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MINI-SYMPOSIUM

The role of interferons type I, II and III in myositis: A review

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Abstract

The classification of idiopathic inflammatory myopathies (IIM) is based on clinical, serological and histological criteria. The identification of myositis-specific antibodies has helped to define more homogeneous groups of myositis into four dominant subsets: dermatomyositis (DM), antisynthetase syndrome (ASyS), sporadic inclusion body myositis (sIBM) and immune-mediated necrotising myopathy (IMNM). sIBM and IMNM patients present predominantly with muscle involvement, whereas DM and ASyS patients present additionally with other extramuscular features, such as skin, lung and joints manifestations. Moreover, the pathophysiological mechanisms are distinct between each myositis subsets. Recently, interferon (IFN) pathways have been identified as key players implicated in the pathophysiology of myositis. In DM, the key role of IFN, especially type I IFN, has been supported by the identification of an IFN signature in muscle, blood and skin of DM patients. In addition, DM-specific antibodies are targeting antigens involved in the IFN signalling pathways. The pathogenicity of type I IFN has been demonstrated by the identification of mutations in the IFN pathways leading to genetic diseases, the monogenic interferonopathies. This constitutive activation of IFN signalling pathways induces systemic manifestations such as interstitial lung disease, myositis and skin rashes. Since DM patients share similar features in the context of an acquired activation of the IFN signalling pathways, we may extend underlying concepts of monogenic diseases to acquired interferonopathy such as DM. Conversely, in ASyS, available data suggest a role of type II IFN in blood, muscle and lung. Indeed, transcriptomic analyses highlighted a type II IFN gene expression in ASyS muscle tissue. In sIBM, type II IFN appears to be an important cytokine involved in muscle inflammation mechanisms and potentially linked to myodegenerative features. For IMNM, currently published data are scarce, suggesting a minor implication of type II IFN. This review highlights the involvement of

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different IFN subtypes and their specific molecular mechanisms in each myositis subset.

KEYWORDS

anti-synthetase syndrome, dermatomyositis, interferon, myositis, sporadic inclusion body myositis

1 | INTRODUCTION

Inflammatory idiopathic myopathies (IIM) are a group of rare autoimmune diseases involving muscle tissue but also other organs. They are categorised into four dominant subsets: dermatomyositis (DM), antisynthetase syndrome (ASyS), inclusion body myositis (IBM) and immune-mediated necrotising myopathy (IMNM) based on clinical manifestations, the presence of myositis-specific autoantibodies (MSA) and myopathological findings.

DM and ASyS are systemic diseases associated with skin, joints and/or lung involvement, in addition to myositis (1). DM is associated with a characteristic skin rash (Gottron's papules/signs with heliotrope rash) as well as myopathological findings such as perifascicular atrophy and the presence of perivascular inflammatory infiltrates (2). Both muscular and cutaneous biopsies display vasculopathic features (loss of capillaries, C5b-9 immunoreactivity on capillaries) (3, 4).

ASyS also harbours a perifascicular pathology characterised by perifascicular necrosis, and the majority of patients have interstitial lung disease while this latter feature rarely occurs in DM patients. ASyS is defined by the presence of one of the anti-histidyl RNA-t-synthetase antibodies (5, 6), and DM is associated with a different set of MSAs: anti-Mi2 (7), anti-SAE1 (8), anti-TIF1 γ (9), anti-NXP2 (10) and anti-MDA5 (11–13).

IBM and IMNM are muscle-specific diseases as they are not substantially associated with extramuscular features. Myopathological features of IBM are characterised by the combination of muscle inflammation with endomysial CD8⁺ T cell infiltrates and distinctive degenerative muscle features (14–17). In contrast, inflammatory infiltrates are usually mild in IMNM (18).

Transcriptomic and myopathological studies have demonstrated that interferons (IFNs) play important roles in the pathophysiology of IIM (19), but the IFN subtypes involved and their molecular mechanisms differ in each myositis subset.

Therefore, our objective was to review, clearly layout and summarise the role of IFNs in myositis. First, we describe the IFN family, including induction and signalling pathways, and then we discuss the concept of genetic interferonopathies in order to highlight the pathogenic role of IFNs in myositis. Finally, we focus on the different IFN pathways involved in each myositis subset: DM, ASyS, IBM and IMNM.

2 | INTERFERONS

2.1 | IFN subtypes

IFNs have been discovered in 1957 by Isaacs and Lindenmann in a study entitled 'virus interference', where they identified a new factor able to prevent the virus from entering the cells and called it 'interferon' (20). To date, three types of IFNs have been identified, based on their specific receptors. In humans, the type I IFN (IFN-I) family consists of 13 subtypes of IFN α and one subtype of IFN β , ϵ , κ , ω each (21); type II IFN (IFN-II) includes IFN γ only (22, 23), and type III IFN (IFN-III) includes IFN- λ 1 (IL-29), IFN- λ 2 (IL28A), IFN- λ 3 (IL-28B) and IFN- λ 4 (24).

The IFN-I receptor, the IFN α receptor (IFNAR), is a heterodimeric transmembrane receptor composed of IFNAR1 and IFNAR2 subunits. IFNAR1 is constitutively associated with the tyrosine kinase 2 (TYK2) and IFNAR2 with the Janus kinase 1 (JAK1). IFN γ binds to a distinct cell surface receptor composed of the IFNGR1 subunit, associated with JAK1 and the IFNGR2, constitutively associated with JAK2. Type III IFNs- also signal through a heterodimeric receptor composed of the IFN λ R1 (high-affinity chain) and IL-10R2, a subunit shared with other cytokine receptors (24).

2.2 | IFN-secretion

2.2.1 | IFN-I

In humans, there is no, or only minimal, constitutive secretion of IFN-I. Production of IFN-I is induced by danger signals (pathogen-associated molecular patterns; PAMP) released by viruses or microbial products. These PAMPs can either stimulate membrane receptors localised at the cellular or endosomal membranes (Toll-Like Receptors, TLRs) (25), or cytosolic pattern recognition receptors including nucleotide sensors such as retinoic acid-inducible gene I (RIG-I)-like receptors (RLRs) or DNA sensors to induce IFN production (26).

Endosomal TLRs recognise nucleic acids (double-strand RNA and single-strand RNA or double-strand DNA by TLR3, 7/8, 9, respectively) (27). TLR activation leads to an activation of IFN response factor (IRF) (25, 28) and IFN production (e.g. IRF5 and IRF7 in

plasmacytoid dendritic cells (pDCs) or IRF3 in monocytes) (29).

TLR-independent pathways involve RIG-I, melanoma differentiation-associated protein 5 (MDA5) and stimulator of interferon genes protein (STING). RIG-I and MDA5 recognise endogenous RNA (respectively single and double-strand) (30) and induce IFN by IRF3 and IRF7 activation by a mitochondrial antiviral-signalling protein (MAVS)-dependent mechanism. STING is activated by the synthesis of cyclic GMP-AMP (cGAMP) triggered by endogenous cytosolic double-strand DNA (dsDNA) leading to IFN production by IRF3 activation (31).

While TLR-dependent and -independent pathways are different, both lead to upregulation of IFN-stimulated genes (ISG) including the IFN gene itself.

All cells harbour IFNR and may produce IFNs, but some are specialised cells like pDCs (32) or monocytes, and others like fibroblasts, dendritic cells and epithelial cells (33).

2.2.2 | IFN-II

IFN γ is a cytokine that is primarily produced by cells of the innate immune system, such as monocytes, macrophages and natural killer (NK) cells (23). IFN-II can be induced by cytokines or following activation of pattern recognition receptors. In addition, CD4⁺ Th1 T lymphocytes and CD8⁺ cytotoxic T cells are able to produce IFN γ (34). IFN-II has a key role in macrophage polarisation but also has homeostatic and anti-tumoural effects (23).

2.2.3 | IFN-III

As type IFN-I, also IFN-III can be expressed in hematopoietic cells. Among nonhematopoietic cells, epithelial cells may also produce IFN-III. Both IFN-I and IFN-III induce the same ISG, but IFN-III primarily targets mucosal epithelial cells and protect against viral attacks (24).

2.3 | IFN signalling, IFN-stimulated genes and IFN effects

IFN receptors transduce the signal mainly through JAK-signal transducer and activator of transcription (JAK-STAT) signalling pathways (31, 35). There are four JAK (JAK 1, JAK2, JAK3 and TYK2) and seven different STAT proteins (STAT1, STAT2, STAT3, STAT4, STAT5a, STAT5b and STAT6) in humans (36). JAK activation enhances phosphorylation, dimerisation and nuclear translocation of STAT resulting in the upregulation of hundreds of ISGs (35). Even though IFN-I, -II and -III have different receptors, their activation

leads to induction of mostly the same set of ISGs (22). Nevertheless, some ISG promoters include a specific sequence: interferon-stimulated response element that is only recognised by the ISGF3 complex (STAT1/STAT2/IRF9), which is only secreted by IFN-I but not by IFN-II (37).

ISGs have three major effects. First, ISGs have indirect anti-viral effects. This major function was the first one described (20). Second, IFNs promote immune responses with an indirect anti-viral effect by activation of inflammatory mediators (e.g. MxA, OAS, RNASEL) (31). Third, ISGs are also involved in immune regulation by downregulating the inflammatory responses (e.g., by suppression of cytokine signalling [SOCS]) (31,38).

3 | THE PATHOGENIC ROLE OF IFNS: GENETIC INTERFERONOPATHIES SHARE COMMON FEATURES WITH DM

3.1 | In humans

The monogenic forms of type I interferonopathies represent a recent group of diseases combining the characteristics of Mendelian transmission with a constitutive activation of the IFN-I signalling pathway (39). Aicardi-Goutières syndromes (AGS), chilblain lupus in the juvenile form, STING-associated vasculopathy with onset in infancy (SAVI), PSMB8-related diseases (like Chronic Atypical Neutrophilic Dermatitis with Lipodystrophy and Elevated temperature [CANDLE]) and Singleton-Merten syndrome (SMS) (39, 40), all of which basically occurring in children, are the main syndromes related to constitutive activation of IFN-I signalling.

Five different genetic mechanisms leading to an overproduction of IFNs have been identified, so far (41). The first one is an abnormal accumulation of nucleic acids linked to a nuclease defect. For example, loss of function mutations in the exonuclease TREX1 (which normally downregulates STING activation), in the RNase H (2A, 2B and 2C) or the RNase SAMHD1, have been linked with AGS and familial chilblain lupus (39, 41). The second mechanism is a constitutive activation or enhanced sensitivity of an innate immune sensor or adaptive molecule. Gain of function mutations in *IFIH1* (coding for MDA5), in *DDX58* (coding for RIG-I) or *TMEM173* (coding for STING) were reported in AGS, SMS and SAVI, respectively (40, 42). A defect of IFN-I regulation can lead to an interferonopathy. For example, loss of function mutations in *ISG15* that normally decreases ISG activity by glycation (43) lead to interferonopathy (*ADARI* mutation found in some AGS). Finally, loss of function mutations in immunoproteasome subunits (e.g. *PSMB8* for $\beta 5i$ subunit) impair its multicatalytic function causing cellular stress leading to sustained IFN-I production (41, 44).

TABLE 1 Overlapping features between DM and genetic interferonopathies

	AGS	Familial lupus	SAVI	CANDLE	Singleton-merten
Gene/pathway	<i>TREX1, RNASEH2, SAMHD1, ADAR, IFIH1</i>	<i>TREX1, SAMHD1</i>	<i>TMEM173</i>	<i>PSMB8</i>	<i>IFIH1 DDX58</i>
Cutaneous manifestations					
Chilblain-like lesions	+	+	+	+	+
Digital necrosis	+	+	+	+	–
Panniculitis	Occasional	–	–	–	+
Heliotope rash	–	–	–	+	–
Telangiectasia			+		
Interstitial lung disease	–	–	+	–	–
Myositis	–	–	+	+	–
Arthritis	–	–	+	+	–
Constitutional fever	+	–	+	–	+

Abbreviations: AGS, Aicardi Goutières syndrome, SAVI, sting-associated vasculopathy with onset in infancy, CANDLE, Chronic Atypical Neutrophilic Dermatitis with Lipodystrophy and Elevated temperature.

Several clinical features observed in genetic interferonopathies are very similar to those observed in DM (Table 1).

3.1.1 | Skin manifestations

Chilblain and ulceration are present in AGS, and distal erythema and ulceration are observed in one-third of patients with CANDLE (44). Erythematous oedema of the eyelids or panniculitis may be observed in genetic interferonopathies, which are also features encountered in some forms of DM. The genetic interferonopathy associated with the cutaneous phenotype having the most similarities with anti-MDA5⁺ DM is SAVI. Indeed, telangiectatic erythema of the light-exposed areas and skin ulcerations are a constant finding. Patients may also have distal ulcerative lesions. These lesions reflect underlying cutaneous vasculopathy and share similarities with the type of skin vasculopathy encountered in many patients with anti-MDA5⁺ DM (45).

3.1.2 | Muscle manifestations

Muscle inflammation has also been reported in genetic interferonopathies. Myositis may be observed in 80% and 25% of patients with CANDLE and SAVI, respectively (44, 45) although the precise characteristics may differ gradually and qualitatively.

Lung and joint involvements may occur both in DM and genetic interferonopathies. SAVI patients frequently develop interstitial lung disease and non-erosive arthritis that may be severe (45). CANDLE patients may also have non-erosive polyarthritis (44). Among DM, interstitial lung disease and synovitis are features commonly observed in anti-MDA5⁺ DM (46).

3.2 | Animal models

In vivo models of genetic interferonopathies (loss or gain of function mutations) confirm the presence of overlapping features with DM.

3.2.1 | Skin manifestations

Cutaneous manifestations have been demonstrated in *Trex1*-deficient mice, which develop inflammation of the skin (47–49). In normal condition, *Trex1* degrades and metabolises endogenous DNA and is a specific negative regulator of STING-dependent signalling. *Trex1*^{−/−} mice develop specific multiorgan inflammation with T cell infiltrates in the skin. STING antagonism attenuates pathological features of the autoinflammatory disease in these mice (49). Skin changes with vasculopathy were also demonstrated in a mouse model with a *TMEM173* gain of function mutation that leads to the activation of STING. Two animal models (knock-in with a V154M substitution and N153S) have been developed to reproduce SAVI (50). In the N153S model, mice developed skin ulceration with vasculopathy and interstitial lung disease. In the V154M knock-in model, mice also developed interstitial pulmonary disease reproducing the phenotype of SAVI, similar to that in anti-MDA5⁺ DM patients.

3.2.2 | Muscle manifestations

Myositis and myocarditis have been demonstrated in some animal models. In *Trex1*^{−/−} mice, histological analysis showed T and B cells in striated and cardiac muscles (48). In *IFNGR*^{+/-} mice with knockout of the IFN pathways regulator SOCS, histological analysis showed massive

infiltration by T cells, macrophages and eosinophils in the skeletal muscle and in the heart (51).

4 | INTERFERON PATHWAY ACTIVATION IN DM

4.1 | IFN pathway and DM-specific autoantibodies

Antigenic targets of DM-specific autoantibodies are involved in the IFN pathways. The most striking example is the anti-MDA5 antibody (occurring in 10% of Caucasian DM patients (1)) associated with a subset of amyopathic DM with potentially severe interstitial lung disease (46). As mentioned above, MDA5 is an RLR playing a key role in IFN secretion (30).

Anti-TIF1 γ antibodies (present in 40% of DM (52)) are associated with DM and a high risk of malignancy in adults (53). TIF1 γ (TRIM33) plays a critical role in downregulating IFN- β production by macrophages (54). TRIM33 is involved in sumoylation (55), a process involved in post-translational modification by activation of small-ubiquitin-like modifiers (SUMO) (8). Sumoylation has an important role in IFN regulation. In the animal model, sumoylation deficiency causes increased levels of IFN- β in skin and muscle tissues (55).

Small ubiquitin-like modifier activating enzyme (SAE) activates small-ubiquitin-like modifier-1 (8) in the process of sumoylation. SAE is targeted by DM-specific autoantibodies in 10% of patients (1). Mi-2/NuRD (56) complex and NXP-2 (57) are both inhibitors of SUMO. Anti-Mi-2 and anti-NXP-2 autoantibodies are observed in 30-35% and 20% of DM patients, respectively (1).

4.2 | IFN pathway is activated in DM

In 1986, Isenberg et al. were the first to report the presence of IFN-I and IFN-II in muscle biopsies of patients with inflammatory myopathies (58).

4.2.1 | Muscle tissue

First evidence of the involvement of an IFN pathway in DM was demonstrated by Emslie-Smith et al. in juvenile DM (59). The authors showed an upregulation of ISG in juvenile DM muscle biopsies. Later Greenberg et al. reported on the activation of IFN pathways in muscle biopsies from adult DM patients (60). This IFN signature was not observed in other subsets of inflammatory myopathies such as polymyositis (60). To date, several studies confirm the presence of an IFN signature in the blood, muscle and skin of DM patients (43, 60–65).

Muscle transcriptomic analyses showed a similarly upregulated IFN signature in all serological subsets of DM (19) including muscle biopsies from amyopathic anti-MDA5 DM patients (66).

Detection of IFN-stimulated proteins by immunostaining in DM muscle biopsies have permitted to demonstrate the translation of some ISG. Sarcoplasmic expression of ISG15, RIG-I and Myxovirus resistance A (MxA) were observed in the perifascicular area of DM muscle biopsies (43, 67–70). Of note, based on these findings, MxA immunostaining is now considered the most specific myopathological diagnostic marker of DM (68, 69).

4.2.2 | Skin tissue

Of the 25 most highly overexpressed genes in the skin of DM patients, 21 were ISGs (3). Moreover, the intensity of the IFN signature is mainly correlated with serum levels of IFN-I (IFN- β) and not with IFN-II (IFN γ) (3). A similar topography of damage is seen in keratinocytes and myofibres in DM (71). Ono et al. showed a correlation between IFN signature in blood and skin and vasculopathy features. Furthermore, they confirmed IFN pathway involvement at a protein level with MxA staining in the blood vessels and interstitial fibroblasts of DM patients. Proteomic analysis of skin biopsies derived from DM patients showed an IFN signature leading to a reduction of collagen by dermal fibroblasts (72).

4.2.3 | Blood

IFN pathway activation is not restricted to the muscular and skin compartments. IFN signature has been found in the peripheral mononuclear blood cells of DM patients (61, 64, 73).

Among the highly overexpressed ISGs, there are common genes overexpressed in DM and interferonopathies such as *IFI27*, *IFI44L*, *IFIT1*, *ISG15*, *RSAD2* and *SIGLEC1* (74). It should be noted that DM with anti-MDA5 antibodies, which has a similar phenotype with SAVI, also has a very similar IFN signature. Nevertheless, the intensity of the IFN signature is less than in genetic interferonopathies (75).

Moreover, the level of IFN pathway activation (i.e. IFN score) correlates with disease activity (61, 73, 76, 77). Accordingly, blood IFN- β levels correlate with skin disease activity (62, 63), and IFN-I α levels correlate with disease severity in anti-MDA5⁺ DM (78, 79).

4.3 | IFN production in DM

The triggers of IFN pathway induction as well as the main sources of IFNs in dermatomyositis remain unclear.

4.3.1 | Immune cells and IFN production

pDCs are known to produce large amounts of IFN-I (32). IFN scores correlate with the pDCs density in the skeletal muscles and the skin of adult and juvenile DM (60, 80) (Figure 1). These cells can be found in the perifascicular and perivascular areas of muscle and skin biopsies (60, 81–83).

4.3.2 | Muscle cells

In muscle tissue, RIG-I activation is responsible for an IFN- β secretion by human myotubes, resulting in an autocrine loop of activation (67). It has been shown that myogenic precursor cells from juvenile DM muscle produce higher levels of IFNs compared to those of healthy controls (84). Previous reports also showed that regenerating muscle fibres in addition to pDCs may produce IFNs in DM (85).

4.3.3 | Keratinocytes

Keratinocytes can also produce IFN-I and IFN-III in DM *in vitro* (86). The IFN secretion in skin is associated

with the recruitment of CXCR3⁺ lymphocytes and the expression of two ISGs, CXCL9 and CXCL10, correlating with capillary loss in juvenile DM (87). Another study showed a key role of immunoproteasome subunit PSMB9 that is up-regulated in DM skin by proteomic analysis and could decrease collagen secretion when treated by IFN *in vitro* (72).

4.4 | IFNs induce muscle damage

Ladislau et al. showed a pathogenic role of IFN-I by impairing myoblast differentiation and by inducing atrophy of myotubes *in vitro* (88). Myotubes demonstrated an upregulation of two genes involved in the atrophy process, *FBXO32* (*ATROGIN1*) and *TRIM63* (*MURF1*).

IFN-I pathway activation also disrupts the capillary network *in vitro*. Interestingly, the administration of JAK inhibitors improve IFN-I-induced damage. Additionally, Meyer et al. showed that IFN- β -induced mitochondrial dysfunction contributes to the impairment of muscle function and persistent inflammation in DM (89).

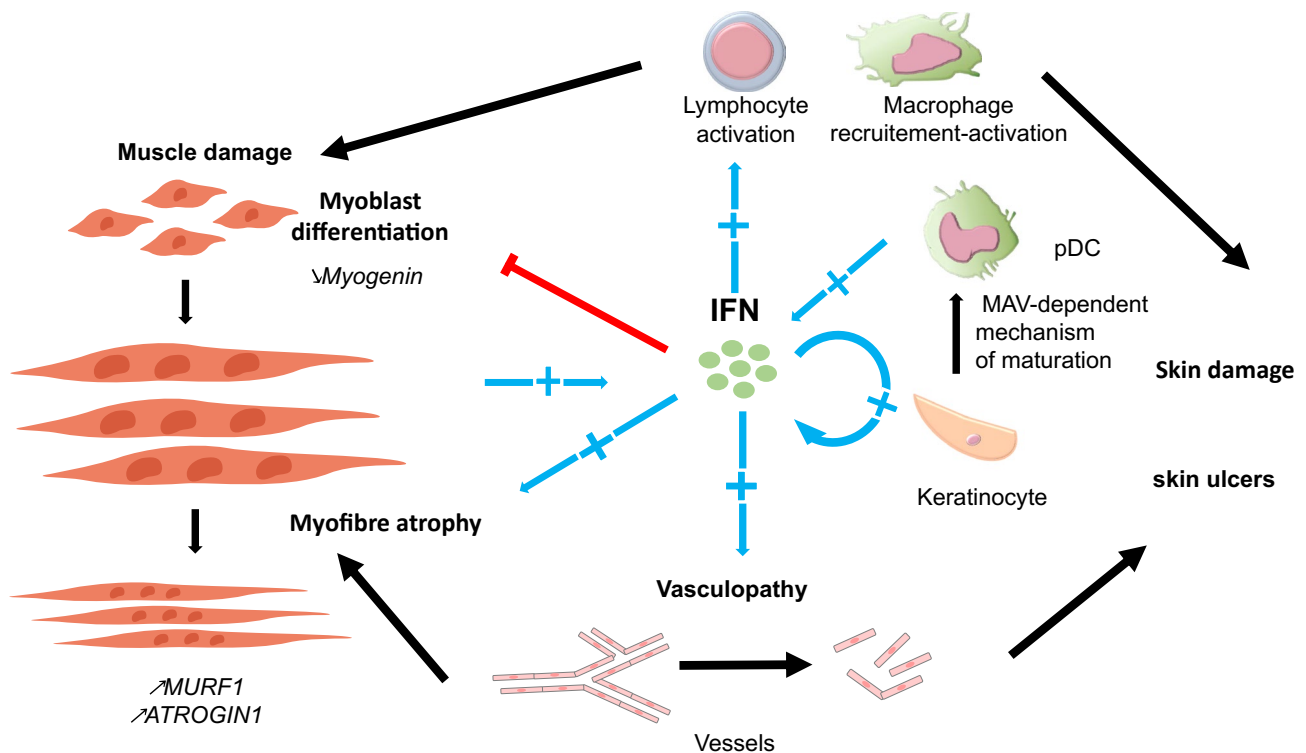


FIGURE 1 Pathologic effects of interferon in dermatomyositis. *In vitro*, type I interferon (IFN-I) induces muscle damage. IFN-I inhibits myoblast differentiation and decreases *myogenin* expression. IFN-I induces myofibre atrophy and overexpression of muscle atrophy genes, *ATROGIN1* (*FBXO32*) and *TRIM63* (*MURF1*). *In vitro*, IFN-I impairs angiogenesis of endothelial cells and disrupts the vascular network. Vasculopathy in dermatomyositis has been linked to indirect muscle pathology features and skin damage like ulcers. *In vitro*, IFN-I-stimulated keratinocytes produce IFN-I and IFN-III and activate plasmacytoid dendritic cells (pDC) as IFN-I producers. pDCs are present in perifascicular and perivascular areas in muscle and skin tissues, but macrophages and muscle fibres may also participate in IFN-I production. IFNs lead to the recruitment and activation of both innate and adaptive immune cells

Altogether, these data highlight the key role of the IFN pathways, especially IFN-I, in the pathophysiology of DM.

5 | INTERFERON PATHWAY ACTIVATION IN ASyS

DM and ASyS have previously been considered as part of the same disease spectrum as both conditions may present with similar skin features and a perifascicular pathology. Indeed, Mozaffar and Pestronk were the first to describe the myopathological features in anti-Jo-1⁺ ASyS patients demonstrating septal inflammation, and myofibre lesions restricted to the perifascicular area referred to as an “immune myopathy with perimysial pathology” (IMPP) (90). More recent studies, have demonstrated that ASyS muscle biopsies display perifascicular necrosis (91–93), while DM is associated with perifascicular atrophy (91), suggesting distinctly separate pathomechanisms.

5.1 | IFN-II pathway in muscle of ASyS

Overexpression of IFN-I related proteins, such as MxA, ISG15 and RIG-I, are absent in ASyS (69, 94), and perifascicular MHC class II (MHC-II) expression is prominently observed in ASyS but not in DM (93, 95). Interestingly, in muscle biopsies from patients with ASyS, CD8⁺ T cells are located in close vicinity to MHC class-II⁺ myofibres, suggesting the involvement of the IFN-II pathway (96). Along that line, *in vitro* studies showed that IFN-II can upregulate the MHC-II expression on primary human myoblasts (97, 98).

Accordingly, RNA sequencing data from a large number of myositis muscle biopsies showed prominent IFN-II activation in patients with ASyS (19), and conversely, IFN-I-induced genes were only mildly expressed in ASyS (19).

5.2 | IFN-II pathway in lungs of ASyS patients

The presence of histidine RNA-Synthetase(HisRS)-reactive CD4⁺ T cells in the bronchioalveolar lavage fluids as reported in ASyS patients (99). Along that line, shared oligoclonal CD4⁺ and CD8⁺ T cells were reported in the lungs and skeletal muscles associated with different myositis subsets (100).

In the bronchioalveolar lavage fluids, these cells display a Th1 phenotype and produce high levels of IFN γ (99). In addition, these Th1 cells are characterised by the expression of C-C chemokine receptor type 5 (CCR5), involved in T cell homing. Interestingly, it was demonstrated that HisRS has chemotactic properties inducing tissue homing of CCR5-positive cells (lymphocytes and antigen-presenting cells) (101).

5.3 | IFN pathways in blood of ASyS patients

In the peripheral blood of ASyS, the presence of HisRS-reactive CD4⁺ T cells was also reported (99) even though the Th1 involvement was less significant in the lungs. Along that line, increased levels of IFN γ -induced chemokines CXCL9 and CXCL10 are observed in the serum of patients with anti-Jo-1 antibodies (102).

Besides, the addition of ASyS patients' serum induces the production of IFN in the blood from healthy donors (64). Indeed, it has been demonstrated that anti-Jo-1 antibody complexed with necrotic cells is able to stimulate peripheral blood mononuclear cells (PBMCs) from healthy donors to produce IFN-I α (103, 104).

5.4 | Animal model of HisRS-induced myositis-associated IFN pathway

Mouse models have been developed and highlight the role of IFN pathway in ASyS (105).

Experimental autoimmune myositis induced by a single intramuscular immunisation with HisRS spontaneously resolved in few weeks. Persistent muscle inflammation and prolonged anti-Jo-1 production were observed only in mice stimulated by HisRS in the presence of a TLR7/8 agonist, but absent in IFN $\alpha\beta$ R-null mice (105).

Altogether these data support the role of IFNs in ASyS. Myopathological findings and transcriptomic studies suggest the preferential involvement of IFN-II, rather than of IFN-I.

6 | INTERFERON PATHWAY ACTIVATION IN sIBM

Sporadic inclusion body myositis (sIBM) is a slowly progressive skeletal muscle disease with degenerative features including rimmed vacuoles and related myonuclear degeneration, mitochondrial pathology and cytoplasmic protein aggregates within myofibres. Novel muscle histochemical degenerative biomarkers, such as p62, LC3 and TDP43, are involved in unfolded protein response, endoplasmic reticulum stress and altered autophagy. One of the pathological hallmarks of sIBM is the predominance of endomysial CD8⁺ T cells (and macrophages) surrounding and occasionally invading MHC-I-positive myofibres.

6.1 | IFN-II pathway in muscle tissue of sIBM

CD8⁺ T cells in sIBM muscles show a highly differentiated effector phenotype (106–108). Greenberg et al. found that muscle fibres are invaded by CD8⁺CD57⁺T

cells (106). The proportion of invaded myofibres correlated positively with the degree of muscle weakness (106). Microarray data from a large series of muscle biopsies identified killer cell lectin-like receptor G1 (KLRG1) as a biomarker of these subpopulations (107). Accordingly, IFN-II gene expression is higher in myofibres invaded by CD8⁺ T cells compared to expression in non-invaded myofibres (109). These cytotoxic CD8⁺ T cells produce high levels of cytotoxic molecules such as perforin and granzymes (110, 111). Along that line, Th-1 cytokines and chemokines such as IFN-II, CXCL9, CCL5 and IL32 are increased in sIBM muscle tissue (107).

Analysis of proteomic profiling from muscle biopsies showed that 39% of upregulated proteins are involved in IFN-I or mixed IFN-I and IFN-II pathways (112).

Accordingly, immunohistochemistry studies confirmed the presence of IFN-related proteins such as ISG15 and IRF8 (112). Moreover, high expressions of CD74 and STAT1, as key molecules of IFN macrophage activation, were detected in sIBM (112). It was also demonstrated that IFN-II can upregulate cell surface MHC-I expression on myoblasts (113).

6.2 | IFN-II pathway in blood of sIBM

Expansion of highly differentiated T cells was also detected in blood of sIBM patients. sIBM patients have increased levels of differentiated populations of T cells such as CD8⁺CD28null (114), CD8⁺Tbet⁺ (115), CD8⁺CD57⁺ (106) and CD8⁺KLRG1⁺ T cells and high levels of Th1 chemokines and cytokines (CXCL9, CXCL10 and IL-12) (114, 116). This specific T-cell population is prone to produce higher levels of IFN-II in sIBM compared to levels in healthy controls (114, 116). Moreover, this population is clonally expanded and shares the same oligoclonal repertoire that muscle-infiltrating CD8⁺ T cells have (108). Greenberg et al. found that 58% of sIBM patients have abnormal populations of circulating large granular lymphocytes meeting diagnostic criteria of T cell large granular lymphocytic leukaemia (LGLL) (106). Highly differentiated clonal T cells in LGLL are resistant to apoptosis (117).

6.3 | IFN-II pathway and muscle degeneration

A growing body of evidence suggests that causality flows from autoimmunity to degeneration (15, 16). Abnormal sarcoplasmic accumulation of β -amyloid may also be explained by the effect of inflammatory cytokines on muscle cells. Beta-amyloid production may be induced in human primary myotubes by exposure to IL-1 β and IFN-II (118). However, overexpression of β -amyloid precursor protein in mouse muscle did not lead to any muscle inflammation (119).

Thus, analysis of the link between IFNs and protein aggregation in muscle cells is crucial to understand the pathogenesis of sIBM, and the immunoproteasome is a key element (Figure 2). Indeed, several studies showed that elements of the immunoproteasome are detected in muscle tissue of sIBM (107, 113). The immunoproteasome is an alternative isoform of the constitutive proteasome induced by inflammatory cytokines, in particular by IFN γ . The main role of the immunoproteasome is to process antigens for presentation on MHC-I molecules to CD8⁺ T cells (120). Specific inhibitors of the immunoproteasome reduced surface expression of MHC-I in myoblasts induced by IFN γ *in vitro* (113). Nakajo-Nishimura syndrome (NNS) is a rare autosomal recessive autoinflammatory disorder caused by a homozygous mutation in *PSMB8* that encodes the immunoproteasome subunit β 5i. NNS is clinically characterised by a pernio-like skin rash, periodic fever and myositis (121). Interestingly, myopathological analysis of small case series of NNS shows sarcolemmal MHC-I expression, CD4⁺ and CD8⁺ T cell infiltrations, p62 and TDP43 aggregations and rimmed vacuoles in myofibres (122) similar to certain pathological features of sIBM muscle. Myeloid cell lines derived from pluripotent stem cell of NNS patients show dysfunction of proteasome activity and overproduction of inflammatory cytokines and chemokines such as IL-6, CCL2 and CXCL10 with upregulation of reactive oxygen species under IFN γ and TNF α stimulation (123).

Altogether these data support the role of IFNs, in particular of IFN-II, in sIBM. IFN-II may be secreted by specific highly differentiated CD8⁺ T cells that are detectable in muscle tissue and blood of sIBM patients. Myodegenerative features in sIBM may be related to IFN-II as well.

6.4 | IFN and IMNM

IMNM represent another IIM subset associated with distinct clinical, serological and histological features. Different from sIBM, the presence of certain pathognomonic MSAs (anti-SRP or anti-HMGCR antibodies) are a major feature of the disease. Recent data point towards a direct pathogenic effect of these autoantibodies, potentially through the activation of the classical complement cascade (124).

Although the IFN signalling pathway does not seem to be the central mechanism of IMNM, transcriptomic data showed a modest overexpression of IFN-inducible genes in muscle tissue compared to those in other IIM (19). The IFN signature appears more driven by IFN-II rather than IFN-I with *IFI30* as the most upregulated IFN-inducible gene (19). This predominant IFN-II signature has been inconsistently identified with *q*RT-PCR assays (96, 116, 125).

Indirect arguments are consistent with a potential role of IFN-II. Although the inflammatory infiltrate is usually scarce in the muscle biopsies of IMNM,

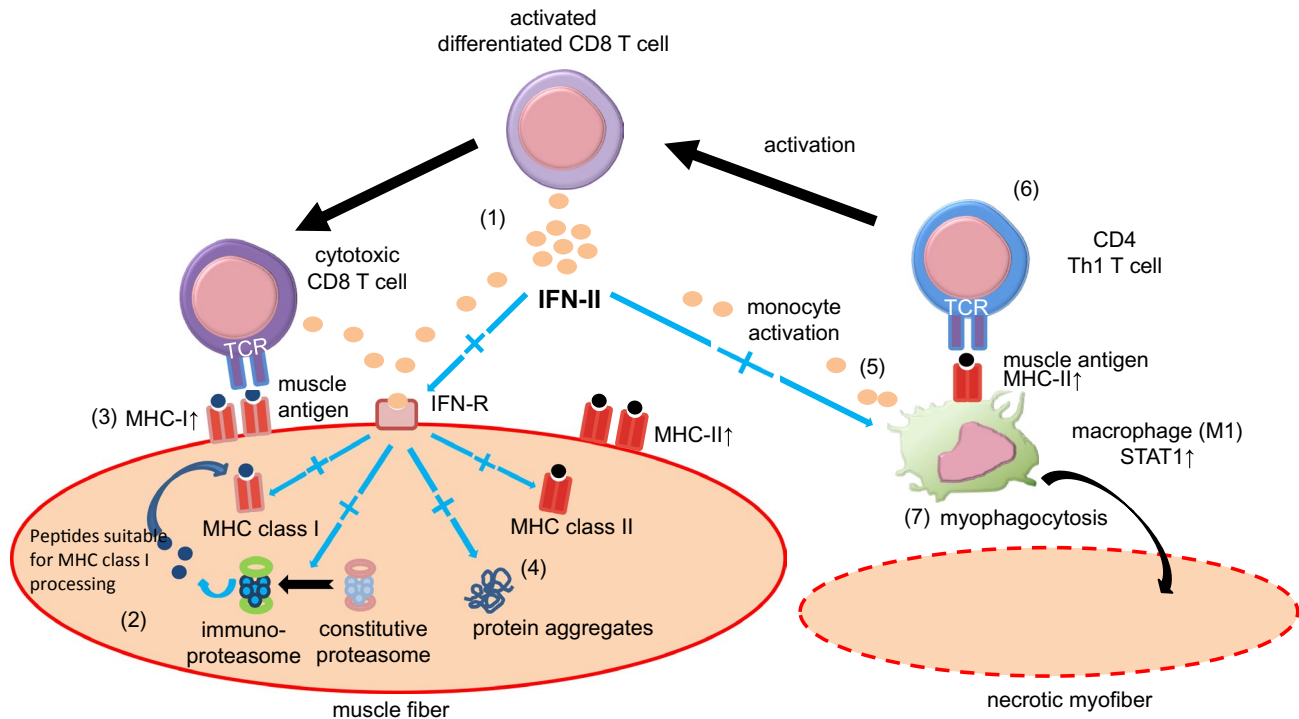


FIGURE 2 IFNs and IFN-II in Inclusion body myositis. (1) Highly differentiated T cells secrete IFN-II. (2) IFN-II stimulates the expression of muscle antigens on MHC class I via the immunoproteasome in muscle fibres (3). Cytotoxic CD8⁺ T cells can be stimulated by the complex of muscle antigens and MHC class I and produce IFN-II (4). IFN-II may induce protein aggregates (5). IFN-II also activates macrophages and MHC class II expression with muscle autoantigen (6). Activated macrophages stimulate CD4⁺ Th1 T cells and (7) are involved in myophagocytosis of necrotic muscle fibres

a decent amount of macrophages and T cells have been described in skeletal muscle tissues independent from myophagocytoses and necrosis. Macrophages harboured an M1 phenotype in the context of Th1 environment with upregulation of *STAT1*, *IFNG*, *TNFA* and *IL12* (126). Double-positive macrophages CD68⁺STAT1⁺ and CD68⁺NOS2⁺ were identified in muscle tissues by immunostainings confirming an M1-polarised macrophage profile (125).

Moreover, mild to moderate sarcolemmal MHC-I expression but the absence of sarcolemmal MHC class II is a characteristic myopathological feature observed in IMNM. As mentioned above, IFN-II induces MHC-I cell surface expression on myoblasts (109).

Only a few studies evaluated the IFN signalling pathway in IMNM suggesting a modest role of IFN, especially by IFN-II.

6.5 | Conclusion

Although IIMs are a heterogeneous group of autoimmune diseases, the IFN cytokine family plays an important role in its pathophysiology. DM is the subset where IFNs, and especially IFN-I, have been highlighted as key pathogenic cytokines leading to consider DM as an acquired interferonopathy in analogy to the monogenic interferonopathies. In ASyS, both IFN-I and II have been involved but transcriptomic analyses and pathological data showed a

predominant role of IFN-II. The key role of terminally differentiated CD8⁺ T cells in sIBM supports IFN-II as an important cytokine, not only involved in muscle inflammation but also in muscle degenerative features, presumably mediated via the immunoproteasome. The role of IFN appears minor in IMNM compared to that in other myositis subsets, yet IFN-II remains important for macrophages signalling.

CONFLICT OF INTEREST

The authors have declared no conflicts of interest for this article.

DATA AVAILABILITY STATEMENT

Not applicable.

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