Hemophagocytic lymphohistiocytosis in an adult kidney transplant recipient successfully treated by plasmapheresis

A case report and review of the literature

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Abstract

Rationale: Hemophagocytic lymphohistiocytosis (HLH) is a life-threatening disease entity primarily described in children, but not less relevant in adults. It is characterized by a misdirected activation of the immune system, resulting in an uncontrolled cytokine release from macrophages and cytotoxic T-cells (CTLs). Primary HLH relies on a genetic predisposition, whereas secondary HLH develops in the context of infections, malignancies or autoimmune diseases. However, the awareness and therapeutic knowledge for HLH in adulthood is limited. Most therapy protocols are almost exclusively validated in pediatric cohorts and for primary HLH. Their transferability to adult individuals with mostly secondary HLH is doubtful. Especially the high liver and bone marrow toxicity of applied etoposide-based protocols is discussed controversially and connected to overwhelming infections and death.

Patient concern: A 51-year old, male, kidney transplant recipient was admitted to our center suffering from diarrhea, fever, nausea, hyponatremia, kidney graft failure, disorientation, progressive hemodynamic instability, and multiorgan failure.

Diagnoses: Clinical and laboratory findings resembled those of a septic shock. Ferritin and soluble interleukin-2 receptor (sCD25) levels were disproportionally elevated. Only a mild hepatosplenomegaly was diagnosed in a CT scan. A T2-weighted, fluid-attenuated inversion recovery MRI showed marked, bilateral and periventricular white matter hyperintensities. The cerebrospinal fluid (CSF) analysis showed a moderately elevated protein content and cell count. There was no evidence of any bacterial, viral, or parasitic infection. The diagnosis of HLH was made.

Interventions & Outcomes: The patient was successfully treated by a combined approach consisting of plasma exchange (PE), corticosteroids, anakinra, and cyclosporine (CsA).

Lessons: HLH is an important differential diagnosis in critically ill patients. Its unspecific clinical picture complicates an early diagnosis and may be misclassified as sepsis. A combination of plasma exchange (PE), corticosteroids, anakinra, and cyclosporine (CsA) may be a promising and less toxic approach for HLH therapy in adults.

Abbreviations: ALT = alanine aminotransferase, AOSD = adult onset Still’s disease, APC = antigen presenting cell, AST = aspartate aminotransferase, BKV = BK-virus, CMV = cytomegalovirus, CNS = central nervous system, CRP = C-reactive protein, CsA = cyclosporine A, CSF = cerebrospinal fluid, CTLs = cytotoxic T-lymphocytes, DIC = disseminated intravascular coagulation, EBV = Ebstein–Barr virus, FHL = familial hemophagocytic lymphohistiocytosis, HAV = hepatitis A virus, HBV = hepatitis B virus, HCV = hepatitis C virus, HEV = hepatitis E virus, HHV6 = human herpes virus 6, HIV = human immune-deficiency virus, HLH = hemophagocytic lymphohistiocytosis, HSV = herpes simplex virus, IFNγ = interferon gamma, IL = interleukin, IVG = intravenous immunoglobulin, JCV = JC-virus, LDH = lactate dehydrogenase, MAS = macrophage activation syndrome, NK cells = natural killer cells, PCR = polymerase chain reaction, PE = plasma exchange, PML = progressive multifocal leukoencephalopathy, PRES = posterior reversible encephalopathy syndrome, sCD25 or sIL-2r = soluble interleukin-2 receptor, SJIA = systemic juvenile idiopathic arthritis, SLE = systemic lupus erythematosus, SNP = single nucleotide polymorphisms, TLR = toll-like receptors, TNFα = tumor necrosis factor alpha, VZV = varicella zoster virus.
1. Introduction

Hemophagocytic lymphohistiocytosis (HLH) is a life-threatening disease characterized by massive cytokine production from activated blood monocytes, macrophages (histiocytes), and cytotoxic T-lymphocytes (CTLs). The ubiquitous cellular organ infiltration and cytokine release evoke an unspecific and often sepsis-like clinical picture. Refractory and prolonged fever, hepatosplenomegaly, hemophagocytosis in the bone marrow, and several laboratory findings such as cytopenia, very high ferritin levels, low or absent natural killer (NK) cell activity, elevated soluble interleukin-2 receptor (sIL-2r), hypertriglyceridemia and/or low fibrinogen are considered as typical HLH features. These parameters form the widely applied HLH-2004 diagnostic criteria. In general, one has to distinguish between primary and secondary HLH. Primary HLH is either of genetic origin, also called familial HLH (FHL), or associated with genetic immunodeficiency syndromes (Table 1). Secondary or acquired HLH occurs mostly in the context of infections, malignancies, and autoimmune diseases. In addition, cases of acquired HLH are described in patients receiving immunosuppressive therapy after solid organ transplantation. The term “macrophage activation syndrome” (MAS) is particularly used for autoimmune-related secondary HLH. Secondary HLH can occur at any age, whereas FHL manifests mainly during infancy or early childhood. The epidemiological data for HLH are limited, especially in adulthood. Thus, its true incidence is probably unknown. The best data for primary HLH or HLH in childhood comes from three studies, indicating a yearly incidence of 1.2 per million children in Sweden and of 7.5 and 3.3 per 10000 hospitalized children in Turkey and the United States of America, respectively. Only one epidemiological study exists for HLH in adults, reporting an incidence of 3.6 per million for malignancy associated HLH. The overall mortality is high and ranges between 45 and 60% for FHL and 5 and 30% for autoimmune-related MAS in children. The situation in adults is even worse. Recent data suggest an overall mortality of 41%. Thus, early diagnosis and initiation of appropriate measures are essential to improve outcomes and quality of life.

Unfortunately, its nonspecific clinical presentation and sepsis-like appearance makes the diagnosis challenging and suggests a large number of undetected cases with potentially fatal outcomes in adult critically ill patients. A major problem is thereby the limited awareness for HLH, leading at least in part to the high mortality in adults. In addition, most clinical guidelines, diagnostic criteria, and treatment protocols are developed and validated in pediatric patients. It is unclear to what extent these approaches are transferable into an adult patient population. Concerns exist especially with regard to the use of the cytotoxic topoisomerase II inhibitor etoposide that is widely applied during pediatric disease manifestations. Herein, we report for the first time on a HLH case in an adult kidney transplant recipient, who was successfully treated by a less toxic approach consisting of plasma exchange (PE), cyclosporine (CsA), anakinra, and corticosteroids.

2. Case report

In April 2017, a 51-year old, male, kidney transplant recipient was admitted to our center in poor general condition due to excessive diarrhea and dehydration along with fever, nausea, hyponatremia (126 mEq/L), and kidney graft failure. There were no signs for rush or polyarthralgia. The patient had a living donor kidney transplantation in 2002 due to unknown primary renal disease. On admission, the patient was on an immunosuppressive therapy with tacrolimus, mycophenolic acid, and low dose methylprednisolone. An empirical, antibiotic therapy with ciprofloxacin and metronidazole was started and the patient developed progressive tachypnea, hypotonia, and disorientation. An arterial blood gas analysis revealed a metabolic acidosis with partial, respiratory compensation (pH 7.35, pCO2 15mm Hg, pO2 161mm Hg, sHCO3 15.6, sodium 135mEq/L, chloride 114mEq/L).

### Table 1

| Hereditary gene defects predisposing for primary HLH. |
|---|
| **Disease** | **Protein/Gene** | **Function/Specials** |
| Familial HLH (FHL) | | |
| FHL 1 | Unknown/unknown | Unknown |
| FHL 2 | Perforin/PRF | Induction of apoptosis, typically early onset. |
| FHL 3 | MUNC13-4/UNC13D | Vesicle priming, early onset, CNS frequently affected. |
| FHL 4 | Syntaxin 11/STX11 | Vesicle transport, rare, onset beyond first months, typically in Turkish or Arabic patients. |
| FHL 5 | MUNC18-2/STXB2 | Vesicle transport, late onset typically with hypogammaglobulinemia, platelet adhesion defect. |
| Immunodeficiency syndromes associated with HLH | | |
| C22q deficiency | C22q7/C22q7 | Impairs cellular and humoral immunity, EBV-associated, lymphoma risk. |
| Chediak-Higashi Syndrome | LYST | Vesicle transport, late onset, partial albinism, retardation. |
| Griscelli syndrome 2 | RAB27A | Vesicle transport, late onset, partial albinism, CNS frequently affected. |
| IXF deficiency | ITK/ITX | Reduced number of NK cells, EBV-associated. |
| XPL1 deficiency | SAP/HOD1A | Transduction and activation of lymphocytes, X-linked, mostly induced by EBV, severe form, risk of lymphoma, hypogammaglobulinemia. |
| XPL2 (XAP deficiency) | XAP/BRIC1 | X-linked, mild HLH, frequently EBV-associated, colitis. |

CNS = central nervous system, EBV = Epstein–Barr virus, NK = natural killer.
Under the assumption of hyperchloremic metabolic acidosis (normal anion gap) due to severe gastrointestinal bicarbonate and fluid losses and a septic clinical picture, the patient was transferred to our intensive care unit. The antibiotic regime was changed to intravenous meropenem and immunosuppressive therapy was reduced accordingly to a methylprednisolone (20 mg) monotherapy. Typical viral (norovirus, rotavirus, adenovirus, astrovirus cytomegalovirus [CMV]) and bacterial (clostridium difficile, shigella, yersenia, and campylobacter species) pathogens of gastrointestinal infection were excluded and multiple blood cultures remained negative. The neurological condition worsened and C-reactive protein (CRP) levels increased further to 120.9 mg/L. T2-weighted, fluid-attenuated inversion recovery (FLAIR) MRI images showed a marked, heterogeneous white matter hyperintensity of the bilateral periventricular regions with occipital, parieto-frontal, and especially dextral accentuation (Fig. 1).

In the context of immunosuppressive therapy in this patient, a diagnosis of progressive multifocal leukoencephalopathy (PML) due to JC-virus (JCV) reactivation, any other virus-associated encephalitis or posterior reversible encephalopathy syndrome (PRES) was suspected. The cerebrospinal fluid (CSF) analysis showed a moderately elevated spinal fluid protein content (480 mg/L), reduced lactate levels (1.6 mmol/L) and an elevated leucocyte count \(12 \times 10^6/L\), consistent with inflammatory CSF alterations. Antiviral therapy with acyclovir and cidofovir was initiated, but the neurological condition worsened. The patient developed hemodynamic instability together with liver and renal failure and needed mechanical ventilation and vasopressor therapy (norepinephrine 23 μg/kg/min). A CT scan of the thorax and abdomen showed only mild hepatosplenomegaly without further pathologies. The most striking laboratory findings at this time were as follows: ferritin 23623 mg/L, serum creatinine 6.6 mg/dL, platelet count \(66 \times 10^9/L\), leucocyte count \(10.79 \times 10^9/L\), hemoglobin 91 g/L, CRP 160 mg/L, total bilirubin 4.6 mg/dL, aspartate aminotransferase (AST) 6106 U/L, alanine aminotransferase (ALT) 1421 U/L, and lactate dehydrogenase (LDH) 2032 U/L. On the basis of therapy-refractory fever, hepatosplenomegaly, massive hyperferritinemia, progressive thrombocytopenia and anemia (though not fulfilling HLH-2004 criteria), central nervous system (CNS) involvement and a sepsis-like clinical picture without any evidence of a microbiological pathogen, the suspected diagnosis of secondary HLH was established.

At this time, only 3 of the HLH-2004 criteria (fever, splenomegaly, and hyperferritinemia, Table 2) were fulfilled, but the clinical picture was highly suspicious for HLH. Thus, a
Table 2
HLH-2004 diagnostic criteria.[3]

| Features | Cutoffs |
|----------|---------|
| Diagnosis of HLH can be made if either a molecular diagnosis consistent with HLH or five out of the following eight criteria are fulfilled |
| Fever (Hepato-)Splenomegaly | ≥ 2 cell lines |
| Cytopenia | <90 g/L (neonates < 100 g/L) |
| -Hemoglobin | < 100 x 10⁹/L |
| -Platelets | < 1.5 x 10⁹/L |
| -Neutrophils | Hypofibrinogenemia or Fasting hypertriglyceridemia |
| Hemophagocytosis | Bone marrow or tissue |
| Additional new diagnostic criteria (HLH-2004) | Hyperferritinemia ≥500 μg/L |
| Elevated soluble CD25 ≥2400 U/mL |
| Low or absent NK cell activity |
| Additional features (not included in HLH-2004 diagnostic criteria) | Elevated bilirubin, transaminases |
| Elevated lactate dehydrogenase |
| Elevated D-dimers |
| Elevated CSF protein, CSF cells |

CSF = cerebrospinal fluid, HLH = hemophagocytic lymphohistiocytosis, NK = natural killer.

Table 3
Overview of laboratory findings on admission, prior plasma exchange, Peak values for HLH-classification and ambulant follow-up after 2 months.

| Parameters | Admission | Prior peak (d9) | Peak values | Follow-up |
|------------|-----------|----------------|-------------|-----------|
| Creatinine, mg/dL | 4.37 (BL 2.99) | 6.2 | 6.57 (d10) | 7.05 |
| Urea, mg/dL | 119 | 184 | 237 (d11) | 196 |
| LDH, U/L | 136 | 2032 | 2032 (d9) | 220 |
| AST, U/L | 28 | 6106 | 6106 (d9) | 17 |
| Bilirubin, total mg/dL | 0.4 | 4.6 | 4.6 (d9) | 0.3 |
| Triglycerides, mg/dL | n.s. | 210 | 366 (14) | 187 |
| CRP, mg/L | 30.4 | 160.2 | 160.2 (d9) | < 2 |
| WBC, ×10⁹/L | 5.52 | 10.78 | 2.04 (d18) | 7.08 |
| Hemoglobin, g/L | 100 | 91 | 65 (d13) | 91 |
| Platelets, ×10⁹/L | 95 | 66 | 39 (d12) | 175 |
| Fibrinogen, g/L | n.s. | 2.64 | n.s. | n.s. |
| Ferritin, μg/L | 23623 | 23623 (d9) | 1508 |

AST = aspartate transaminase, BL = baseline creatinine, CRP = C-reactive protein, d = days after admission, LDH = lactate dehydrogenase, n.s. = not specified, PE = plasma exchange, WBC = white blood cells.

Table 4
HLH-2004 treatment protocol.[3]

| Substance | Dosing |
|-----------|--------|
| Dexamethasone IV | 10 mg/m² 2 weeks, 5 mg/m² 2 weeks, 2.5 mg/m² 2 weeks, 1.25 mg/m² 2 weeks, tapering over 8 weeks |
| Etoposide IV | 150 mg/m² twice weekly for the first 2 weeks, then weekly during initial therapy |
| Cyclosporine A | Start 6 mg/kg daily (divided in 2 doses), aiming levels around 200 μg/L |
| Intrathecal therapy* | < 1 year 6 mg, 1–2 years 8 mg, 2–3 years 10 mg, >3 years 12 mg |
| Methotrexate | < 1 year 4 mg, 1–2 years 6 mg, 2–3 years 8 mg, >3 years 10 mg |
| Prednisolone | Prophylactic cotrimoxazol (week 1 and onward) |
| Supportive therapy | NIS 0.5 g/kg once every 4 weeks |
| Oral antifungal (1–9 weeks) | |

CSF = cerebrospinal fluid, IV = intravenous, MG = intravenous immunoglobulin.

* Only if progressive neurological symptoms or no improvement of abnormal CSF, maximum of 4 weekly doses.
therapy with 250 mg prednisolone and PE were initiated on day 9 after admission. The laboratory findings prior to initiation of PE are displayed in Table 3. The decision for PE was based on data retrieved from children with autoimmune related MAS that was successfully treated by PE.\textsuperscript{23,24} A therapy with etoposide and CsA according to the HLH-2004 protocol (Table 4) was avoided due to high liver toxicity of etoposide and known association of CsA with suspected PRES and PML development.

Before steroid therapy was started, an active viral infection of the CSF (JCV, varicella zoster (VZV), herpes simplex virus (HSV)) was ruled out by polymerase chain reaction (PCR). During the following 5 days, the patient developed relevant hypertriglyceridemia (366 mg/dL), progressive thrombocytopenia (39 to 10^9/L), anemia (nadir of 6.5 g/dL) and leukocytopenia (3.37 to 10^9/L), before all cell lines recovered under PE therapy. The diagnosis of hemophagocytosis was confirmed in a bone marrow biopsy, but admittedly of minor severity. The diagnosis of HLH could be made according to the HLH-2004 criteria 5 days after initiation of therapeutic measures. In total, 18 plasmaphereses were performed and with initiation of plasma exchange the patient’s condition improved impressively, vasoressor therapy was terminated after the first procedure and the neurological impairment as well as the CNS lesions were regressive in a control MRI only 4 days after initiation of therapy (Fig. 1). The laboratory parameters in relation to PE procedures are shown in Figure 2. Anakinra, a recombinant interleukin-1 (IL-1) receptor antagonist, was started as maintenance therapy on day 19 with 200 mg every other day based on 3 recent publications, which showed a beneficial effect of IL-1 blockade via anakinra in adults suffering from HLH and MAS-like syndrome.\textsuperscript{25–27} After improvement of CNS lesions during PE and establishment of a definite diagnosis of HLH, PRES, and PML were considered as unlikely and a CsA therapy at 3 mg/kg per day was added to the maintenance therapy on day 27. Target

![Figure 2. Laboratory findings in relation to plasma exchange. *Plasma exchange (one plasma volume). AST = aspartate aminotransferase, CRP = C-reactive protein, LDH = lactate dehydrogenase.](image-url)
CsA blood concentration was 150 μg/L. All cell lines recovered further and the PE therapy was terminated on day 31. The prednisolone dose was tapered from 250 mg for 2 days to 125 mg for 5 days to a maintenance dose of 20 mg/day. During the further clinical course a ventilator-associated sepsis with pseudomonas aeruginosa in blood cultures and tracheal secretion was successfully treated with pipericillin/tazobactam and ciprofloxacin that was changed to meropenem according to the antibiotic resistance pattern.

After termination of plasma exchange the ferritin levels remained stable at 2000 to 2500 μg/L during maintenance therapy with CsA, anakinra and low-dose prednisolone. After 48 days in the ICU, the patient was transferred to the general ward and improved further. A potential infectious trigger for HLH and the reason for the initial gastrointestinal symptoms could not be identified. A virologic screening for BV-virus (BKV), CMV, Ebstein–Barr virus (EBV), enterovirus, hepatitis virus A/B/C/E (HAV, HBV, HCV, HEV), HSV, human herpes virus 6 (HHV6), human immune-deficiency virus (HIV), JC-virus (JCV), parainfluenza virus, parvovirus B19, and VZV showed no evidence of an active viral infection. The patient was discharged 75 days after the initial admission.

2.1. Concept of secondary HLH

Secondary or acquired HLH can occur at any age, but generally manifests in older ages, is more frequent than primary HLH, and appears mostly in the context of infectious, malignant or autoimmune diseases.[1] One of the most common factors that trigger the disease are viruses of the herpes family (CMV, HSV, HHV8), especially EBV.[22,28] In addition, other viral (HAV, HBV, HCV, measles, dengue, enterovirus, parvovirus B19, etc.), bacterial (campylobacter, chlamydia, staphylococcus, salmonella, tuberculosi(s), fungal (aspergillus, candida) and parasitic (leishmanial, malaria, toxoplasma) pathogens can serve as a trigger.[29] Malignant disease, most notably leukemia, lymphoma and especially t-cell driven disease entities, are also able to trigger HLH.[31]

The term “Macrophage activation syndrome” (MAS) is particularly used for autoimmune-related, secondary HLH[3] and by some authors also for HLH in adults.[29] The highest incidence of MAS in pediatric rheumatology is found in patients with systemic juvenile idiopathic arthritis (SJIA), systemic lupus erythematous (SLE), and Kawasaki syndrome,[10] whereas adult-onset Still’s disease (AOSD) and SLE play a major role in adulthood.[18] About 7 to 11% of children with SJIA seem to develop MAS[17,31] and according to newer reports up to 30 to 40% may suffer from subclinical MAS.[32] Epidemiological data on secondary HLH/MAS in adults are almost absent. Studies are needed to generate a better understanding of the epidemiology of secondary HLH, especially in adults and nonautoimmune diseases.

2.2. Diagnostic criteria

The first guidelines for the diagnosis of HLH were published by the Histiocyte Society in 1991[36] and in revised version in 2004 (HLH-2004).[3] Today, at least 6 relevant diagnostic guidelines exist,[3,32–36] (Table 5), but only 2 are proposed for use in adult patient population.[35,36] All other guidelines either focus on primary HLH or HLH secondary to autoimmune diseases in children.[1,3,12–34] Thus, validated scores for a reliable and early HLH/MAS diagnosis in adults are still missing apart from autoimmune or malignant diseases (Table 5). Furthermore, the differentiation between sepsis and HLH/MAS remains a major problem in clinical practice, especially when liver, kidney, and multigran failure occur.[17]

2.3. Pathogenesis

Central to the pathogenesis of HLH is a massive production of pro-inflammatory cytokines (“cytokine storm”) through an uncontrolled activation of macrophages and CTLs.[30] An impairment of cytotoxic activity of NK cells and CTLs (mainly cytotoxic CD 8⁺ T-cells) is considered a hallmark of most HLH entities.[22,38] The general understanding is that on the basis of impaired cytotoxic cell function, NK cells, and CTLs fail to sufficiently eliminate antigen-presenting cells, especially macrophages, leading to an inability to terminate immune responses.[10] As a consequence, uncontrolled expansion of macrophages and CTLs with production of large amounts of cytokines occurs with IL-1, IL-6, IL-18, tumor necrosis factor alpha (TNFα), interferon gamma (IFNγ) being the driving forces.[10] The result is an ubiquitous inflammation and cellular organ infiltration, causing the diverse, syndrome-like disease pattern.[28,39] This hypothesis applies to primary HLH, where genetic mutations cause a dysfunction of perforin-mediated cytotoxicity in 97% of FHL patients.[38] However, in patients suffering from secondary HLH the postulated NK cell deficiency is only seen in 22%.[38] Thus, the precise pathogenesis for secondary HLH where normal NK cell function is found remains speculative. In this setting, an acquired immunodeficiency on the basis of a malignancy, a HIV infection,[39] an iatrogenic immunodeficiency due to immunosuppressive drug therapy (solid organ- and stem cell transplantation)[40–42] or any other reason is still discussed as key role in adult HLH pathogenesis[28,42] However, in otherwise healthy and immune competent individuals, it remains unclear how viral infections, autoimmune diseases and malignancies induce secondary HLH. Single nucleotide polymorphisms in cytokine genes, which lead to an aggravated and prolonged immune response are just as considered[43,44] as repeated stimulation of toll-like receptors due to genetic predisposition.[45–47] In summary, the general pathophysiology in acquired HLH remains incompletely understood, especially when NK cell deficiency is absent.

2.4. Clinical picture & classic laboratory findings

HLH has a syndrome-like appearance with characteristic clinical and laboratory findings. Particularly during the disease onset, symptoms are rather unspecific and patients develop a clinical picture with fever, hemodynamic instability, hepatosplenomegaly, CNS affection, and an increase in leukocytes and CRP. Unfortunately, early diagnosis is often hampered by delayed appearance of the characteristic laboratory findings.[5] Fever is induced by a misguided cytokine release, unresponsive to antibiotic treatment and often of subacute, prolonged and recurrent character.[48] It is one of the most consistent symptoms, but certainly not specific for HLH diagnosis.[2,49] CNS symptoms were initially described in FHL,[49,51] but manifest as well in adults suffering from secondary HLH.[52] They range from minor alterations of consciousness, headache, memory loss and disorientation to Guillain–Barré-like syndromes, nerve palsy, meninigismus, and seizures. The CSF in HLH patients usually shows an elevated spinal fluid leukocyte count and a mild to moderate elevation of protein content.[49,52] MRI scans typically display polymorphic, bilateral and especially periventricular white matter hyperintensities in T2-weighted, FLAIR images.[51,53] Therefore,
## Table 5
Relevant HLH/MAS diagnostic criteria[^33-36^].

| Diagnostic features | HLH-94 | HLH-2004 | MAS 05 | HScore | MAS 16 | Tamamyan |
|---------------------|--------|----------|--------|--------|--------|----------|
| Molecular diagnosis | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Fever               | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| (Hepato-)Splenomegaly| ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Cytopenia           | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Hypo(hyper)gammaglobinemia | ✔️ | ✔️   | ✔️   | ✔️   | ✔️   | ✔️       |
| Hypertriglyceridemia | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Hemophagocytosis    | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Hyperferritinemia ≥500 mg/L | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ |
| Elevated sCD25 ≥ 2400 U/mL | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ |
| Low or absent NK cell activity | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ |
| Coagulopathy        | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Elevated LDH        | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Elevated AST        | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Elevated D-Dimers   | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Decreased albumin   | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Monocytosis         | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Elevated B2 microglobulin | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ | ✔️ |
| Immunosuppression   | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| CNS dysfunction     | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |
| Renal failure       | ✔️     | ✔️       | ✔️     | ✔️     | ✔️     | ✔️       |

| Validation cohort   | Mainly FLH | FHL and sHLH in childhood | SJIA-associated MAS | HLH in adults | SJIA-associated MAS | Malignancy-associated HLH in adults |
|---------------------|-------------|---------------------------|---------------------|---------------|---------------------|-------------------------------------|
| Diagnosis           | All required | 5/8 required or molecular diagnosis | SJIA+ ≥2 criteria | Likelihood for HLH | SJIA+ferritin >684 µg/L≥2 criteria | High likelihood for sHLH when ≥5 criteria |

[^33-36^]: AST = aspartate aminotransferase; CNS = central nervous system; FHL = familial or primary HLH; LDH = lactate dehydrogenase; MAS = macrophage activation syndrome; SJIA = systemic juvenile idiopathic arthritis; sHLH = secondary HLH.
HLH patients with CNS affection may often be misdiagnosed with viral or other inflammatory CNS diseases.

2.4.1. Hyperferritinemia. Hyperferritinemia is generally considered as one of the most characteristic findings in HLH. According to a recent study, median and maximal ferritin levels in adults range between 2367 and 25,720 ng/mL, and 6067–684,000 ng/mL, respectively. However, there are also patients, who are diagnosed with HLH with significantly lower ferritin levels (181–2424 ng/mL).[30] Several studies investigated the significance of ferritin levels for HLH diagnosis in children. One study showed a 84% sensitivity for FHL diagnosis for ferritin levels above 500 ng/mL,[3] whereas 2 other studies exhibited a 70% sensitivity and 68% specificity and a 90% sensitivity and 96% specificity for concentrations over 2000 ng/mL and 10,000 ng/mL, respectively.[8,44] The diagnostic value of ferritin concentrations in adult HLH is unknown. A study by Schram et al.[55] suggested that ferritin levels beyond 50,000 ng/mL are associated with a variety of disorders (hemochromatosis, hepatocellular injury, hematologic malignancy, solid malignancy, inflammatory conditions, and hemolytic anemia) and are nonspecific.

2.4.2. Cytopenia. Cytopenia is caused by cytokines, especially IFN-γ and TNF-α and usually affects two or more cell lines. Hemophagocytosis, however, seems to play a minor role.[48] Median thrombocyte counts appear to be lower in adults than in children (6–32 × 10^9/L vs. 44–69 × 10^9/L) and are reduced below 100 × 10^9/L in 86 to 93% of all HLH cases. Median hemoglobin (Hb) levels range from 6.7 to 11.4 g/dL in adults.[2] Hemolysis, however, has to be ruled out in this context. Haptoglobin measurement and exclusion of auto-antibody-mediated hemolysis via Coombs-test are obligatory. The degree of neutrophilcytopenia differs in the literature. While severe agranulocytosis is described in most but not all pediatric cases,[49] neutrophilcytopenia does not represent a characteristic finding in adults.[56]

2.4.3. Hypertriglyceridemia and hypofibrinogenemia. Hypertriglyceridemia is caused by inhibition of the lipoprotein lipase through TNF-α. It occurs in 30 to 89% of HLH patients. Fibrinogen concentrations differ substantially in literature and studies have to be analyzed together with standard coagulation parameters and platelet counts, especially to exclude other coagulopathies of critically ill patients like disseminated intravascular coagulation (DIC).[157]

2.5. Special features

The characteristic clinical picture and the classic laboratory findings are supplemented by special diagnostics, which play a major role in diagnostic criteria of HLH and MAS: hemophagocytosis and sCD25.

2.5.1. Hemophagocytosis. Despite the fact that the disease owes its name to this feature, hemophagocytosis is rather of minor relevance as diagnostic criterion. Hemophagocytosis occurs in many inflammatory conditions such as sepsis, bacterial, and viral infections (influenza, malaria, leishmaniosis), rheumatologic diseases and after blood transfusions.[58,59] It manifests late in the disease course and was found in only 32% of children on admission, but in 85% at the time of diagnosis.[60] In other reports, hemophagocytosis was present in 59 to 100% of children and in 52 to 100% of adults. Although one study claims a high sensitivity of hemophagocytosis for HLH diagnosis of 83%, the specificity is rather low and additional criteria need to be fulfilled to establish a definite diagnosis.

2.5.2. Soluble interleukin-2 receptor (sCD25). The soluble interleukin-2 receptor (sIL-2Rα or sCD25) is a surrogate parameter for T-cell activation[61] and suggested as a sensitive marker for HLH detection. However, the primary data on its sensitivity is derived from only 3 pediatric patient cohorts.[62] On the basis of one report, which showed an 93% sensitivity and 100% specificity for values ≥ 2400 U/mL, this threshold was incorporated in the 2004 HLH diagnostic criteria.[8,62] sCD25 levels are supposed to correlate with higher disease activity, worse clinical response to therapy and worse clinical outcomes, but fail to reliably distinguish between subtypes of HLH/MAS.[62]

In general, median sCD25 values lie between 2963 an d21,500 U/mL, but data in adults are limited. Two studies in adults showed absolute values between 1891 an d206567 U/mL[63,64] and in a recent review 79% of 775 HLH cases had sCD25 concentrations > 2400 U/mL and 38% > 10,000 U/mL.[18] Prospective clinical trials are needed to clarify the diagnostic relevance of sCD25 in adults and nonautoimmune disease entities.

2.6. Therapy/treatment

The first prospective treatment protocol was established in 1994 by the Histioocyte Society[33] but revised in 2004.[3] The HLH-2004 protocol now recommends a combined, early chemo-immunotherapy approach consisting of CsA and pulse therapy with dexamethasone and etoposide for at least 8 weeks.[3] Nevertheless, the severe side-effects of etoposide and especially the high liver toxicity and bone marrow suppression remain a general concern.[10] In this context, deaths due to overwhelming infections are reported in primary and secondary HLH in childhood[11,12,65] Therefore, most clinicians start with an intravenous dexamethasone pulse therapy and added CsA (2–7 mg/kg per day) if there is no rapid response to steroid monotherapy. In most of these patients, CsA induces quick disease control and allows for rapid steroid reduction,[116] but one has to be aware of neurological and renal toxicity. In the end, the HLH-2004 protocol is reserved for more severe and therapy-refractory cases and efforts exist to establish less toxic therapy alternatives. Previous attempts on the basis of PE, intravenous immunoglobulins (IVIG), and methylprednisolone are under investigation.[23,24,66,67]

However, treatment studies in adults, mostly include <20 patients and are of retrospective character.

Furthermore, the studies are extremely heterogenic and drugs were applied in various doses and combinations.[18] Thus, treatment decisions are rather based on clinical expertise, than on reliable data. The situation in patients who develop HLH under immunosuppressive therapy (see also our case report) is even more complex when HLH may be triggered by malignancies or overwhelming infectious diseases. In these patients, the underlying triggering event must be treated but HLH specific therapy may also be needed.

2.6.1. Plasma exchange. In the 1980s, PE was shown to induce transient clinical remission in FHL patients.[66] However, PE was later replaced by chemo-immunotherapy. Lately, PE is regaining attention for acquired HLH as a less toxic therapeutic alternative to the HLH-2004 protocol.[58,67] Especially in autoimmune-related MAS in children, PE combined with corticosteroids has proven therapeutic efficacy by showing an effective disease suppression and improved survival compared to etoposide-based regimens.[23,24,69] Furthermore, in case reports PE seems to be beneficial as salvage therapy, when a CsA and steroid regime fail
to control autoimmune-related MAS on the basis of SJIA or AOSD.\textsuperscript{12,36,70} In adults and apart from autoimmune diseases, there exist only a few case reports about PE as HLH therapy.\textsuperscript{71–73} Most of them showed fatal outcomes, although PE was initially capable of stabilizing the patient’s condition. Taken together, PE may be a promising less toxic therapeutic option, especially to control disease onsets of secondary HLH. However, the data are mostly derived from case reports and after termination of PE; the choice of maintenance therapy has to be discussed. A congeneric therapeutic option may be a highly effective cytokine clearance via high cutoff membranes (cytosorb),\textsuperscript{74} but up to now, there is no report of a use in the context of HLH therapy.

### 2.6.2. Interleukine-1-directed therapy

Independent from the underlying pathogenesis, a common feature of HLH entities is a massive cytokine release. Thus, cytokine-directed therapy, targeting cytokines that are typically involved in HLH pathogenesis, is a rising therapeutic alternative. According to the current knowledge, especially IL-1 may be one promising target.\textsuperscript{30}

IL-1 is a pro-inflammatory cytokine produced by peripheral mononuclear cells.\textsuperscript{75} It leads to leukocyte activation and production of other cytokines like IL-6 and is considered as a relevant molecule in the pathogenesis of SJIA.\textsuperscript{47,76} This is supported by data showing an IL-1 related gene expression profile in the blood of patients with newly diagnosed SJIA.\textsuperscript{46,76}

In addition, serum of SJIA patients is capable of activating IL-1 related genes in monocytes of healthy individuals.\textsuperscript{47} Anakinra, a recombinant IL-1 receptor antagonist, is an established treatment agent in SJIA, inducing a long-lasting remission in more than half of SJIA patients\textsuperscript{77,78} and represents a major therapeutic option in AOSD.\textsuperscript{79} The contribution of IL-1 to HLH development is, however, unclear. Several studies showed a beneficial effect of anakinra in SJIA-associated MAS after inadequate response to steroids and cyclosporine,\textsuperscript{80,81} as well as in AOSD complicated by MAS.\textsuperscript{26,82} The data of anakinra in autoimmune independent HLH entities is rather limited. To our knowledge three studies reported a beneficial effect of anakinra in a heterogeneous adult HLH cohort (after transplantation, autoimmune disease, acute lymphoblastic leukemia, EBV infection).\textsuperscript{27} AOSD related MAS\textsuperscript{26} and a sepsis-related MAS-like syndrome, defined by hepatobiliary dysfunction and DIC,\textsuperscript{25}

### 3. Conclusions

To our knowledge this is the first report of an adult kidney transplant recipient suffering from severe, acquired HLH years after transplantation, who was successfully treated by an initial combination of PE and steroids, followed by a maintenance therapy with anakinra, CsA and low dose steroids. HLH has been reported in kidney transplant recipients, mostly weeks after renal transplantation. In only a few patients HLH occurred years after surgery.\textsuperscript{83} Late occurrence of HLH was seen in patients with neoplasia (T-cell lymphoma, angiosarcoma, karpso sarcoma), parasitic infection (leishmaniosis, toxoplasmosis, babesiosis, histoplasmosis), parvovirus B19 and histoplasmosis.\textsuperscript{84} In our patient, HLH occurred 15 years after living donor kidney transplantation and both solid and hematological malignancies were excluded by means such as multiple CT scans, bone marrow aspiration and flow cytometry. Although, there was no evidence of a viral, bacterial or parasitic infection on admission, the patient presented with severe gastroenteritis, diarrhea, dehydration, and nausea suggesting an infectious HLH trigger. This in line with previous reports where HLH in kidney transplant patients was associated with viral infections such as CMV, EBV, HHV6, HHV8, BKV or with bacterial infections like tuberculosis.\textsuperscript{44}

In our patient, the sepsis-like appearance delayed the diagnosis of HLH and the initiation of adequate therapeutic measures. The patient suffered from hepatic and renal transplant failure, severe neurologic dysfunction and relevant hemodynamic instability. Until the time of initiation of PE therapy, none of the 6 known HLH diagnostic criteria were fulfilled and characteristic features like hypertriglyceridemia and relevant pancytopenia according to the HLH 2004 criteria occurred late.\textsuperscript{3} Additionally, hemophagocytosis was of only minor severity in an early bone marrow aspiration and could rather be attributable to any other inflammatory condition like sepsis. This observation is consistent with reports from other authors, who found that hemophagocytosis lacked specificity for HLH, when not supported by other criteria.\textsuperscript{85} sCD25 is considered as a parameter with higher sensitivity.\textsuperscript{86} A retrospective analysis of a sample that was drawn in our patient 3 days before initiation of PE, confirmed a significantly increased sCD25 level of 5450 U/mL. Thus, although we did not measure NK cell activity, our case report suggests that the HLH diagnostic criteria (HLH-2004) may not be sufficient to diagnose the disease in adults at early stages.

High ferritin together with persistent fever, progressive thrombocytopenia or anemia (even when not fulfilling HLH 2004 cutoffs) and CNS impairment pointed to the suspected diagnosis of HLH in our case. But once the diagnosis was considered the question came to therapeutic measures. In our patient, several contraindications existed for the HLH-2004 protocol (Table 4). Other causes of the CNS symptoms such as PRES, PML or viral infection have not been ruled out completely during MRI scan and CSF analysis and the patient suffered from progressive pancytopenia and severe hepatic failure, so that the application of immunosuppressive therapy or etoposide were not reasonable at that time. Thus, we considered cytokine clearance via a PE and corticosteroid-based regime as less toxic therapy alternative, especially to account for the mentioned contraindications. It has already been shown in children that secondary HLH can be successfully treated with a combination of PE, IVIG and methylprednisolone. This approach led to improved survival compared to etoposide- or CsA-based regimes.\textsuperscript{84} In adults, however, treatment with PE had only been reported in the form of case reports\textsuperscript{71–73} with mostly fatal outcomes. The PE frequency was low, ranging from one procedure at all\textsuperscript{71,73} to twice per week\textsuperscript{72} and maintenance therapies were diverse including etoposide- and CsA-based regimes,\textsuperscript{71} or corticosteroid monotherapy,\textsuperscript{73} suggesting an insufficient concept for disease control after PE termination.

In our case, a high frequency PE led to a tremendous improvement of the patient’s general condition as well as the neurological status. After the occurrence of further characteristic HLH symptoms (hypertriglyceridemia, relevant pancytopenia) over the next days, PRES, viral disease and PML were considered as unlikely and a maintenance therapy with anakinra and CsA was started. PE was terminated after 18 procedures. All laboratory findings turned to normal, the disease activity was sufficiently suppressed under a maintenance therapy with anakinra and low dose corticosteroids and the patient could be discharged 75 days after admission. Several follow-ups showed a sufficient disease control. The kidney graft, however, did not recover and we started with intermittent hemodialysis two months after discharge.
In summary, this case report suggests high frequency PE in combination with steroids as an effective and less-toxic alternative to the HLH-2004 protocol for severe HLH onsets in adults or as a bridging approach as long as contraindications to more aggressive therapies exist. Furthermore, the maintenance therapy with CsA, anakinra and low dose corticosteroids showed excellent disease control after termination of PE. The relevance of PE as a first line treatment option for severe HLH in adulthood, as well as the here applied combination of CsA and IL-1-directed maintenance therapy by anakinra has to be assessed in prospective future studies.

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