Clinical manifestations of Kaposi sarcoma herpesvirus lytic activation: multicentric Castleman disease (KSHV–MCD) and the KSHV inflammatory cytokine syndrome

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Soon after the discovery of Kaposi sarcoma (KS)-associated herpesvirus (KSHV), it was appreciated that this virus was associated with most cases of multicentric Castleman disease (MCD) arising in patients infected with human immunodeficiency virus. It has subsequently been recognized that KSHV–MCD is a distinct entity from other forms of MCD. Like MCD that is unrelated to KSHV, the clinical presentation of KSHV–MCD is dominated by systemic inflammatory symptoms including fevers, cachexia, and laboratory abnormalities including cytopenias, hypoalbuminemia, hyponatremia, and elevated C-reactive protein. Pathologically KSHV–MCD is characterized by polyclonal, IgM-lambda restricted plasmacytoid cells in the intrafollicular areas of affected lymph nodes. A portion of these cells are infected with KSHV and a sizable subset of these cells express KSHV lytic genes including a viral homolog of interleukin-6 (vIL-6). Patients with KSHV–MCD generally have elevated KSHV viral loads in their peripheral blood. Production of vIL-6 and induction of human (h) IL-6 both contribute to symptoms, perhaps in combination with overproduction of IL-10 and other cytokines. Until recently, the prognosis of patients with KSHV–MCD was poor. Recent therapeutic advances targeting KSHV-infected B cells with the anti-CD20 monoclonal antibody rituximab and utilizing KSHV enzymes to target KSHV-infected cells have substantially improved patient outcomes. Recently another KSHV-associated condition, the KSHV inflammatory cytokine syndrome (KICS) has been described. Its clinical manifestations resemble those of KSHV–MCD but lymphadenopathy is not prominent and the pathologic nodal changes of KSHV–MCD are absent. Patients with KICS exhibit elevated KSHV viral loads and elevation of vIL-6, homolog of human interleukin-6 and IL-10 comparable to those seen in KSHV–MCD; the cellular origin of these is a matter of investigation. KICS may contribute to the inflammatory symptoms seen in some patients with severe KS or primary effusion lymphoma. Additional research is needed to better define the clinical spectrum of KICS and its relationship to KSHV–MCD. In additional, research is needed to better understand the pathogenesis and epidemiology of both KICS and KSHV–MCD, as well as the optimal therapy for both of these disorders.

Keywords: Kaposi sarcoma-associated herpesvirus, human herpesvirus 8, multicentric Castleman disease, KSHV inflammatory cytokine syndrome, interleukin-6, human immunodeficiency virus

KAPOSI SARCOMA-ASSOCIATED HERPESVIRUS AND ASSOCIATED DISEASES

Kaposi sarcoma (KS)-associated herpesvirus (KSHV) is a gamma herpesvirus that is established as the causative agent of a several human illnesses, including KS, primary effusion lymphoma (PEL), and, in the setting of human immunodeficiency virus (HIV) infection, almost all cases of multicentric Castleman disease (MCD; Chang et al., 1994; Cesaran et al., 1995; Soulier et al., 1995). In KS, the predominant cell type is the spindle cell, which expresses endothelial markers, while the cell of origin in PEL and KSHV–MCD is of B-cell lineage. KS and KSHV–MCD are polyclonal in nature, while PEL is considered to be monoclonal. Like other herpesviruses, KSHV infection can exhibit both latent and lytic phases, distinguished by their viral gene expression patterns (Moore and Chang, 1998; Staskus et al., 1999; Sun et al., 1999). KSHV can establish latency in various cell types including endothelial cells and B-lymphocytes (Monini et al., 1999; Wu et al., 2006; Della Bella et al., 2008). Gene expression in latent phase is extremely limited (Jenner et al., 2001), with expressed genes primarily being directed to enhance cell survival, inhibit apoptosis, and evade the host immune response (Moore and Chang, 1998; Wen and Damania, 2010). Following activation of the lytic switch gene, RTA, encoded by open reading frame (ORF) 50, the virus enters the lytic phase, wherein the full complement of viral encoded genes is expressed and the host cellular machinery is redirected to the manufacture and assembly of progeny virions. The triggers of lytic activation in vivo remain unclear. It is speculated that intermittent lytic cycles may play a
role in maintaining viral persistence in the host (Gregory et al., 2009).

The KSHV genome is notable for its molecular piracy of cellular genes (Moore et al., 1996; Ganem, 2010). These appear to promote viral survival, and there is evidence that they contribute to the pathogenesis of KSHV-associated diseases through interactions with the human cellular machinery and immune system. These gene mimics include viral homologs of interferon regulatory factors (IRF), ORF K9/vIRF-1, and ORF K11.5/vIRF-2; a viral homolog of human interleukin-6 (hIL-6) called viral IL-6 (vIL-6; encoded by ORF K2); and a constitutively active G-protein coupled receptor (GPCR) with pro-angiogenic and immunomodulatory functions encoded by ORF74 (Aoki et al., 2000; Jenner et al., 2001; Yarchoan, 2006). While vIL-6 is commonly referred to as a lytic gene, in some circumstances it may be expressed in the absence of full lytic activation (Chatterjee et al., 2002). KSHV also encodes at least 12 small non-coding miRNAs, whose functions are still to be comprehensively determined (Ganem and Ziegelbauer, 2008; Umbach and Cullen, 2010). There is evidence that these also interact with host proteins in ways that favor viral survival and may also induce expression of other host inflammatory proteins including IL-10 (Abend et al., 2010; Lei et al., 2010; Lu et al., 2010; Qin et al., 2010).

Viral IL-6 exhibits a number of important differences from its human counterpart. Their sequence homology is approximately 25% (Moore et al., 1996; Neipel et al., 1997). vIL-6 has a signaling potency 100- to 1000-fold lower that that seen with hIL-6 in in vitro models (Aoki et al., 2000). However, unlike hIL-6, vIL-6 is able to signal by forming a tetrameric complex with the IL-6 receptor superfamily gp130 component, which is widely expressed in human tissues, without requiring binding to the classical gp80 IL-6 receptor (Heinrich et al., 1998; Boulanger et al., 2004; Meads and Medveczky, 2004; Hu and Nicholas, 2006). As a result, vIL-6 may be able to initiate signaling in a wider variety of cell types than does hIL-6 (Osborne et al., 1999; Meads and Medveczky, 2004). In addition, there is in vitro evidence of vIL-6 autocrine action in lytically infected cells, though the function and significance of this activity is as yet uncertain (Chen et al., 2009a,b).

In the absence of HIV infection or other causes of immunodeficiency, most infection with KSHV is clinically silent, even when intermittent lytic activation cycles occur. However, as T cell immunity declines, KSHV-infected patients become increasingly likely to develop KSHV-associated tumors, particularly KS. It is not unusual for patients to manifest more than one KSHV-associated tumor. Each of these tumors has also been reported in other immunosuppressed and older individuals (Teruya-Feldstein et al., 1998). In KS tissue, KSHV is found predominantly in latent phase as assessed by immunohistochemical and genetic analysis of tumor tissue, while there is limited expression of KSHV viral genes, especially vIL-6, in PEL cells (Teruya-Feldstein et al., 1998; Jones et al., 1999; Staskus et al., 1999; Parravicini et al., 2000). By contrast, and unusually among herpesvirus-associated tumors, KSHV–MCD is associated with lytically active KSHV in a substantial proportion of KSHV-infected cells (Oksenhendler et al., 1996; Staskus et al., 1999; Parravicini et al., 2000). This lytic activation, and particularly the production of vIL-6, is thought to underlie its pathogenesis (Oksenhendler et al., 2000; Aoki et al., 2001a; Polizzotto et al., 2011).

MULTICENTRIC CASTLEMAN DISEASE

There are several forms of Castleman disease, which historically have been distinguished by clinical findings (based on whether involved nodes are singular or multiple), pathologic findings, and more recently etiology (idiopathic or KSHV-associated; Cronin and Warnke, 2009). The idiopathic forms, as originally described, include a hyaline vascular form (usually unicentric) and a plasmacytoid form (usually multicentric). The KSHV-associated form is characteristically multicentric, with distinct pathologic features mirroring its unique etiology, as described below (Oksenhendler et al., 1996). Nearly all Castleman disease arising in the setting of HIV infection is KSHV-associated MCD (KSHV–MCD), while KSHV–MCD also rarely occurs in non-HIV infected individuals, particularly the immunocompromised. KSHV–MCD is considered to be rare, although its incidence is not well defined. By contrast to KS, there is evidence that MCD has become more common since the advent of combination anti-retroviral therapy (cART; Powles et al., 2009).

KSHV–MCD is characterized by intermittent flares of inflammatory symptoms, including fever, fatigue, and cachexia, and edema, together with lymphadenopathy and/or splenomegaly (Oksenhendler et al., 1996; Stebbing et al., 2008). Gastrointestinal and respiratory symptoms are common. Flares are often severe and can be fatal. Common laboratory abnormalities include anemia, thrombocytopenia, hypoalbuminemia, hyponatremia, and elevated inflammatory markers such as C-reactive protein (CRP; Oksenhendler et al., 2000; Oksenhendler, 2009). The clinical course waxes and wanes, but until recently, has generally been fatal within 2 years of diagnosis, with patients succumbing to the severe inflammatory syndrome, concurrent infections, or progressing to lymphoma (Oksenhendler et al., 2000; Stebbing et al., 2008). The differential diagnosis of fever and adenopathy, even with other laboratory abnormalities, in the HIV infected individuals is broad. As a result, KSHV–MCD may be difficult to diagnose and is often missed. Also, while there are few reports of KSHV–MCD in Africa, there is recent evidence that it may be substantially underdiagnosed in this region (Ayers et al., 2011).

The diagnosis of KSHV–MCD is made pathologically by a lymph node biopsy demonstrating hypocellular germinal centers with a vascularized core and by a proliferation of polyclonal, IgM λ-restricted plasmacytoid cells in the interfollicular area (Oksenhendler et al., 1996; Du et al., 2001; Chadburn et al., 2008). A proportion of these cells are KSHV-infected, as demonstrated by expression of the KSHV latency associated nuclear antigen (LANA-1), and of these a subset express KSHV lytic genes, especially vIL-6 (Du et al., 2001; Figure 1). Areas of KS spindle cell proliferation in affected nodes are also not uncommon (Naresh et al., 2008). However, the majority of the cellular burden within affected nodes is comprised of uninfected plasmacytoid B-lymphocytes (Chadburn et al., 2008; Naresh et al., 2009).

As noted, the symptom complex of KSHV–MCD is associated with inflammatory dysregulation attributable to KSHV lytic activation. Both production of vIL-6 and induction of host production of hIL-6 have separately been postulated to contribute
to symptoms, perhaps in combination with enhanced production of human interleukin-10 (IL-10) and other human cytokines (Oksenhendler et al., 2000; Aoki et al., 2001a,b; Bower et al., 2009). Viral and human IL-6 over-expression in mouse models can each produce phenotypes resembling that of KSHV–MCD (Brandt et al., 1990; Aoki et al., 2001b). Symptomatic episodes (“flares”) are also associated with high KSHV viral loads (Oksenhendler et al., 2000; Polizzotto et al., 2011). Histopathologic evaluation of nodes involved by KSHV–MCD suggests that the cellular origins of vIL-6 and hIL-6 are largely distinct: involved nodes show vIL-6 production within the periphery of affected germinal center, in the lytic subset of the relatively small number of KSHV-infected cells. By contrast, the larger number of KSHV-uninfected cells in the core of affected nodes can produce hIL-6 but not vIL-6 (Chadburn et al., 2008).

KSHV INFLAMMATORY CYTOKINE SYNDROME

In addition to these well-recognized KSHV-associated diseases, our group has recently described what appears to be a novel inflammatory symptom complex associated with KSHV infection (Uldrick et al., 2010a). In six patients with HIV and KSHV co-infection, but without any pathological evidence of MCD, we observed an inflammatory syndrome resembling that seen in MCD. These patients presented with abnormalities including fevers, sweats, fatigue, wasting, cytopenias, hypoalbuminemia, and hyponatremia, associated in some cases with lymphadenopathy or effusions (Table 1). Notably, four of the six patients also had severe KS. Each patient underwent at least one tissue biopsy to rule out KSHV–MCD as a cause of symptoms. None had pathologic evidence of KSHV–MCD (or KSHV-associated lymphoma), although abnormal KSHV-infected mononuclear cells were sometimes detected in biopsy specimens. None developed KSHV–MCD on follow-up, and one patient who died had no evidence of KSHV–MCD or lymphoma at autopsy (Uldrick et al., 2010a).

We hypothesized that while these symptoms did not appear to be caused by KSHV–MCD, they might nonetheless similarly result from KSHV lytic activation and consequent cytokine production. On investigation, this cohort indeed exhibited KSHV viral burdens and cytokine profiles similar those seen with KSHV lytic activity in MCD flares, including substantial elevation of vIL-6 and hIL-6 (Figure 2; Uldrick et al., 2010a). Patients also exhibited markedly elevated KSHV viral loads in peripheral blood mononuclear cells. While the number of patients is small, the large differences in the immunologic and virologic milieu in these patients compared to KS patients without inflammatory symptoms was statistically significant (Figure 2). Our findings in this small cohort provide evidence that patients with lyrically active KSHV may manifest systemic clinical symptoms even in the absence of the pathologic changes diagnostic of MCD. These features are distinguishable from the chronic immune activation of HIV infection by the presence both of markedly elevated KSHV VLs in peripheral blood and by the detection of vIL-6 in KSHV inflammatory cytokine syndrome (KICS). As seen in Figure 2, neither of these features are evident in patients with KSHV and HIV co-infection who have KS but do not have systemic inflammatory symptoms. While in the original series all patients were HIV infected, we have subsequently observed this syndrome in patients without HIV.

Based on this original cohort, we have proposed naming this putative clinical syndrome KICS. The assessment of this small retrospective cohort of patients suggest that clinical manifestations of KICS are protean, and none are specific to the syndrome. Rather, they likely reflect common endpoints of proinflammatory

Table 1 | Key clinical and laboratory findings in patients with KICS and KSHV–MCD.

| Parameter       | KICS   | KSHV–MCD |
|-----------------|--------|----------|
| Number of patients | 6      | 8        |
| CD4* [cells/μL; median, range] | 255 (28–492) | 189 (21–364) |
| HIV VL* [copies/mL; median, range] | 4650 (<50–110,000) | <50 (<50–664) |
| Sodium (mEq/L; median, range) | 132 (127–139) | 130 (127–136) |
| Albumin (g/dL; median, range) | 2.6 (2.0–3.9) | 2.7 (1.7–3.9) |
| Hemoglobin (g/dL; median, range) | 8.7 (7.0–14.8) | 10.2 (7.4–12.7) |
| Leukocytes (10^3 cells/μL; median, range) | 4.5 (2.3–6.4) | 5.7 (2.7–10.1) |
| Platelets (10^3 cells/μL; median, range) | 158 (62–231) | 126 (11–262) |

*From Uldrick et al., 2010a, used with permission. All patients in both cohorts were HIV-infected.
cytokine deregulation in the setting of KSHV lytic replication and pathologic expression of KSHV encoded genes such as vIL-6. These symptoms may also be observed in other inflammatory states, and at times can be confused clinically with sepsis. KICS is therefore a diagnosis of exclusion, requiring pathologic exclusion of MCD; careful evaluation to exclude alternate explanations for the clinical manifestations, such as serious intercurrent infection; demonstration of systemic inflammation; and demonstration of KSHV viral activity. Based on the initial case series, we have proposed a working case definition for KICS (Table 2), incorporating clinical, laboratory, and radiographic manifestations together with evidence of KSHV activation and systemic inflammation, and a requirement that KSHV–MCD be excluded on pathological examination. A clinical protocol to study this syndrome prospectively is underway at the National Cancer Institute, National Institutes of Health, based on this working definition (NCT01419561).

The precise relationship of KICS to KSHV–MCD is unclear, and it is possible that KICS is a heterogeneous condition. Pathological evaluation and perhaps functional imaging with 18-[F]-fluorodeoxyglucose positron emission tomography (FDG-PET) may shed additional light on this question, and perhaps assist in identifying lymph nodes to biopsy that may reveal KSHV–MCD. We have preliminary evidence to show that in KSHV–MCD, pathologically involved nodes are detectable by FDG-PET, and that PET findings are distinct from those of suppressed HIV or intercurrent lymphoma (Polizzotto et al., 2010). Comparison of PET findings in KICS and KSHV–MCD may also provide a non-invasive means to assess disease activity and shed light on syndrome pathogenesis, including the possible cellular sites of disease activity. It is
also possible that some patients with KICS may represent a pro-
drome to KSHV–MCD or an unusual or limited presentation of
KSHV–MCD, and may eventually evolve into frank KSHV–MCD.
Alternatively, certain patients may have another cellular source
apart from lymphocytes responsible for the excess vIL-6 cytokine
production, perhaps extensive KS spindle cells or KSHV-infected
monocytes or macrophages. Notably, while several patients with
KICS in the original cohort had severe KS, several other patients in
this cohort with severe KS had less severe laboratory abnormali-
ties suggestive of KICS based on the clinical criteria used
for the initial case series. Similarly, PEL patients can exhibit severe
inflammatory symptoms with disease activity, and may exhibit
elevated KSHV viral loads (Uldrick et al., 2011b). Taken together,
these observations raise the possibility that KSHV lytic activa-
tion and associated inflammatory clinical manifestations may be
more common than appreciated in patients with other KSHV-
associated diseases. It is possible that significant clinical overlap
exists between KICS, KS, and PEL.

Among patients with KSHV and HIV co-infection, KICS may be
an important cause of morbidity and mortality. KICS was
associated with 50% mortality in our small series (which likely
captured relatively severe cases). We also hypothesize that KS may
be relatively refractory to therapy in patients with KICS because
the cytokine excess can help drive KS spindle cell proliferation.
In a small number of patients our group has seen with KICS and
KS that was difficult to treat, strategies to reduce KSHV burden
and intercurrent inflammation (including rituximab with cyto-
toxic chemotherapy) appear to have led to improvements in KS.
One can thus hypothesize that KICS may in some cases be associ-
ated with the worsening of KS that can be observed after starting
cART [KS immune reconstitution syndrome (IRIS)], although
KICS has not been formally studied in this setting. Notably, some
investigators report elevated hIL-6 levels in IRIS patients without
KS, though KSHV and viral cytokines have not, to our knowledge,
been assessed in relation to systemic symptoms (Stone et al., 2001).

**RATIONAL THERAPEUTIC APPROACHES IN KSHV–MCD AND
KICS**

There is no standard therapy for KSHV–MCD, and most articles
describing therapy have consisted of case reports or very small series.
Some of the approaches that have shown activity include interferon-alpha, surgical resection of the spleen, cytotoxic
chemotherapy (either single agents or combinations), ganciclovir,
glucocorticoids, and rituximab (Boulanger et al., 2004; Waterston
and Bower, 2004; Dispenzieri and Gertz, 2005; Neuville et al., 2005;
Bower et al., 2007; Mylona et al., 2008; Bower, 2010). Our group has
explored selective targeting of KSHV–MCD plasmablasts based on
their expression of KSHV lytic genes that phosphorylate certain
drugs (Davis et al., 2007). One KSHV lytic gene, ORF36, encodes
a phosphotransferase that activates ganciclovir to a cytotoxic
triphosphate moiety, and another, ORF21, encodes a thymidine
kinase that phosphorylates zidovudine (AZT), again to a cytotoxic
moiety (Cannon et al., 1999; Gustafson et al., 2000). Members of
our group have shown that PEL cells in which KSHV was
lytically activated produced increased amounts of triphosphate
moieties of AZT and ganciclovir and that AZT and ganciclovir
had synergistic toxicity, at doses attainable in patients, against
these PEL lines with activated KSHV (Davis et al., 2007). No
comparable cytotoxicity was seen in non-KSHV-infected Burkitt
lymphoma cell lines. Based on these in vitro results, we explored the
combination of high dose zidovudine and valganciclovir, an
oral pro-drug of ganciclovir, in 14 patients with HIV infection
and KSHV–MCD (Uldrick et al., 2011a). This regimen delivered symp-
tomatic control and normalization of correlates of lytic KSHV
activity (including KSHV viral load, vIL-6, and hIL-6) in a high
proportion of MCD patients. Responses were evaluated based on

### Table 2 | Proposed working case definition of KSHV inflammatory cytokine syndrome.

| 1. CLINICAL MANIFESTATIONS* | b. Laboratory abnormalities |
|-----------------------------|---------------------------|
| a. Symptoms                | Anemia                    |
| Fever                      | Thrombocytopenia           |
| Fatigue                    | Hypoalbuminemia            |
| Edema                      | Hyponatremia               |
| Cachexia                   |                           |
| Respiratory symptoms       |                           |
| (including cough, dyspnea, |                           |
| airway hyperreactivity)    |                           |
| Gastrointestinal disturbance|                           |
| (including nausea, anorexia, abdominal discomfort, altered bowel habit) | |
| Arthralgia and myalgia     |                           |
| Altered mental state       |                           |
| Neuropathy with or without |                           |
| pain                       |                           |
|                         |                           |
| 2. EVIDENCE OF SYSTEMIC INFLAMMATION |                   |
| Elevated C-reactive protein|                           |
|                         |                           |
| 3. EVIDENCE OF KSHV LYTIC ACTIVITY |                   |
| Elevated KSHV viral load in peripheral blood mononuclear cells (> 100 copies/10^6 cells) | |
|                         |                           |
| 4. NO EVIDENCE OF KSHV-ASSOCIATED MULTICENTRIC CASTLEMAN DISEASE |                   |
| Exclusion of MCD requires pathologic assessment lymph node, bone marrow, or spleen | |

The working case definition of KICS requires the presence of at least two clinical manifestations drawn from at least two categories (1a, b, and c), together with each of the criteria in 2, 3, and 4. Clinical manifestations for the working definition are drawn from the initial case series and from findings commonly seen in KSHV–MCD.
newly established criteria (NCI–HAMB criteria) for improvement in clinical symptoms, laboratory parameters (including hemoglobin, platelets, CRP, albumin, and sodium), and radiographic parameters (Uldrick et al., 2011a). Overall, 12 patients (86%) attained major clinical responses by these criteria, with seven patients (50%) attaining clinical complete responses, three (21%) attaining symptom free disease, and two (14%) attaining partial clinical response. In addition, seven patients (50%) attained major biochemical responses, with three (21%) attaining a complete response, four (29%) attaining a partial response. Finally, five patients (36%) attained a major radiographic responses including four (29%) with complete radiographic responses. Relapses were relatively common with this regimen, and the progression-free survival was 6 months. However, patients overall did well, and with 43 months of median follow-up, the overall survival was 86%.

Several recent studies have reported that KSHV–MCD can respond to therapy with the anti-CD20 monoclonal antibody rituximab (Corbellino et al., 2001; Marcelin et al., 2003; Ocio et al., 2005; Bower et al., 2007). It is postulated that in this setting, it acts by targeting the B-lymphocyte KSHV reservoirs, though it should be noted that the level of expression (weak or absent) of CD20 on lytic KSHV-infected B-lymphocytes themselves is contentious. Alternatively, it may target IL-6 production by other B cells that are stimulated by the KSHV-infected plasmablasts. However, different investigators have reported variable results with this agent, and two studies noted that its use was associated with a worsening of KS (Neuville et al., 2005; Bower et al., 2007). Because of this concern, and because many patients with MCD either have KS or are at risk of KS, we have explored the combination of rituximab and liposomal doxorubicin (Doxil) in KSHV–MCD (R/Dox). Liposomal doxorubicin has a high affinity for the pathogenic spindle cell in KS lesions, and, when available, is currently the standard of care for first line therapy of KS (Northfelt et al., 1998; Cianfrocca et al., 2010). It was also possible that it could help target the KSHV-infected MCD plasmablasts. In an ongoing study, we have treated 12 patients with this R/Dox regimen (Uldrick et al., 2010b). R/Dox was highly effective even in heavily pretreated patients. All achieved a clinical CR by NCI–HAMB criteria (after median two cycles therapy, range 1–5); nine (75) achieved a biochemical CR, one (8%) biochemical PR, and two (17%) biochemical SD; while best radiographic response (nodes and spleen) was CR in six (50%) and PR in six (50%). Toxicity was not marked, comprising predominantly infusion reactions and neutropenia.

Until recently, the expected survival of most patients with KSHV–MCD was 2 years or less (Oksenhendler et al., 2000). With the recent advances in therapy, along with the use of highly active anti-retroviral therapy (HAART) for those patients with HIV-associated KSHV–MCD, the overall prognosis of patients with KSHV–MCD has improved markedly, and a number of patients can now attain long remissions. Nonetheless, patients with flares of KSHV–MCD can rapidly become critically ill, with a clinical presentation similar to patients with sepsis, and may require careful monitoring, sometimes in an intensive care unit, during the initial treatment. Our experience is that some patients may benefit from short-term glucocorticoids during initial KSHV–MCD treatment, although patients should be probably weaned as soon as possible as glucocorticoids may mask intercurrent infections and can exacerbate KS (Gill et al., 1989). It should be noted, however, that there is no standard therapy for KSHV–MCD and it may be worthwhile to consider a variety of approaches, depending on disease severity and other parameters. Our group has been exploring the use of consolidation therapy after patients achieve remissions with several months of either interferon-alpha or high dose zidovudine with valganciclovir. Additional studies will be needed to explore new therapies, to establish the best therapy for KSHV–MCD, and perhaps through a combination of treatments, to establish the best way to optimize therapy for different patients with the goal of durable long-term remission.

Because of the small number of patients with KSHV–MCD, there is little commercial interest in developing drugs specifically for this disease, and it is unlikely that a therapeutic antibody to vIL-6 will be developed in the near future. However, an antibody to the IL-6 receptor is in clinical use for rheumatoid arthritis, while a different antibody against IL-6 is being evaluated in the setting of MCD not associated with KSHV. Because hIL-6 is markedly elevated and believed to contribute to disease pathogenesis in KSHV–MCD, antibodies targeting IL-6 signaling may have activity, and we are currently conducting a clinical study of tocilizumab, an anti-IL-6 receptor antibody in patients with symptomatic KSHV–MCD (NCT01441063).

Kaposi sarcoma-associated herpesvirus inflammatory cytokine syndrome is a newly described syndrome, and there is little clinical experience to guide therapy. In those cases where KICS accompanies KS or PEL, it is reasonable to hypothesize that treatment of the underlying tumor may help reduce the KSHV-associated cytokines. However, it should be noted that in the original series describing this syndrome KS was often quite difficult to treat, perhaps because of stimulation by the IL-6 or other cytokines. It may be worthwhile to consider treatment approaches to KICS similar to those used for KSHV–MCD. These may include, for example, antiviral drugs such as ganciclovir with activity against KSHV, the use of high doses of zidovudine or ganciclovir that can be activated by KSHV to toxic moieties as described above, the use of rituximab to kill B cells that may harbor KSHV or be producing cytokines, or the use of liposomal doxorubicin to kill KS spindle cells. In the original series of six patients of KICS, one patient received high dose zidovudine and valganciclovir, one patient received valganciclovir alone, and two received liposomal doxorubicin alone (Uldrick et al., 2010a). The patient who received zidovudine/valganciclovir demonstrated an initial improvement in laboratory parameters, though his clinical course was later complicated significantly by intercurrent illness, while one patient who received valganciclovir monotherapy had a transient clinical response. The two patients who received liposomal doxorubicin (both with KS requiring therapy) also had transient clinical responses.

**SUMMARY AND FUTURE CONSIDERATIONS**

Soon after the discovery of KSHV in 1994, it became apparent that KSHV-associated PEL and KSHV–MCD were separate disease entities. Since that time, we have learned much about the
pathogenesis of these diseases, and for KSHV–MCD in particular, therapeutic advances have led to substantial improvement in survival. At the same time, many questions remain concerning KSHV–MCD and its relationship to other KSHV-induced diseases. More recently, KICS has been described as a new syndrome caused by KSHV; but there is much to be learned about this entity. What cells are primarily responsible for the cytokine overproduction in KICS, and what pathophysiologic mechanisms are responsible for its overproduction? What is the spectrum of disease, and what is its incidence and prevalence in different populations? How is it best managed? It is not unreasonable to hope that answers to many of these questions will be forthcoming in the next decade and that this research will help lead to improved tools to diagnose and treat these entities.

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