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important future direction. Although future work will be required to fully answer these questions, targeting phosphatases offers an interesting and promising new direction for anti-TB HDT.

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Lighting the way to host-directed immunotherapeutics
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Developing broad-spectrum, host-directed antiviral therapeutics can be adapted to combat emerging viruses. In this issue of Cell Chemical Biology, Maarifi and colleagues implement a Nano luciferase reporter-based protein complementation assay to screen for small molecules and identify Gilteritinib, which enhances interferon induction and antagonizes virus replication.

Drug development, from discovery to clinical application, is a lengthy process that creates a barrier to treat emerging viruses. The limited information known about novel viruses upon their emergence further precludes timely identification and introduction of virus-targeted therapeutics. A potential strategy to circumvent the delay in the development of virus-specific antivirals is the identification of host-directed therapeutics. Targeting the host’s immune responses could be an effective strategy to combat emerging viruses with various viruses as well as reduce the likelihood of escape mutant evolving. The type I interferon (IFN-I) response is a logical target. IFN-I induction and signaling pathways are innate signaling cascades critical for mounting an effective antiviral response. In this issue of Cell Chemical Biology, Maarifi et al. (2022) identify Gilteritinib as a broad-spectrum antiviral that facilitates IFN-I induction.

In response to virus infection, pathogen-associated molecules stimulate the activation of the IFN-I pathway. Viral nucleic acids commonly act as agonists for host pathogen-recognition receptors (PRRs), including the cytoplasmic retinoic acid-inducible gene I (RIG-I)-like receptors (RLRs) and endosomal Toll-like receptors (TLRs) (McNab et al., 2015). After the PRRs bind to their viral agonist, a signaling cascade is triggered. Adaptor molecules transduce the signal facilitating the assembly of complexes that promote TANK-binding kinase 1 (TBK1) and IkB kinase ε (IKKe)-mediated phosphorylation and activation of the IFN regulator factor (IRF)3 and/or IRF7 transcription factors (Figure 1). Phosphorylated IRF3 and/or IRF7 translocate to the nucleus and bind to the positive regulatory domain I and III regions of IFN-stimulated regulator elements in the promoter of IFN-I genes to induce their expression.
expression (Honda et al., 2006). For optimal IFN-β expression, other transcription factors, including nuclear factor-κB and AP-1 (a heterodimer of activating transcription factor 2 and c-Jun) are recruited to PRDs II and IV, respectively. In addition, depending on the cell type and the viral agonist, other IRFs (e.g., IRF-1, IRF-5) can be activated to induce optimal IFN-I levels. The assembly of these transcription factors at the IFN-I gene promoters enhances their expression through the recruitment of histone acetyl transferases including CREB binding protein (CBP). Once synthesized, IFN-I binds to the IFN-I receptor (IFNAR) to activate an antiviral transcriptional repertoire comprised of IFN-stimulated genes (ISGs). ISGs are the effector proteins that restrict virus growth at various stages of their replication cycles. IFN-I signaling also induces the expression of negative feedback regulators such as suppressors of cytokine signaling (SOCS), which can antagonize both IFN-I induction and signaling (Yoshimura et al., 2005) to minimize IFN-I-associated damage to the host. An important consideration when developing host-directed therapeutics targeting the IFN-I pathways is how to
specifically activate the antiviral factors without coincidentally enhancing the expression of pro-inflammatory factors.

Maarifi et al. (2022) apply a reporter protein complementation strategy to screen for kinase inhibitors that induce IRF-induced gene expression to identify host-directed antivirals. Specifically, they adapted their luminance-based Alpha Centauri assay, first developed to identify inhibitors of the human immunodeficiency virus-1 (HIV-1) integrase (Fernandez et al., 2021), to monitor IRF nuclear translocation and localization to gene promoters. They fused the 13 amino acid ε component of Nano luciferase to IRFs and the complementary fragment, Cen, to CBP, so that the enzymatically active luciferase would assemble when the IRF and CBP co-localize. The luminescence resulting from the Nano luciferase-mediated cleavage of furimazine is the readout. The use of CBP as the complementary protein, versus a nuclear localization signal, increases the specificity of the assay beyond IRF nuclear localization and toward the formation of an active transcriptional complex. To validate this method, they over-expressed RIG-I-2CARD, a constitutively active form of RIG-I that contains only the caspase recruitment domains, to stimulate IFN-I induction and monitor the recruitment of the different IRFs to their promoter. Treatment with MRT67307 to block IRF kinases TBK1 and IKKe; prevents IRF3 phosphorylation and nuclear translocation, and they observed diminished luminescence signal following RIG-I-2CARD expression. The authors also investigated which IRFs are induced following infection with a variety of viruses. They found that different groups of IRFs engage with CBP at their promoters following virus infections. For example, they found that infection with Sendai virus activates IRFs 1, 3, and 7, Influenza A virus induces IRFs 3 and 7, but severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) does not induce translocation of any IRFs. The authors hypothesized that virus-encoded innate immune antagonists, such as the papain-like protease (PLP), prevent the activation of IRFs. In support of this, treatment with PLP inhibitor GRL-0617 during SARS-CoV-2 infection allowed for the activation of IRFs 3, 5, and 7.

Because the strongest luminescence signals detected during SARS-CoV-2 infection resulted from IRF7-ε-CBP-Cen interaction, this experimental set-up was used to screen a panel of 21 kinase inhibitors for immunomodulatory activity. Using this assay, the authors identified the AXL and fms-like tyrosine kinase 3 (FLT3) inhibitor Gilteritinib (Dhillon, 2019), which enhances IRF7-CRB interaction in response to infection with SARS-CoV-2, stimulates expression of IFNα4, IFNβ, and CXCL10, and inhibits SARS-CoV-2 replication. Importantly, they found that Gilteritinib specifically stimulated the antiviral arm without altering expression of pro-inflammatory IFNα-γ, IL-6, or TNF-α. In the absence of virus infection, Gilteritinib had no immunostimulatory effect, implying that the molecule does not directly activate IRF7. Treatment with an IFNAR antibody that blocks IFN-I signaling prevented immunostimulation by Gilteritinib and partially reversed its antiviral effects during SARS-CoV-2 infection. Using gene knockdown experiments, they showed that Gilteritinib’s activity was mainly mediated through blocking AXL but not its other target FLT3.

The partial retention of antiviral activity in the absence of IFN-I signaling and following AXL knockdown suggests that Gilteritinib may also have an IFN-I signaling- and AXL-independent mechanism for the inhibition of SARS-CoV-2. AXL is a member of the TAM (Tyro3, Axl, and Mer) receptor tyrosine kinase (RTK) family. This group of RTKs typically functions to facilitate the phagocytosis of apoptotic cells and/or membranes with detectable phosphatidylserine (PS) (Lemke, 2013). In order to recognize apoptotic cells, growth arrest-specific protein 6 (Gas6) binds to PS and the extracellular immunoglobulin domains of TAMs, mainly AXL, to trigger their activation (Figure 1). Several enveloped viruses exploit TAMs to facilitate virus entry when PS is incorporated into their membrane. Further work is needed to determine whether SARS-CoV-2 can similarly hijack AXL and/or the other TAMs to facilitate entry.

In addition to a potential role for facilitating virus entry, blocking AXL’s enzymatic activity could bolster IFN-I signaling. Investigation in dendritic cells has demonstrated that AXL negatively regulates both IFN-I and pro-inflammatory cytokine production (Rothlin et al., 2007; Sharif et al., 2006). AXL signaling facilitates the expression of transcriptional repressors Twist 1 and 2 following IFN-I signaling to antagonize the expression TNFα, IL-8, and COX2 (Sharif et al., 2006). Although Maarifi et al. did not elucidate the precise molecular mechanism by which Gilteritinib promotes the IFN-mediated antiviral response, it could involve cross-talk between AXL and the IFN receptor (Figure 1). Studies have shown that AXL and IFNAR1 interact following IFN-I signaling (Rothlin et al., 2007). The AXL-IFNAR crosstalk results in the formation of STAT1 homodimers, which stimulates gene expression of SOCS1 and SOCS3 that negatively regulate TLRs, RLRs, and IFNAR through blocking critical adaptor molecules such as TRAFs 3 and 6 (activated by TLR agonists) and JAK1 (activated downstream the IFN-I receptor) (Rothlin et al., 2007). Because of AXL’s key role as a negative regulator of pro-inflammatory pathways and in the maintenance of tissue homeostasis through the clearance of apoptotic cells to prevent autoinflammatory diseases (Lemke, 2013), it will be important to evaluate the antiviral therapeutic potential for Gilteritinib in phagocytic cells and characterize the role of AXL in regulating innate antiviral immune responses in multiple cell types.

The Alpha Centauri approach taken by Maarifi and colleagues could potentially be applied beyond therapeutics screening. An important finding of their study was the distinct IRF activation profiles following infection with different virus species. A potential permutation of their approach could be fusing the components of a split enzyme proximity labeling system (Qin et al., 2021) to identify transcriptional regulators recruited to IFN-I gene promoters in response to unique viruses. This type of technology could be applied to emerging viruses to aid in understanding which innate immune pathways are viable targets for host-directed therapeutics. Although this approach does present some limitations, especially considering that it needs over-expression or transduction of the Alpha Centauri components, the system could be adapted to elucidate how IRFs and other transcription factors differentially regulate cytokine production in a cell type specific manner. Overall, Alpha Centauri could provide an accelerated approach for the identification of host-targeted antivirals and at the same time...
providing a tool to study mechanisms of action.

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DECLARATION OF INTERESTS

The authors declare no competing interests.

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