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Immunopathogenesis Based Adamantiades- Behcet Disease Vasculitis Treatment

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1. Introduction

All immune cell lineages arise from the multipotent stem cells in the bone marrow. The development and regulation of immune system cells (immune homeostasis) depends on the programmed appearance of specific cell surface molecules (e.g. receptors) and responsiveness to certain cytokines and chemokines. After maturation, the immune system cells include antigen presenting cells (APCs), other phagocytic cells like neutrophils, eosinophils, basophils and lymphocytes which include T lymphocytes, B lymphocytes and natural killer (NK) cells. Depending on an initial stimulus and the microenvironmental conditions at a given time, each lymphocyte lineage can be subdivided to different populations with discrete specialized functions. Subpopulations of crucial importance are CD4⁺ T cells (mainly Th1, Th2, Th17 and T regulatory cells) and CD8⁺ T cells, CD16⁻ and CD16⁺ NK cells and CD5⁺ and CD5⁻ B cells.

As a structure, the immune system consists of central locations of immune cell production and differentiation (bone marrow and thymus) and peripheral organs, where encounter with antigen and response to it occurs, that is the spleen, lymph nodes, tonsils and mucosa-associated lymphoid tissue (MALT). Mucosal surfaces and the skin provide the main primary access for foreign antigens to meet cells of the immune system, where they first react primarily with APCs.

Innate immunity includes skin and mucosal barriers, as well as the innate immune cells that are triggered and react in minutes after they encounter 'foreign' material. The innate immune system, as it represents the first line of host defense against foreign entities, is regarded as a nonspecific system and its main role is to engulf and eliminate pathogens, trigger pro-inflammatory responses and present antigens to adaptive immune cells, which are subsequently activated. More recently it has been shown that the innate immune system has a necessary degree of specificity that enables it to discriminate between self and foreign particles (e.g. microorganisms). Germline encoded and constitutively expressed *pattern recognition receptors (PRRs)*- like Toll receptors on innate immune cells, help them to discriminate highly conserved *pathogen associated molecular patterns (PAMPs)*, which activate specific signaling pathways that lead to robust but well-defined innate immune responses.

Research on innate immunity is progressively uncovering molecular and cellular mechanisms by which it recognizes environment and reacts to it, leading to protective or

pathological and harmful immune responses. Understanding innate and adaptive immunity pathways in detail will have a great impact on revealing the immunopathology of immune-mediated diseases like Adamantiades-Behcet Disease (ABD) and developing more targeted immunotherapeutic agents to manipulate, even life-threatening, ABD vasculitis.

2. General considerations in ABD immunopathogenesis

ABD owes its name to Benediktos Adamantiades (Greek ophthalmologist), who had described it first in 1930 and Hulusi Behcet (Turkish dermatologist), who described hypopyon, iritis and orogenital aphthosis in 1937 (Adamantiades, 1931; Behcet, 1937). However, the first description of the disease dates back, 2500 years, to Hippokrates, to the 5th century BC, in the Epidemion book, case 7, where it is described as aphthous ulceration, genital ulceration and iridocyclitis.

There were other forms of fever...
Many developed aphthae, ulcerations...
Many ulcerations about the genital parts...
Watery ophthalmies of a chronic character, with pains...

Since the early descriptions, additional clinical features have been described and added to disease spectrum. ABD can be manifested with different organ involvement, articular, neurological, pulmonary and gastrointestinal, besides the classical clinical triad proposed by the International Study Group for Behcet’s Disease.

Nowadays, ABD diagnosis rests upon a set of established criteria, where no single clinical or laboratory feature is pathognomonic (International Study Group for Behcet’s Disease, 1990, Table 1).

Recurrent oral ulcerations plus two	Recurrent genital ulcers Skin lesions (erythema nodosum, acne, folliculitis) Ocular lesions (uveitis) Positive pathergy test
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Table 1. Diagnostic Criteria for ABD

From early descriptions, it was shown that patients with ABD cluster according to a geographic pattern that coincides with the ancient “Silk Route”. It has a higher incidence (100 times more common) in the countries of the Mediterranean Basin, Middle East and Far East, which includes Greece, Turkey, Saudi Arabia, Iran, Korea ,China and Japan while it is rare in North America. Although sporadic cases are described, there is an increased frequency among relatives (familial aggregation), mainly in HLA-B51 class I gene positive relatives. Genetic susceptibility factors seem to be connected with more severe disease expression. Particularly, in Asia, up to 80% of patients are HLA-B51 positive. Recent studies focus on new genetic markers that could amplify immune responses responsible for disease expression (HLA and non-HLA genes), probably under an environmental challenge e.g. an infectious agent (the infectious model), (Fietta, 2005; Kaklamani et al, 1998; Verity et al, 2003; Yazici H et al, 1999).

ABD etiopathogenesis still remains a mystery to be elucidated. There is substantial evidence for an infectious background participating in disease initiation, especially, in genetically predisposed individuals. A number of viruses like HSV, Parvovirus B19 or bacteria like

uncommon serotypes of *Streptococcus* have been implicated. At the moment there is no concrete proof of the existence of certain microorganisms in the etiology of ABD and the initiation of the immunological process. Molecular mimicry between bacterial (HSV, streptococci) and human heat shock proteins (HSP), due to significant peptide homology between bacterial and human HSPs has been widely accepted, might be the basic link between infection and (auto)immunity in predisposed subjects, and could initiate and augment T and B cell immune responses. Besides, the option of exposure to chemicals was not supported for long. However, although immunological abnormalities concerning innate and adaptive immunity in ABD have been extensively studied, there is no adequate proof that ABD is either a typical autoimmune disease- it does not fulfill substantial criteria- or an auto-inflammatory disease- it is a very weak hypothesis at the moment (Direskeneli & Saruhan-Direskeneli, 2003; Kapsimali et al, 2010; Mumcu et al, 2007; Zouboulis & May, 2003).

ABD is a systemic vasculitis which may affect arteries and veins of all sizes in any organ and underlies disease manifestations. To a great extent, disease pathology points to a vascular inflammation. Biopsies of affected organs usually reveal vasculitis, where intense infiltrations of neutrophils predominate and CD4+ T lymphocytes are present in the early stages.

Venous vascular involvement and thrombosis is the characteristic feature of the disease, probably responsible for most of the tissue damage seen. Inflammatory changes in large arteries are typical of vasculitis, lack wall thickening and granuloma formation and have a tendency towards aneurysm formation. It is more severe in males. ABD lacks an increased risk of atherosclerosis. The term 'vasculo-Behcet' has been adopted to describe cases where vascular features dominate in disease expression. After initial vascular lesion (usually venous thrombosis), other vascular lesions may follow and lead to progressive, multifocal vascular disease with an unfavorable prognosis (Melikoglu et al, 2008).

Vascular inflammation is mainly due to complex innate-immune, pro-inflammatory mechanisms which activate and perpetuate adaptive/specific immune responses against antigens not identified clearly. Endothelial dysfunction, partly due to anti-endothelial autoimmune reactions and non-specific coagulation/fibrinolytic abnormalities, has also been demonstrated and participates in disease pathophysiology to some extent. In such cases aggressive treatment consisting of immunosuppressive drugs is recommended (Calamia et al, 2011; Calamia & Kaklamanis, 2008).

Management decisions are often difficult to take and there is a lack of controlled trials. A sequential therapeutic approach, based on ABD immunopathology and the immunomodulatory action achieved by immunosuppressive drugs is discussed.

2.1 Innate immunity cells abnormalities in ABD

Neutrophil involvement in ABD pathogenesis represents a challenging issue, as neutrophil hyperactivation seems to play a pivotal role in the inflammatory vasculitic characteristics of disease lesions. Neutrophils are the dominant subpopulation of the inflammatory infiltrates in the vessel wall (Hayasaki et al, 2004).

Neutrophil hyperactivity is a major contributor to the well-known pathergy skin reaction, which might be positive during periods of active disease. Pathergy reaction is a non-specific skin hyper-reactivity to minor trauma, such as a needle prick (Boura et al, 2007; Ozdemir et al, 2007).

At the tissue level, neutrophils were described to be present in the anterior chamber of the eye and the corneal epithelium, as well as in the ciliary body and chorioid tissue (Mendoza-Pinto et al, 2010).

These cells are reported to be functionally impaired in the disease. Eksioglu-Demiralp et al described specific defects concerning neutrophil activation, oxidative burst and phagocytic activity (Eksioglu-Demiralp et al, 2001). On the other hand, increased synthesis of reactive oxygen species (ROS) has been reported in ABD. ROS-mediated oxidative stress seems to play an important role in pathogenesis, as advanced oxidation protein products were described to correlate with disease activity (Yazici C et al, 2004). This contradiction might mean that functional activity of neutrophils might correlate to disease activity or other, poorly defined, parameters.

Additionally, it has been postulated that endothelial cells cooperate with neutrophils and monocytes in the vasculitic process. Endothelial cell activation and subsequently increased expression of cell adhesion molecules is a characteristic feature of the disease, which can trigger neutrophils and monocytes to initiate natural immune responses on the vessel wall. Recently, a novel mechanism by which neutrophils may interact with cytokine-activated endothelial cells in the presence of anti-endothelial cell antibodies has been described. Main receptors were Fc-gamma-RIIa and CXCR1/2. Moreover, endothelium activation leads to excessive thrombin formation and impaired fibrinolytic function, thus contributing to the generation of a prothrombotic state in ABD (Florey et al, 2007; Lee MT et al, 2007; Probst et al, 2004).

Furthermore, neutrophils are primed by overproduced pro-inflammatory cytokines, secreted by antigen presenting cells (APCs) and the endothelium (e.g. TNF- α , IL-1, IL-6, IL-8 etc). Under such conditions, they secrete other cytokines that prime themselves, cause Th1 polarization and maintain a non-specific, pro-inflammatory microenvironment to retain enhanced activity (Carletto et al, 1997; Katsantonis et al, 2000; Zouboulis et al, 2000).

Recently, the role of matrix metalloproteinases, particularly MMP-2 and MMP-9 was underlined in ABD. Pay et al found increased levels of these molecules in the serum of ABD patients and this was strongly correlated to vasculo-Behcet disease, especially aneurysm formation. Other investigators confirmed the elevated MMP-9 levels in the cerebrospinal fluid of ABD patients, suggesting that they might be involved in the pathogenesis of neuro-ABD (Hamzaoui K et al, 2009; Pay et al, 2007).

It is reported that neutrophils are more resistant to spontaneous apoptosis (via Fas/Fas ligand interactions) during the remission of ABD uveitis, while resistance was restored in the active phase of the disease (Fujimori et al, 2008).

Monocytes are also involved in ABD pathogenesis. Recent studies support the possible role of bacterially-induced innate immune responses through monocyte Toll-like receptor (TLR) activation. It was reported that TLR-2 was decreased after stimulation with LPS and HSP-60, or increased after bacterial lipoteichoic acid stimulation. TLR-6 expression was found to be enhanced after stimulation with HSP-60 and *Streptococcus sanguis* (Neves et al, 2009; Yavuz et al, 2008). As mentioned above, HSP-60 and *Streptococcus sanguis* represent candidate antigens in ABD pathogenesis.

Antigen-presenting cells are reported to be activated in ABD patients, since ex vivo cultured monocytes were able to produce increased amounts of TNF- α , IL-6 and IL-8 (Mege et al, 1993). Moreover, there is strong evidence suggesting that APCs may be primarily involved in ABD immunopathogenesis. Early data comes from pathergy reaction histology, where

Saito et al showed that Langerhans cells were increased in the dermis layer shortly after needle prick (Saito et al, 1980).

However, it has not been clarified whether APC hypereactivity is a primary defect of these cells or the result of disease related conditions. Nevertheless, IL-12 and IL-18, cytokines mostly produced by APCs, are known to strongly promote Th1 polarization of immune responses. Both these cytokines have been found in increased levels during active disease, implying a central role for APCs in ABD pathogenesis (Hamzaoui K et al, 2002).

Finally, Pay et al described decreased numbers of peripheral plasmacytoid dendritic cells, along with decreased levels of IFN- β and suggested that this cell subset may also participate in disease pathogenesis (Pay et al, 2007).

2.1.1 Gamma-delta ($\gamma\delta$) T cells

These specialized T cells (TCR comprising of γ and δ heterodimer) have been increasingly investigated during recent years. They are thought to play a critical role in mucosal immunity, being the first line of defense against invading micro-organisms, as well as the link between innate and adaptive immune response (Casetti et al, 2008).

A subset of $\gamma\delta$ T cells (Vgamma9+Vdelta2+) was found in increased numbers in the blood of active ABD patients. These cells were reported to express IL-2 receptor and HLA-DR surface molecules, indicating a high activation status. The Vgamma9+Vdelta2+ T cells were described to infiltrate target organs, such as the eye, while other studies found that they strongly proliferate after stimulation with bacterial extracts of the oral flora (Bank et al, 2003; Verjans et al, 2002; Yamashita et al, 1997).

The critical role of gammadelta T cells in ABD is underlined in the study of Treusch et al, who found a decrease in their numbers after treatment with interferon-alpha-2a (Treusch et al, 2004).

Later, it was shown that expanded CD56+ and CD8+ gammadelta T cells in ABD were capable of secreting IFN- γ and TNF- α . Data is further supported by the fact that these cells seem to be polyclonally rather than oligoclonally activated in ABD patients with oral ulcers. Probably because of repeated inflammation of the oral mucosa, due to a wide variety of antigenic stimuli, activation and proliferation of these cells might play a role in disease clinical heterogeneity (Freysdottir, 1999, 2006).

2.1.2 NK and NKT cells

Early studies reported an increase in NK cell numbers in ABD patients. However, others described a decrease in their cytotoxic activity in the active phase of the disease, which was restored in remission periods. However, recent studies reported that, although increased in active disease, NK cells do not seem to exert impaired cytotoxic activity, as they were shown to adequately suppress IFN- γ production by CD4+ T cells, thus regulating a Th1 response in ABD (Onder et al, 1994; Suzuki et al, 1992; Yamaguchi et al, 2010).

Natural killer T cells (NKT) comprise a sublineage of T cells that share characteristics of conventional T cells and NK cells and bridge innate and adaptive immunity. Their activation is restricted by the MHC molecule CD1d. Early studies reported an increase of CD4+CD16+ and CD4+CD56+ T cells in ABD. Other investigators confirmed the presence of increased CD56+ T cells in the periphery of patients with active ABD uveitis, in contrast with other forms of immune-mediated uveitis (Eksioglu-Demiralp et al, 1999; Yato & Matsumoto, 1999).

In the tissue level, NKT cells were found to be increased in the aqueous humor of uveitis patients. Additionally, in neuro-ABD patients, NKT cells were reported to be decreased in the periphery and increased in the cerebrospinal fluid, thus suggesting a recruitment of these cells in the site of inflammation. It should be mentioned that these cells were strongly Th1 polarized. Further studies demonstrated that CD8+CD56+ T cells represent potent cytotoxic effectors in ABD uveitis, as they were found to express both surface FasL and preformed intracellular perforin (Ahn et al, 2005; Hamzaoui K et al, 2006; Yu et al, 2004).

2.2 Adaptive immunity cells abnormalities in ABD

Although the exact pathogenetic mechanisms behind ABD remain to be fully elucidated, much evidence has been accumulated regarding the involvement of adaptive immune response in the disease. Recent studies have revealed the central role of T-cell mediated immunity; in parallel, histopathological studies have demonstrated, mainly, T-cell-dominated perivascular infiltrates in skin, oral lesions and retina, as well as in choroids and periretinal scar tissue (Charteris et al, 1992a, 1992b; Gul et al, 1995, Poulter & Lehner, 1989).

2.2.1 T cell abnormalities

CD4+ T helper cells differentiate into, at least, four distinctive functional subsets, namely Th1, Th2, Th17 and T regulatory cells (Mucida & Cheroutre, 2010). For all of these lymphocyte subpopulations quantitative and/or qualitative abnormalities have been shown in ABD (Chi et al, 2008; Hamzaoui K et al, 2006, Ilhan et al, 2008; Mantas et al, 1999).

Genetic and environmental factors may have an influence independently or in association with other parameters, such as the ligation of T-cell receptor (TCR), the activation of co-stimulatory molecules, the predominance of a given cytokine in the microenvironment and others (Direskeneli et al, 1999).

Many investigators suggest that a Th1 polarization of the immune response is present, at least, in active ABD. Th1 type proinflammatory cytokines, such as IL-2 and IFN- γ , are reported not only to be increased in peripheral blood of patients with ABD, but also to correlate with disease activity. Additionally, IL-12, which drives Th1 immune response from naïve T cells, is also reported to be elevated in ABD. Interleukin-12 is suggested to play a substantial role in ABD; when recombinant IL-12 was added to peripheral blood lymphocytes from patients with active disease, it prevented spontaneous and Fas-induced apoptosis, resulting to proliferation of autoreactive Th1 lymphocytes that could contribute to a prolonged autoimmune inflammation (Esin et al, 1997; Frassanito et al, 1999; Nara et al, 2008; Sugi-Ikai et al, 1998).

Recently, IL-15 was reported to be increased in ABD, either in serum or in the aqueous humor of uveitis patients; IL-15 is able to induce pro-inflammatory cytokines, such as TNF- α and IL-1 β , and facilitate maintenance of auto-reactive CD8+ T cells (Curnow et al, 2008; Waldman, 2004).

The Th1 cells were proven to expand with high individual variability of the V β TCR, suggesting that there is no single antigen responsible for T cell activation (Direskeneli et al, 1999).

Histopathological studies also point to a Th1 type tissue infiltration in ABD, possibly driven by the highly expressed IL-12 and IL-23 in intestinal and cutaneous lesions. Interestingly, both streptococcal antigens and auto-antigens, such as aB-crystallin also drive an IL-12

mediated response in ABD (Imamura et al, 2005; Kulaber et al, 2007; Lew et al, 2008; Yanagihori et al, 2006).

Concerning Th2 type of immune response, both, cells and signature cytokines, such as IL-4, IL-10 and IL-13 are reported to be unaffected or decreased in ABD by most researchers in the serum or in the aqueous humor. However, other studies reported a significant expression of IL-4 in oral ulcers from ABD patients, suggesting the presence of a more complex antigenic stimulus during the disease (Ahn et al, 2006; Dalghous et al, 2006, Mantas et al, 1999).

Th17 cells represent a relatively new functional subset of T helper cells, which mainly produce IL-17A-F, IL-22, IL-23 and TNF α . Interleukin-6 and TGF- β are considered to be the major cytokines responsible for the differentiation of these cells from naïve T cells. Recent studies have shown a possible role for these cells in the majority of systemic and organ-specific autoimmune diseases (Mesquita et al, 2009).

Concerning ABD, active disease is characterized by high serum levels of IL-6 and IL-17, compared to patients in remission, while serum levels of IL-23 and IL-17 are reported to be elevated in active uveitis. However, an increased Th17 response is not confirmed by all studies (Chi et al, 2008; Ferrante et al, 2010; Saruhan-Direskeneli et al, 2003).

Recent papers underline Th17 involvement in disease pathogenesis in the context of their ratio to Th1 cells. Kim et al reported an increased peripheral Th17/Th1 ratio, especially in uveitis and folliculitis patients, while there were no significant differences concerning Th1/Th2 and Th17/Th2 ratios (Kim et al, 2010). Histopathological studies, however, did not confirm these results in the tissue level (Canete et al, 2009).

Cytotoxic T cells are also reported to be activated in ABD, probably as a response to a profound inflammation. CD8⁺ T cells were also found in increased numbers in the aqueous humor in uveitis patients. Furthermore, recent studies support that these cells play a fundamental role in ABD flares, as they were found in increased numbers in active disease, along with $\gamma\delta$ T cells. Additionally, CD8⁺ T cells were shown to express the CD69 molecule (early activation marker), while their numbers and function returned to normal after successful management of the disease flare (Houman et al, 2004; Yasuoka et al, 2008).

2.2.2 Immune regulation in ABD

Firstly described by Sakaguchi et al in the '70s, T regulatory cells represent potent regulators of the immune response. These cells are divided into two broad categories, namely the "naturally occurring" and the "inducible" T regulatory cells (Sakaguchi et al, 1995).

Interest for immune regulation in ABD intensified as early as 1982, when Victorino et al and Sakane et al independently reported abnormal T cell suppressor activity (Sakane et al, 1982; Victorino et al, 1982). Recent studies were not able to definitely shed light in this issue. Paradoxically, CD4⁺CD25⁺FoxP3⁺ T regulatory cells were reported to be increased in active ABD compared to inactive disease and healthy controls, while they could suppress their T effector cell counterparts. The authors suggested that this may be due to the need for inflammation suppression in ABD (Hamzaoui K et al, 2006).

On the other hand, other investigators showed that T regulatory cells were significantly decreased before ocular attack in ABD patients, suggesting that they may serve as a predictive marker in ocular disease, thus, allowing treatment intensification before relapse (Nanke et al, 2008).

The exact interactions between T regulatory cells and other immune cells in ABD are not well understood. Hamzaoui suggested a possible cross talk between these cells and NKT cells, as these subpopulations can modulate each other through cell-to-cell contact and IL-2 dependent mechanisms, respectively (Hamzaoui K, 2007). Furthermore, other cells with possible regulatory activity, such as gammadelta-T-cells were reported to be incapable of maintaining a regulatory profile in ABD (Clemente et al, 2010).

T regulatory cells represent a promising therapeutic target for suppressing inflammation in ABD. Recently, Sugita et al reported T-reg expansion after in vitro stimulation with infliximab and significant suppression of bystander T effector cells in active ABD uveitis (Sugita et al, 2011). Additionally, other investigators reported that vitamin D is an important promoter of T cell regulation in vivo in ABD patients, as its levels were strongly correlated to T regulatory cells (Hamzaoui K et al, 2010).

2.2.3 B cell abnormalities and autoantibodies in ABD

Although not a typical autoimmune disease, ABD is characterized by B cell activation, despite the absence of disease-specific autoantibodies. Early studies demonstrated that the total numbers of B cells were normal during the disease; however, an increased expression of activation markers, such as CD13, CD133, CD80 and memory marker CD45RO was reported. Despite the presumed B cell activation, plasma cells were found in low levels, in contrast to other well-defined autoantibody-mediated disorders. These results suggest a modified B cell function in ABD, although it is not known if it is related to a weak stimulus by an unknown external antigen or it represents an epiphenomenon in the context of the general dysfunction of the immune system (Eksioglu-Demiralp et al, 1999).

After the identification of the multiple roles of B cells in the immune response, such as antigen presentation and cytokine release, newer studies reported some interesting results. B cells (along with plasma cells) were found in histopathological studies in the synovium, the central nervous system and the vessel wall of patients with ABD (Hirohata, 2008; Hirohata & Kikuchi, 2009; Suh et al, 2001).

Furthermore, it was recently reported that B-cell activating factor (BAFF) receptor was upregulated on B cells in patients with ABD vasculitis, as well as serum BAFF levels and mRNA in skin biopsies. Later, Hamzaoui A et al showed that BAFF, along with IL-6 and IL-13, were, also, increased in the broncho-alveolar lavage (BAL) fluid from ABD patients with active pulmonary involvement; All these factors are known to actively participate in the maturation and activation of B cells (Hamzaoui A, 2010; Hamzaoui K, 2008).

Consequently to B cell activation, various autoantibodies have been detected in the sera of ABD patients; however, their role in disease pathogenesis remains to be defined. Antinuclear antibodies (ANA) and rheumatoid factors are typically absent. On the other hand, autoantibodies against cell surface antigens, such as anti-endothelial cells antibodies (AECA), or against mucosal antigens have repeatedly been reported. Recent studies, based on proteomics, have revealed the presence of antibodies against more specific antigens, such as α -enolase, α -tropomyosin, kinectin, selenium-binding protein, esterase-D and carboxy-terminal subunit of Sip-1 (Delunardo et al, 2006; Direskeneli et al, 1995; Lee KH et al, 2003; Lu et al, 2005; Mahesh et al, 2005; Michelson et al, 1985; Mor et al, 2002; Okunuki et al, 2007, 2008).

Concerning AECA, their prevalence in ABD is reported to vary from 18-50%; however, difficulties in method standardization led to significant discrepancy between the different studies and raised concerns about their clinical utility. Furthermore, other studies confirmed that AECA were not quite specific for the disease; instead, they are frequently detected in other systemic vasculitides and/or autoimmune diseases. Despite their inspecificity, AECA were shown to mediate enhanced polymorphonuclear adhesion to activated endothelial cells, thus promoting tissue injury during systemic inflammation. An association between IgG-AECA and central nervous system involvement in ABD was reported (Dinc et al, 2003; Souza et al, 2007; Zeng et al, 2004).

Alpha-enolase represents a target antigen of IgM AECA, reported to be detectable in approximately half of ABD patients, especially in those with overt vascular involvement; however, its specificity is not satisfying (Lee JH et al, 2009).

3. Coagulation and fibrinolytic abnormalities in ABD

Although no specific defect in the coagulation cascade has so far been detected, ABD is characterized by thrombotic manifestations in 7-38% of the patients. In most studies, both coagulation and fibrinolytic pathways seem to be activated (Leiba et al, 2004).

Various pro-coagulant conditions, such as deficiencies of protein C, protein S, antithrombin III and factor V Leiden and prothrombin G20210A mutations have been described in ABD patients (Ates et al, 2003; Caramaschi et al, 2010). Additionally, higher levels of homocysteine, factor VIII, thrombin-antithrombin III complexes (TAT) and prothrombin fragment 1+2 maintain the intravascular generation of thrombin in these patients (Espinosa et al, 2002; La Regina et al, 2010). Despite all these data, it seems that thrombophilic factors do not play a major role in the thrombotic tendency of ABD. Leiba et al, in a tightly designed study, showed that the only factor that was strongly correlated to thrombosis in these patients was hypertriglyceridemia (Leiba et al, 2004). Other studies reported that active ABD patients were characterized by hypertriglyceridemia, low HDL and high Lp(a) levels (Musabak et al, 2005).

Regarding the fibrinolytic cascade, most studies have revealed defective fibrinolysis with lower levels of tissue plasminogen activator (t-PA) and increased thrombin activatable fibrinolysis inhibitor (TAFI) and thrombomodulin. Other studies involve the plasminogen activator inhibitor (PAI), as this molecule appears to be increased in ABD, resulting in defective tPA/PAI complex formation (Donmez et al, 2010; Ozturk et al, 2004). Furthermore, the endothelial dysfunction due to immune-mediated vasculitis may also play an important role in the thrombotic tendency of the disease.

Another important thrombogenic factor in ABD is the presence of antiphospholipid antibodies. Indeed, anticardiolipin (ACA) and anti- β 2-glycoprotein 1 (anti- β 2-GPI) antibodies, as well as lupus anticoagulant (LA) have been described in selected ABD patients (Bonnet et al, 2004; Kandolf-Sekulovic et al, 2005). Additionally, newer epitopes, such as antibodies against the phosphatidylserine-prothrombin complex (anti-PS/PT) or annexin V, have also been reported to be associated with thrombosis in these patients (Aslan et al, 2004; Kawakami et al, 2009). The significance of these antibodies in thrombotic manifestations of ABD remains to be fully elucidated.

In Table 2, a synopsis of the main abnormalities of the immune cells in ABD is presented.

Neutrophils	Hyperactivation Cytokine and MMPs secretion Resistance to apoptosis Tissue infiltration
Monocytes/Macrophages	Altered TLR expression Secretion of IFN- γ , IL-1, IL-8, IL-12, TNF- α
Gammadelta T cells	Vgamma9+Vdelta2+ expansion Secretion of IFN- γ , TNF- α
NK cells	Polyclonal proliferation Increased numbers Impaired cytotoxic activity?
NKT cells	Abnormal KIR repertoire Recruitment to the site of inflammation High expression of FasL and perforin Interaction with T regulatory cells?
CD8+ T cells	Increased numbers Highly activated (CD69)
CD4+ T cells (Th1 polarized)	Secretion of IL-2, IL-12, IFN- γ Resistance to apoptosis Tissue infiltration
CD4+ T cells (Th17 polarized)	Disease initiation? Secretion of IL-17, IL-23 Increased Th17/Th1 ratio
T regulatory cells	Probably low numbers during active disease Impaired suppressive activity?
B cells	Normal numbers, but activated Increased expression of BAFF receptor Autoantibody production (AECA and others)

Table 2. Main abnormalities in immune cells in ABD

A proposed model for ABD vasculitis pathogenesis

(schematic representation -1, -2, -3)

In genetically predisposed individuals, due to an exogenous stress, e.g. an infectious insult, a systemic or local non-specific immune response is mediated via innate immune cells and the endothelium.

Respiratory burst products and proinflammatory cytokines and chemokines secreted at the site of the initial insult further attract monocytes, neutrophils and dendritic cells around the area and the blood-endothelial barrier breaks down (schematic representation -1).

Initial insult is neutralized, but HSPs derived from bacterial load, as well as HSPs from affected tissue/s give rise to a second, more specific T-cell-mediated immune response, according to molecular mimicry and/or epitope spreading model. These events might be augmented and spread out in case of repeated antigenic stimuli (second hit). Endothelial-innate cell contact and activation perpetuates the proinflammatory milieu and along with

local tissue injury produces a ‘danger model’ situation, where APCs, through PAMP-PRR cross-talk, are activated and induce downstream Th1 and Th17 polarization of the specific immune response. Main cytokines mediating this process are IL-1, TNF- α , IL-6, IL-8, IL-12, IL-18 and others, (schematic representation -2).

T cells, through this antigen-driven activation, are committed and clonally expanded against certain candidate peptides via well known mechanisms of molecular cross-reactivity. The orchestrated immune reaction involves B cells activated by CD4+ T cells to produce autoantibodies (AECA?) that are able to induce endothelial damage and contribute to neutrophil migration as well.

Gammadelta-T-cells probably contribute to the pathogenetic process, by cytokine secretion which augments Th1 polarization. The role of cell-mediated cytotoxicity is not clear at the moment. Probably, CD8+, NK and NKT cells are recruited in the site of inflammation and mediate a cytotoxic immune response.

Hyperactivated neutrophils, primed by cytokines and the activated endothelium (overexpressing cell adhesion molecules), migrate into the vessel wall and produce reactive oxygen species and MMPs, thus contributing to vasculitic lesions and aneurysm formation.

Additionally, low numbers or defective function of T regulatory cells probably cannot adequately regulate the hyperactivation status of effector cells, (schematic representation -3).

Finally, during disease perpetuation, tissue damage may produce new epitopes (damage-associated molecular patterns, DAMPs) that can further augment immune cells activation and contribute to the clinical heterogeneity of the disease.

This pathogenetic model is briefly presented in Figure 1.

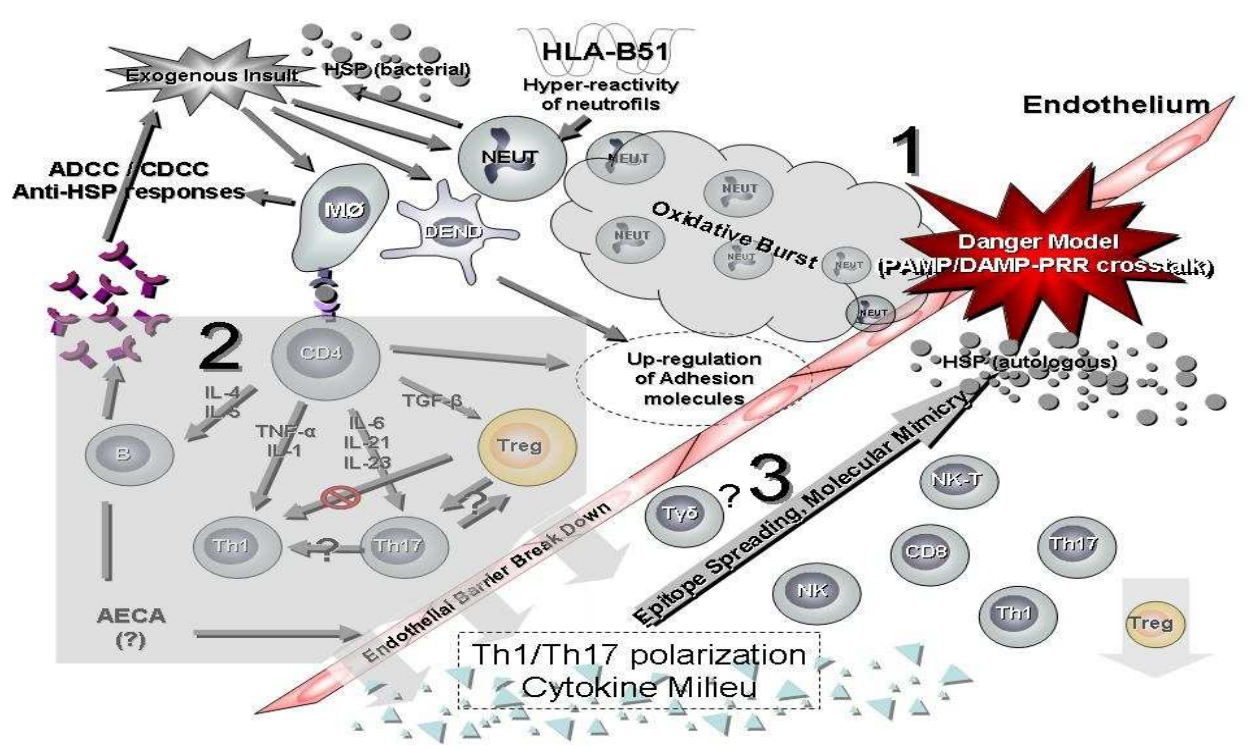


Fig. 1. Proposed pathogenetic model of ABD vasculitis

4. Therapeutic approach to ABD vasculitis

4.1 General principles

Although treatment of ABD was reviewed recently, management of complicated cases, especially those with vascular involvement, still poses a great challenge to experienced physicians. Nowadays, the recently published EULAR (European League Against Rheumatism) recommendations are considered to be the accepted guidelines for the management of the disease. Nonetheless, treatment concerning vascular involvement, either major vessel or visceral disease (medium-small vessel vasculitis) is largely based on expert opinion and uncontrolled evidence from open trials and observational studies. So the need for further properly designed controlled clinical trials is obvious (Hatemi et al, 2008).

In general, topical therapy should be applied in mucocutaneous lesions and should be the only treatment for mild symptoms. Although antiseptics seem to prevent secondary infections and reduce pain intensity, antibiotics face antigen burden and they are important in identifying disease relapses as well. Drugs with anti-inflammatory action are used as adjuvants but, beyond this, it is imperative to emphasize that the physician must be vigilant for alarming clues of systemic vasculitis and ready to implement immunomodulating regimen (Alpsoy, 2005; Alpsoy et al, 2007).

Treatment decisions depend on the age, sex, comorbidities, patient's willingness to cooperate and other parameters, such as the site and severity of involvement, as well as the frequency of recurrences. Fortunately, a part of disease burden can be managed with mild systemic immunosuppression. Early and aggressive immunosuppression is mandatory in severe or life-threatening cases. Below, the most commonly used immunomodulating drugs for the management of major vessel disease or visceral involvement will be discussed. (Table 3)

4.2 Glucocorticosteroids (C/S)

It is well known that glucocorticosteroids modify immune response to diverse stimuli by suppressing inflammatory reactions in general. They have profound metabolic effects and they affect the replication, movement and activity of virtually all cells involved in inflammatory process. Although C/S exert cytotoxic effects to activated T cells at high doses, their immunomodulating effects are probably mostly attributed to their ability to modify cellular immune functions when low doses are used orally. At the cellular level C/S bind to their intracellular receptor forming an active complex. The complex moves to the nucleus, where it binds to DNA sequences modifying chromatin structure and altering the transcription machinery. Another mechanism of action is that glucocorticosteroids-glucocorticosteroids receptor complex directly increases I κ B (inhibitory factor κ B) expression. I κ B inhibits the actions of transcription nuclear factor NF κ B. It is well known that NF κ B promotes gene induction of cytokines, chemokines and adhesion molecules, thus amplifying lymphocyte proliferation and leading to the perpetuation of the inflammatory process (Bruton et al, 2005; Rhen & Chidlowski, 2005).

Through these mechanisms C/S suppress cellular immunity, both Th17 and Th1 responses. This action explains their therapeutic use in ABD. Although more detailed trials are needed, worldwide clinical experience has shown that C/S are the cornerstone of treatment for many ABD complications. High oral doses, up to 100mg/day for a long period of time, or pulse treatment with 1gr/day for 3 consecutive days or more are given, in serious or life-threatening cases, like aneurysm formation or cavernous sinus thrombosis (Hatemi et al,

2008). It was demonstrated that in large-vessel arterial disease (arterial occlusion or aneurysm) high dose glucocorticosteroids alone did not improve occlusive disease and mortality (Li Thi Huong et al, 1995). On the contrary, other studies have shown that C/S in combination with other immunosuppressants had a significant beneficial effect in vascular complications of ABD (Alpsoy & Alkman, 2009, Sharaf & Yazici, 2009). In another recently conducted randomized trial, low dose intramuscular depot C/S was used and proved to be beneficial in milder manifestations, like erythema nodosum, but no for genital ulcers. The trial used the drug in depot form for intramuscular injection, so no conclusions can be drawn for the efficacy of the usual oral or intravenous regimen (Mat et al, 2006).

Adverse effects such as osteoporosis, metabolic disturbances, psychosis must be taken into account. EULAR committee recommends the use of C/S as a first line therapy for managing all ABD manifestations. Different level of evidence for every case is probably attributed to data availability. Based on immunopathology and clinical data, we suggest the administration of high dose intravenous corticosteroids for the achievement of remission in large vessel or CNS vasculitis, followed by a slow tapering schedule to the lowest effective dose which sustains remission, under close clinical and laboratory follow up (Bruton et al, 2005; Hatemi et al, 2008; Li Thi Huong, 1995; Sharaf & Yazici, 2009).

4.3 Azathioprine

Azathioprine is a prodrug of 6-mercaptopurine, containing an imidazole ring and acts like an antimetabolite. Azathioprine and mercaptopurine appear to induce immunosuppression by interfering with purine nucleic acid metabolism. The drug intervenes with the process of cell division at steps that are required for lymphoid cell proliferation following antigenic stimulation. Stimulated lymphoid cells incapable of entering cell mitotic phase undergo apoptosis, establishing a relevant state of immunosuppression. Although RNA synthesis is blocked at the same time, these agents appear to have less effect on cell function if implemented after antigen exposure than on nucleic acid synthesis in proliferating, antigen encountering, T cells (Bruton et al, 2005).

The efficacy of azathioprine in ABD is well established. The usual dose is 1-2.5 mg/kg/day (not exceeding 200 mg/day) and an average of 3 months is required for its effect to be established. This slow immunomodulating action limits the use of azathioprine as a sole agent of therapy in acute vasculitis presentation. However, it can be used additionally in the acute phase as a corticosteroid sparing agent, as well as prophylactically. Yazici H et al propose azathioprine as a first line therapy for non-life-threatening cases of neurological complications (Yazici H et al, 1990). According to the EULAR recommendations, azathioprine should be used in any ABD patient having inflammatory eye disease with posterior segment involvement (Hatemi et al, 2008). Its efficacy and safety was evaluated in a randomized, double-blind, placebo-controlled trial. In this trial, there were fewer new cases of ocular involvement in azathioprine plus corticosteroids group comparative to corticosteroids alone (one versus eight). Additionally, the patients taking azathioprine had less frequent oral ulcers, genital ulcers and arthritis (Yazici H et al, 1990). Contemplating the pathophysiology of ABD (see above), immunosuppressants like azathioprine may play a role in the management of vessel thrombosis through anti-inflammatory action on vessel wall. Results from Sharaf et al confirm the clinical experience that by dampening immune system activation, pro-thrombotic diathesis can be, at least partially, reversed (Sharaf & Yazici, 2009).

In general, azathioprine could be used as a first line treatment in combination with corticosteroids in almost all serious manifestations of the disease and is preferable for maintenance therapy after remission induction. Common side effects are gastrointestinal disturbances, liver toxicity and bone marrow suppression but rarely do they lead to drug discontinuation. In parallel, due to B cell impairment and mild hypoglobulinemia, humoral immunity can be weakened by these cytotoxic agents as well.

4.4 Cyclophosphamide (CyP)

Cyclophosphamide is a highly active alkylating agent, which impairs DNA duplication and cell division. Although it is not a cell specific drug, the greater effect is observed upon rapidly proliferating cells, such as T cell clones after their encounter with the antigen. Initially, cyclophosphamide has proven a reliable agent in management of lupus nephritis and ANCA associated vasculitis and, eventually, has found a distinguished place in ABD vasculitis (Bruton et al, 2005).

It has been known that pulse CyP, combined with intravenous C/S regimen, help to achieve remission in ABD vascular and/or visceral involvement (Du et al, 1990; Hamza et al, 1992). According to EULAR recommendations, large vessel vasculitis, such as pulmonary and/or peripheral aneurysms, need to be treated with CyP for at least 2 years (Hatemi et al, 2008). The drug may be administrated either intravenously, as pulse treatment in doses 400-700mg/m², or orally in doses of 1-3 mg/day. The exact dose depends upon the severity of the disease and considerations of renal and hepatic impairment. Deep immunosuppression following CyP administration has a life-saving effect. Also in a double-blind cross over trial, CyP has proven beneficial in patients with intractable uveitis (Davatchi et al, 1999).

Due to its severe toxicity (bone marrow suppression, mucositis, neurotoxicity, syndrome of inappropriate ADH secretion, cancer), this drug must be used in life-threatening or non-responding cases. We prefer to administer CyP intravenously, as monthly or 3-weekly pulses of 500-1000mg bolus, in order to partially avoid bladder toxicity. For safety reasons, mesna (chelate agent) and adequate fluids must be administered concurrently.

4.5 Cyclosporine

Cyclosporine is a cyclic polypeptide, which forms a complex with cyclophilin, an intracellular factor mediating signaling pathways. This complex binds to calcineurin and inhibits Ca²⁺-stimulated dephosphorylation of the cytosolic component of the transcription factor NFAT. Activated cytoplasmic NFAT translocates to the nucleus and binds to nuclear components required for complete T-cell activation, blocking of IL-2 and other Th1 cytokine producing genes (Th1 to Th2 switch). Besides this, calcineurin also possess peptidyl-prolyl cis-trans-isomerase enzymatic activity. The peptidyl-prolyl cis-trans-isomerase domain of calcineurin facilitates binding to CD147, also known as an inducer of extracellular-matrix metalloproteinases (MMPs). This binding causes CD147 to translocate to the cell surface, where it plays a critical role in stimulating matrix-metalloproteinase activity, leading to matrix degradation. Thus, this agent inhibits the enzymatic activation of MMPs, which leads to vessel wall thinning (Weintraub, 2009). Cyclosporine is effective against T-cell-dependent immune reactions and a suitable drug for immune intervention in the pathophysiological process of ABD vasculitis (see above). Additionally, cyclosporine, in synergy with C/S, downregulates the activity of NKT cells controlling inflammatory process (Bruton et al, 2005; Chi et al, 2010).

The efficacy of cyclosporine in ABD has been evaluated primarily in patients with ocular, mucocutaneous and articular manifestations. Approximately 50% of patients showed clinical remission when treated with cyclosporine. Two well designed trials proved that cyclosporine protects ocular degeneration and visual acuity better than conventional regimen containing prednisolone (Atmaca, 1994; Ernakova, 2003). A direct comparison of the therapeutic potential of cyclosporine to oral cyclophosphamide was conducted by Ozyazgan Y et al in patients with ocular inflammation. Therapeutic superiority concerning visual acuity was seen in the cyclosporine group at six months, but similar results were noted for both agents at two years (Ozyazgan et al, 1992). Similarly, comparison of cyclosporine with glucocorticoids or chloramboucil indicates its superiority to control ocular symptoms. EULAR recommendations suggest cyclosporine or infliximab in combination with azathioprine and corticosteroids for refractory eye involvement and major vessel disease as well (strength of recommendation C) (Hatemi et al, 2008).

Side effects are hypertension, nephrotoxicity and predisposition to infections. Furthermore, the role of cyclosporine in vascular neurological involvement in ABD has not been clarified yet and its use is discouraged (Bruton et al, 2005).

4.6 Colchicine

Colchicine exerts a variety of pharmacological effects, especially in cellular immunity. The drug arrests cell division in G1 phase by interfering with microtubule and spindle formation. This anti-mitotic effect is greatest on cells with rapid turnover (e.g. neutrophils and expanding lymphocyte clones). Colchicine also renders cell membranes more rigid and decreases the secretion of chemotactic factors from activated neutrophils, thus inhibiting inflammation in many ways (Actulga et al, 1980; Bruton et al, 2005).

Clinical experience and randomized trials support that the use of colchicine, alone or in combination with penicillin, is beneficial in reducing the number of episodes of aphthous mucositis and arthritis (Al Waiz et al, 2005). According to the authors' opinion and EULAR recommendations, colchicine has no role in severe large vessel vasculitis management and in acute relapses. Slow onset of action and lack of T cell specificity render this drug improper for use in severe cases. Beyond that, there is strong recommendation (category Ib) for the management of cases where the dominant lesion is erythema nodosum (medium size vasculitis) (Hatemi et al, 2008).

Side effects of colchicine such as nausea, vomiting, diarrhea and hematologic disturbances are well controlled with stepwise use of the drug.

4.7 Methotrexate (MTX)

Methotrexate competitively inhibits dihydrofolate reductase (DHFR), an enzyme that participates in the tetrahydrofolate synthesis. Methotrexate acts specifically during DNA and RNA synthesis and, thus, it is cytotoxic during the S-phase of the cell cycle. MTX, by inhibiting the enzymes involved in purine metabolism, and leads to intracellular accumulation of adenosine and suppression of intercellular adhesion molecule expression by T cells. These two mechanisms provoke T cell inhibition (Bruton et al, 2005; Johnston et al, 2005).

Concerning ABD vasculitis, literature for MTX is relatively poor. However, Borhani Haghighi A proposes MTX as a first line therapy in neurological complications (Borhani Haghighi, 2009). The relatively low cost and the acceptable safety profile make MTX a preferable choice (Davatchi et al, 2003; Kikuchi et al, 2003). EULAR committee recommends

MTX for CNS involvement (strength of recommendation C), but no distinction is made between vascular and parenchymal involvement (Hatemi et al, 2008). Common side effects are liver toxicity and macrocytosis.

4.8 Biologic agents

4.8.1 Interferon

Although interferons (alpha, beta, and gamma) were identified by their antiviral activity, these agents also have important immunomodulatory functions. Interferons bind to membrane receptors type I and II and produce their immunomodulating effects through a cascade of gene induction and transcriptional changes. The final result is enhancement of macrophage phagocytic activity and augmentation of specific cytotoxicity by CD8+T lymphocytes. How these changes modify immune system reaction in ABD has not been clarified. Possible mechanisms include reduction of viral antigen load and antiproliferative properties on lymphocytes and damaged tissue cells. Another probable mechanism of action is the up-regulation of sTNF- α -RII (soluble TNF- α receptor) and IL-1ra (interleukin 1 receptor antagonist). It should be emphasized that other studies entangle IFN- α action in the induction and propagation of autoimmune process (Kotter et al, 2004; Theofilopoulos et al, 2005).

In a systematic review by Kotter I et al, the authors concluded that nearly all patients with mucocutaneous symptoms and/or arthritis and/ or uveitis responded to IFN- α administration in a dose related manner (Kotter et al, 2004). Additionally, in other studies including cases of refractory uveitis, IFN- α had similar positive results. These reports were confirmed by Tugal-Tutkun et al, although they reported lower rates of complete response (Tugal-Tutkun et al, 2006). EULAR committee justifies the use of IFN- α with or without corticosteroids for eye disease, instead of cyclosporine or azathioprine. On the contrary, EULAR recommendations do not consider the use of IFN- α for major artery vasculitis due to insufficient data from large studies (Hatemi et al, 2008). We consider that potent, traditional immunosuppressants are preferable in these cases and that administration of an agent with parallel immunostimulant properties in full-blown inflammation is ambiguous (Boura et al, 2006). Side effects of IFN- α include psychosis, nausea, vomiting, diarrhea, hematologic changes, liver toxicity and leucoencephalopathy (Bruton et al, 2005).

4.8.2 Anti – TNF- α (tumor necrosis factor) agents

TNF- α , as a pro-inflammatory cytokine, promotes immune reactions and inflammation. Infliximab is a chimeric anti-TNF- α monoclonal antibody, which binds with high affinity to TNF- α and prevents the cytokine from binding to its receptors. A number of studies (uncontrolled, open labeled, case series and reports) indicate that infliximab control mucocutaneous, joint, neural, gastrointestinal and eye manifestations of disease, inducing remission (Boura et al, 2006; Sfrikakis et al, 2001, 2004). Additionally, in these studies, infliximab proved adequate to sustain remission in cases refractory to azathioprine and/or cyclosporine. Sfrikakis et al propose infliximab as a second line therapy for ABD relapses when usual therapy fails or as a first line regimen in severe sight-threatening uveitis (Sfrikakis et al, 2004). Our experience is in agreement.

Etanercept contains the ligand binding portion of a human TNF- α receptor fused to the Fc portion of human IgG1, and binds to TNF- α , thus preventing it from interacting with its receptors. Melikoglu M et al conducted a controlled study using etanercept in ABD.

Etanercept was superior to placebo in controlling oral ABD manifestations, whereas the rate of genital ulcers healing, joint symptoms and papulopustular skin lesions were not statistically different between the two groups (Melikoglu et al, 2005). Different results were reported from the Study Group on Autoimmune Diseases (GEAS), who published an impressive rate of response in ABD using etanercept (96%), (Ramos-Casals et al, 2008). These contradictory results indicate that more studies need to be undertaken in order to assess etanercept in ABD.

Like infliximab and etanercept, adalimumab binds to TNF- α , preventing it from activating TNF- α receptors. There is an increasing number of reports that adalimumab is promising on patients with symptoms refractory to other agents (Bawazeer et al, 2010). EULAR recommendations include anti TNF- α agents in possible therapeutic regimens for gastrointestinal and CNS involvement, but no mention is made for major vessel inflammation (Hatemi et al, 2008).

Side effects (infections, cancer) and cost must be taken into account. TNF- α antagonists could be reserved as a second line therapy in intractable posterior uveitis and CNS vascular disease as well.

4.8.3 Anakinra (anti-interleukin-1)

IL-1 mediates its pro-inflammatory actions through IL-1 receptor binding on the surface of immunocompetent cells. For homeostatic reasons there is a naturally occurring IL-1 receptor antagonist (IL-1ra) in serum, which competes with IL-1 for receptor binding, blocks IL-1 activity and ameliorates its pro-inflammatory actions. The balance between IL-1 and IL-1ra may contribute to the extent of an inflammatory response. Anakinra, a recombinant non-glycosylated version of human IL-1 receptor antagonist, may have a therapeutic role in ABD. Although literature is limited, a case report of a patient with severe vascular and gastrointestinal ABD, refractory to conventional treatment, who responded only to anakinra has been published (Botsios et al, 2008). However, later reports question the long term efficacy of anakinra in preventing secondary complications, such as amyloidosis (Bilginer et al, 2010). These ambiguous results confirm the view that further studies are needed.

4.9 Other treatments

There are limited data available regarding the use of alternative therapeutic options in ABD. Intravenous immunoglobulins have been tested either in a small number of patients or in a heterogeneous group of patients with eye inflammation (Seider et al, 2001).

IL-6 plays a central role in disease pathophysiology. IL-6 levels have been reported to be elevated in the cerebrospinal fluid of patients with active CNS disease. Probably anti-IL-6 treatment could be useful in vascular manifestations of ABD. In fact, Borhani Haghighi A proposed tocilizumab (anti-interleukin-6 agent), as an additional weapon to the therapeutic armamentarium (Borhani Haghighi & Safari, 2008).

Lim SH and de Cata A reported resolution of ABD manifestations after non myeloablative hematopoietic stem cell transplantation (autologous or derived from allogeneic bone marrow or umbilical cord blood). There is an increasing number of studies proposing this practice as an upcoming treatment for refractory ABD. The rationale is the depletion of activated immune cells from the body (De Cata et al, 2007; Lim et al, 2009)

All these therapeutic modalities have not been tested adequately and are not mentioned in current guidelines.

5. Specific considerations for vascular inflammation in ABD

5.1 Major vessel involvement

Although large artery disease is uncommon in ABD, it can lead to significant morbidity and mortality. The percentage of ABD patients with arterial damage varies from 5 to 18%. The two main presentations of arterial involvement is lumen stenosis, or more likely, aneurysm formation.

Although pathogenesis is yet unclear, the hypothetical model already proposed for other large vessel vasculitides might be of value. According to this model, IL-1 and TNF- α secretion from dendritic cells, macrophages and neutrophils results in the production of radical oxygen species and metalloproteinases with subsequent elastic fiber destruction, thinning of media layer and expansive remodeling of the artery (Weyand & Goronzy, 2003). Chronic uncontrolled inflammation also leads also to arterial occlusions due to intimal thickening. These two forms of vessel remodeling may be detected in the same patient (Buggage et al, 2005).

Aneurysm formation and/or occlusive lesions may require combined medical and surgical intervention under the supervision of an experienced clinical immunologist. Urgent surgery is mandatory in case of enlarging or ruptured aneurysms or due to organ-threatening ischemia. Equally significant is the implementation of postoperative immunosuppressive therapy, to prevent relapse and common complications after arterial bypass surgery, such as graft occlusion and new aneurysm formation at the site of anastomosis (Calamia et al, 2011; Tuzun et al, 1997).

The first goal is to achieve remission with immunosuppressive drugs. We suggest aggressive therapy with methylprednisolone 1000mg x 3 or more consecutive days, followed by oral C/S and CyP in intravenous pulse treatment (4-8mg/kg) until clinical and laboratory remission.

A number of patients with major-vessel arterial disease were studied in a retrospective manner. This study illustrated that C/S, even in high doses, did not improve occlusive disease when fibrosis of intima was irreversible, emphasizing the importance of early immune intervention (Le Thi Huong, 1999).

Anticoagulation is employed successfully in several cases, but EULAR committee do not recommend anticoagulants or antiplatelet drugs due to lack of evidence from randomized controlled trials and possible danger of bleeding in sites of aneurysmal rupture (Hatemi et al, 2008). (see algorithm 1)

5.2 Medium - small vessel involvement

Although large vessel involvement is responsible for deaths in ABD, small and medium vessel vasculitis maintains the main characteristic in disease pathophysiology. Pathological findings compatible with vasculitis from different anatomical sites have been reported. More specifically, the histopathological findings range from minor neutrophilic vascular infiltration to true leukocytoclastic (oral lesions) or lymphocytic (genital lesions) vasculitis. Although such interesting clues indicate a variance in the pathophysiologic process, there are no data to suggest different therapeutic approaches in patients with different type of lesions (Chun et al, 1990; Nazarro et al, 1966).

Retinal vasculitis is a serious finding in posterior uveitis of ABD patients. EULAR committee recommends a treatment regimen that includes azathioprine and systemic corticosteroids for the management of the first attack. Doses of prednisone up to 1 mg/kg/day for one

month with tapering thereafter as tolerated proved to be adequate in most of the cases. Furthermore, long term follow-up of patients proves that local and/or systemic corticosteroids and azathioprine are beneficial for maintenance treatment as well (Hatemi et al, 2008; Iscan et al, 2005; Yazici et al, 2010).

In severe relapsing or refractory eye inflammation, initial pulse therapy with intravenous methylprednisolone (1 g/day for three days or more) is needed. A number of observational studies have suggested efficacy of infliximab for the treatment of inflammatory eye disease. Combination of the above agents with cyclosporine or IFN- α can be used for intractable eye inflammation (EULAR recommendations 1 and 2), although the level of evidence is low (grade of evidence III) (Hatemi et al, 2008; Yazici et al, 2010).

Alternatively, the addition of cyclophosphamide pulse treatment prior to infliximab in refractory ocular ABD could be used. There is a possible synergistic effect of cyclophosphamide and infliximab in controlling ocular inflammation when used consecutively (submitted for publication).

In general, neurological complications could be treated in the same manner as posterior uveitis. The combination of high-dose C/S (3–7 pulses of intravenous methylprednisolone 1 g/day) with another immunosuppressive agent (preferably azathioprine) is required. Cyclophosphamide, methotrexate and TNF- α antagonists represent quite useful alternatives in severe parenchymal disease. For dural sinus thrombosis, short pulses of intravenous C/S are usually adequate. In neurological involvement, cyclosporine should not be introduced due to its potential neurotoxicity (strength of recommendation C) (Borhani Haghighi, 2009; Borhani-Haghighi & Safari, 2010; Hatemi et al, 2008; Kotter et al, 2006).

Serious renal lesions are rare and range from minor IgA deposits in glomerulus to catastrophic crescentic glomerulonephritis. In case of clinically important glomerulonephritis, treatment varies according to other organ involvement. A reasonable approach is cyclophosphamide 750-1000 mg /month plus glucocorticosteroids 500-1000 mg /day for three days and 60-100 mg oral prednisolone in tapering doses until clinical remission (Altıparmak et al, 2001). (see algorithm 2)

5.3 Venous thrombosis

Deep vein thrombosis (DVT) is the most frequent vascular manifestation of ABD. DVT is believed to result from endothelial inflammation leading to thrombosis through pro-thrombotic alterations in the vessel wall, upregulation of adhesion molecules and endothelial dysfunction. DVT can occur in any site of the vascular tree, most commonly in lower extremities, but also in hepatic veins, cerebral venous sinuses, superior and inferior vena cava (Ames et al, 2001).

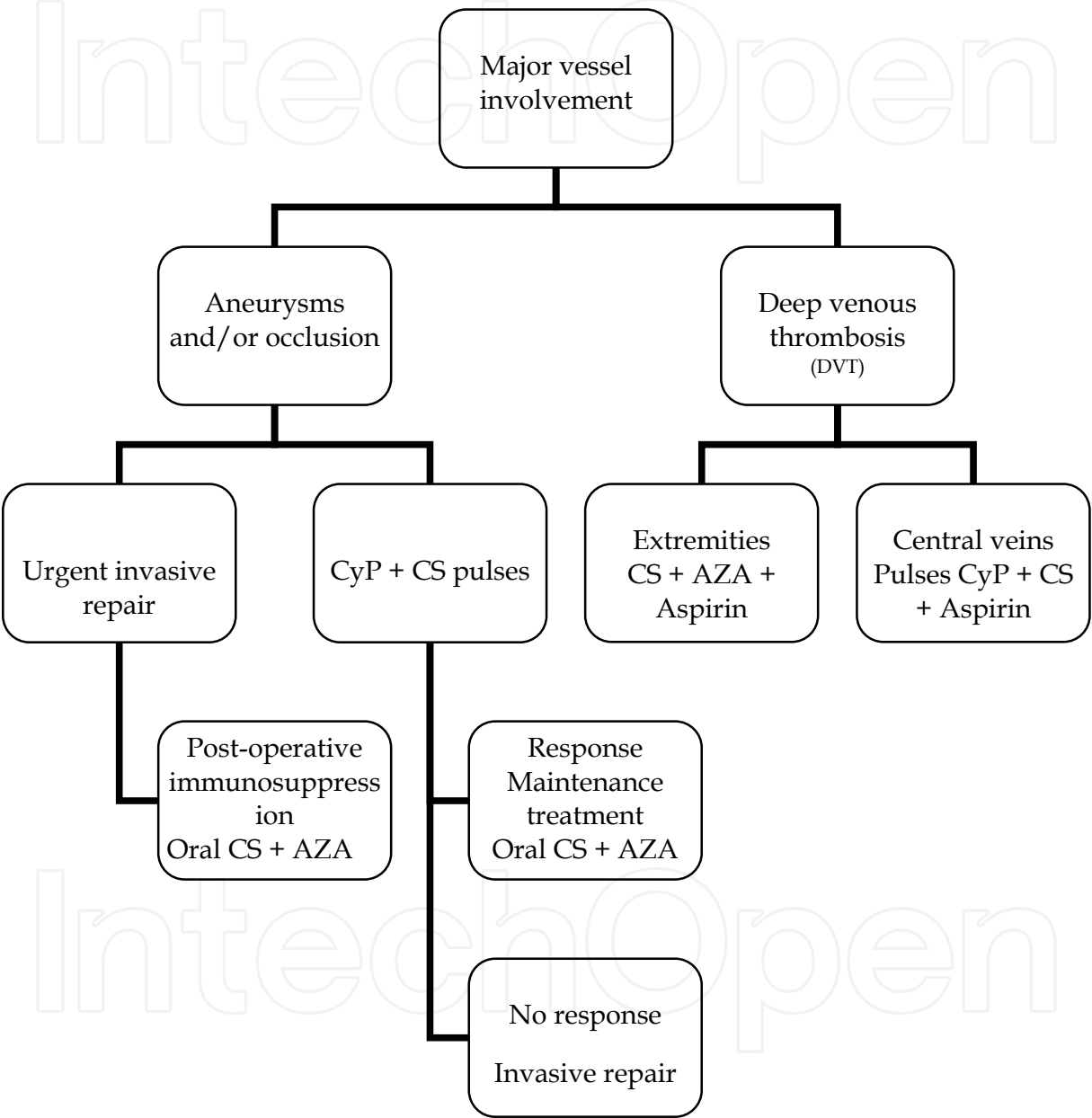
Thrombotic events are prevented by ameliorating systemic inflammation, rather than by the institution of primary anticoagulation (Kuzu et al, 1994; Mader et al, 1999; Sarica-Kucukoglu et al, 2006). EULAR does not recommend the use of antiplatelet drugs, anticoagulants or fibrinolytic agents for thrombotic events in ABD, due to insufficient data from large studies (Hatemi et al, 2008).

The authors suggest that venous thrombosis must be faced on case-by-case basis, through cooperation with appropriate subspecialists. When anti-thrombotic intervention is considered, aspirin is a rational choice. Acetylsalicylic acid interferes with the transcriptional activation of the interferon- γ gene, a mechanism of action that may be

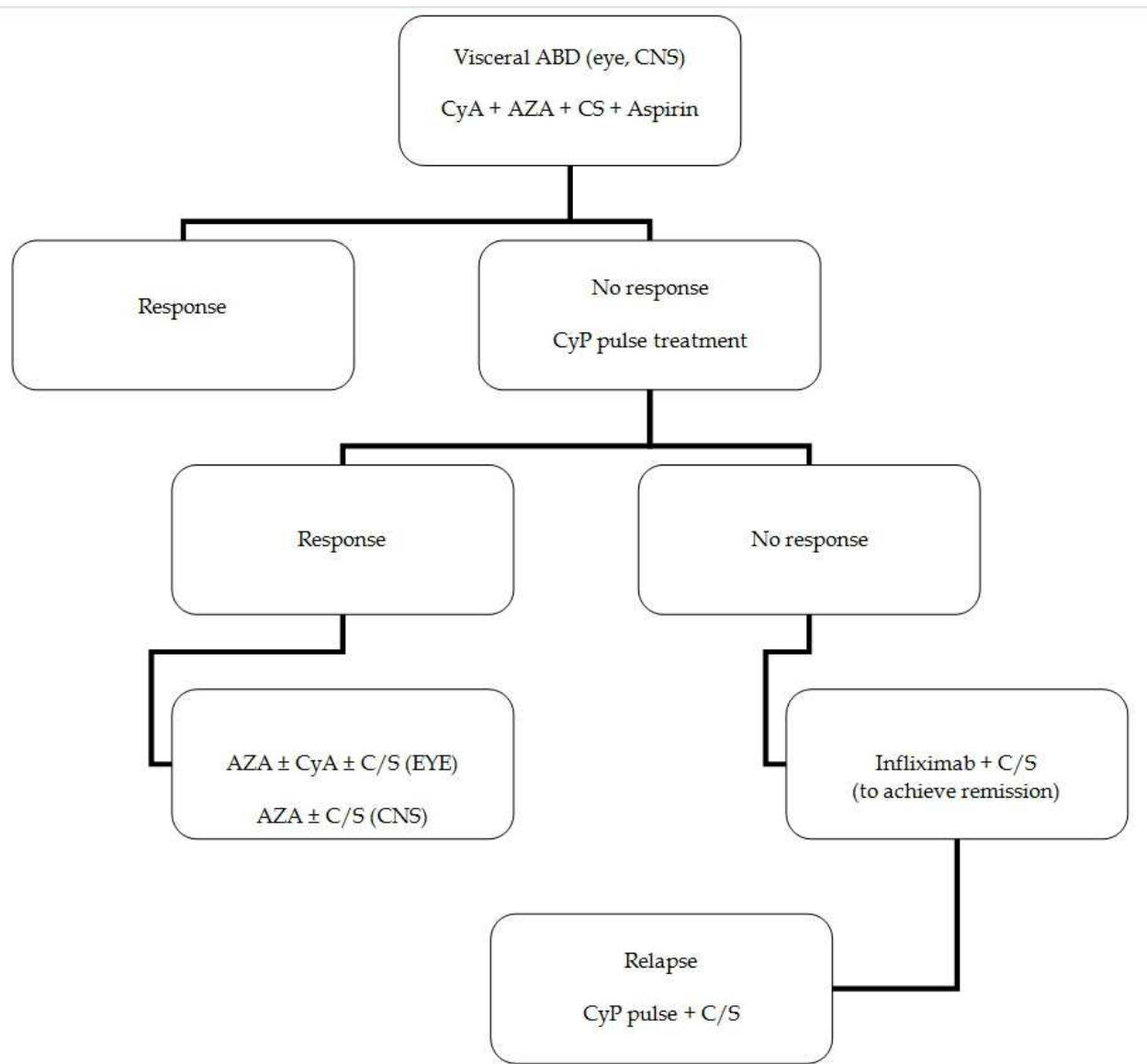
particularly useful in inhibiting T-cell function in ABD. Furthermore, from the pathophysiological models of giant cell arteritis, we know that IFN- γ favor intimal hyperplasia and endothelial dysfunction. (see algorithm 1)

Target	Therapeutic Goals	Current or emerging approach
Interleukin-1	Suppression of acute phase response Inhibition of macrophages	Anakinra, C/S
Interleukin-6	Suppression of acute phase response Inhibition of macrophages	Tocilizumab, C/S
Interferon- γ or interferon- γ inducing genes	Disruption of adaptive T-cell response	acetosalicylic acid, C/S
Metalloproteinases	Inhibition of matrix degradation- -inhibition of expanding remodeling	Cyclosporine
Innate immune responses	Inhibition of nuclear factor k- B	C/S, Interferons
Adaptive immune responses	Inhibition of T cell clones activation	Cyclophosphamide, Cyclosporine
Vascular thrombosis	Inhibition of prothrombotic changes of endothelium	acetosalicylic acid, anticoagulants, immunosuppressants
TNF- α	Inhibition of TNF- α mediated immune response	etanercept, infliximab, adalimumab

Table 3. Therapeutic targets for pathogenic pathways in Adamantiades- Behcet disease and indicative approaches



Algorithm 1. Therapeutic intervention for major vessel involvement in ABD.



Algorithm 2. Therapeutic intervention for visceral involvement (eye, CNS) in ABD.

6. Conclusions

The multi-level pathogenetic process of visceral ABD implies that inflammation control by one single agent is seemingly impossible; indeed, various combinations of immunomodulating agents are proposed. Early administration of immunosuppressive therapy is critical before permanent damage is established. Invasive approaches must be implemented when needed. Urgent intervention is mandatory in life threatening circumstances. Although there is an increasing amount of data from controlled trials regarding specific issues in ABD therapy (targeting management of early ABD and particular disease mechanisms), current guidelines are largely based on the opinion of experts.

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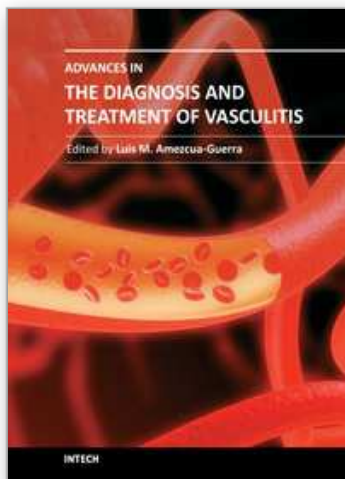
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This book represents the culmination of the efforts of a group of outstanding experts in vasculitis from all over the world, who have endeavored to draw themselves into this volume by keeping both the text and the accompanying figures and tables lucid and memorable. The book provides practical information about the screening approach to vasculitis by laboratory analysis, histopathology and advanced image techniques, current standard treatment along with new and more specific interventions including biologic agents, reparative surgery and experimental therapies, as well as miscellaneous issues such as the extra temporal manifestations of "temporal arteritis" or the diffuse alveolar hemorrhage syndrome. The editor and each of the authors invite you to share this journey by one of the most exciting fields of the medicine, the world of Vasculitis.

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