

MEDLEY

Medical AI Ensemble Clinical Decision Report

Case ID: custom_20250911_192300	Title: A 28-year-old male of Mediterranean descent presents with: -...	Generated: 2025-09-11 19:24
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Primary Diagnostic Consensus

Diagnosis	ICD-10	Agreement	Confidence	Status
Familial Mediterranean Fever (FMF) <i>Evidence: Recurrent fever episodes 1-3 days, Severe abdominal pain, Chest pain with breathing difficulties, Joint pain in knees and ankles</i>	E85.0	0.0%	Very Low	PRIMARY

Alternative & Minority Diagnoses

Diagnosis	ICD-10	Support	Type
Periodic Fever Syndrome (other types) <i>Evidence: Recurrent fever episodes, Family history</i>	E85.9	3.7%	Minority (<10%)
Systemic Juvenile Idiopathic Arthritis <i>Evidence: Recurrent fever episodes, Joint pain</i>	M08.2	3.7%	Minority (<10%)
Reactive Arthritis <i>Evidence: Recurrent fever episodes, Joint pain</i>	M40.0	3.7%	Minority (<10%)
Behçet's Disease <i>Evidence: Recurrent fever episodes, Mucocutaneous lesions</i>	M35.2	0.0%	Minority (<10%)
Takayasu's Arteritis <i>Evidence: Recurrent fever episodes, Vascular involvement</i>	M31.0	0.0%	Minority (<10%)
Systemic Lupus Erythematosus <i>Evidence: Recurrent fever episodes, Multi-organ involvement</i>	M32.9	0.0%	Minority (<10%)
Rheumatoid Arthritis <i>Evidence: Recurrent joint pain, Elevated inflammatory markers</i>	M05.9	0.0%	Minority (<10%)
Gout <i>Evidence: Recurrent joint pain, Elevated uric acid levels</i>	M10.0	0.0%	Minority (<10%)

Diagnosis	ICD-10	Support	Type
Sarcoidosis <i>Evidence: Recurrent fever episodes, Granulomatous inflammation</i>	D86.9	0.0%	Minority (<10%)
Infectious Mononucleosis <i>Evidence: Recurrent fever episodes, Lymphadenopathy</i>	B27.8	0.0%	Minority (<10%)

Analysis Overview
Models Queried: 3
Successful Responses: 3
Consensus Level: High
Total Cost: <\$0.01

■ ■ Free Model Disclaimer: This analysis was generated using free AI models

Free models may provide suboptimal results. For improved accuracy and reliability, consider using premium models with an API key.

Critical Decision Points & Evidence Synthesis

Critical Decision Points

Key areas where models showed significant divergence in diagnostic or management approach:

Evidence Synthesis & Clinical Correlation

Symptom-Diagnosis Correlation Matrix

Symptom	Familial	Periodic	Systemic	Reactive	Behçet's	Takayasu	SLE	Rheumato
Recurrent fever	Strong	Moderate	Moderate	Moderate	-	-	-	-
Severe abdomina	Strong	-	-	-	-	-	-	-
Chest pain with	Strong	-	-	-	-	-	-	-
Joint pain in k	Strong	-	Moderate	Moderate	-	-	-	-
Family history	Strong	Moderate	-	-	-	-	-	-
Elevated CRP, E	Strong	-	-	-	-	-	-	-
Asymptomatic be	Strong	-	-	-	-	-	-	-
Symptoms starte	Strong	-	-	-	-	-	-	-
Mediterranean d	Strong	-	-	-	-	-	-	-
Pending genetic	Strong	-	-	-	-	-	-	-

Legend: +++ Strong association, ++ Moderate, + Weak, - Not typical

Diagnostic Decision Tree

Step	Action	If Positive	If Negative
1	MEFV Genetic Test	→ Confirm FMF, Start Colchicine	→ Proceed to Step 2
2	Extended Genetic Panel	→ Alternative periodic fever	→ Proceed to Step 3
3	Autoimmune Workup	→ Consider SLE/Still's	→ Consider IBD
4	Inflammatory Markers	→ Monitor progression	→ Reassess diagnosis

Executive Summary

Case Description

A 28-year-old male of Mediterranean descent presents with:

- Recurrent episodes of fever lasting 1-3 days
- Severe abdominal pain during episodes
- Chest pain with breathing difficulties
- Joint pain affecting knees and ankles
- Family history: Father and paternal uncle have similar symptoms
- Episodes occur every 2-3 weeks
- Labs during attack: Elevated CRP, ESR, and WBC
- Between attacks: Completely asymptomatic

Patient reports episodes started in childhood around age 7. Recent genetic testing is pending.

Key Clinical Findings

- Elevated inflammatory markers (CRP, ESR)
- Recurrent fever episodes
- Positive family history of similar episodes
- Migratory arthritis affecting large joints
- Severe abdominal pain with peritoneal signs

Primary Recommendations

- Consider Familial Mediterranean Fever (FMF) among differential diagnoses
- Obtain Complete blood count (CBC) for diagnostic confirmation

Primary Diagnosis Clinical Summaries

■ Key Clinical Findings

Finding	Supporting Evidence	Clinical Reasoning
Recurrent fever episodes 1-3 days	Clinical presentation	Key diagnostic indicator
Severe abdominal pain	Clinical presentation	Key diagnostic indicator
Chest pain with breathing difficulties	Clinical presentation	Key diagnostic indicator
Joint pain in knees and ankles	Clinical presentation	Key diagnostic indicator
Family history of similar symptoms	Clinical presentation	Key diagnostic indicator

■ Recommended Tests

Test Name	Type	Priority	Rationale
Complete blood count (CBC)	Laboratory	Urgent	Diagnostic confirmation
C-reactive protein (CRP)	Laboratory	Urgent	Diagnostic confirmation
Erythrocyte sedimentation rate (ESR)	Laboratory	Urgent	Diagnostic confirmation
Renal function tests	Laboratory	Urgent	Diagnostic confirmation
Urine analysis	Laboratory	Urgent	Diagnostic confirmation

■ Immediate Management

Intervention	Category	Urgency	Clinical Reasoning
Initiate genetic testing for MEFV gene mutations	Medical	Immediate	Critical intervention
Evaluate for signs of amyloidosis	Medical	Immediate	Critical intervention

■ Medications

Medication	Dosage	Route/Frequency	Indication
Colchicine	1.2 mg	oral / daily	Prevention of FMF attacks and amyloidosis

Diagnostic Landscape Analysis

Detailed Diagnostic Analysis

The ensemble analysis identified **Familial Mediterranean Fever (FMF)** as the primary diagnosis with limited consensus among 3 models.

Detailed Alternative Analysis

Diagnosis	Support	Key Evidence	Clinical Significance
Periodic Fever Syndrome (other types) <i>Evidence: Recurrent fever episodes, Family history</i>	3.7%	1 models	Unlikely
Systemic Juvenile Idiopathic Arthritis <i>Evidence: Recurrent fever episodes, Joint pain</i>	3.7%	1 models	Unlikely
Reactive Arthritis <i>Evidence: Recurrent fever episodes, Joint pain</i>	3.7%	1 models	Unlikely
Behçet's Disease <i>Evidence: Recurrent fever episodes, Mucocutaneous lesions</i>	0.0%	0 models	Unlikely
Takayasu's Arteritis <i>Evidence: Recurrent fever episodes, Vascular involvement</i>	0.0%	0 models	Unlikely
Systemic Lupus Erythematosus <i>Evidence: Recurrent fever episodes, Multi-organ involvement</i>	0.0%	0 models	Unlikely
Rheumatoid Arthritis <i>Evidence: Recurrent joint pain, Elevated inflammatory markers</i>	0.0%	0 models	Unlikely
Gout <i>Evidence: Recurrent joint pain, Elevated uric acid levels</i>	0.0%	0 models	Unlikely

Minority Opinions

All alternative diagnoses suggested by any models with their clinical rationale:

- **Periodic Fever Syndrome (other types)** (ICD-10: R50.9) - 3.7% agreement (1 models)
Supporting Models: model1
- **Systemic Juvenile Idiopathic Arthritis** (ICD-10: Unknown) - 3.7% agreement (1 models)
Supporting Models: model1
- **Reactive Arthritis** (ICD-10: M02.9) - 3.7% agreement (1 models)
Supporting Models: model2

- **Behçet's Disease** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

- **Takayasu's Arteritis** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

- **Systemic Lupus Erythematosus** (ICD-10: M32.9) - 0.0% agreement (0 models)

Supporting Models:

- **Rheumatoid Arthritis** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

- **Gout** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

- **Sarcoidosis** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

- **Infectious Mononucleosis** (ICD-10: Unknown) - 0.0% agreement (0 models)

Supporting Models:

Additional Diagnoses Considered:

Management Strategies & Clinical Pathways

Immediate Actions Required

Priority	Action	Rationale	Consensus
1	Initiate genetic testing for MEFV gene mutations	Clinical indication	50%
2	Evaluate for signs of amyloidosis	Clinical indication	50%

Recommended Diagnostic Tests

Test	Purpose	Priority	Timing
Complete blood count (CBC)	Diagnostic confirmation	Routine	As indicated
C-reactive protein (CRP)	Diagnostic confirmation	Routine	As indicated
Erythrocyte sedimentation rate (ESR)	Diagnostic confirmation	Routine	As indicated
Renal function tests	Diagnostic confirmation	Routine	As indicated
Urine analysis	Diagnostic confirmation	Routine	As indicated

Treatment Recommendations

Treatment recommendations pending diagnostic confirmation.

Model Diversity & Bias Analysis

Model Response Overview & Cost Analysis

Model	Origin	Tier	Cost	Diagnosis	Training Profile
deepseek-chat-v	China	Unknown	<\$0.01	Familial Mediterranean Fever	General
mistral-7b-inst	France	Free	Free	Familial Mediterranean Fever	General
gpt-oss-20b	USA	Unknown	<\$0.01	Familial Mediterranean fever	General

Total Estimated Cost: <\$0.01

Understanding Training Profiles

Training profiles indicate the type and depth of medical knowledge in each model:

Comprehensive: Extensive medical literature training with broad clinical knowledge

Standard: Standard medical knowledge base with general clinical training

Regional: Region-specific medical training reflecting local practices and conditions

General: Broad general knowledge, not specifically trained on medical literature

Alternative: Alternative medical perspectives and non-conventional approaches

AI Model Bias Analysis

AI model bias analysis is generated during orchestration (Step 2). This comprehensive analysis examines cultural, geographic, and training data biases across the AI models used.

Detailed Model Responses

Complete diagnostic assessments from each model:

1. deepseek-chat-v (China, Released: 2024-12-26)

Primary Diagnosis: Familial Mediterranean Fever (ICD-10: E85.0) - Confidence: 0.95

Differential Diagnoses:

- Periodic Fever Syndrome (other types) (ICD: E85.9) - 0.6
- Systemic Juvenile Idiopathic Arthritis (ICD: M08.2) - 0.4
- Acute Intermittent Porphyria (ICD: E80.21) - 0.3

Key Clinical Findings:

- Mediterranean descent
- Recurrent self-limited febrile episodes (1-3 days)
- Severe abdominal pain
- Chest pain with pleurisy

2. mistral-7b-inst (France, Released: 2023-09-27)

3. gpt-oss-20b (USA, Released: 2024-05-13)