

Evans syndrome: clinical perspectives, biological insights and treatment modalities

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Abstract: Evans syndrome (ES) is a rare and chronic autoimmune disease characterized by autoimmune hemolytic anemia and immune thrombocytopenic purpura with a positive direct anti-human globulin test. It is classified as primary and secondary, with the frequency in patients with autoimmune hemolytic anemia being 37%–73%. It predominates in children, mainly due to primary immunodeficiencies or autoimmune lymphoproliferative syndrome. ES during pregnancy is associated with high fetal morbidity, including severe hemolysis and intracranial bleeding with neurological sequelae and death. The clinical presentation can include fatigue, pallor, jaundice and mucosal bleeding, with remissions and exacerbations during the person's lifetime, and acute manifestations as catastrophic bleeding and massive hemolysis. Recent molecular theories explaining the physiopathology of ES include deficiencies of CTLA-4, LRBA, TPP2 and a decreased CD4/CD8 ratio. As in other autoimmune cytopenias, there is no established evidence-based treatment and steroids are the first-line therapy, with intravenous immunoglobulin administered as a life-saving resource in cases of severe immune thrombocytopenic purpura manifestations. Second-line treatment for refractory ES includes rituximab, mofetil mycophenolate, cyclosporine, vincristine, azathioprine, sirolimus and thrombopoietin receptor agonists. In cases unresponsive to immunosuppressive agents, hematopoietic stem cell transplantation has been successful, although it is necessary to consider its potential serious adverse effects. In conclusion, ES is a disease with a heterogeneous course that remains challenging to patients and physicians, with prospective clinical trials needed to explore potential targeted therapy to achieve an improved long-term response or even a cure.

Keywords: Evans syndrome, autoimmune cytopenias, low-dose rituximab, IVIG, systemic lupus erythematosus, HSCT, sirolimus, mofetil mycophenolate, HSCT

Introduction

Evans syndrome (ES) is an uncommon autoimmune disease that was defined by Robert Evans in 1951 when he studied the relationship between autoimmune hemolytic anemia (AIHA) and immune thrombocytopenic purpura (ITP). He described the first diagnosis criteria of ES, including the presence of anemia, reticulocytosis, increased blood bilirubin and fecal urobilinogen, no family history of hemolytic diseases, evidence of antibodies against erythrocytes at 37°C, hemolysis of transfused erythrocytes, the presence of purpura, prolonged bleeding time, bone marrow aspiration with normal or increased number of megakaryocytes and the absence of exogenous toxic agents or a baseline disease. Evans divided five groups of patients and demonstrated the presence of a serum agglutinating factor in patients with ITP¹ based on previously reported

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observations in guinea pigs and rabbits.² Some authors modified the definition of Evans as a destruction of at least two hematological blood lineages, as observed in patients with neutropenia, hemolytic anemia and thrombocytopenia.^{3,4} Importantly, AIHA is a complex syndrome with a warm, cold, mixed and biphasic thermal behavior of antibodies, but only warm ones characterize ES.

The typical course of ES is characterized by an heterogeneous chronic disease with clinical variability at onset, spontaneous remissions and exacerbations.^{1,5,6} Its worldwide frequency is unknown; however, research of ES in AIHA is available in some cohorts that report an incidence of about 37%–73% in this setting.^{6,7} A higher rate in female gender has been reported in 60%–70% of ES patients.^{5,8–10}

Diagnosis

First, an exhaustive clinical history must be taken to determine the risk factors for developing ES, such as infections,

malignancies, autoimmune diseases, recent vaccinations, drugs or a family history of immune disorders,¹¹ complemented with a thorough physical examination focused on signs of anemia or thrombocytopenia. It is important to mention that primary ES is a diagnosis of exclusion and secondary ES must be searched for to determine a baseline disease because treatment and responses considerably differ between primary and secondary ES.

Characterization of laboratory features requires a complete blood count and direct examination of peripheral blood; anemia, thrombocytopenia, reticulocytosis, poikilocytosis, mainly due to the presence of spherocytes,^{5,9} increased indirect bilirubin and lactate dehydrogenase and a positive direct anti-human globulin test (DAT) test confirming ongoing immune hemolysis¹² are to be expected. The main laboratory characteristics are summarized in Figure 1.

Additional studies can include the detection of antibodies against platelets (PA-IgG), documented in 35% of individu-

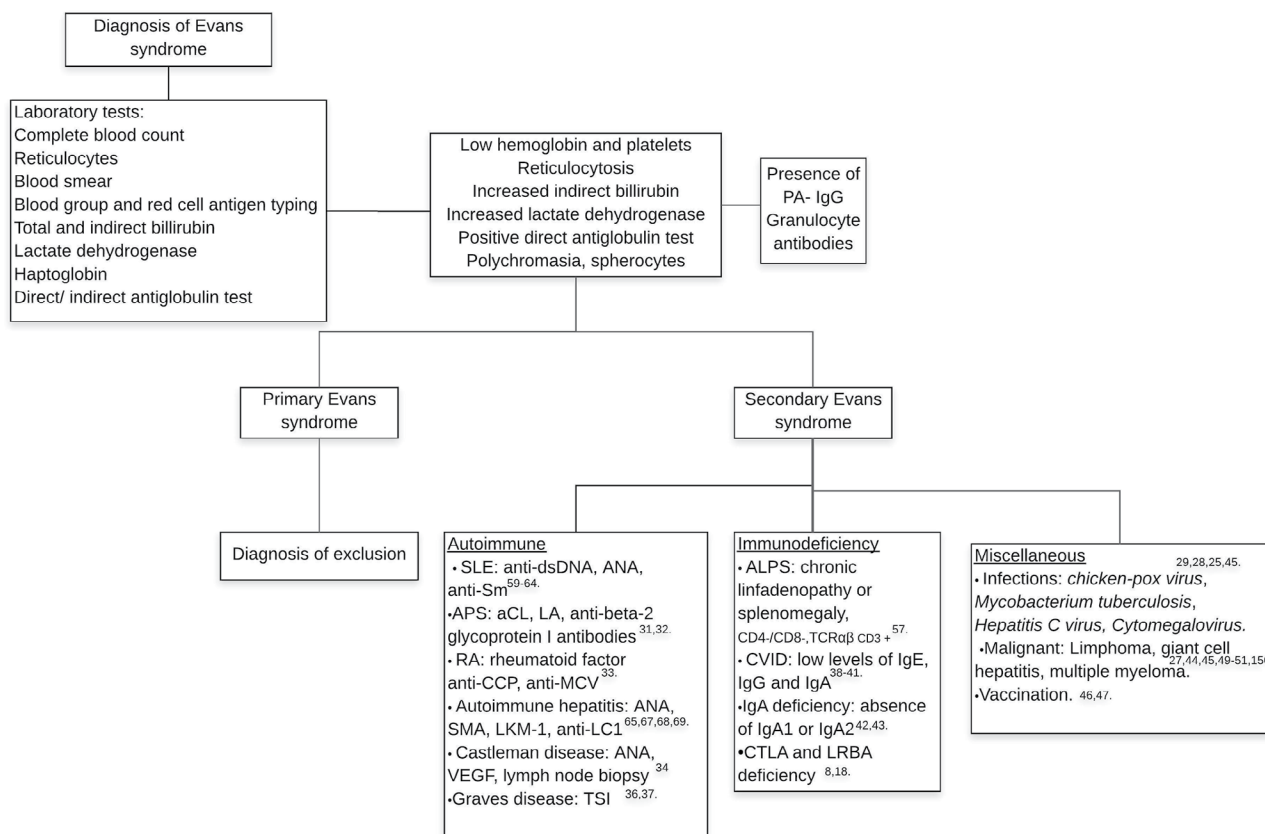


Figure 1 Diagnostic approach for Evans syndrome.

Abbreviations: aCL, anticardiolipin; ALPS, autoimmune lymphoproliferative syndrome; ANA, antinuclear antibodies; anti-CCP, anti-cyclic citrullinated peptide; anti-LC1, anti-liver cytosol antibody; anti-MCV, anti-mutated citrullinated vimentin; anti-Sm, anti-Smith antibody; APS, antiphospholipid syndrome; CVID, common variable immunodeficiency; LA, lupus anticoagulant; LKM-1, liver kidney microsomal type I antibodies; PA-IgG, platelet-associated IgG; RA, rheumatoid arthritis; SLE, systemic lupus erythematosus; SMA, smooth muscle antibody; TSI, thyroid stimulating immunoglobulin; VEGF, vascular endothelial growth factor.

als, mostly type anti-IIb-IIIa.⁹ Anti-granulocyte antibodies can be detected in some patients, but a lack of these antibodies does not exclude neutrophil involvement. Other antibodies must be considered to identify main autoimmune diseases associated with ES. The diagnostic work up is shown in Figure 1.

Clinical features

Clinical features are associated with anemia and thrombocytopenia including pallor, weakness, fatigue, jaundice, petechiae, ecchymosis, gingivorrhagia and epistaxis.^{5,8,10} Due to the prolonged immunosuppressive therapy and/or the associated underlying immune deficit, there is a risk of 66.6% of patients developing respiratory tract infections.⁸

Differential diagnosis of ES

There are some diseases with similar characteristics to ES. Main differential diagnoses are as follows: hemolytic uremic syndrome, characterized by microangiopathic hemolytic anemia, thrombocytopenia, leukocytosis, negative DAT test, clinically manifested as diarrhea, vomiting, fever, pallor, acute renal failure in children predominantly secondary to infections with *Escherichia coli*, *Shigella dysenteriae* or *Streptococcus pneumoniae*;¹³ thrombotic thrombocytopenic purpura, a disease associated with a functional deficit of ADAMTS 13 presenting as microangiopathic hemolytic anemia, thrombocytopenia, fever and neurological or kidney impairment;¹⁴ Kasabach–Merritt syndrome, a rare disease predominant in early infancy with a high mortality rate of about 30% characterized by microangiopathic hemolytic anemia, thrombocytopenia, low fibrinogen levels, increment of D-dimer and giant hemangioma;^{15,16} and paroxysmal nocturnal hemoglobinuria, a chronic life-threatening disease with hemolytic anemia, pancytopenia and thrombosis of vital

organs, mainly in adults. These pathologies must be excluded because treatment is considerably different in each.

Pathophysiology

Several authors have proposed different disease pathways, summarized by the presence of immune dysregulation with antibodies against erythrocytes, platelets and/or granulocytes¹⁷ and decreased CD4:CD8 ratio. Mechanisms currently proposed for explaining ES are shown in Table 1; the pathophysiology of ES remains unknown. In the following section, immunologic alterations observed in this syndrome are exposed in the larger context of autoimmunity and other relevant data are theoretically extrapolated to the disease.

Deficiency of CTLA-4 (CD152) and LRBA

CTLA-4 or CD152 is an inhibitory transmembrane receptor at the surface of regulatory T-cells that bind with a high affinity to CD80/CD86 molecules in antigen-presenting cells with subsequent endocytosis and downregulation contributing to immune homeostasis. This deficiency has been related to ES;¹⁸ on the other hand, LRBA is an intracellular protein that binds to CTLA-4 cytoplasmic fraction in regulatory T-cells following its endocytosis, avoiding its degradation. CTLA-4 deficiency as a cause of alteration in T-cell homeostasis has been documented in an animal model.¹⁹ Recently, a gene mutation in LRBA and CTLA-4 proteins has been related to patients with ES;⁸ also, this immune dysregulation has been found in recurrent severe opportunistic infections in humans and fatal lymphoproliferative disease in mice.^{8,20}

Deficit of TPP2

A recent clinical trial revealed that tripeptidyl peptidase 2 (TPP2) is a molecule with an important role in aging, immu-

Table 1 Principal mechanisms proposed to explain the pathogenesis of Evans syndrome

Studies	Design	Population	Patients	Mechanism
Besnard et al, 2018 ⁸	Retrospective cohort	Children with ES	18	CTLA-4, LRBA, KRAS, STAT3 gain-of-function mutations are associated with ES secondary to a loss of T-cell homeostasis.
Stepensky et al, 2015 ²¹	Clinical trial	Children with ES	2	TPP2 is an essential molecule for cell survival under stress; deficiency is associated with higher levels of autoantibodies, ABCs, perforins and MHC I, with a consequent immunosenescence; there is an increased risk for viral infections and development of autoimmunity.
Karakantza et al, 2000 ²³	Short report	Children with ES	1	Increased levels of Th1 cytokines, especially INF- γ , explains the presence of autoreactive B-cells against platelets and erythrocytes, confirmed by the lower levels of IFN- γ after splenectomy.

Abbreviations: ABCs, age-associated B cells; ES, Evans syndrome; INF, interferon; MHC I, major histocompatibility complex I; CTLA-4, cytotoxic T-lymphocyte antigen 4; LRBA, lipopolysaccharide responsive beige-like anchor protein; TPP2, tripeptidyl peptidase 2.

nosenescence, autoimmunity and tumorigenesis. Its deficiency is associated with ES.²¹ Serological analysis shows that the deficit of TPP2 is related to the presence of anti-nucleolar, anti-cytoplasmic, anti-nuclear antibodies and an increased level of age-associated B cells (ABCs), CD11c⁺, with a higher expression of major complex histocompatibility I molecules. This may indicate the presence of antigens on their surface, explaining the association between ABCs and autoimmune diseases.²¹ ABCs are usually present in the elderly and have been demonstrated in patients with autoimmune diseases including systemic lupus erythematosus (SLE), rheumatoid arthritis or scleroderma and systemic sclerosis.²² Further studies are needed to clearly understand the specific role of these cells in ES and autoimmune disease pathophysiology.

Decreased CD4/CD8 ratio

Abnormalities of immune regulation have been demonstrated in ES patients with a decreased level of T helper and increased T cytotoxic cells with a low CD4:CD8 ratio compared with healthy controls. Excessive production of T cytotoxic lymphocytes or a diminished activity of T helper cells could avoid the production of immunoglobulins (Ig) M and G, as observed in common variable immunodeficiency (CVID) patients.¹⁷ It has been recently shown in vitro that after CD8+ T-cell activation, there are increased levels of interferon gamma with a subsequent stimulation of autoreactive B-cells against erythrocytes and platelets.²³ It could be that this increased activity of T-CD8+ cells favors deficiency of CTLA-4, which regulates inflammatory cytokine production.²⁰ These mechanisms could plausibly cause a disruption of T-cell homeostasis with an elevated concentration of interferon gamma and subsequent activation of B-cells against erythrocytes and platelets.

Classification

Classification of ES includes primary/idiopathic, with this being an exclusion diagnosis with no underlying condition, and secondary in the presence of a baseline disease. It has been demonstrated that secondary disease responds better than primary ES.²⁴ Multiple miscellaneous causes of secondary ES have been documented.^{25–52}

Secondary ES

Autoimmune lymphoproliferative syndrome

Previously named Canale–Smith syndrome,⁵³ autoimmune lymphoproliferative syndrome (ALPS) is a disorder characterized by a Fas gene mutation with an alteration in T-cell apoptotic pathways causing lymphoproliferation manifested

as hepatomegaly, splenomegaly, lymphadenopathies, an increased risk of Hodgkin lymphoma and immune cytopenia, mainly AIHA and ITP, with the presence of antibodies against erythrocytes and platelets,⁵⁴ features similar to those observed in autoimmune multilineage cytopenia.¹⁰ Diagnostic criteria for ALPS were established in 1999 by the National Institutes of Health and require two necessary criteria, such as the presence of chronic lymphoproliferation, increased levels of double-negative T-cells and one primary accessory criterion with a deficit of apoptosis in vitro or genetic mutation in FAS, FASL and/or CASP 10.⁵⁵ Some authors suggest ruling out this diagnosis in all pediatric patients with AIHA and ES.⁵⁴ Others consider ALPS as a cause of secondary ES, as reported in a center with a higher frequency of concomitant ALPS and ES in 47%–50% of patients, confirmed by the presence of elevated double-negative T-cells.^{56,57} It is important to diagnose ALPS before starting treatment, since it is not recommended to administer rituximab to these patients except in specific severe unresponsive cases because of an association with developing hypogammaglobulinemia, which has been reported. CVID has also been considered as a cause of secondary ES.^{38,40} A recent study documented the increased expression of Fas in patients with ES and CVID.⁴¹ Interestingly, an association between ALPS and ES has been documented only in children.^{56,57}

Systemic lupus erythematosus

Several studies have demonstrated the relationship between SLE and ES;^{9,10,58} prevalence of secondary ES in SLE has been reported in 1.7%–2.7% of cases.^{24,59} Hemorrhagic manifestations are frequently observed in this group, including petechiae, ecchymosis, epistaxis, gingivorraghia, gastrointestinal bleeding and hematuria.^{5,59} Spontaneous splenic rupture as an initial manifestation of ES has been documented,⁶⁰ confirming the heterogeneous course of the disease. Steroids are the suggested first line of therapy.⁶¹ In refractory cases, rituximab at standard doses has proven effective with a reduction in PA-IgG, DAT strength,⁶² decreased antinuclear antibodies (ANA) titer, and anti-CD19, -CD21, -B3, -TNF- α ,⁶³ 1–6 months after treatment.^{62,63} Autologous hematopoietic stem cell transplant in a refractory case with SLE, antiphospholipid syndrome and ES led to clinical remission, negative ANA and DAT tests, plausibly explained by a graft vs immunity effect.⁶⁴ Interestingly, secondary ES has a longer remission than the primary disease.²⁴

Autoimmune hepatitis

The relationship between ES and autoimmune hepatitis has been reported by few centers, with main clinical features

including pallor, fatigue and splenomegaly.⁶⁵ Diagnosis for autoimmune hepatitis and ES is confirmed by the presence of elevated hepatic enzymes, ANA, anti-smooth muscle antibody, DAT, PA-IgG and a confirmatory biopsy with rosette formation.⁶⁶ Initial treatment with prednisolone and rituximab is recommended to prevent refractoriness as observed with some immunosuppressants such as azathioprine, mycophenolate mofetil (MMF) and intravenous immunoglobulin (IVIG). Improvement in cytopenias and normalization of hepatic enzymes have been observed, mainly with rituximab, in a year of follow-up.^{65,67–69}

Chronic lymphocytic leukemia

CLL is a worldwide common cause of hematologic cancer in the elderly.⁷⁰ It is frequently associated with autoimmune cytopenias.^{70,71} The frequency of ES in CLL is 2.9%.⁷² A diagnosis of secondary ES in CLL requires the presence of AIHA and ITP at onset or within 1 month of the baseline disease. Determination of genetic mutations is necessary to establish prognosis in CLL. A 13q14 deletion is the most common chromosomal alteration and is associated with a good prognosis;⁷³ however, poor prognostic factors are predominant in patients with ES and CLL, such as a 17p13 deletion, presence of ZAP-70 and a mutation of TP53,⁷² explaining the lower survival rate in these cases, compared to patients with only CLL.

ES during childhood

The incidence of ES in children is unknown due to a lack of published studies, with the few available consisting of case reports or retrospective cohorts.^{6,12,59,74–76} From the largest cohorts of autoimmune cytopenias, it can be concluded that ES is more frequent during childhood; in a study, 11 (4.1%) of 179 children had ES⁶ vs 6 (0.78%) of 766 adults.⁷⁷ It is important to note that autoimmune cytopenias in children are frequently associated with primary immunodeficiencies and in this group, they should always be ruled out.⁷⁸

A referral center of autoimmune cytopenias in France has been operating since 2004, and children with ITP, AIHA and ES have been prospectively evaluated in the OBS'CEREVANCE cohort.^{7,79} A report from this prospective cohort included 156 children from 26 different centers; median age was 5.4 years (0.2–17.2) with a male:female ratio of 1.44. Sixty-nine percent of children required more than one line of treatment, including rituximab, azathioprine, splenectomy, cyclosporine and MMF.⁸⁰

A large multicenter retrospective study in 21 hospitals included 42 children with a median age of 7.7 years. These patients received 1–12 modalities of treatment, mostly

steroids plus IVIG, whereas others underwent splenectomy and/or received cyclosporine, vincristine, danazol, azathioprine, cyclophosphamide and plasmapheresis, with no optimal response. Twenty-four (58%) patients had complications – 12 hemorrhagic and 12 suffered severe infections, mainly sepsis, pneumonia, meningitis, abscess and osteomyelitis.³

In pediatric ES cases lack of response or relapse can be severe, even after splenectomy or the use of multiple immunosuppressive agents. In addition, no large clinical trials assessing outcomes for different treatment strategies and long-term responses in children have been published.

ES during pregnancy

ES during pregnancy is not frequent and usually the diagnosis is established previously; main differential diagnoses in this circumstance are HELLP syndrome, thrombotic thrombocytopenic purpura and uremic hemolytic syndrome. There is a passive immune transfer of maternal IgG antibodies via placenta to the fetal circulation, which explains the transient thrombocytopenia or hemolytic anemia previously reported in newborns or fetus of women with autoimmune cytopenia.⁸¹ In general, pregnant women with ES have a good outcome if appropriate treatment is administered opportunely; however, there is an increased risk of developing abruptio placentae and postpartum hemorrhage. Fetuses usually have a bad prognosis associated with high-risk complications occurring such as severe hemolytic anemia or hemorrhages secondary to significant thrombocytopenia, mainly intracranial bleeding with intra-extrauterine death or neurological impairments. Hematological therapy is challenging because of drugs' teratogenic effects; treatment of choice is steroids plus IVIG. Its action mechanisms include a decreased level of antibodies crossing the placenta⁸² and of maternal IgG antibodies by downregulation, with a reduction of these in fetal circulation.⁸³ Last treatment options are azathioprine or splenectomy, exclusively in refractory third trimester cases.⁸⁴

Treatment

There are no clinical trials available for ES treatment and indications for starting therapy have not been established by evidence-based studies. Few retrospective reports and small cohorts have documented responses which are summarized in Table 2.

It is reasonable to treat patients with clinically significant decreased platelet and hemoglobin levels; nevertheless, as observed in ITP, the decision to start therapy in non-symptomatic patients varies in each case depending on physician experience.¹¹

Table 2 Second-line therapeutic options for patients with Evans syndrome

Drug	Dose	Route of administration	Results
Rituximab	N/A ⁹	N/A	CR: 5; PR: 4; NR: 2
	375 mg/m ² /week/4 doses ⁹⁸	N/A	Remission: 1/1
	375 mg/m ² /week/3–4 doses ⁷⁴	N/A	Remission: 1/3
	375 mg/m ² /week/4 doses ⁹⁹	N/A	Good response
	375 mg/m ² /week/4 doses ¹⁰⁰	N/A	Remission: 1/1
	375 mg/m ² /week/4 doses ¹⁰¹	IV	Remission: 1/1
	375 mg/m ² /week/4 doses ¹⁰²	IV	Good response: 1/1
	375 mg/m ² /week/4 doses ³⁴	IV	NR: 1
	375 mg/m ² /day/8 doses ¹⁰³	N/A	NR: 1
	375 mg/m ² /week/3 doses ⁹⁶	IV	Response: 5/5
	375 mg/m ² /week/4 doses ¹⁰⁴	IV	Remission: 1/1
	375 mg/m ² /week ⁹⁷	IV	Patient 1: NR to AIHA, but CR for ITP; Patient 2: NR; Patient 3: CR for AIHA, NR for ITP; Patient 4: NR to AIHA
	375 mg/m ² /3 weeks/5 doses ¹⁰⁵	IV	Good response: 1/1
	375 mg/m ² /week ¹⁰⁶	IV	Remission: 1/1
375 mg/m ² /week/6 weeks ¹⁵⁷	IV	Remission: 2/2	
375 mg/m ² /week/4 weeks ⁶⁹	IV	Response: 1/1	
Alentuzumab	10 mg/day/10 days ¹³⁴	IV	Response: 2; transient response: 1
Mycophenolate mofetil	650 mg/m ² /b.d. ¹⁰⁸	N/A	Response: 1/4
	50 mg/kg/day ¹¹¹	N/A	Complete remission: 1/1
	50 mg/kg/day ¹¹⁰	N/A	Sustained remission: 1/1
	500 mg/day, then 1–3 g/day ¹⁵⁸	N/A	Response: 1/0, one with ES
Cyclosporine	500 mg/day/b.d., then 1 g/day/b.d. ¹⁰⁷	N/A	Response: 4
	N/A ¹¹⁴	N/A	Relapse
	6 mg/kg/day ¹¹⁵	N/A	Response: 1/1
	N/A ¹¹⁹	N/A	CR: 16/18
	10 mg/kg/day/8 months ¹¹⁶	N/A	Remission: 1/1
	5 mg/kg/day/b.d. for 6 days, then 3 mg/kg/day ¹¹⁸	N/A	PR: 1/1
Cyclophosphamide	5–6 mg/kg/day divided b.d. ¹²⁰	Oral	Response: 5
	10 mg/kg/day, alternative days ¹¹⁷	N/A	Response
	1,000 mg ¹²⁹	IV	CR: 6/8
	50 mg/kg/day/4 days ¹⁵⁹	N/A	Remission: 1, NR: 1
	500 mg/m ² /4 weeks ¹³⁰	N/A	PR: 1/1

Abbreviations: AIHA, autoimmune hemolytic anemia; b.d., twice daily; CR, complete response; ES, Evans syndrome; ITP, immune thrombocytopenic purpura; IV, intravenous; N/A, not available; NR, no response; PR, partial response.

The main objective consists of achieving a long-term complete response. There is no therapeutic regimen established. Steroids with and without IVIG are recommended as front-line therapy.^{11,12,85} Red blood cell/platelet transfusion is indicated only in severe symptomatic patients due to the risk of exacerbations.¹¹

First-line treatments

Steroids

It is postulated that steroids inhibit the ability of macrophages for clearing platelets and erythrocytes.⁸⁶ Prednisolone or prednisone at 1–2 mg/kg/day^{11,12} must be administered in

all cases, and in patients with severe clinical manifestations, an initial dose of 4–6 mg/kg/day within the first 72 hours is recommended. The initial response rate documented for patients who receive steroids at 1–2 mg/kg/day is about 82%–83%.⁹ A regimen including a prednisolone megadose of 30 mg/kg/day for 3 days, then 20 mg/kg/day for 4 days, followed by progressive tapering to 10, 5, 2, 1 mg/kg/week has been administered and most patients have shown complete response.^{6,11,75} Three weeks after starting treatment patients need a re-evaluation; if complete response is achieved, slow tapering within 6 months is necessary to avoid adverse effects. In patients with a partial response, the initial

dose of prednisolone must continue for two more consecutive weeks.¹¹ If no response is achieved, it is mandatory to start second-line therapy.¹² A loss of response is mainly associated with tapering the steroid dose or viral infections.⁶ IVIG may be added if ITP is prevalent.⁸⁷ In cases where thrombocytopenia predominates, the presence of hepatomegaly has been related to a good response.⁸⁸

Intravenous immunoglobulin

Concentrated IgG from human plasma donors acts by blocking the FC γ receptor on the macrophages,⁸⁹ which remains a controversial therapy. In some centers, it is recommended as a first-line treatment together with steroids, and in others as a complementary drug in severe thrombocytopenic patients. A Canadian guideline recommends IVIG only in patients with predominance of ITP symptoms; in cases where AIHA symptoms are more conspicuous, other treatments are preferred.⁹⁰ IVIG was first indicated in 1950 for primary immune deficiencies, but it is currently an important therapy for autoimmune and hematologic diseases such as acquired hemophilia, acquired hypogammaglobulinemia, acquired von Willebrand disease, aplastic anemia, AIHA, autoimmune neutropenia and ES.⁹⁰ This is an important life-saving resource in uncontrolled patients;¹¹ the most common dose administered is 0.4 g/kg/day for 4 days and 5 g/kg in the case of predominance of AIHA.⁹¹ Before using this therapeutic modality, it is important to consider its expense which is about 4,000 Canadian dollars per dose.⁹⁰ A high risk of presenting adverse effects including fever, headache, fatigue, nausea, palpitations and myalgias occurs in 20%–30% of patients.^{92,93} Severe complications have been documented, consisting mainly of acute kidney injury, arterial and venous thromboembolic events, aseptic meningitis, hemolytic anemia and infections.⁹⁴

Second-line treatment

Rituximab

Rituximab is a chimeric anti-CD20 targeted drug that has been increasingly used as the second-line treatment in steroid-refractory or relapsing ES. It has an immunosuppressive effect with B-cell depletion in peripheral blood after one infusion.⁹⁵

A pediatric multicentric prospective study analyzed the efficacy of rituximab at 375 mg/m² in 15 patients with refractory AIHA. All patients previously received two or more courses of immunosuppressive agents (steroids, azathioprine, IVIG, cyclosporine and azathioprine) and 5/15 had ES; of these 5 patients, 3 responded with an increment in platelet

levels from 27 to 140 \times 10⁹/L, 2 months after starting rituximab.⁹⁶ Another multicenter cohort retrospectively evaluated 68 adults with ES; eleven received rituximab at the standard dose with an initial response of 82% after a mean of 12 months, and two of the eleven patients relapsed.⁹

A retrospective research investigated the use of rituximab at the standard dose plus prednisone in 17 refractory children with ES. An overall response of 76% was achieved; the main side effects during rituximab administration were vomiting, facial edema and rash, and at 1–2 weeks after infusion, transient neutropenia and pneumonia developed in one patient.⁷⁴ This suggested that adding rituximab to steroids can be effective in refractory cases, mainly as a second-line treatment, and emphasized the fact that repeated doses are efficient in relapsing patients. The effects of rituximab in 14 patients with immune cytopenias including 4 ES cases were reported. No patient had a concomitant response in both AIHA and ITP, suggesting that rituximab can be effective only in some patients with refractory ES.⁹⁷ There are ten case reports of single patients treated with rituximab, eight adults and two children; nine of these ten patients achieved remission.^{34,98–106} These studies suggest that rituximab is a good option in refractory patients, and that repeated courses are necessary to achieve long-term responses. We have previously reported the outcome of six patients with ES treated at our institution, four women and two men with an age range of 10–42 years. Three received steroids as the first-line treatment and low doses of rituximab, 100 mg/week/4 weeks, as the second-line treatment with a good clinical response; after second relapse, however, two patients required splenectomy.¹⁰

Mycophenolate mofetil

The 2-morpholinoethyl ester of mycophenolic acid is a selective, competitive immunosuppressant prodrug that reduces lymphocyte proliferation by inhibition of inosine monophosphate dehydrogenase.¹⁰⁷ Some authors have shown its efficacy in autoimmune cytopenias.^{12,107} A recent retrospective multicenter study evaluated the use of MMF in 40 and 16 children with ITP and ES, respectively. Most patients previously received one to three lines of treatment; 13/16 cases achieved a complete response and only 1 patient had nausea as a secondary effect.¹⁰⁸ The excellent response in that study encourages further clinical trials as it proved that MMF is safe and effective for ES, confirming its administration as a second–third line therapy.^{11,109} Effectiveness of MMF has been associated with a sustained complete remission as observed in two children; the first case was refractory to steroids and IVIG¹¹⁰ and the second to steroids, IVIG, rituximab,

vincristine and cyclophosphamide.¹¹¹ Both patients received MMF at 50 mg/kg/day, while the second patient also received cyclosporine A.

The use of MMF has been related to a higher risk of developing lymphoma. Nevertheless, in a prospective study of 200 patients, main adverse effects were epigastric pain and headache, with no manifestations suggestive of lymphoma.¹¹² A dose of 600 mg/m² of MMF twice a day followed by an evaluation 3 months after the initial treatment is recommended because of the risk of relapse and scarce information on the long-term adverse effects.¹⁰⁹ These results suggest that MMF can be a useful and safe second-line treatment for ES patients refractory to steroids or IVIG.

Cyclosporine

Cyclosporine has been used in small series of patients with ES since 1994. It is an immunosuppressive drug whose mechanism of action is that it inhibits the activation of T-cells.¹¹³ A total of 28 patients have been treated with this modality in different studies.^{114–120} Most were refractory to steroids, IVIG or another immunosuppressant. A response was observed in 26/28 refractory patients; thus, cyclosporine can achieve an excellent response in these circumstances. Nevertheless, it is important to carefully select patients considered for this drug in cases with no response to less-toxic drugs. If administered, continuous monitoring of serum concentrations of cyclosporine is mandatory due to the severe adverse effects associated with its administration, including malignancy and nephrotoxicity, which have been observed after administration of low doses of cyclosporine (6 mg/kg/day) after 20 years of treatment.¹¹⁵

Splenectomy

Splenectomy was the only option in refractory relapsing patients before the introduction of rituximab, being classically considered as a second-line treatment in patients with autoimmune cytopenias, achieving a response of 60%–75% in AIHA and ITP,¹⁰¹ while in ES, the responses are documented to be considerably heterogeneous, between 0% and 66%.^{3,6,121,122}

Recently, splenectomy has been replaced by rituximab due to the risks of the surgical procedure. It is important to mention that in children <6 years of age, it is not a recommended option because of a higher risk of septicemia/meningitis with *S. pneumoniae*, *Neisseria meningitidis* or *Haemophilus influenzae* with both classic and laparoscopic techniques.^{11,12,123} Also, splenectomy is contraindicated in patients with ALPS due to an elevated risk of sepsis and recurrent multilineage cytopenias.¹²⁴

Reported complications after splenectomy in patients with hematologic diseases include bleeding, infections and thromboembolic events.¹²⁵ Recently, a case report with fatal pulmonary embolism as a consequence of splenectomy in a patient with ES was published.¹²⁶ Laparoscopic splenectomy is an option that could avoid some secondary adverse effects, as observed in five patients with ES who underwent laparoscopic splenectomy. Two cases achieved normalization of the platelet count at a mean of 18 months and did not require additional treatment. One patient had a good initial response, but required additional medical treatment after 18 months, and two cases did not respond.¹²¹

Third-line treatments

Cyclophosphamide

Cyclophosphamide is a powerful alkylating agent with anti-mitotic and immunomodulatory effects, is used in cancer and autoimmune diseases.¹²⁷ It is able to suppress an ongoing immune response. Although few centers have analyzed the use of cyclophosphamide in ES, a slow but progressive response has been described.¹²⁸ In a report, 48 patients with autoimmune cytopenias associated with CLL were evaluated. Eight had ES and were treated concomitantly with rituximab, dexamethasone and cyclophosphamide, achieving an overall response of 75%. The duration of response of CLL patients was shorter in cases of ES compared to AIHA or pure red cell aplasia.¹²⁹ In another center, cyclophosphamide was administered in a refractory child at a dose of 500 mg/m² intravenously (IV), achieving a good response with improvement of hemoglobin and platelet levels.¹³⁰ It is known that the use of this drug is associated with a higher frequency of adverse effects, including infections and secondary malignancies.¹²⁸ Current results suggest that cyclophosphamide can induce remission in ES patients and it could be administered as a third–fourth-line option in refractory cases.

Alemtuzumab

A humanized monoclonal antibody against CD52 has recently been tested for AIHA and ITP and was found to show a good response.^{131,132} This antibody exerts its action through depleting lymphocytes; recently, we studied this therapeutic option in refractory ITP and AIHA at 10 mg on days 1 and 3 administered subcutaneously in combination with subcutaneous rituximab at 100 mg/week/month, and the response was 100%.¹³³ A prospective cohort of 21 patients with autoimmune cytopenias received alemtuzumab IV at 20 mg/day for 10 days; 3 patients had ES, 2 responded, whereas the third had a transient response. The first patient achieved

a response within 1 week, but relapsed at 3 months and received cyclosporine and a second course of alemtuzumab. The second patient responded 3 days after administration of treatment, but relapsed at 3 months and received a second course of alemtuzumab; however, this patient died due to a cerebral hemorrhage. The last patient achieved a transient response and died because of lung cancer.¹³⁴

Thrombopoietin-receptor agonists

Second generation of thrombopoietin receptor agonists, romiplostim and eltrombopag, had a potential effect stimulating thrombopoiesis as observed in ITP,¹³⁵ and recently in ES, as documented in a patient refractory to steroids, splenectomy, IVIG and rituximab who received eltrombopag, romiplostim plus steroids with a duration of remission of 40 days at the last follow-up, suggesting that the combination of drugs is effective in decreasing platelet destruction and increasing the proliferation and maturation of megakaryocytic cells, as observed with steroids and thrombopoietin receptor agonists, respectively; this could be a highly effective double-targeted therapeutic approach.¹³⁶ Another center reported a patient refractory to rituximab who received romiplostim at 1–2 µg/kg/week/4 doses and after splenectomy was performed, with a remission duration of 153 days since the first romiplostim administration.¹³⁷

Hematopoietic stem cell transplantation

Scarce reports have documented the use of stem cell transplantation in ES. The main benefit of allogeneic HSCT is that it currently constitutes the only curative treatment option for ES by resetting the immune system,¹³⁸ thus, this therapy could be a valuable alternative for patients with multiple relapses, with severe complications affecting their quality of life and/or if a lack of response to immunosuppressive drugs exists.

Allogeneic HSCT

In 1997 the first case report of a child with ES treated with ten different therapeutic immunosuppressive lines without a good response was published. The patient suffered severe bleeding (intracranial, gastrointestinal) and considerable neurological impairment. Due to his unresponsive disease, an allogeneic cord blood transplant was performed with a conditioning regimen of 225 cGy daily for 6 days plus IV cyclophosphamide, 60 mg/kg for 2 days. This patient achieved remission; nevertheless, at day 286 posttransplant, the patient developed streptococcal pharyngitis and, subsequently, liver failure, coma and died.¹³⁹ An adult with refractory ES treated with multiple interventions received an allogeneic HSCT with a

conditioning regimen of cyclophosphamide, 200 mg/kg plus anti-thymocyte globulin at 90 mg/kg. Three months after transplant, antibodies against erythrocytes and platelets were negative with 100% chimerization. He maintained complete remission for the 30 months of follow-up.¹⁴⁰

Autologous HSCT

A prospective non-randomized study analyzed the use of cyclophosphamide at 50 mg/kg/day with autologous depleted peripheral blood stem cells for ITP in five cases with ES. They received 6–14 lines of treatment prior to transplant. Three achieved complete response and two did not respond. All cases presented bacteremia and one died after developing multiple myeloma.¹⁴¹

In another case, a female adult received an autologous HSCT. She suffered SLE plus antiphospholipid syndrome and suddenly developed florid ES. The conditioning regimen was cyclophosphamide at 50 mg/kg/day, anti-T lymphocyte globulin 10 mg/kg/day and 6-methylprednisone 1 g/day. The patient achieved a good response with normal blood counts.⁶⁴

A child with refractory ES received a double autologous stem cell transplant and achieved a short-term response. On relapse, unrelated umbilical cord blood transplantation was performed with a conditioning regimen of busulfan, thiotepa, etoposide and antithymocyte globulin. After cord blood transplant, the patient achieved a good response.¹⁴²

The largest retrospective study of autoimmune cytopenias treated with HSCT included 36 patients, wherein 7 had ES. Five received an allogeneic HSCT and one achieved remission, two died and two were not evaluated. This suggests that auto-HSCT can be an alternative in severe autoimmune cytopenias.¹⁴³

Graft vs host disease has an important effect in eliminating auto-antibodies against platelets and erythrocytes, with a higher rate and prolonged remission in secondary cases of ES. The high risk of morbidity has restricted the use of HSCT for treating ES and other autoimmune diseases. Thus, results suggest that it is important to start therapy with medical treatments. If there is no response or the patient has a chronically relapsing course and if a human leukocyte antigen-identical sibling donor is available, allogeneic HSCT is preferred. In older patients with significant comorbidities who lack a human leukocyte antigen-identical donor, autologous HSCT could be performed.¹⁴⁴

ES secondary to solid organ or HSCT

Solid organ and bone marrow transplantation has been associated to the development of ES in some centers.^{145–151} The

main mechanism behind this association is still unknown; immunosuppression secondary to chemotherapy of maintenance regimen could be implicated to an immune dysregulation of T-cells.¹⁴⁵ In 19 infants with genetic diseases, a mismatched cord blood transplantation was performed with a conditioning regimen of busulfan, cyclophosphamide and antithymocyte globulin; 10 cases developed chronic graft vs host disease with autoimmune cytopenias and 3 had ES.¹⁴⁶ A child with recurrent Hodgkin lymphoma underwent an autologous blood stem cell transplantation, after 15 months the patient developed ES refractory to steroids and IVIG; cyclosporine A was effective in this case.¹⁴⁷ A female patient with recurrent Hodgkin disease treated with an autologous bone marrow transplant with a conditioning regimen of cyclophosphamide, etoposide and carmustine developed ES.¹⁴⁸ A patient with non-Hodgkin lymphoma with multiple relapses received a peripheral blood stem cell transplant with a conditioning regimen of etoposide, carboplatin and cyclophosphamide; 49 days after transplantation the patient developed ES, non-Hodgkin lymphoma relapsed and died.¹⁴⁹ Also, development of refractory ES in a patient with multiple myeloma treated with an autologous stem cell transplantation has been documented.¹⁵⁰

Additional potential supporting interventions

Anecdotal reports suggest that immunological benefits can be gained in patients with autoimmune diseases including SLE and rheumatoid arthritis from simple measures, including oral honey drink and *Nigella sativa* oil¹⁵² or powder as nutritional support measures. It is reported that wet cupping therapy (Al-Hijamah) can clear patients' blood from autoantibodies, inflammatory mediators and immunological noxious substances.^{153,154}

Perspectives in ES

New treatment approaches for patients suffering from ES should lead to long-lasting remissions, less-severe relapses and an increased opportunity for cure. These scenarios will be possible as knowledge of the molecular landscape in this intriguing disease deepens, thus allowing the design of sophisticated biological therapy and/or improved HSCT strategies for severe cases, particularly in children.

A Phase II active clinical trial (Identifier: NTC00392951)¹⁵⁵ recently evaluated sirolimus at 3 mg/m² and then at 2.5 mg/m² depending on its serum levels in refractory autoimmune cytopenias including ES. A total of 30 patients were evaluated, including 8 with ES. There were three complete responses, three partial responses and two patients with no

response. Main objectives were to document toxicities of sirolimus, including mucositis, elevation of triglycerides and cholesterol and to assess the efficacy of this approach in patients with refractory ES.¹⁵⁶

Conclusion

ES is a rare disease with a heterogeneous course characterized by multiple relapses during lifetime, despite multimodal treatment. A combination of genetic and epigenetic factors appears to be involved in its pathogenesis, and the knowledge of its precise etiology will help to design specific targeted therapies with less-severe adverse effects, significantly improving patients quality of life. Results with currently available therapeutic modalities are unsatisfactory and the disease runs a chronic relapsing course, eventually complicated by catastrophic clinical events that can lead to death. Prospective studies and clinical trials derived from knowledge on its pathophysiology are needed to improve the long-term response and even cure individuals suffering ES.

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Disclosure

The authors report no conflicts of interest in this work.

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