity on chromosomes 1p and 19q are characteristic for oligodendroglioma. They rarely exhibit mutations in the *TP53* gene or amplification of the *EGFR*. Information is compiled from the literature in particular from the [1, 7–9, 62].

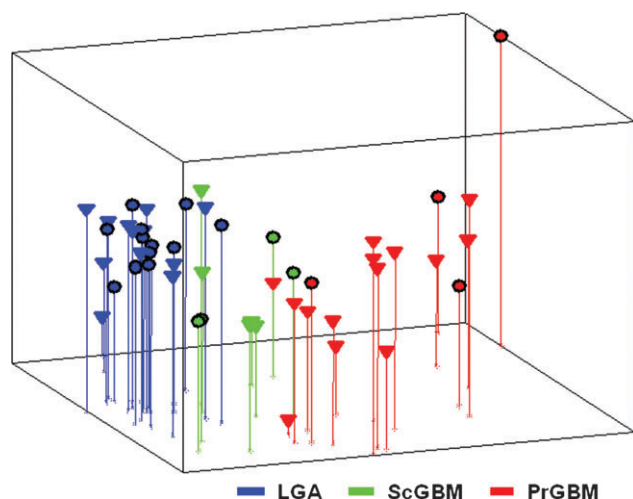


Figure 2. Gene expression profiles differentiate glioma subtypes. MDS (multidimensional scaling) based on overall gene expression (1185 genes) of 51 astrocytic gliomas. The color code indicates the tumor subtype according to histological and clinical criteria. LGA, low grade astrocytoma; ScGBM, secondary glioblastoma; PrGBM, primary glioblastoma. With permission from [19].

Recent comprehensive molecular profiling efforts provided evidence for the existence of molecular subtypes of glioblastoma that may differ in their clinical behavior [19–28]. Several prognostic factors emerged from these and other retrospective studies. However, the insights gained will require validation in independent data sets and prospective clinical trials to determine their predictive value for response to specific therapeutic interventions.

prognostic molecular markers improve diagnostic precision

Tumor histology and malignancy grade guide treatment decisions while taking into account clinical parameters such as age, performance status and extent of tumor resection, are all important prognostic factors. Identification of molecular prognostic markers will be crucial for the design of future trials. However, many markers may just reflect a different natural history and not necessarily contribute to decisions for individual patient management. Nevertheless, molecular markers may greatly improve diagnostic precision and characterize tumor entities with distinct biological behavior.

The best example is the identification of oligodendroglioma with loss of heterozygosity on chromosomes 1p and 19q (LOH 1p/19q). This tumor type is associated with high response rates to chemotherapy and a prolonged survival [8, 29].

Recently, the outcome of anaplastic oligoastrocytoma and oligodendroglioma treated within two independent randomized trials with a similar design was reported by the Radiotherapy Oncology Group (RTOG) and the European Organization of Research and Treatment of Cancer (EORTC) [30, 31]. In these trials PCV-chemotherapy was administered either before (RTOG) or after radiotherapy (EORTC), respectively. Combined losses on chromosomes 1p and 19q represented a favorable prognostic marker and characterized a distinct glioma entity independent of whether initial treatment included PCV chemotherapy or not. Patients with oligodendroglioma with LOH 1p/19q had a much better outcome than patients without LOH 1p/19q. Chemotherapy prolonged progression-free survival in all patients but failed to improve overall survival, even the subset of patients considered the most sensitive to

chemotherapy with LOH 1p/19q. Hence we need to conclude that LOH 1p/19q is primarily a prognostic factor describing a distinct pathologic entity. Its determination improves the accuracy of morphologic diagnosis overcoming subjective interobserver differences that have severely limited comparability between institutions and treatments [32]. Consequently, the ongoing European and Canadian trial (EORTC 22033/26033) for low grade glioma stratifies the patients for 1p deletions in the tumor.

targeted treatments

As pathogenetic factors are diverse, multiple treatment strategies are possible. The integration of clinical and molecular information has provided a list of mechanisms that may be targeted. Aspects of selected treatment strategies are highlighted below. Otherwise reference is given to recent comprehensive reviews of the respective fields.

DNA-repair-driven therapeutics

Epigenetic silencing of the *MGMT* gene encoding a DNA repair enzyme was recently found to be of predictive value for benefit from the alkylating agent temozolomide (TMZ) in a randomized clinical trial for newly diagnosed glioblastoma [33]. The trial had shown that the addition of concomitant and adjuvant (maintenance) TMZ to radiotherapy (RT) improved the 2-year survival rate from 10% in the RT group to 26% for TMZ/RT, setting a new standard of care for glioblastoma patients [3]. Subsequent analysis of the patient's tumor tissues revealed that benefit of the addition of TMZ chemotherapy was basically confined to patients whose tumors had a methylated *MGMT* promoter [33]. At 2 years, 46% of the patients treated with TMZ/RT and whose tumors were *MGMT* methylated survived, compared to only 14% for the patients with unmethylated tumors. High expression of the *MGMT* repair enzyme is known to reverse part of the treatment effect of alkylating agents by rapidly repairing O⁶-methyl guanine, the biologically most important lesion for triggering a cytotoxic response [34, 35]. Since the *MGMT* protein is a suicide repair enzyme that gets inactivated when the alkyl group from the lesion is transferred to the active site of the enzyme, depletion of the enzyme from the tumor cells may be used as a therapeutic strategy to improve sensitivity to alkylating agents. One approach uses a non-cytotoxic substrate of *MGMT*, such as O⁶-BG (O⁶-benzylguanine) or PaTrin-2 [O(6)-(4-bromothienyl)guanine] [36] that may deplete cells of the repair enzyme. Phase I clinical trials have evaluated O⁶-BG in association with TMZ or carmustine (BCNU) [37, 38]. An alternative strategy is to use an intensified, dose-dense TMZ administration schedule aiming at depleting *MGMT* by continued exposure. This approach is currently undergoing clinical testing within a large randomized international Intergroup trial (RTOG0525/EORTC26052–22053) [39]. Patients are stratified prior to randomization by their *MGMT* methylation status. [For additional information, see <http://www.rtog.org> or <http://www.eortc.org> or contact the study chairs: Mark Gilbert (mrgilbert@mdanderson.org) or Roger Stupp (roger.stupp@chuv.ch).] The cytotoxic property of the

O⁶-methyl guanine lesion that persists in the DNA in absence of functional *MGMT*, depends on an intact mismatch repair (MMR) system that eventually triggers the signaling pathways leading to cell death [40, 41]. Thus, best response is expected in patients whose tumor is deficient for *MGMT*, but proficient for MMR. In adult glioblastoma there is no evidence for lack of MMR, in contrast to pediatric glioblastoma [13]. However, additional resistance factors exist, and may include abrogation of apoptosis pathways and deregulated survival signaling that need to be identified and subsequently overcome.

Other approaches aim at rescuing or increasing sensitivity to alkylating agent therapy by inactivating base excision repair (BER) [42]. Approaches have been to use methoxyamine that inhibits apurinic/apyrimidinic endonuclease (APE)-mediated cleavage by binding to AP-sites; while inhibition of Poly(ADP-ribose)polymerase-1 (PARP) impairs the recruitment of BER proteins [43]. This approach has been effective in breast cancer models with either defective *BRCA1* or *BRCA2* that are both involved in repair of double strand breaks [44]. In glioblastoma cell lines or respective xenograft models PARP inhibitors proved to be particularly effective when MMR was deficient [45, 46]. The redundancy in the targeted pathway, in this case repair, has to be overcome in order to achieve a successful treatment response. Hence, targeting repair processes might be a promising strategy for improving efficacy of alkylating agent chemotherapy.

small molecule inhibitors

Over the last years small molecule drugs have been developed specifically inhibiting aberrantly activated signaling pathways. These pathways may not only be activated in the tumor cells themselves, but also in the tumor stroma and tumor-associated endothelial cells [47]. Thus targeting of respective pathways operative in different compartments, or inhibitors of multiple tyrosine kinases, such as PTK-787, may have synergistic effects and thereby enhance treatment efficacy. Evaluation of these small molecule drugs, including peptides, polypeptides and antibodies specifically targeting signaling pathways aberrantly activated in gliomas will be central to future clinical trials (recently reviewed in [48]) (Figure 3). Based on molecular profiles of glioblastoma, the EGFR- and PI3K-pathway and their downstream partners represent particularly attractive targets [49, 50]. Promising are also strategies that particularly aim at angiogenesis [47] such as inhibitors of vascular endothelial growth factor (VEGF), vascular endothelial growth factor receptor (VEGFR), or inhibitors of integrins [51]. However, based on modest responses in trials with single agents, and encouraging synergistic effects in preclinical studies, combination treatments are warranted, combining cytotoxic agents plus one or several targeted drugs (or molecules targeting multiple pathways) [52, 53]. Many new targeted drugs, or combinations thereof, are currently tested or have been tested in phase I or II trials [53, 54]. One limitation in the clinic is the frequent interaction with the commonly used enzyme-inducing anti-epileptic drugs.

Since these strategies aim (mainly) at specific molecular targets it is mandatory to establish molecular profiles of the

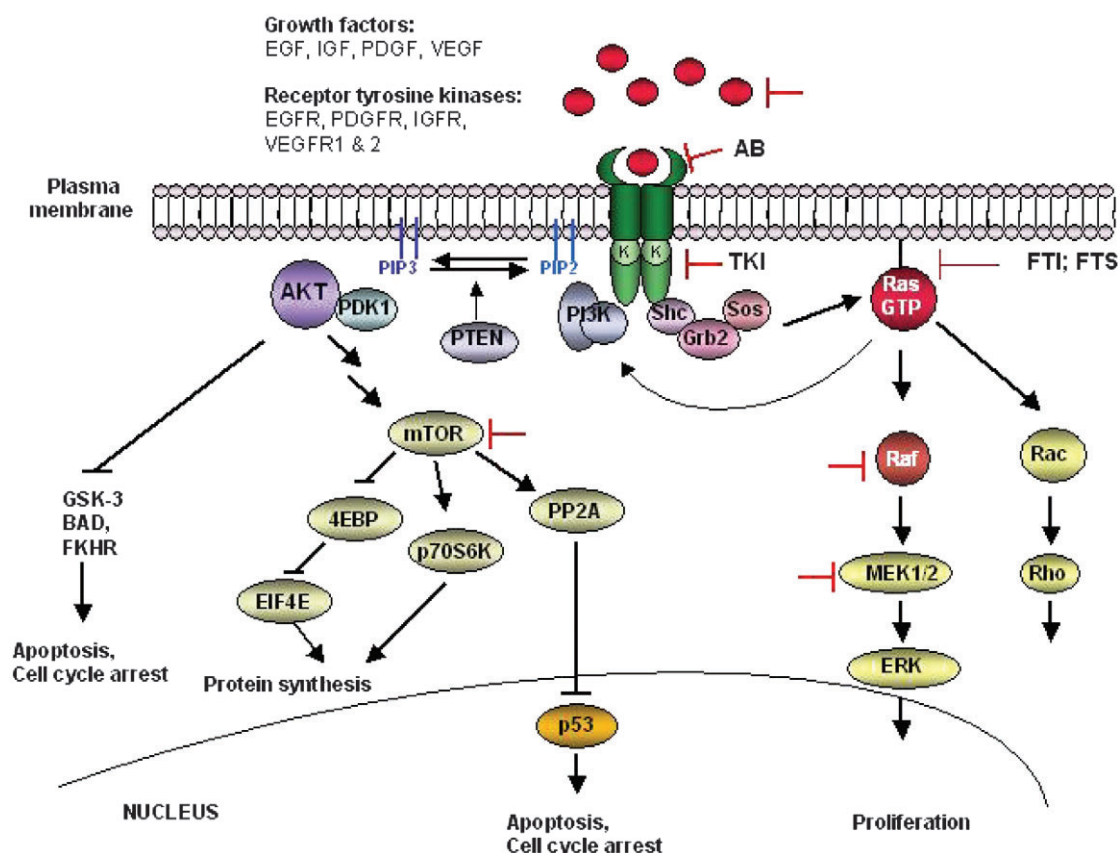


Figure 3. Signaling pathways and their inhibitors. Simplified diagram for signaling pathways operative in glioma or in tumor epithelial cells and respective inhibitors thereof (see also Table 1). The red trait signifies site of specific inhibition. EGF, epidermal growth factor; EGFR, epidermal growth factor receptor; IGF, insulin-like growth factor; IGF1R, insulin-like growth factor receptor; PDGF, platelet-derived growth factor; PDGFR, platelet-derived growth factor receptor; VEGF, vascular endothelial growth factor; VEGFR1 & 2, vascular endothelial growth factor receptor 1 and 2; PI3K, phosphatidylinositol 3-kinase; PTEN, phosphatase and tensin homolog; PDK1, phosphatidylinositol-dependent kinase; AKT, v-akt murine thymoma viral oncogene homolog; mTOR, mammalian target of rapamycin; 4EBP, eukaryotic initiation factor 4E (EIF4E)-binding protein; p70S6K, ribosomal protein, S6 kinase 70kD; PP2A, protein phosphatase 2A; GRB2, growth factor receptor-bound protein 2; SOS, Ras; Raf, MEK, mitogen-activated ERK kinase; and ERK, extracellular-regulated kinase; TKI, tyrosine kinase inhibitor; AB, specific monoclonal antibodies; FTS, Farnesyltransferase inhibitors.

tumors for evaluation of correlations with treatment response. Since the EGFR represents a particularly attractive target it has been extensively evaluated in malignant glioma. Retrospective correlative studies suggested that response to EGFR tyrosine kinase inhibitors did not depend solely on the presence of the target, an activated EGFR [54]. Mellinshof et al. [55] proposed that co-expression of EGFRvIII mutant and PTEN in the tumor were crucial for response to EGFR inhibitors, while in a study by Haas Kogan et al. [56] none of the responders expressed EGFRvIII, but the response was associated with EGFR amplification and expression, and low levels of PKB/Akt phosphorylation. Of note, most of these molecular studies were performed on tumor tissue obtained at the initial diagnosis and may not reflect the molecular pattern at recurrence after multiple prior treatment. Molecular analysis of few tumor tissue samples, available after resection performed during and after prior therapy with the EGFR inhibitors (erlotinib or gefitinib), seemed to suggest inefficient inhibition of EGFR phosphorylation [57]. Thus, the molecular signature relevant for response to this treatment remains to be elucidated and will need more in-depth molecular analysis to elucidate the

molecular mechanisms responsible for these unexpected, additional treatment resistances.

It follows that prospective trials for targeted treatments need to be designed with an integrated translational research component allowing for future molecular selection of patients potentially benefiting from a certain agent or treatment regimen and identifying the relevant pathways that may need to be targeted in addition, in order to improve antitumor activity. Thus, common to all ongoing or planned trials is the absolute necessity of availability of tumor material for molecular profiling (paraffin-embedded or ideally fresh-frozen) in order to establish molecular criteria for the choice of individually tailored treatment approaches in the future.

outlook: gene signature guide drug choice

Novel approaches have recently been described identifying gene expression signatures indicative of oncogenic pathways that may be specifically targeted for therapy [58, 59]. Using this

Table 1. Targeted treatments^a

Target	Name	
Tyrosine kinase inhibitors		
EGFR	Gefitinib	ZD1839, Iressa
EGFR	Erlotinib	OSI-774, Tarceva
EGFR	Cetuximab	C225, Erbitux
EGFR		EKB569
EGFR & ErbB-2	Lapatinib	GW-572016
EGFR & VEGFR		AEE788
VEGFR & EGFR		ZD6474
PDGFR & c-kit	Imatinib	ST1571, Gleevec
	mesylate	
PDGFR & VEGFR		PTK787
PDGFR & VEGFRs, c-kit, Flt-3	Sunitinib	SU11248
PDGFR & VEGFRs, c-kit, Flt-3		AG-013736
PDGFR		SU101
PDGFR		MLN518
Raf, VEGFRs, PDGFR	Sorafenib	Bay 43-9006
VEGFR		AZD2171
Kinase inhibitors		
MEK		CI-1040
MEK		UO126
MEK		PD0325901
VEGF inhibitors		
	Bevacizumab	Avastin
	VEGF Trap	
	Pegabtanib	
Farnesyl transferase inhibitors		
	Tipifarnib	R115777, Zarnestra
	Lonafarnib	SCH66336, Sarasar
mTOR inhibitors		
	Temsirolimus	CCI-779
	Everolimus	RAD001
	Sirolimus	Rapimmune; Wyeth AP23573
Histone deacetylase inhibitors		
	Depsipeptide	FK228
	Suberoylanilide hydroxamic acid	
DNA repair		
MGMT		O ⁶ -benzyl guanine
MGMT	PaTrin-2	O(6)-(4-bromothienyl)guanine
PARP		INO-1001
BER	AP-sites	methoxyamine
Other		
integrins avb3 & avb5	Cilengitide	EMD 121974
Hsp90	17-allylamino-geldanamycin	
Proteasome inhibitor	Bortezomib	PS-341, Velcade
PKC inhibitor	Tamoxifen	

^aCompilation of the literature, references as mentioned in the text, in particular the review by Kesari et al. [48].

concept Solit et al. identified B-raf mutations as predictive for sensitivity to MEK-inhibition in tumor xenograft models, while mutations in ras that are further upstream and signals in addition to other growth promoting pathways led only to partial response. Thus, gene expression profiles may not only identify activated oncogenic pathways but also indicate the presence of redundant pathways that could be responsible for unexpected resistance to treatment with small molecule drugs, despite confirmed presence of the activated target. In the future it might become feasible to deduce, from the tumor derived gene expression profile of a patient, which oncogenic pathways are activated and to devise a respective rational combination treatment. This would conclude the current 'one fits all' treatment strategies based on subjective categorization of morphologic tumor features and some patient characteristics. Further advances may come from new mouse glioma models that recapitulate aberrant molecular pathways relevant in human glioma, or xenograft models of human primary tumors that may prove to be very useful for testing new small molecule drugs preclinically [60, 61].

At present, the test for the *MGMT*-methylation status represents the only predictive factor for benefit from treatment that has been evaluated in a clinical trial, but requires confirmation in ongoing prospective clinical trials. In the individual management of glioblastoma patients this information may direct the choice for alkylating agent chemotherapy, or suggest the addition of alternative treatment modalities, not depending on *MGMT* processing. Hopefully, new predictive factors will become available within the next years to stratify patients according to their individual molecular profiles to respective targeted therapies.

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