REVIEW

Strategically targeting MYC in cancer [version 1; referees: 2 approved]

Valeriya Posternak¹, Michael D. Cole¹,²

¹Department of Pharmacology and Toxicology, Geisel School of Medicine at Dartmouth, Norris Cotton Cancer Center, Lebanon, NH, USA
²Department of Genetics, Geisel School of Medicine at Dartmouth, Norris Cotton Cancer Center, Lebanon, NH, USA

Abstract

MYC is a major driver of cancer cell growth and mediates a transcriptional program spanning cell growth, the cell cycle, metabolism, and cell survival. Many efforts have been made to deliberately target MYC for cancer therapy. A variety of compounds have been generated to inhibit MYC function or stability, either directly or indirectly. The most direct inhibitors target the interaction between MYC and MAX, which is required for DNA binding. Unfortunately, these compounds do not have the desired pharmacokinetics and pharmacodynamics for in vivo application. Recent studies report the indirect inhibition of MYC through the development of two compounds, JQ1 and THZ1, which target factors involved in unique stages of transcription. These compounds appear to have significant therapeutic value for cancers with high levels of MYC, although some effects are MYC-independent. These approaches serve as a foundation for developing novel compounds to pharmacologically target MYC-driven cancers.

This article is included in the F1000 Faculty Reviews channel.

Corresponding author: Michael D. Cole (mcole@dartmouth.edu)

How to cite this article: Posternak V and Cole MD. Strategically targeting MYC in cancer [version 1; referees: 2 approved] F1000Research 2016, 5(F1000 Faculty Rev):408 (doi: 10.12688/f1000research.7879.1)

Copyright: © 2016 Posternak V and Cole MD. This is an open access article distributed under the terms of the Creative Commons Attribution Licence, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Grant information: The author(s) declared that no grants were involved in supporting this work.

Competing interests: The authors declare that they have no competing interests.

First published: 29 Mar 2016, 5(F1000 Faculty Rev):408 (doi: 10.12688/f1000research.7879.1)
**Introduction**

The MYC protein plays a crucial role in a variety of cellular processes, including cell proliferation and differentiation, cell cycle progression, metabolism, and apoptosis\(^{1,2}\). MYC is a pleiotropic transcription factor that regulates a variety of functions by promoting activation or repression of genes on a global scale\(^{3-5}\). As a transcription factor, MYC heterodimerizes with MAX and directly binds to a consensus sequence on DNA, CACGTG\(^6\). MYC-mediated transcriptional activation involves an interaction between MYC and a nuclear cofactor, transformation/transcription domain-associated protein (TRRAP), through a conserved domain on MYC, MYC Box II (MBII)\(^7\). TRRAP is in complex with histone acetyltransferases that acetylate histones around gene promoters, inducing an open chromatin conformation, making it possible for RNA polymerase II (RNA Pol II) recruitment and productive transcription\(^8,9\).

MYC expression is tightly regulated under normal circumstances and is increased in response to extracellular stimuli, such as growth factors\(^{10-11}\). Chromosomal translocation, gene amplification, and mutations in signaling pathways promote MYC overexpression independently of growth factor stimulation, which leads to unrestrained cell proliferation and tumorigenesis\(^1\). MYC is deregulated in approximately 70% of human cancers\(^1\), and many studies have observed that MYC inhibition can result in tumor regression and cell differentiation in a host- and cell-dependent manner\(^1\). Widespread activation of MYC in a range of tumors and the reversibility of MYC-induced tumorogenesis have made MYC an appealing target for cancer therapy. However, MYC lacks innate enzymatic function and small-molecule interactions that facilitate most pharmacological strategies. Furthermore, as a transcription factor, MYC is localized in the nucleus and hence is inaccessible to any antibody-based therapies. For these reasons, MYC is widely considered ‘undruggable’, a frustrating limitation for such a well-established driver of cancer. Nevertheless, numerous strategies have been employed to target MYC at various stages of biological and pathological development, and there have been significant advances in understanding the MYC dependence of cancer and developing novel approaches to targeting MYC activity in the past five years (Figure 1). To date, directly targeting MYC’s interaction with MAX by using compounds like 10058-F4 has proven unsuccessful in vivo, although a biological agent, Omomyc, has proven informative. However, a new library screen resulted in the identification of a potent in vivo MYC-MAX inhibitor, KJ-Pyr-9, that has some efficacy\(^12\). More recent studies have demonstrated that indirect approaches using compounds designed to inhibit key factors involved in transcriptional initiation and elongation seem selective for the MYC oncogenic pathway\(^{15-18}\). These new developments in therapeutic targeting of MYC in cancer have broad implications in a challenging field where inhibition of MYC has been shown to result in tumor regression but has proven problematic to execute because of difficulties in delivery or specificity.

**Targeting the MYC and MAX interface**

Since dimerization with MAX is essential for MYC DNA-binding activity\(^1\), disruption of the MYC/MAX interaction by using small molecules is an obvious strategy of targeting MYC functionality. A number of selective low molecular weight inhibitors that disrupt the interaction between MYC and MAX have been developed\(^19\). One of these is 10058-F4, a molecule that prevents heterodimerization and is capable of penetrating cells with low non-specific toxicity\(^{20,21}\). The compound has demonstrated the ability to inhibit mammalian cell growth, cell cycle progression, and expression of MYC target genes in vitro. A number of studies have reported that short-term pharmacological inhibition of MYC using 10058-F4 or more potent analogs leads to tumor regression in vivo. More recently, KJ-Pyr-9, a compound identified in a pyridine library screen, was identified as a potent inhibitor of the MYC/MAX interaction and it displays the correct pharmacokinetic properties necessary for in vivo administration\(^{41}\). Although these compounds have shown specificity for the MYC/MAX interaction, targeting a bHLH-LZ domain is inherently inefficient and potentially non-specific since many other proteins contain these motifs. Nevertheless, 10058-F4 and KJ-Pyr-9 appear to have differential efficacy in vivo, depending on tumor type, differential metabolism of the compounds, and tumor model\(^{42,43-45}\). Taken together, these data suggest that direct inhibition of MYC through disruption of the MYC/MAX interaction is promising but requires further experimentation to establish specificity and efficiency in humans.

A second strategy to inhibit MYC/MAX dimerization is Omomyc, a mutant basic helix-loop-helix domain that acts as a potent dominant negative molecule by sequestering MYC and preventing its binding to MAX and DNA\(^{46,47}\). Under normal circumstances, MYC is unable to homodimerize, but Omomyc is a MYC homolog that contains four amino acid substitutions augmenting homodimerization and non-functional heterodimerization with MYC. Although Omomyc cannot penetrate human tumors and hence is ineffective as a cancer therapeutic, it has proven useful to explore the consequences of MYC inhibition in vivo. Omomyc can stimulate MYC-induced apoptosis of NIH3T3 cells in a MYC-dependent manner in vitro and of MYC-overexpressing tumor cells in a mouse model of K-Ras-driven lung adenocarcinoma\(^{36,37}\). Recent studies have demonstrated that the bHLH-LZ domain of MAX (MAX\(^*\)) can be transduced across cell membranes through endocytosis and is able to translocate to the nucleus, suggesting that compounds mimicking the bHLH-LZ domain may be efficacious in vivo\(^\text{26}\). Another recent study extended the efficacy of MYC inhibition as a therapeutic strategy (using Omomyc) in the treatment of human glioma in a mouse model of astrocytoma, human glioblastoma cell lines, and patient-derived tumors in vitro and in vivo\(^\text{11}\). Interestingly, general inhibition of MYC activity is tolerated in the mouse, albeit with severely reduced proliferation in the skin, testes, gastrointestinal tract, and hematopoietic lineages\(^17\). Remarkably, the proliferation defects were fully reversible, suggesting that anti-MYC therapy could be used to treat human disease since tumor cells often apoptosis upon MYC inhibition whereas normal cells simply fail to proliferate.

**Indirectly targeting MYC through BRD4 bromodomain inhibition**

Recent findings suggest that MYC promotes gene expression by global transcriptional amplification, although there have been other interpretations of the data\(^{48,49,50}\). The transcriptional amplification model proposes that MYC binds to virtually all active promoters in any cells and enhances transcriptional elongation. These studies have established a positive correlation between MYC levels and phosphorylation of serine-2 (S2) on the carboxy-terminal...
domain (CTD) of RNA Pol II, which is linked to transcriptional elongation. Phosphorylation of S2 on the CTD is catalyzed by P-TEFb (positive transcription elongation factor b), which can be activated by binding to the bromodomain protein BRD4. Bromodomains bind to acetylated lysines (Ac-K) on histones and other proteins, and the binding of BRD4 to P-TEFb results in recruitment to promoters and productive transcriptional elongation. BRD4 is a member of the BET family of proteins and by itself is a key mediator of an aggressive squamous cancer, NUT midline carcinoma. Small-molecule screens have identified compounds that inhibit the binding of the BRD4 bromodomain to Ac-K. The most extensively characterized compound developed for this purpose is JQ1, a powerful inhibitor of BRD4. JQ1 binds to the Ac-K-binding site of BET bromodomains and effectively displaces BRD4 from chromatin, preventing transcriptional elongation. Treatment with JQ1 results in cell differentiation of NUT cells and attenuates growth of BRD4-dependent carcinomas in vivo. Efficacy of JQ1 in a number of myeloid-derived tumors, such as acute myeloid leukemia (AML) and multiple myeloma, has been demonstrated. Notably, these studies have revealed that the effect of JQ1 on tumor regression appears to be specifically mediated by downregulation of MYC itself, its downstream targets, and inflammatory signals. The link between JQ1 and MYC expression is not totally clear but may involve the dependence of MYC on multiple enhancers and ‘super-enhancers’ that are highly dependent on BRD4. These findings have led to a number of potential combination therapies in conjunction with JQ1 that synergistically result in tumor regression. These therapies include indirectly targeting MYC in combination with the PI3K pathway, mechanistic target of rapamycin (mTOR), or histone deacetylases (HDACs) for the treatment of T-cell acute lymphoblastic leukemia, pancreatic ductal adenocarcinoma, and osteosarcoma, respectively. BET inhibitors have also been shown to induce apoptosis of osteosarcoma cells independently of MYC downregulation and display synergistic effects when combined with CDK inhibitors, indicating that this strategy could be employed in the treatment of osteosarcoma.
The use of bromodomain-binding compounds has very recently been developed into a new strategy to target BRD4 and subsequently MYC. dBET1 is a novel compound developed to target BRD4 for protein degradation, in contrast to JQ1, which inhibits the bromodomain of BRD4\cite{14,18}. dBET1 is a bivalent compound composed of JQ1 and thiadiazole that creates a link between BRD4 and cereblon (CRBN), a component of a cullin-RING ubiquitin ligase that catalyzes proteasomal degradation\cite{3}. Together, these data provide a rationale for targeting BRD2, BRD3, and BRD4 for degradation. As with JQ1, the MYC protein and its transcriptional pathway appear to be the most strongly affected. Treatment with dBET1 produces an improved apoptotic response at lower concentrations in AML and lymphoma cell lines, accompanied by a decrease in MYC levels compared with JQ1. This strategy can be exercised to target a wide variety of proteins that may have no innate enzymatic function as long as high-affinity ligands are available.

**Targeting CDK7 as an indirect inhibitor of MYC**

Another very recent study suggests a second indirect approach to target MYC. TFIIH, a complex involved as a basal factor in transcriptional initiation, is composed of a number of proteins, including the catalytic subunit cyclin-dependent kinase 7 (CDK7)\cite{45,46}. CDK7 phosphorylates serine-5 (S5) on the CTD of RNA Pol II, which induces transcriptional initiation, production of nascent mRNA, mRNA capping and methylation, and promotes proximal pausing\cite{51,52}. THZ1 was developed as a novel covalent inhibitor of CDK7, and its high selectivity for CDK7 results from chemical linkage to a cysteine residue that resides outside of the canonical kinase domain\cite{53}. Interestingly, THZ1 specifically downregulates MYC in MYCN-driven neuroblastomas compared with normal cells, and this effect is attributed to the presence of super-enhancers upstream of the MYCN gene\cite{54}. Although the mechanism accounting for MYC specificity requires further investigation, targeting CDK7 in tumors addicted to super-enhancer-associated transcription factors provides a novel platform for targeting multiple aberrant genes with a single agent. Therapeutically, THZ1 was shown to be highly effective in killing MYC-driven tumors, including neuroblastoma, small cell lung cancer, and triple-negative breast cancer\cite{55,56,57}. Treatment with THZ1 leads to a substantial reduction in tumor volume by suppressing cell proliferation and inducing apoptosis. THZ2, an analog of THZ1, was developed to overcome the instability of THZ1 \textit{in vivo} and demonstrated improved pharmacokinetics with an amended half-life and high potency for CDK7\cite{58}. Together, these data provide a rationale for targeting CDK7 in tumors that are dependent on high levels of MYC for transcription.

**Synthetic lethal interactions with MYC**

Although MYC itself is difficult to drug, tumor cells often exhibit ‘oncogene addiction’ or changes in gene expression and physiology that make them extremely dependent on a specific oncogenic pathway for growth or survival or both. This dependence theoretically can be exploited to search for a tumor cell’s Achilles heel (that is, pathways that become rate-limiting for the growth/survival of tumor cells but not their normal counterparts). An early study identified AMPK (AMP-dependent kinase) as critical for the survival of cells with high levels of MYC\cite{60}. Synthetic lethality has also been observed in MYC-overexpressing cells when spliceosome core factors or metabolic pathways are targeted for inhibition\cite{11,15}. A more general approach has been taken to uncover new therapeutic strategies for cancer by interrogating the connection between genomic aberrations and response to a wide panel of anti-cancer drugs. Bioinformatic tools were used to identify a synthetic lethal relationship between MYC overexpression and sensitivity to dasatinib, a multikinase inhibitor. This platform sets a framework for the discovery of novel combination therapies to target MYC-driven tumors.

**Conclusions and Future directions**

A large number of direct and indirect MYC inhibitors have been developed in the last decade, and some are more efficacious and specific than others. Although direct inhibitors of MYC, precisely those targeting the interaction between MYC and MAX, are more specific for MYC itself, they target a bHLH-LZ domain conserved between many transcription factors. Chemical inhibition of a domain present in MYC alone would provide a more targeted approach for MYC inhibition. The mechanisms by which indirect inhibitors of MYC, such as JQ1 and THZ1, act remain to be well characterized. Furthermore, additional experimentation is required to determine the efficacy of these compounds in human cancer. Ultimately, it may be necessary to strategically target MYC from a multitude of angles, taking advantage of its well-established role as a master regulator of transcription in cancer cells.

**Competing interests**

The authors declare that they have no competing interests.

**Grant information**

The author(s) declared that no grants were involved in supporting this work.

---

**References**

1. Meyer N, Penn LZ: Reflecting on 25 years with MYC. Nat Rev Cancer. 2008; 8(12): 976–90. [PubMed Abstract] [Publisher Full Text]

2. Tormay A, Relaiz Y, Oskarsson T, et al.: c-Myc regulates mammalian body size by controlling cell number but not cell size. Nature. 2001; 414(6865): 769–73. [PubMed Abstract] [Publisher Full Text] [F1000 Recommendation]

3. Dang CV, O'Donnell KA, Zeller KI, et al.: The c-Myc target gene network. Semin Cancer Biol. 2006; 16(4): 259–64. [PubMed Abstract] [Publisher Full Text]

4. Lin CY, Loven J, Rahl PB, et al.: Transcriptional amplification in tumor cells with elevated c-Myc. Cell. 2012; 151(1): 56–67. [PubMed Abstract] [Publisher Full Text] [Free Full Text] [F1000 Recommendation]

5. Nie Z, Hu G, Wei G, et al.: c-Myc is a universal amplifier of expressed genes in lymphocytes and embryonic stem cells. Cell. 2012; 151(1): 68–79. [PubMed Abstract] [Publisher Full Text] [Free Full Text] [F1000 Recommendation]
11. Kelly K, Cochran BH, Stiles CD, et al.: Cell-specific regulation of the c-Myc gene by lymphocyte myelons and platelet-derived growth factor. Cell. 1983; 39(Pt 2): 603–10. PubMed Abstract | Publisher Full Text

17. Hart JR, Gamer AL, Yu J, et al.: Inhibitor of MYC identified in a Kröhnke pyridine library. Proc Natl Acad Sci U S A. 2014; 111(34): 12556–61. PubMed Abstract | Publisher Full Text | F1000 Recommendation

18. Truong N, Dao T, Borrello I, et al.: In vivo imaging of MYC protein localization in pancreatic ductal adenocarcinoma. Mol Cancer. 2014; 13: 189. PubMed Abstract | Publisher Full Text

21. Nilson KA, Guo J, Turek ME, et al.: Modelling Myc inhibition as a cancer therapy. Nature. 2008; 455(7213): 679–83. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

22. Schubbert S, Cardenas A, Chen H, et al.: Targeting the MYC and PI3K pathways eliminates leukemia-initiating cells in T-cell acute lymphoblastic leukemia. Cancer Res. 2014; 74(23): 6704–13. PubMed Abstract | Publisher Full Text

24. Brodin J, Keel M, Wranne B, et al.: BET inhibitors induce apoptosis through targeted inhibition of c-Myc. Proc Natl Acad Sci U S A. 2015; 112: 5395–400. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

26. Baker EF, Taylor S, Gupta A, et al.: BET inhibitors induce apoptosis through a BYC independent mechanism and synergize with CDK inhibitors to kill osteosarcoma cells. Sci Rep. 2015; 5: 10120. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

28. Heidemann M, Hintermair C, Voß K, et al.: Activation and repression by oncogenic c-Myc shape tumour-specific gene expression profiles. Nature. 2014; 511(7510): 488–92. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

29. Montagne M, Beaudoin N, Forlin D, et al.: The Max b-HLH-LZ can transduce signals into cells and inhibit c-Myc transcriptional activities. PLoS One. 2012; 7(2): e32172. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation
Co-transcriptional Capping and Pausing. Mol Cell. 2015; 59(4): 576–87.

53. Kwiatkowski N, Zhang T, Rahl PB, et al.: Targeting transcription regulation in cancer with a covalent CDK7 inhibitor. Nature. 2014; 511(7511): 616–20. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

54. Christensen CL, Kwiatkowski N, Abraham BJ, et al.: Targeting transcriptional addictions in small cell lung cancer with a covalent CDK7 inhibitor. Cancer Cell. 2014; 26(6): 909–22. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

55. Wang Y, Zhang T, Kwiatkowski N, et al.: CDK7-dependent transcriptional addiction in triple-negative breast cancer. Cell. 2015; 163(1): 174–86. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation

56. Liu L, Ulbrich J, Müller J, et al.: Deregulated MYC expression induces dependence upon AMPK-related kinase 5. Nature. 2012; 483(7391): 608–12. PubMed Abstract | Publisher Full Text | F1000 Recommendation

57. Hsu TY, Simon LM, Neill NJ, et al.: The spliceosome is a therapeutic vulnerability in MYC-driven cancer. Nature. 2015; 525(7569): 384–8. PubMed Abstract | Publisher Full Text | F1000 Recommendation

58. Shim H, Chun YS, Lewis BC, et al.: A unique glucose-dependent apoptotic pathway induced by c-Myc. Proc Natl Acad Sci U S A. 1998; 95(4): 1511–6. PubMed Abstract | Publisher Full Text | Free Full Text

59. Martins MM, Zhou AY, Corella A, et al.: Linking tumor mutations to drug responses via a quantitative chemical-genetic interaction map. Cancer Discov. 2015; 5(2): 154–67. PubMed Abstract | Publisher Full Text | Free Full Text | F1000 Recommendation
Open Peer Review

Current Referee Status:  

Editorial Note on the Review Process

F1000 Faculty Reviews are commissioned from members of the prestigious F1000 Faculty and are edited as a service to readers. In order to make these reviews as comprehensive and accessible as possible, the referees provide input before publication and only the final, revised version is published. The referees who approved the final version are listed with their names and affiliations but without their reports on earlier versions (any comments will already have been addressed in the published version).

The referees who approved this article are:

1. **David L Levens**, Center for Cancer Research, National Cancer Institute, Bethesda, Maryland, USA  
   *Competing Interests*: No competing interests were disclosed.

2. **Chi Dang**, Abramson Cancer Center, University of Pennsylvania, Philadelphia, Pennsylvania, USA  
   *Competing Interests*: No competing interests were disclosed.