

Medical Al Ensemble Clinical Decision Report

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Primary Diagnostic Consensus

Diagnosis	ICD-10	Agreement	Confidence	Status
Familial Mediterranean Fever	E85.0	60.0%	Moderate	PRIMARY

Alternative & Minority Diagnoses

Diagnosis	ICD-10	Support	Туре
Systemic Juvenile Idiopathic Arthritis	Unknown	40.0%	Strong Alt (≥30%)
PFAPA Syndrome	M04.8	40.0%	Strong Alt (≥30%)
Inflammatory Bowel Disease	K50.9	40.0%	Strong Alt (≥30%)
Periodic Fever Syndrome (other types)	E85.8	20.0%	Alternative (10-29%)
Adult-Onset Still's Disease	M06.1	20.0%	Alternative (10-29%)
Hyper-IgD Syndrome (MKD)	E85.8	20.0%	Alternative (10-29%)
TNF Receptor-Associated Periodic Syndrome (TRAPS)	Unknown	20.0%	Alternative (10-29%)
Acute Intermittent Porphyria	E80.21	20.0%	Alternative (10-29%)
Vasculitis (e.g., Polyarteritis Nodosa)	Unknown	20.0%	Alternative (10-29%)

Analysis Overview Models Queried: 5 Successful Responses: 5 Consensus Level: 0.95 Total Estimated Cost: <\$0.01

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/Finding	Familial Med	Systemic Juv	PFAPA Syndro	Inflammatory
Chronic Inflamm	-	-	-	?
Gi Symptoms	+	-	++	?
Fever	+++	+++	+++	?
Abdominal Pain	-	-	-	?
Joint Symptoms	-	-	-	?

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	ightarrow 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

- · Positive family history of similar episodes
- Migratory arthritis affecting large joints
- Recurrent fever episodes

Primary Recommendations

- Moderate consensus (60.0%) suggests Familial Mediterranean Fever
- Seek immediate medical attention
- Consult a healthcare professional
- Obtain Genetic testing for FMF for diagnostic confirmation

Primary Diagnosis Clinical Summaries

Diagnostic Landscape Analysis

Detailed Diagnostic Analysis

The ensemble analysis identified **Familial Mediterranean Fever** as the primary diagnosis with 60.0% consensus among 3 models.

Detailed Alternative Analysis

Diagnosis	Support	Key Evidence	Clinical Significance
Systemic Juvenile Idiopathic Arthritis	40.0%	2 models	Worth investigating
PFAPA Syndrome	40.0%	2 models	Worth investigating
Inflammatory Bowel Disease	40.0%	2 models	Worth investigating
Periodic Fever Syndrome (other types)	20.0%	1 models	Less likely
Adult-Onset Still's Disease	20.0%	1 models	Less likely
Hyper-IgD Syndrome (MKD)	20.0%	1 models	Less likely
TNF Receptor-Associated Periodic Syndrome (TRAPS)	20.0%	1 models	Less likely
Acute Intermittent Porphyria	20.0%	1 models	Less likely

Minority Opinions

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

Additional Diagnoses Considered:

- Systemic Juvenile Idiopathic Arthritis (ICD-10: Unknown) 40.0% (2 models)
- PFAPA Syndrome (ICD-10: M04.8) 40.0% (2 models)
- Inflammatory Bowel Disease (ICD-10: K50.9) 40.0% (2 models)
- Periodic Fever Syndrome (other types) (ICD-10: E85.8) 20.0% (1 models)
- Adult-Onset Still's Disease (ICD-10: M06.1) 20.0% (1 models)
- Hyper-IgD Syndrome (MKD) (ICD-10: E85.8) 20.0% (1 models)
- TNF Receptor-Associated Periodic Syndrome (TRAPS) (ICD-10: Unknown) 20.0% (1 models)
- Acute Intermittent Porphyria (ICD-10: E80.21) 20.0% (1 models)
- Vasculitis (e.g., Polyarteritis Nodosa) (ICD-10: Unknown) 20.0% (1 models)

Diagnostic Confidence Analysis

Management Strategies & Clinical Pathways

Immediate Actions Required

Priority	Action	Rationale	Consensus
1	Seek immediate medical attention	Clinical indication	50%
2	Consult a healthcare professional	Clinical indication	50%

Recommended Diagnostic Tests

Test	Purpose	Priority	Timing
Genetic testing for FMF	Diagnostic confirmation	Routine	As indicated
Inflammatory markers	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
gemma-2-9b-it	USA	Free	Free	Not specified	General
gemma-3-12b-it	USA	Unknown	<\$0.01	Periodic Fever Syndrome (likely Familial Mediterranean Fever - FMF)	General
mistral-7b-inst	France	Free	Free	Familial Mediterranean Fever	General
shisa-v2-llama3	Japan/USA	Free	Free	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

Al Model Bias Analysis

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

Primary Diagnosis Bias Factors:

- Cultural: Models from 6 countries with Western dominance may miss cultural factors. Chinese models (100.0%) provide alternative perspective.
- Geographic: Western model dominance (500.0%) creates strong bias toward Western medical paradigms. High Western medical paradigm influence expected
- Training Data: English-dominant training data creates systematic bias against non-Western medical practices and symptom presentations.

Alternative Diagnoses Bias:

• Missed: Traditional Medicine Conditions - Western model dominance may miss traditional medicine diagno...

• Missed: Socioeconomic-Related Conditions - Homeless status bias may cause dismissive attitudes and miss...

Bias Mitigation Recommendations:

- Socioeconomic Bias: Consider cultural context in diagnosis interpretation
- Geographic/Cultural Bias: Incorporate diverse cultural perspectives in diagnosis

Detailed Model Responses

Complete diagnostic assessments from each model:

- 1. deepseek-chat-v (China, Released: 2024-12-26)
- 2. gemma-2-9b-it (USA, Released: 2024-06-27)
- 3. gemma-3-12b-it (USA, Released: 2024-12-11)
- 4. mistral-7b-inst (France, Released: 2023-09-27)
- 5. shisa-v2-llama3 (Japan/USA, Released: 2024-12-20)