

### Reportable Parameters

28 reportable parameters: CRP, Hs CRP, SAA, WBC, Neu#, Lym#, Mon#, Eos#, Bos#. Neu%, Lym%, Mon%, Eos%, Bas%, RBC, HGB. HCT, MCV, MCH, MCHC, RDW-SD, RDW-CV, PLT, MPV, PDW, PCT, P-LCR, P-LCC

Research Parameters 6 research parameters ALY#, ALYX Lie#, UC%. NRBC#. NRBC%

Graph 13D Scattergram, 3 2D Scattergrams and 3 histograms

Throughput Up to 90 tests per hour

Sample Volume 20pL

Test Mode CBC, CBC+DtFF. CBC+DIFF+CRP, CBC+DIFF+SAA, CRP, SAA, CRP+SAA, CBC+DIFF+CRP+SAA

Sampling Mode Closed in auto loader and open in vial

Blood Mode Whole blood .capillary blood and prediluted modes

Dimension (W\*H\*D) <595mm\* <600mm\* < 700mm

Weight <72kg

Diagnosis

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**Redefine Infection** 



## **DM79X** 5-Part Hematology + CRP +SAA Joint Analyzer

### The Clinical Significance of WBC, CRP & SAA

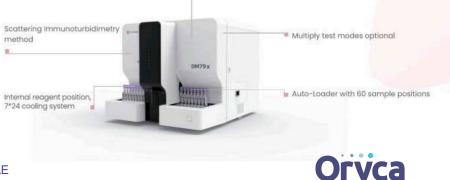
ltem	Ref. Range	Bacterial Infection	Viraf infection	Mixed infection
WBC	4-10*10* /L	Neutrophil Increase	Lymphocyte increase	Might increase
CRP	<10mg/L	Significantly Increase	Slightly or none increase	Increase
SAA	<10mg/L	Increase faster than CRP but drop faster	Significantly increase	Significantly increase

### The Clinical Alert of CRP & SAA Joint Detection during Acute Phase of Infection

SAA	CRP	Clinical alert
<10 mg/l	Within normal range	No acute inflammatory
10 mg/L-100 mg/l	Normal or slightly increase	Virus Infection (mild)
100 mg/L-500 mg/L	> 50 mg/L	Bacterial infection, dynamically monitor SAA and CRP to evaluate the therapeutic effect
*500 mg/L	Increasebut <50 mg/L	Viral infection (severe), bacterial infection, need 12-24 h review

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Triditional method two instrument multiply steps