Primary vasculitides that affect children are a challenging and complex group of disorders that may involve any system of the body and lead to significant morbidity and mortality. In recent years, there have been significant advances in the field of childhood vasculitides, including the development of classification criteria and outcome assessment. Although some forms of vasculitis occur in both children and adults, considerable differences exist between childhood and adult vasculitides; we review childhood vasculitides, thus highlighting their differences with the adult forms of the disease. We will also discuss monogenic forms of vasculitis, such as deficiency of adenosine deaminase type 2 (DADA2) and haploinsufficiency of A20 (HA20).

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Introduction

Primary vasculitides are a rare group of disorders characterized by inflammation of the blood vessels. The clinical features depend on the size of the vessel involved. Although most forms of primary vasculitis are more common in adults than in children, some types of vasculitis such as Henoch-Schönlein purpura (HSP)/Immunoglobulin A vasculitis (IgAV) and Kawasaki disease (KD) tend to occur predominantly in childhood. In this review, we aim to summarize pediatric vasculitides, with emphasis on the main differences and similarities in clinical features and classification between children and adults.
Epidemiological and clinical differences of vasculitides in children

Pediatric vasculitides account for 1%–6% of pediatric diseases [1,2]. A study from the United Kingdom (UK) reported the estimated annual incidence of primary vasculitis to be approximately 23 per 100,000 children [3]. The distribution varies in different ethnic groups [1,2,4]. For example, in the authors’ experience, KD and Behçet’s disease (BD) are more common in Asian and Turkish children, respectively, although we lack strong precise epidemiological data. In adults, the most common primary vasculitis is giant cell arteritis (GCA), whereas GCA is never seen in childhood [5,6]. Takayasu arteritis (TA) is a large-vessel vasculitis that affects the aorta and/or its major branches. It is rare but may occur even in infancy. The median age of disease onset in children is approximately 11 years [7]. It is more prevalent among Japanese and Asian children [8]. Among pediatric patients with TA, the most common findings are arterial hypertension (82%), headache (31%), fever (29%), dyspnea (23%), and weight loss (22%) [9–11]. Musculoskeletal involvement is more common, and ocular findings are less common in children than in adults [9–11], whereas bruits and claudication are more common in adults than in children [12].

In national registries, IgA-V (HSP) and KD are the most common forms of childhood vasculitis. In most parts of the world, IgA-V (HSP) is the most common type of pediatric vasculitis with an annual incidence of 13–20/100,000 [3,13]. Recently, a French population-based study reported an annual incidence of IgA-V (HSP) of 18.6/100,000 in children [14]. However, in adults, IgA-V (HSP) is less common, with an annual incidence of 0.1–1.8/100,000 [15]. IgA-V (HSP) occurs most commonly between the age of 5 and 15 years in children [15]. Upper respiratory infection may be a predisposing factor in pediatric and adult patients with IgA-V (HSP), whereas malignancy and drug exposure emerge as predisposing factors almost exclusively in adults [16]. There are important clinical differences between pediatric and adult patients. Adult patients tend to have a more severe course of disease, with more significant renal involvement: 8–68% of adult IgA-V (HSP) has been reported to be complicated by chronic renal failure, whereas it occurs in only 1.8–15% of pediatric patients with IgA-V (HSP) [17–19]. Pediatric patients more commonly present with abdominal pain before the appearance of purpuric rash than adult patients with the disease, whereas diarrhea is more common in adult patients with IgA-V (HSP) with abdominal involvement than in pediatric patients [17,20]. However, there are no prominent differences regarding the distribution of purpura, joint involvement, or laboratory findings [21].

Antineutrophil cytoplasmic antibody (ANCA)-associated vasculitides (AAV) are a group of vasculitides consisting of granulomatosis with polyangiitis (GPA) (formerly known as Wegener’s granulomatosis), eosinophilic GPA (EGPA) (previously known as Churg–Strauss syndrome, CSS), and microscopic polyangiitis (MPA) [22]. AAV are rare in childhood, with an incidence of 2.4 cases per million per year; GPA is the most common form [3]. GPA occurs around adolescence, with a median disease onset of 11.6 years and an estimated incidence of 0.1 per 100,000 [23]. In adults, GPA has a peak incidence in the sixth decade and is associated with a slight male predominance. The reported frequency of GPA has increased during the past few decades in adults. The annual incidence of GPA is 2.1–15 cases per million in Europe [24]. AAV are associated with significant morbidity, with a higher frequency of relapse, accrue damage in childhood [25]. Pediatric patients with AAV have more fever, more subglottic stenosis, more ischemic abdominal pain, more nasal cartilage damage, and less myalgia and peripheral neuropathy than adult patients with AAV [25]. EGPA is a rare necrotizing vasculitis. In pediatric patients with EGPA, cardiopulmonary involvement such as cardiomyopathy (42% in children versus 24% in adults) and pulmonary infiltrates (88% in children versus 59% in adults) are more common, whereas peripheral neuropathy is less common in these patients than in adult patients with EGPA (39% compared with 69% adult patients) [26]. Approximately 69% of pediatric patients with GPA are anti-PR3 ANCA (c-ANCA) positive, and approximately 21% of them are antmyeloperoxidase (MPO) ANCA (p-ANCA) positive [27]. Almost all the patients with GPA are ANCA positive in the active phase of the disease [28]. The presence of MPO-ANCA in pediatric and adult patients with EGPA is 25% and 38%, respectively [26].

Polyarteritis nodosa (PAN) is a primary systemic necrotizing vasculitis that affects predominantly medium-sized vessels [29]. The actual incidence of the disease is unknown in childhood, but it is estimated to be the third most common vasculitis among children [4]. In adults, it has an annual
incidence of 2-9 cases per million, with a peak age of 25 and 50 years and male predominance [3]. The disease onset age is around 9-10 years in children without gender predominance. When pediatric and adult patients with PAN are compared, weight loss and renal and neurologic involvement are more common in adults than in children, whereas arthralgia/arthritis and skin involvement are more common in children than in adults [30]. Childhood PAN tends to have a more benign course (with less renal and neurologic involvement) than adult-onset PAN [30].

Although KD is a predominantly childhood vasculitis, some very rare case reports of adult patients with KD have been published. It is a self-limiting vasculitis; however, the risk of coronary artery disease significantly increases in untreated cases [31]. It usually affects children under the age of 5 years. It has a higher prevalence among Japanese children with an annual incidence of 360 cases per 100,000, whereas the incidence in the UK is 8.1 cases per 100,000 cases per year [3]. A patient may be classified as having KD when she/he has fever lasting for more than 5 days (which cannot be explained by other causes) together with at least four of the following clinical findings: (1) bilateral conjunctival injection, (2) changes in oropharyngeal mucous (fissured lips, strawberry tongue, or injected pharynx), (3) changes in peripheral extremities (erythema, edema of hands and feet, or desquamation), (4) polymorphous rash, and (5) cervical lymphadenopathy [26]. Of note, atypical features such as arthritis, aseptic meningitis, pneumonitis, urethritis, diarrhea, hydrops of gallbladder, and uveitis may occur. Infants tend to have an atypical course with fewer classical signs. Some laboratory findings such as anemia, decreased albumin levels, elevation of alanine aminotransferase, and increased platelet levels after the seventh day may help to detect incomplete cases. Echocardiography should be performed in all patients with three or more suspected findings even if there are not four classical clinical criteria [31].

BD and Cogan syndrome are defined as variable vessel vasculitis [22]. BD is the only variable vessel vasculitis that occurs during childhood. BD manifests with recurrent oral and/or genital aphthous ulcers; additionally, cutaneous, ocular, articular, gastrointestinal, and/or central nervous system inflammatory lesions may occur [22]. Children account for 5.4%-7.6% of all patients with BD [32,33]. The disease has high prevalence along the Silk Road, from Japan to the eastern edge of the Mediterranean Sea, including areas of the former Ottoman Empire. The prevalence of BD along the Silk Road is 77-100 cases per 100,000 cases compared to only 0.1-15.9 cases per 100,000 in Western Europe [34-38]. There are differences in BD between pediatric and adult patients. Family history as well as neurologic and gastrointestinal involvement are more common in pediatric patients than in adult patients [39]. Uveitis is more common and more severe in boys than in girls [40-44], and genital ulcers are more common in girls than in boys [41-45].

**How to define and classify pediatric vasculitides**

Vasculitides are defined according to the predominant size of the involved vessels. With the improvement of knowledge of the etiopathogenesis, the CHCC nomenclature system for defining the different forms of vasculitis was updated in 2012 [22]. The eponyms used in vasculitides were replaced by descriptive names, whenever possible [22]. Thus, Wegener’s granulomatosis, CSS, and HSP were renamed as GPA, EGPA, and IgAV, respectively. In addition, new subcategories including single-organ vasculitis (which includes isolated central nervous system vasculitis), vasculitis associated with systemic disease (such as systemic lupus erythematosus), and vasculitis associated with a probable etiology (such as Hepatitis B virus-associated vasculitis) were added to the definition of vasculitides. Furthermore, CHCC 2012 has divided small-vessel vasculitides into two groups depending on the pathogenesis as ANCA-associated vasculitides and those associated with the presence of immune complexes.

Owing to the lack of any pathognomonic tests to diagnose these rare diseases, classification criteria are used to define homogeneous groups of vasculitis for research purposes. In 1990, the American College of Rheumatology (ACR) created a set of classification criteria for adult patients (Table 1) [46,47]. The Ankara 2008 Consensus Conference criteria were endorsed by European League Against Rheumatism, Pediatric Rheumatology International Trials Organization, and Pediatric Rheumatology European Society (EULAR/PRINTO/PRES). These organizations have developed new sets of criteria for childhood vasculitides, including criteria for childhood HSP/IgAV, PAN, GPA, and TA (Table 1) [48,49]. When the current update of the ACR/EULAR criteria for adults with vasculitis is completed (from the
Table 1
Comparison of the American College of Rheumatology (ACR) 1990 [47] and Ankara 2008 (EULAR/PRES/PRINTO) classification criteria [48,49] *(adapted from Refs. [47–49]).*

<table>
<thead>
<tr>
<th>Henoch–Schönlein purpura (HSP)/Immunoglobulin A vasculitis</th>
<th>ACR criteria</th>
<th>Ankara 2008 criteria</th>
<th>Main differences between pediatric-specific criteria and adult criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>≥2 of the following:</td>
<td>Purpura or petechia (mandatory) with lower limb predominance plus one of the following:</td>
<td>Palpable purpura became a mandatory criterion. Histopathologic findings were specified as predominance of IgA deposit Arthritis/arthralgia and renal involvement were included.</td>
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<tr>
<td>● ≤20 years of age at disease onset</td>
<td>● Abdominal pain</td>
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<tr>
<td>● Palpable purpura</td>
<td>● Histopathology (predominant IgA deposit in a biopsy)</td>
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<tr>
<td>● Acute abdominal pain</td>
<td>● Arthritis or arthralgia</td>
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<tr>
<td>● Biopsy showing granulocytes in the wall of small arterioles/venules</td>
<td>● Renal involvement</td>
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<tr>
<td>Granulomatous polyangiitis</td>
<td>≥2 of the following:</td>
<td>ANCA-positivity was included</td>
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<td></td>
<td>At least three of the following:</td>
<td>Computerized tomography findings were included to define pulmonary abnormalities, and more definitive items were added for upper and lower airway involvement</td>
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<tr>
<td></td>
<td>● Histopathology (granulomatous inflammation)</td>
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<td></td>
<td>● Upper airway involvement</td>
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<td></td>
<td>● Laryngo-tracheo-bronchial stenosis</td>
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<td></td>
<td>● Pulmonary involvement (chest X-ray or CT showing the presence of nodules, cavities, or fixed infiltrates)</td>
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<tr>
<td></td>
<td>● ANCA positivity</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>● Renal involvement</td>
<td></td>
<td></td>
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<tr>
<td>Polyarteritis nodosa</td>
<td>≥3 of the following 10 criteria:</td>
<td>The presence of histopathological or angiographic abnormalities became a mandatory criterion. The presence of testicular pain/tendency, weight loss, and evidence of hepatitis B infection were removed.</td>
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<tr>
<td></td>
<td>● Granulocyte or mixed leukocyte infiltrate in an arterial wall on biopsy</td>
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<tr>
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<td>● Arteriographic abnormalities</td>
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<td></td>
<td>● Livedo reticularis</td>
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<td></td>
<td>● Myalgia</td>
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<td></td>
<td>● Diastolic blood pressure &gt;90 mmHg</td>
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<td></td>
<td>● Mono- or polyneuropathy</td>
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<td></td>
<td>● Elevated blood urea nitrogen or creatinine</td>
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<td></td>
<td>● Testicular pain/tenderness</td>
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<td></td>
<td>● Hepatitis B reactants</td>
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<td></td>
<td>● Weight loss &gt;4 kg</td>
<td></td>
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<tr>
<td>Takayasu arteritis</td>
<td>≥3 of the following:</td>
<td>Angiography (conventional, CT, or MRI) of the aorta or its major branches and pulmonary arteries showing aneurysm/dilatation, narrowing, occlusion, or thickened arterial wall (mandatory) plus one of the following:</td>
<td>Angiographic findings became a mandatory criterion</td>
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<td></td>
<td>● Arteriographic evidence of narrowing or occlusion of the entire aorta, its primary branches, or large arteries in the proximal, upper, or lower extremities</td>
<td>● Pulse deficit or claudication</td>
<td>Pulse deficit and claudication became a single criterion</td>
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<td></td>
<td>● Decreased brachial artery pulse</td>
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<td>● Claudication of an extremity &gt;10 mmHg difference in systolic blood pressure between arms</td>
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<tr>
<td></td>
<td>● A bruit over subclavian arteries or the aorta</td>
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<td></td>
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<td></td>
<td>● Age at disease onset ≤40 years</td>
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</table>

* The table only includes pediatric vasculitides for which validated criteria have been developed. The table excludes giant cell arteritis, Kawasaki disease, and microscopic polyangiitis.
Diagnostic and Classification of Vasculitis Study, DCVAS), some criteria introduced for adults may need to be adapted for use in children.

In pediatric practice, the diagnosis of IgA-V (HSP) is a clinical one; biopsies are rarely performed. However, the presence of IgA in the biopsy sample is specific for the disease. Thus, the presence of predominant IgA deposition in biopsies of affected children is one of the new pediatric criteria for IgA-V (48). The revised pediatric criteria require the presence of palpable purpura (predominantly seen on the lower limbs) together with at least one of the following: abdominal pain, IgA deposition on histology, arthritis/arthralgia, or renal involvement [48]. This criteria set had a higher sensitivity (100% vs. 87.1%) and a similar specificity (87% vs. 87.7%) than the ACR criteria when applied to children [48].

Recently, a group of adult rheumatologists have evaluated the performance of Ankara 2008 pediatric criteria in adult patients with IgA-V (HSP) and found a higher sensitivity and specificity (99.2% vs. 86.8% and 86.0% vs. 81%) than the original ACR criteria for HSP [50].

The most significant changes in the Ankara 2008 criteria for childhood PAN were as follows: (1) histopathology or angiographic abnormalities became a mandatory criterion and (2) some criteria such as the presence of testicular pain/tenderness, weight loss, and hepatitis B reactants were removed because of their low specificity [48].

For GPA, the major differences between the pediatric Ankara 2008 criteria and the 1990 ACR adult criteria were the inclusion of ANCA positivity, the addition of computerized tomography abnormalities to pulmonary findings, and more definitive items for upper and lower airway involvement. The sensitivity and specificity of the Ankara 2008 criteria for childhood GPA were 93.3% and 99.2%, respectively [48]. For TA, the most marked changes in Ankara 2008 criteria were as follows: (1) angiography findings became a mandatory criterion, (2) age limitation was removed, (3) hypertension and elevated acute phase reactants were included, and (4) pulse deficit and claudication became a single criterion. With these changes, the sensitivity of Ankara 2008 criteria for childhood TA increased to 100% and the specificity to 99.9% [48].

KD classification criteria were not validated in the Ankara 2008 Consensus Conference. Patients are classified as KD according to American Heart Association (AHA) criteria as follows: unexplained fever lasting for more than 5 days together with at least four of the following clinical findings: (1) bilateral conjunctival injection, (2) changes in oropharyngeal mucous (fissured lips, strawberry tongue, or injected pharynx), (3) changes in peripheral extremities (erythema, edema of the hands and feet, or desquamation), (4) polymorphous rash, and (5) cervical lymphadenopathy [31].

An international expert consensus group (the pediatric BD [PEDBD] group) has suggested a new set of criteria to classify pediatric BD [51]. A child can be classified as having BD when presenting with three or more of the following criteria: (1) oral aphthosis (≥3 attacks per year), (2) genital aphthosis (typically with scars), (3) skin involvement (necrotic folliculitis, acniform lesions, or erythema nodosum), (4) neurologic involvement (except isolated headaches), (5) ocular manifestations (anterior uveitis, posterior uveitis, or retinal vasculitis), and (6) vascular features (venous thrombosis, arterial thrombosis, or arterial aneurysms) [51]. Recently, two centers from Turkey and Israel evaluated the performance of the new pediatric criteria and found them to have a sensitivity of 73.5% and a specificity of 98.9% for childhood BD [52].

Outcome and management of pediatric vasculitides

Disease activity and damage are the most important domains to determine the disease outcome. PRES Pediatric Vasculitis Working Party has validated an activity score, called the Pediatric Vasculitis Activity Score (PVAS) [53]. This score has been modified from the activity score for adults (Birmingham Vasculitis Activity Score, BVAS) and comprises the sum of the new or worsening clinical characteristics in nine organs/systems [53]. To date, the Vasculitis Damage Index (VDI) has not been validated in children, although a preliminary version has been produced and tested [54]. Some items such as puberty delay and growth retardation of children have been included in the preliminary pediatric version of the VDI [54].

Apart from the epidemiological, clinical, and classification differences, the outcomes in children are also different from those in adults. IgA-V (HSP) is usually a self-limiting form of vasculitis. However, gastrointestinal involvement during the early phase of the disease and renal involvement during the
later stage are risk factors for a worse long-term outcome. Furthermore, older children have a more severe course with worse renal outcomes and more relapses than younger children [55,56]. Adults with IgA-V (HSP) have worse renal outcomes than children with the disease [17–19].

With regard to the other vasculitides, Belostotsky et al. [57] reported that the mortality rate of pediatric patients with GPA was 12% during a 17 year period. Zwerina et al. [26] demonstrated that EGPA has a higher mortality rate in children than in adults (19% in children versus 5% in adults). Childhood PAN has a benign course, with mortality rates of 1–4% in children [29,58] compared to those of 24.9% in adults [59]. Eleftheriou et al. [7] found that the mortality rate for pediatric TA was 27% in the UK.

In untreated cases of KD, coronary involvement is present in up to 25% of cases. Recently, Lin et al. [60] demonstrated that the severity of coronary artery involvement during the first month of disease was the most important predictor for late coronary outcomes. However, in patients who were given appropriate treatment within the first 10 days of fever, the incidence of coronary artery lesions decreased from approximately 25%–5%.

Long-term morbidities such as Cushing’s syndrome and osteopenia, due to glucocorticoid therapy, are common in pediatric BD [45]. The data regarding long-term outcomes of childhood BD are limited, and the outcome mainly depends on organ involvement.

**Treatment approaches**

Pediatric treatment approaches mainly depend on the recommendations published for adult patients, together with small case series and personal experiences of managing pediatric cases. The SHARE initiative (Single-Hub Access for pediatric Rheumatology in Europe) [61] has been developing evidence-based recommendations for the treatment of pediatric vasculitides (manuscript in preparation). Treatment of IgA-V (HSP) is mainly conservative; studies show that early steroid treatment does not prevent renal involvement [62]. However, in the presence of severe gastrointestinal or skin involvement, systemic glucocorticoids may be required [33]. We hope that the SHARE initiative will provide clinicians with a more standardized approach in IgA-V (HSP) treatment in the future.

Management of AAV in childhood mainly depends on clinical trials conducted in adults [68]. The recommendations endorsed by EULAR have been reviewed elsewhere.

KD is one of the most common vasculitides occurring during childhood. Recently, the AHA updated its guideline for the management and follow-up of KD [26]. It is currently recommended that the measurement of coronary artery size should be made and compared with Z scores to provide a more quantitative assessment [31]. Patients should be treated with a single dose of 2 g/kg intravenous immunoglobulin (IVIG) together with acetylsalicylic acid (ASA). However, the standard dose of ASA at initiation is still controversial [31]. Patients can be classified as IVIG resistant, in the presence of persistent or recrudescent fever for at least 36 h after the end of the first IVIG infusion [31]. Patients with IVIG resistance may be treated with a second dose of IVIG, high-dose pulse steroids, tumor necrosis factor (TNF) inhibitors, cyclosporine A, or anakinra [31]. Patients should be evaluated by a cardiologist at 4–6 weeks after discharge and during the first year. If there is no coronary involvement, routine cardiology follow-up is not indicated after the first year, but patients should be assessed for cardiovascular risk factors [31]. Risk scoring systems to predict IVIG resistance have been suggested [63–65]; however, these scores have not been as successful in patients of Western ethnicity [66,67]. The Kobayashi score [63–65] remains the most widely used scoring system. In the presence of high risk, patients should be treated with corticosteroids combined with IVIG and ASA [31]. Recently, a phase I/IIa trial in children with KD demonstrated the safety and tolerability of anakinra in KD [68].

Classic treatment of PAN usually follows practice in adults [69]. We still lack evidence-based data for the treatment of PAN in childhood. The ongoing “MYPAN trial” is a multicenter, open-label, randomized controlled trial conducted to compare the efficacy of cyclophosphamide with mycophenolate mofetil (a potentially less toxic therapy) for the induction of remission in pediatric forms of PAN.

EULAR recommendations for adult patients with TA have already been established [70], but we lack randomized controlled trials for the treatment of pediatric patients with TA. Promising results with tocilizumab have been published in pediatric patients with TA as a single-center experience [71].
A new group of vasculitides in childhood: the monogenic vasculitides

Deficiency of adenosine deaminase type 2

Recently, a new monogenic disorder, mimicking PAN, has been described: deficiency of adenosine deaminase 2 (DADA2) [72,73]. This is an autosomal recessive inherited disease that manifests with early-onset stroke and vasculopathy, elevated acute phase reactants, fever, peripheral neuropathy, mild immunodeficiency, and fluctuating low titers of antibodies [72,73]. Loss-of-function mutations in the cat eye syndrome chromosome region candidate 1 (CECR1) gene, which encodes adenosine deaminase 2 (ADA2), are associated with the disease [72,73]. Deficiency of ADA2 leads to endothelial damage and reduction in the anti-inflammatory macrophage population [72,73]. However, the pathogenesis of the disease remains unclear. The presence of early-onset stroke, parental consanguinity or familial cases, or unresponsiveness to conventional treatment should be seen as red flags in patients with presumed PAN, which should prompt a genetic analysis for mutations in CECR1. ADA2 enzyme activity is significantly reduced in affected individuals compared to that in healthy controls [74]. ADA2 enzyme replacement therapy may be a treatment option in the future; however, current treatment is usually with TNF inhibitors [72,75].

Autoinflammatory interferonopathies presenting with vasculopathy

STING-associated vasculitis of infancy (SAVI) is an autoinflammatory disease, caused by gain-of-function mutations in the TMEM173 gene [76]. SAVI manifests with severe vasculopathy and interstitial lung disease [76]. Patients present with vasculitic lesions such as violaceous plaques and/or nodules prominent in cold-sensitive acral areas. Gangrene and infarcts of fingers and toes leading to tissue loss are the most remarkable skin involvements of the disease [76]. Skin biopsies of patients show marked vascular inflammation, limited to capillaries without any sign of medium- or large-vessel vasculitis. In addition, microthrombotic vascular changes may be observed in some biopsy samples [76]. Janus kinase (JAK) inhibitors may be a promising therapeutic alternative. A Baricitinib (JAK1 and JAK2 inhibitor) trial is underway [77].

Haploinsufficiency of A20

Haploinsufficiency of A20 (HA20) is a newly described monogenic autoinflammatory syndrome that mimics BD [78]. This disease is caused by high-penetrance heterozygous mutations in TNF α-induced protein 3 (TNFAIP3/A20), which encodes the nuclear factor kappa-B (NF-κB) regulatory protein [78]. A20 suppresses NF-κB signals, and this subsequently leads to increased expression of inflammatory cytokines through its deubiquitinase activity, and a loss-of-function mutation in this gene leads to an increase in inflammatory processes, especially NF-κB signaling [78]. Sixteen patients have been described with early-onset recurrent oral, genital, and/or gastrointestinal ulcers. There is no standard treatment for the disease. Published cases were treated with TNF inhibitors, anti-IL1 receptor (anakinra), or anti-IL6 receptor (tocilizumab) [78–80].

Association of primary immune deficiencies and vasculitides in childhood

Systemic vasculitides related to immunodeficiency have already been reported, including Wiskott–Aldrich syndrome (WAS) and interleukin-12 receptor beta-1 (IL-12Rb1) deficiency [81,82]. In 2016, a new monogenic autosomal dominant immune dysregulatory disease, called COPA Syndrome, was described. Patients manifest with arthritis, renal disease, and interstitial lung disease with pulmonary hemorrhage, resembling AAV [83]. Recently, we described a patient with a hypomorphic mutation in the recombination-activating (RAG) 1 gene, presenting with a severe PAN-like disease [84]. This patient had pulmonary hemorrhage, digital necrosis, and finger amputation. The tissue biopsy of the patient was compatible with PAN. In patients with atypical presentations of vasculitis, genetic defects and immune deficiencies should be considered, especially if there is consanguinity.
Current initiatives in pediatric vasculitis

The pediatric field is improving the recognition and management of patients with vasculitis. International registries such as the PRINTO vasculitis registry have served as the basis for the childhood classification criteria and to define the characteristics of these diseases. Another international registry, ARCHIVE, has served for validation studies and for defining treatment strategies [85]. A number of collaborative studies are currently underway to improve our understanding of the diagnosis, biomarkers, and treatment [86]. Furthermore, through international collaborations, pediatricians have provided management recommendations for two childhood vasculitides: KD and IgA Vasculitis (HSP) (submitted work). Pediatricians have led the work in defining characteristics of some rare vasculitides of childhood, such as the pediatric central nervous system vasculitides, pediatric BD, and vasculitis mimics due to single-gene defects.

Conclusion

Vasculitides in children present distinct features. Collaboration among pediatricians has enabled better characterization of these diseases. Multicenter studies may lead us to classify, treat, and manage this challenging disease group; indeed, new pediatric initiatives are emerging as outlined above.

Summary

New data emerging in childhood vasculitides suggest that the vasculitic syndromes display differences in epidemiology and clinical aspects. These differences and the lack of validation of the adult criteria have led pediatricians to develop separate classification criteria for childhood vasculitides. The adult vasculitis activity score has been adapted for use in pediatric studies with the term “pediatric vasculitis activity score (PVAS).” Treatment strategies mainly depend on adult experience and recommendations because most of these diseases are very rare in childhood. Pediatricians have led the work on monogenic vasculitides because these new diseases often have a childhood onset. Among the new monogenic diseases, ADA2 deficiency and “STING associated vasculopathy of infancy” have been particularly important in our practice.

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Conflicts of interest

The authors declare no conflict of interest.

Practice points

- Systemic vasculitides are a group of disorders that occur during childhood characterized by inflammation of vessel walls, with an increased risk of morbidity and mortality. Thus, early diagnosis and multidisciplinary assessment are very important.
- There are differences in the forms of the disease between children and adults in terms of epidemiology, type of vasculitis, and clinical features.
- Monogenic forms of vasculitis should be considered in children with vasculitis of early onset or with an atypical course, especially if there is parental consanguinity.
Research agenda

- Further epidemiologic studies are needed to define the true incidence and prevalence of these rare diseases.
- We still lack randomized controlled trials in many types of childhood vasculitis.
- Multicenter studies and collaboration may guide us in improving the outcome in these patients.
- Understanding the pathogenesis of the new monogenic forms of vasculitis will not only help us in better management of these patients but also shed light on our understanding of the more common forms of vasculitis.

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