



EDUCATIONAL OBJECTIVE: Readers will consider autoinflammatory syndromes in the differential diagnosis of recurrent fever

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Autoinflammatory syndromes: Fever is not always a sign of infection

ABSTRACT

Autoinflammatory syndromes are a newly understood group of conditions characterized by recurrent episodes of fever, rash, and serositis. Generalists and specialists should know about and consider these syndromes in the differential diagnosis of recurrent fever. This article reviews the genetics, pathophysiology, clinical presentation, and treatment of several of these relatively recently discovered diseases.

KEY POINTS

In many of the autoinflammatory syndromes, genetic abnormalities and consequent disordered regulation of the innate immune system lead to overactivity of pro-inflammatory cytokines and subsequent inflammatory symptoms.

Early recognition and treatment with immunoregulatory agents may improve quality of life and reduce the risk of disease sequelae.

Abnormal regulation of the innate inflammatory pathway has also been implicated in the pathogenesis of conditions as phenotypically diverse as gout, type 2 diabetes, atherosclerosis, and epilepsy.

A 22-YEAR-OLD MAN of Turkish ancestry presents to your office for an urgent visit. One day before the visit, he abruptly developed a fever with temperatures as high as 104°F (40°C), abdominal pain, joint pain, and a red rash on the lower right leg. He has no cough, nasal congestion, rhinorrhea, ear or eye pain, oral ulcers, vomiting, or diarrhea. After reviewing his chart, it becomes apparent that he has experienced similar intermittent, random, and self-limited episodes over the last 4 years.

On examination, he is febrile with diffuse abdominal tenderness and guarding. Bowel sounds are normal, and there is no rebound. The left knee is slightly swollen and limited in range of motion, and there is a large, non-palpable, blanching, erythematous lesion over the anterior lower leg.

While pondering diagnostic possibilities, you remember reading about autoinflammatory syndromes that result in recurrent episodes of fever and multisystemic inflammatory symptoms but cannot recall the evaluation and therapeutic options for these conditions.

These syndromes pose diagnostic challenges for physicians. Although these conditions are uncommon and may mimic malignancy or infection, they should be considered in patients who have recurrent febrile illness. While the autoinflammatory syndrome of familial Mediterranean fever (FMF), the diagnosis in the case above, is well known, our growing understanding of genetics and the immune system has unearthed a growing number of autoinflammatory syndromes.

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Abbreviations used in this article

CAPS	—cryopyrin-associated periodic syndromes
DIRA	—deficiency of interleukin 1 receptor antagonist
FCAS	—familial cold autoinflammatory syndrome
FMF	—familial Mediterranean fever
HIDS	—hyperimmunoglobulin D syndrome
HLA	—human leukocyte antigen
HMG-CoA reductase	—3-hydroxy-3-methyl-glutaryl-CoA reductase
IgD	—immunoglobulin D
IgG	—immunoglobulin G
IL-1	—interleukin 1
IL-6	—interleukin 6
MVK	—mevalonate kinase
MWS	—Muckle-Wells syndrome
NOMID	—neonatal-onset multisystemic inflammatory disorder
NLRP3	— NOD-like receptor family, pyrin domain
PAPA	—pyogenic arthritis, pyoderma gangrenosum, and acne (syndrome)
PFAPA	—periodic fever, aphthous stomatitis, pharyngitis, and adenopathy (syndrome)
TNF	—tumor necrosis factor
TRAPS	—tumor necrosis factor receptor-associated periodic syndrome

A GENETICALLY DIVERSE BUT CLINICALLY SIMILAR GROUP OF CONDITIONS

The autoinflammatory syndromes are a group of genetically diverse but clinically similar conditions characterized by recurrent attacks of fever, rash, serositis, lymphadenopathy, and musculoskeletal involvement. This category of diseases is rapidly expanding and was built on the discovery of the genetics behind FMF, hyperimmunoglobulin D syndrome (HIDS), tumor necrosis factor receptor-associated periodic syndrome (TRAPS), and the cryopyrin-associated periodic syndromes (CAPS). More recent additions to the list include Blau syndrome and the syndrome of pyogenic arthritis, pyoderma gangrenosum, and acne (PAPA).

In autoinflammatory syndromes, genetic mutations lead to dysregulation of the innate

immune system and to episodic manifestations of systemic inflammation. Many patients have first- or second-degree relatives with similar symptoms, reflecting the genetic abnormalities underlying this class of conditions. Unlike patients with other rheumatic diseases, patients with autoinflammatory diseases do not have autoreactive T lymphocytes, and they typically lack pathogenic autoantibodies.

The characterization of genetic autoinflammatory syndromes shows the importance of a well-regulated innate immune system and sheds light on the role of the innate immune system in common medical conditions such as gout and type 2 diabetes (see below).

THE INNATE IMMUNE SYSTEM: OUR FIRST LINE OF DEFENSE

The innate immune system is the first line of immune defense. It is evolutionarily conserved. Unlike the adaptive immune response, the innate immune response is not antigen-specific, and its activation does not produce a memory response. Generally speaking, it is composed of certain white blood cells (neutrophils, dendritic cells, macrophages, natural killer cells), proinflammatory signaling proteins (cytokines), and the complement system. Interleukin 1 (IL-1), interleukin 6 (IL-6), and tumor necrosis factor (TNF) alpha are the critical and most potent proinflammatory cytokines of the innate immune system.

To date, nearly all mutations that have been linked to the autoinflammatory syndromes disrupt regulation of inflammatory signaling within the innate immune system. This disruption generates a proinflammatory state, often leading to a final common pathway ending with activation of the inflammasome.

The inflammasome is a complex of distinct proteins which, when brought together, serve to convert inactive prointerleukin 1 beta to the active proinflammatory cytokine IL-1 beta.¹ Formation of the inflammasome can be mediated by multiple different signals including microbial products, endogenously produced substances such as cholesterol and uric acid, or by proinflammatory cytokines and chemokines (FIGURE 1).

The innate immune system in monogenic autoinflammatory syndromes

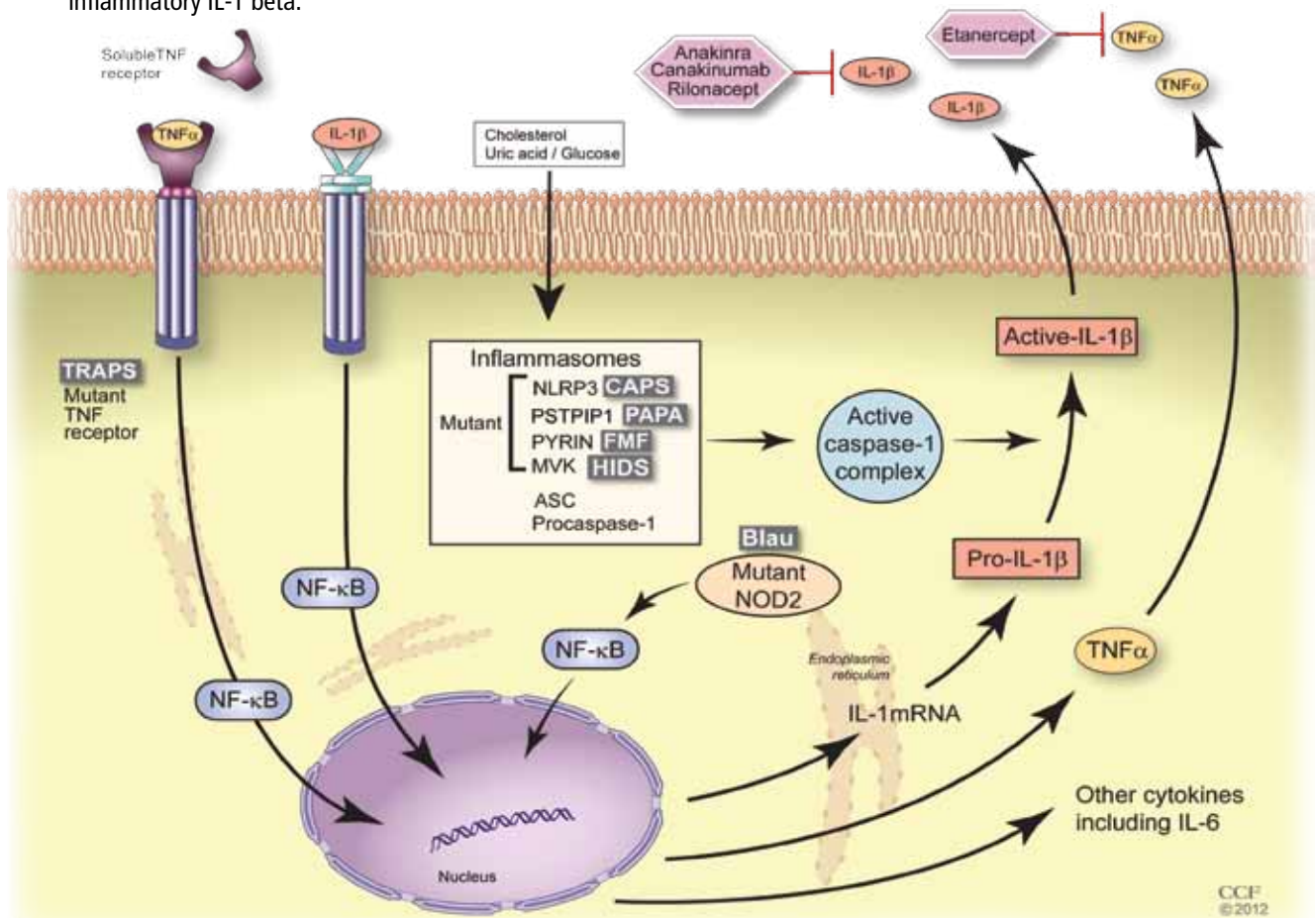
In autoinflammatory syndromes, genetic mutations lead to dysregulation of the innate immune system and to episodic manifestations of systemic inflammation.

CAPS, FMF, HIDS, AND PAPA

In cryopyrin-associated periodic syndromes (CAPS), familial Mediterranean fever (FMF), hyperimmunoglobulin D syndrome (HIDS), and the syndrome of pyogenic arthritis, pyoderma gangrenosum, and acne (PAPA), mutations stimulate inflammasomes, resulting in caspase-1 stimulation and the formation of pro-inflammatory IL-1 beta.

Drug therapy

Anakinra (Kineret), canakinumab (Ilaris), and rilonacept (Arcalyst) inhibit interleukin 1 (IL-1) and have shown efficacy in treating Blau syndrome, CAPS, and PAPA. Titrating doses of the TNF antagonist etanercept (Enbrel) may be efficacious in TRAPS.



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TRAPS

In tumor necrosis factor receptor-associated periodic syndrome (TRAPS), mutant tumor necrosis factor (TNF) receptor is sequestered in the endoplasmic reticulum, leading to production of NF-kappa B and the transcription of proinflammatory cytokines, including IL-1 beta.

Blau

In Blau syndrome, mutant NOD2 leads to increased production of NF-kappa B and to the transcription of proinflammatory cytokines, including IL-1 beta.

FIGURE 1

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FAMILIAL MEDITERRANEAN FEVER

FMF is the most common and well characterized autoinflammatory syndrome. Described in 1949, its etiology was not understood until the genetic mutation that causes it was discovered in 1997.²⁻⁴

The Mediterranean fever gene *MEFV* encodes pyrin, a protein with an important role in controlling IL-1 production. Mutations in *MEFV* affect pyrin-mediated regulation, and IL-1 production increases.

Classically, FMF is described as autosomal recessive, although many patients have only one abnormal allele.⁵ Possibly, the abnormal allele confers an evolutionary advantage in resisting an endemic pathogen, an idea reflected in the carrier frequencies of different *MEFV* mutations in certain Mediterranean and Middle Eastern ethnic populations (Sephardic Jews, Turks, Arabs, Armenians).^{6,7} Also, carriage of mutations in *MEFV* in patients with Crohn disease has been associated with a higher risk of extraintestinal manifestations and colonic stricture,⁸ and their carriage in patients with multiple sclerosis has been associated with a rapid progression of that disease.⁹

Brief episodes of fever and serositis

Although FMF usually presents at ages 5 to 15, about 20% of patients with FMF suffer their first inflammatory attack after age 20 years.

Attacks are characterized by brief episodes of fever with temperatures higher than 102°F (38.9°C), lasting less than 72 hours, accompanied by intense serositis. Abdominal serositis may be severe enough to mimic appendicitis and lead to exploratory surgery.

About 70% of patients experience arthritis (predominantly in the legs), and 40% develop erysipeloid erythema, an intensely erythematous, warm, tender, and plaque-like lesion on the lower extremities. Biopsy of involved skin shows a diffuse, primarily neutrophilic, inflammatory cell infiltrate.

Laboratory examination reveals marked elevation of acute-phase reactants, which may normalize between episodes. The diagnosis can be made using a combination of clinical suspicion, sequencing of the *MEFV* gene, and a positive response to a trial of colchicine (Colcrys).

Without treatment, repetitive attacks of inflammation may result in amyloidosis of the kidneys or liver. The risk of amyloidosis is related to several discrete risk factors, such as country of residence, *MEFV* genotype, and serum amyloid A genotype.¹⁰⁻¹² Patients should be monitored for physical manifestations of amyloidosis at least annually.

FMF patients have also been described who develop vasculitides such as Henoch-Schönlein purpura, polyarteritis nodosa, or Behçet disease.

Colchicine is the mainstay of FMF treatment

Colchicine has been the mainstay of therapy for patients with FMF for almost 40 years.¹³⁻¹⁵ Its benefits in FMF are clear: symptoms cease in nearly 70% of patients treated with colchicine, and an additional 25% have a reduction in the severity and frequency of attacks.

Only 5% to 10% of patients have no response to colchicine; this may be partially due to individual dose limitations imposed by common drug-associated gastrointestinal side effects.¹⁶⁻¹⁸ For these patients, newer biologic drugs that inhibit IL-1 activity, such as anakinra (Kineret) and rilonacept (Arcalyst), offer great promise.

Typically, FMF attacks become less frequent and less severe with age. However, the overall prognosis in FMF is related mainly to the individual's genotype and the associated risk of amyloidosis.¹⁹

HYPERIMMUNOGLOBULIN D SYNDROME

HIDS is another autosomal recessive autoinflammatory syndrome.²⁰

The genetic defect underlying HIDS lies within the mevalonate kinase gene *MVK*.²¹ Mevalonate kinase, an enzyme, plays an important role in the cholesterol biosynthesis pathway, following the initial step by 3-hydroxy-3-methyl-glutaryl-CoA (HMG-CoA) reductase. Mutations are primarily missense mutations in highly conserved areas of protein that result in decreased *MVK* activity (1% to 5% of normal).^{22,23} Decreased production of geranylgeranyl pyrophosphate resulting from disruption in the HMG-CoA reductase pathway subsequently leads to increased release of IL-1 beta from peripheral blood mononuclear cells and triggers inflammatory symptoms.²⁴

**IL-1, IL-6,
and TNF are
the most potent
proinflammatory
cytokines**

Attacks of HIDS begin early in life

HIDS attacks begin early in life, with more than 70% of patients suffering their first attack before age 2, but adult-onset disease has been reported. Patients may report that routine childhood vaccinations triggered attacks, a historical finding unique to HIDS.

Attacks typically last 4 days; a longer duration can help the clinician differentiate HIDS from FMF.

More than 90% of patients have cervical lymphadenopathy, and 80% have an erythematous rash characteristically located on the palms and soles. About 70% of patients have headache, arthritis, and abdominal pain.

During attacks, laboratory examination reveals elevated acute inflammatory reactants. As the name implies, serum levels of immunoglobulin D (IgD) are elevated. However, this finding is not specific to HIDS and may also be found in patients with Still disease or FMF or in those who smoke cigarettes. Serum IgD levels fluctuate throughout life, and the sensitivity of commercially available IgD test kits is variable.

Assessment of mevalonic acid levels in the urine during febrile attacks offers a more sensitive, specific, and reliable diagnostic test for HIDS.²⁵ While genetic sequencing is the gold standard of diagnostic testing, close to 30% of patients meeting clinical criteria for HIDS have no definable mutation.²⁶

Treatment of HIDS can be challenging

Oral corticosteroids are effective in HIDS, but their long-term side effects are undesirable. Patients rarely respond to colchicine, differentiating them from FMF patients.

Etanercept (Enbrel), a fusion protein composed of the soluble TNF receptor and the Fc portion of the human IgG1 protein, has been efficacious in some patients.^{27,28} IL-1 inhibitors have also been used with increasing efficacy in the treatment of HIDS attacks.^{29,30}

Although the frequency of attacks decreases with age, long-term follow-up of 28 Dutch HIDS patients found that their quality of life was still lower than that in country-matched controls.³¹

TUMOR NECROSIS FACTOR RECEPTOR-ASSOCIATED PERIODIC SYNDROME

In 1982, a large multiplex family from Scotland and Ireland was described who had a newly recognized syndrome termed familial Hibernian fever, characterized by recurrent fever, rash, and abdominal pain.³² In 1998, the genetics of this autosomal dominant condition were characterized,³³⁻³⁵ and it is now known by the acronym TRAPS.

TRAPS has a variable presentation owing to a variety of mutations in the gene encoding the cell surface receptor for TNF (*TNFRSF1A*). *TNFRSF1A* mutations affecting conserved cysteine residues important for protein folding correspond to severe disease phenotypes.

The R92Q mutation has an allele frequency of up to 4% of the population. It has no impact on the structure and function of the TNF receptor protein and is associated with a heterogeneous disease course. In contrast, the P46L mutation has an allele frequency of 1% of the population and typically is associated with a milder disease course characterized by older age of onset, shorter episodes, and a low frequency of amyloidosis.³⁶⁻³⁹

The R92Q and T61I mutations, which have low penetrance, have been increasingly reported in adult patients with the autoimmune diseases systemic lupus erythematosus, rheumatoid arthritis, and multiple sclerosis.⁴⁰⁻⁴² Their influence is believed to contribute to proinflammatory responses but not to provide additional genetic susceptibility as provided by human leukocyte antigen (HLA) genotypes for susceptibility for these autoimmune diseases.

TRAPS attacks last longer than FMF and HIDS attacks

TRAPS attacks last 7 days or more, differentiating TRAPS from FMF and HIDS. Patients may present from infancy into adulthood, but more typically present in the toddler period.

Most patients experience intense myalgia as well as abdominal and pleuritic chest pain. A single-center series in 2002 described close to half of patients diagnosed with TRAPS as having had an intra-abdominal surgical procedure; in 10% necrotic bowel was identified,

In FMF, abdominal serositis may be severe enough to mimic appendicitis

Elevated IgD
is not specific
for HIDS

yet the biopsy typically revealed only a serosal mononuclear infiltrate.⁴³

Like FMF and HIDS, TRAPS can cause an erythematous rash. The rash usually appears on an extremity, is centrifugal, and travels proximal-to-distal in concert with symptoms of myalgia. Deep tissue biopsy often demonstrates an intense, neutrophilic fasciitis sparing the underlying musculature. Painful conjunctivitis with periorbital edema also may occur.

Laboratory values reflecting widespread systemic inflammation and elevated acute-phase reactants are encountered during attacks and in some cases may persist between episodes.

Genetic testing can be used to confirm the diagnosis. The probability of finding a mutation in *TNFRSF1A* depends highly on whether the patient has affected relatives. In a series of 28 patients with recurrent inflammatory syndromes and *TNFRSF1A* mutations, 9 (32%) had a family history of recurrent inflammatory syndromes, while in 176 patients with sporadic, nonfamilial “TRAPS-like” symptoms, *TNFRSF1A* mutations were uncommon.^{37,38}

Etanercept is effective for TRAPS

Systemic corticosteroids may be effective for treating TRAPS, but ever-increasing doses are often required.

Etanercept’s ability to bind both soluble and bound TNF explains its relative efficacy in treating TRAPS even though other TNF inhibitors have proven ineffective.^{44,45} With etanercept, the prognosis of TRAPS patients is typically good. Etanercept has even been effective in treating cases of renal amyloidosis from long-standing TRAPS, although it has not been shown to facilitate regression of renal amyloid mass.^{46,47} However, responses to treatment with etanercept may wane with time, and resistant cases have been reported.

IL-1 blockade with anakinra has been shown to be effective in the short term and long term in small case series, providing a reasonable alternative for patients who are difficult to manage.

CRYOPYRIN-ASSOCIATED PERIODIC SYNDROMES

- Perhaps the most clinically diverse hereditary autoinflammatory syndromes are the cryopyrin-associated periodic syndromes

(CAPS). There are three overlapping phenotypes: Familial cold autoinflammatory syndrome (FCAS)

- Muckle-Wells syndrome (MWS)
- Neonatal-onset multisystemic inflammatory disorder (NOMID).

Mutations in *NLRP3*

CAPS symptoms stem from mutations within the *NLRP3* gene (NOD-like receptor family, pyrin domain), which encodes the protein, cryopyrin.⁴⁸ *NLRP3* mutations result in an abnormal cryopyrin structure, abnormal inflammasome activity, and increased IL-1 beta production.^{49,50}

There is poor genotype-phenotype association in CAPS; the same *NLRP3* point mutation can result in variable features, typically of either FCAS and MWS or MWS and NOMID overlapping phenotypes, supporting the hypothesis that modifier genes play a role in phenotypic expression.

Inheritance patterns in CAPS are autosomal dominant, but spontaneous mutations are also common. In fact, approximately two-thirds of patients with mutation-negative NOMID have somatic *NLRP3* mutations, indicating that somatic *NLRP3* mosaicism contributes to the clinical syndrome.⁵¹

Clinical features of the CAPS

The hallmarks of the CAPS include recurrent fevers, urticarial rash, and central nervous system inflammation. Characteristically, CAPS patients present in the neonatal period through early childhood, but adult-onset cases, which may have less typical features, have been reported.

Patients with FCAS develop brief episodes (< 24 hours) of fever, joint pain, and urticarial rash when exposed to sudden drops in ambient temperature.

Patients with MWS have more frequent, prolonged attacks, which may or may not be related to changes in ambient temperature. They also develop fever and urticarial rash and may develop arthritis and headaches from aseptic meningitis.

Patients with NOMID often present with fever and persistent urticarial rash shortly after birth and suffer from chronic aseptic meningitis, which can lead to papilledema and op-

tic nerve atrophy. Frontal bossing of the skull and overgrowth of the epiphyseal regions of long bones with accompanying growth delay are also characteristic of NOMID.

IL-1 antagonists offer relief from CAPS

Many patients with FCAS do not require treatment and may move to a warmer climate to avoid rapid swings in ambient temperature. Otherwise, control of IL-1 beta activity is essential to the successful treatment of CAPS. Patients with MWS and NOMID require treatment with IL-1 antagonists, and the biologic drugs anakinra, rilonacept, and canakinumab (Ilaris) offer the possibility of symptomatic relief and long-term control of the disease.⁵²⁻⁵⁴

Prognosis depends on the phenotype

The overall prognosis for patients with CAPS largely depends on phenotype.

Patients with FCAS generally have progressive improvement in attack frequency and severity over time and are at minimal risk of amyloidosis.

Patients with MWS have a relatively good prognosis when treated with IL-1 antagonists, making them at low risk of amyloidosis and sensorineural hearing loss.

However, patients with NOMID are at high risk of sensorineural hearing loss, growth delay, and amyloidosis unless the condition is recognized and treated early in its course. Mortality rates historically are as high as 20% in untreated patients with NOMID.⁵⁵

■ OTHER AUTOINFLAMMATORY SYNDROMES

More recently, other autoinflammatory syndromes of known genetic etiology have been described.

***NLRP12*-associated autoinflammatory disorders**

A subset of patients with clinical manifestations attributable to CAPS but without mutations at the *NLRP3* locus have mutations in another NLRP family member expressed in peripheral blood mononuclear cells on the *NLRP12* gene. They are therefore labeled as having an *NLRP12*-associated autoinflammatory disorder.^{56,57}

Deficiency of interleukin 1 receptor antagonist

IL-1 receptor antagonist is a naturally occurring antagonist of IL-1 alpha and IL-1 beta. In patients with deficiency of IL-1 receptor antagonist (DIRA), the action of these potent proinflammatory proteins is unopposed, leading to severe pustular rash and osteitis.^{58,59}

Pyogenic arthritis, pyoderma gangrenosum, and acne syndrome

Patients with PAPA syndrome also have increased IL-1 production, in this case due to a mutation in the cytoplasmic adapter protein proline-serine-threonine phosphatase-interacting protein (*PSTPIP1*) gene, leading to the development of the symptoms included in the PAPA acronym.⁶⁰

Majeed syndrome

Majeed syndrome is caused by a mutation in the *LPIN2* gene, resulting in the early onset of chronic recurrent multifocal osteomyelitis, neutrophilic dermatosis, and dyserythropoietic anemia.⁶¹

Blau syndrome

Some patients with Blau syndrome (granulomatosis, arthritis, and uveitis) have *NOD2/CARD15* gene mutations.⁶² Cases of DIRA, PAPA, and Blau syndrome have been reported that responded favorably to treatment with IL-1 antagonists.

Periodic fever, aphthous stomatitis, pharyngitis, and adenopathy syndrome

Although symptoms of the periodic fever, aphthous stomatitis, pharyngitis, and adenopathy (PFAPA) syndrome typically begin in childhood, adult-onset cases have been reported.⁶³

Patients with PFAPA syndrome develop predictable, stereotypic febrile attacks that last on average 5 days and occur approximately every 4 weeks. Between attacks, patients are healthy; during attacks, they may experience oral ulceration (aphthous stomatitis), exudative or nonexudative pharyngitis, and enlarged and tender cervical lymph nodes. Up to 60% of PFAPA patients also experience abdominal pain.

No single genetic mutation has been identified, although it has been shown that

The duration of attacks is one clue in distinguishing different auto-inflammatory syndromes

Many patients with FCAS do not need treatment and may move to a warmer climate to avoid rapid temperature swings

45% of PFAPA patients have a parent or sibling with recurrent fever and 12% have a parent or sibling with a PFAPA-like phenotype, suggesting that the disease has a genetic basis.⁶⁴ Recent studies have demonstrated that T-cell-regulated complement activation and IL-1 production are altered in PFAPA patients, thus supporting the hypothesis that PFAPA is an autoinflammatory syndrome.⁶⁵

Treatment. In view of the syndrome's self-limited nature, treatment is reserved for patients with a severe presentation or for patients whose condition is especially burdensome.

The fever's height may partially respond to nonsteroidal anti-inflammatory drugs, but these drugs have little effect on the duration or frequency of fever.

One or two doses of prednisone (1 mg/kg) within 6 hours of fever onset is effective in aborting the febrile episode in 90% of patients; however, up to 50% of patients may experience an increased frequency of attacks after treatment with systemic corticosteroids.^{66,67}

Additional options include daily colchicine, which may lengthen the time between attacks, and cimetidine (Tagamet), which has been shown to prevent PFAPA attacks in approximately one-third of patients.⁶⁷⁻⁶⁹

The prognosis of PFAPA is quite favorable, and without intervention 40% of patients experience a significant reduction in the severity and frequency of fever attacks within 5 years of diagnosis. To date, there have been no reports of amyloidosis or hearing loss in PFAPA patients.

■ DIAGNOSTIC EVALUATION OF SUSPECTED AUTOINFLAMMATORY DISEASE

The autoinflammatory syndromes pose a true diagnostic challenge for physicians. Tremendous advances have been made in molecular and genetic testing. Nevertheless, the history and careful physical examination can lead the astute clinician to the proper diagnosis when evaluating a patient with a suspected autoinflammatory syndrome.

Critical elements in the history include age at the onset of attacks, duration of attacks, associated symptoms (serositis, adenopathy, myalgias, arthralgias, arthritis, ocular symp-

toms, central nervous system symptoms, rash), family members with similar symptoms, and ethnic background.

Internists should remember that autoinflammatory syndromes are part of the differential diagnosis in adult patients with a recurrent febrile illness. A vigorous search for malignancy and infection (especially tuberculosis) should be conducted in all patients. However, the repetitive, stereotypic nature of autoinflammatory syndromes differentiates them from typical confounders.

The utility of acute-phase reactants in the diagnostic evaluation is limited, as many conditions result in abnormal values. However, serial monitoring of inflammatory markers such as the erythrocyte sedimentation rate and C-reactive protein level in patients with a formally diagnosed autoinflammatory syndrome can be useful in tracking disease activity, identifying flares, and monitoring the efficacy of therapy.

In cases of suspected HIDS, assessment of IgD levels is not recommended, since IgD can be elevated in a number of autoinflammatory and rheumatologic conditions. Instead, preference should be given to testing mevalonic acid levels in the urine in patients with HIDS or suspected HIDS.

Patients with central nervous system symptoms should undergo a thorough examination, including a formal ophthalmologic evaluation, imaging, and possibly lumbar puncture to assess intracranial pressure and inflammatory changes in the cerebrospinal fluid.

Dermatologic manifestations should be examined firsthand and assessed as needed with magnetic resonance imaging to elucidate fascial inflammation or with full-thickness biopsy.

Gross bony abnormalities should be evaluated with plain radiography.

Audiologic testing may be indicated in the diagnostic evaluation of patients with recurrent fever.

Renal or hepatic biopsy may be indicated in the evaluation for amyloidosis; amyloid deposition has been reported in patients with TRAPS and clinical FMF not presenting with fever.^{70,71}

Genetic testing is commercially available for patients with suspected hereditary autoin-

Diagnostic algorithm for patients with suspected autoinflammatory conditions

Patient with recurrent episodes of fever and stereotypic symptoms not explained by infectious or oncologic etiologies

Fever < 3 days, painful serositis, in patient of Jewish, Near-Eastern, or Mediterranean descent?

Yes

Consider familial Mediterranean fever (FMF)
Send for *MEFV* analysis
Consider trial of colchicine (Colcrys)

No

Fever > 7 days, periorbital edema, migratory rash?

Yes

Consider tumor necrosis receptor-associated periodic syndrome (TRAPS)
Send for *TNFRSF1A* analysis
Trial of twice-weekly etanercept (Enbrel)

No

Cold-induced urticaria, sensorineural hearing loss, or other central nervous system symptoms?

Yes

Consider Muckle-Wells syndrome
Send for *NLRP3* analysis
Trial of anakinra (Kineret)

No

Prominent lymphadenopathy, rash on palms and soles, attacks triggered by vaccinations?

Yes

Consider hyperimmunoglobulin D syndrome
Check serum IgD and urinary mevalonic acid
Send for *MVK* analysis

No

Neonatal onset with urticarial rash and bony overgrowth?

Yes

Consider neonatal-onset multisystemic inflammatory disorder
Consider lumbar puncture and fundoscopic examination
Trial of anakinra

No

Consider periodic fever, aphthous stomatitis, pharyngitis, and adenopathy (PFAPA) syndrome (if aphthous stomatitis or cervical adenopathy is present); systemic lupus erythematosus; vasculitis; inflammatory bowel disease

No

Assess for rash, eye inflammation, atypical arthritis or osteomyelitis (deficiency of interleukin 1 receptor antagonist [DIRA], Blau, Majeed, pyogenic arthritis, pyoderma gangrenosum, and acne [PAPA] syndromes)

Treatment of PFAPA syndrome is reserved for severe cases

FIGURE 2

flammatory syndromes. However, clinicians should be cautioned that up to 30% of patients with phenotypic manifestations characteristic of a given autoinflammatory syndrome have normal results on genetic testing. In addition, the results of genetic testing may take several months to return and may cost patients and families up to several thousand dollars, as some insurers refuse to cover this procedure. Genetic testing may ultimately be indicated

for proper counseling of reproductive risk.

Responses to short courses of medications such as colchicine, prednisone, and IL-1 receptor antagonists also represent diagnostic tools.

FIGURE 2 provides a proposed diagnostic algorithm for patients with suspected recurrent fever syndromes. TABLE 1 summarizes clinical and genetic features of the common autoinflammatory syndromes.

TABLE 1

Quick reference guide to common autoinflammatory syndromes

SYNDROME ^a	CLINICAL FEATURES	DURATION OF ATTACKS	MODE OF INHERITANCE	GENE	THERAPY	RISK OF AMYLOIDOSIS
FMF	Serositis Arthritis Rash	< 72 hours	Recessive	<i>MEFV</i>	Colchicine (Colcris) IL-1 inhibition	+++
HIDS	Cervical adenopathy Rash Arthritis Headache Attacks triggered by vaccination	3–5 days	Recessive	<i>MVK</i>	Prednisone Etanercept (Enbrel) IL-1 inhibition Anti-IL-6	–
TRAPS	Myalgia Serositis Periorbital edema Traveling rash	7–10 days	Dominant	<i>TNFRSF1A</i>	Prednisone Etanercept IL-1 inhibition Anti-IL-6	++
CAPS						
FCAS	Urticaria Joint pain	< 24 hours	Dominant	<i>NLRP3</i>	Cold avoidance	–
MWS	Urticaria Arthritis Headache Hearing loss	5–7 days	Dominant	<i>NLRP3</i>	IL-1 inhibition	++
NOMID	Urticaria Arthritis Headache Hearing loss Papilledema Bony changes	Constant	Dominant	<i>NLRP3</i>	IL-1 inhibition	+

^a FMF = familial Mediterranean fever
HIDS = hyperimmunoglobulin D syndrome
TRAPS = tumor necrosis factor receptor-associated periodic syndrome
CAPS = cryopyrin-associated periodic syndromes
FCAS = familial cold autoinflammatory syndrome
MWS = Muckle-Wells syndrome
NOMID = neonatal-onset multisystemic inflammatory disorder

Advances in understanding the innate immune system will lead to improvement in the care we give

NEW INSIGHT INTO MORE COMMON CONDITIONS

Advances in the understanding of the autoinflammatory syndromes have provided new insight into the role of the innate immune system in other, more common conditions.⁷² Indeed, abnormal regulation of the innate inflammatory pathway has been implicated in the pathogenesis of conditions as phenotypi-

cally diverse as gout, type 2 diabetes, atherosclerosis, and epilepsy.^{73,74}

TABLE 2 presents examples of the innate immune system's involvement in the pathogenesis of several common chronic conditions.

Further study of autoinflammatory syndromes will add to our understanding of the innate immune system. These advances will lead to continued improvement in the

TABLE 2

Role of the innate immune system in common chronic conditions

Atherosclerosis

Crystals containing cholesterol are deposited in vessel walls and directly stimulate NLRP3, leading to increased tissue levels of interleukin 1 (IL-1) beta and inflammation^{63,64}

Gout

Monosodium urate crystals trigger localized inflammatory response by activating the NLRP3 inflammasome; blockade of IL-1 beta activity leads to significant improvement in gout symptoms^{65,66}

Osteoarthritis

Calcium phosphate crystals directly stimulate IL-1 beta production via activation of the NLRP3 inflammasome⁶⁷

Type 2 diabetes

Hyperglycemia indirectly activates the NLRP3 inflammasome, leading to increased levels of IL-1 beta; prolonged increased levels of IL-1 beta can lead to insulin resistance, hyperglycemia, and beta cell death^{64,68}

care we give patients, both for the classic autoinflammatory syndromes and for other, more common, genetically complex conditions.

Our 22-year-old patient's fever, abdominal pain (presumed peritonitis), erysipelas-like skin lesion, and arthritis are typical of FMF. Therefore, genetic testing was performed, which revealed a single *MEFV* gene mutation (M694V). Colchicine has been efficacious in preventing flares of his disease. ■

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