BioCLIA® Autoimmune Control Set

RF IgG

RF IgG Assay Controls

Key to Symbols Used

REF	Catalog Number		Expiration Date
IVD	For *In Vitro* Diagnostic Use	LOT	Lot Number
	Store between +2°C and +8°C		Consult Instruction for Use
	Manufacturer	EC REP	Authorized Representative in European Union
		Σ	Contains Sufficient for < *n* > Tests
Control L	Control 1		
Control H	Control 2		Chemical Risk Warning
			Biological Risk Warning

BioCLIA® Autoimmune Control Set, RF IgG

Intended Use

The BioCLIA Autoimmune Control Set, RF IgG is intended for the quality control purposes of the BioCLIA RF IgG performed on the BioCLIA® 1200 and BioCLIA® 6500.

Catalog Numbers

MY00333 (2 X 1 mL)
MY00384 (4 X 1 mL)

Summary and Principles of the Procedure

Rheumatoid factors (RFs) are anti-antibodies specific to the degenerated IgG antibodies caused by infection factors (bacteria, viruses, etc.). B cell clones for expressing RFs exist in rheumatoid arthritis (RA) patients and about 50% of healthy human bodies. A large amount of RFs produced under certain pathologic conditions such as degenerated IgG or EB virus directly stimulating the B cells. [1] Commonly, RFs include IgA, IgD, IgE, IgG and IgM types. [2, 3]

The presence of RFs in serum can indicate the occurrence of suspected autoimmune activity unrelated to rheumatoid arthritis, such as that associated with tissue or organ rejection. In such instances, RF may serve as one of several serological markers for autoimmunity. [4] The sensitivity of RFs for established rheumatoid arthritis is only 60-70% with a specificity of 78%. [5] RFs positive patients are likely to have extra-articular manifestation that includes subcutaneous nodules and vasculitis. Patients with a high concentration of RFs suffer from diseases like rheumatoid arthritis (RA, 50-90%), Sjogren's syndrome (75-95%). [6] In addition, it was also found that RFs are related to systemic lupus erythematosus (SLE, 15-35%), polymyositis (55-10%), systemic sclerosis (20-30%), mixed connective tissue disease (MCTD, 50-60%) or cryoglobulinemia (40-100%). [7, 8] Specifically, determination of RF-IgG has a high specificity for RA diagnosis.

Materials supplied

- **RF IgG Control 1** A tube contains 1mL, ready to use reagent. Control 1 contains anti-RF IgG antibodies in MOPS (PH6.0-7.0) buffer (Low).

RF-G	Control	L

Preservatives: 5-Bromo-5-Nitro-1, 3-Dioxane (BND) < 1%.

- **RF IgG Control 2** A tube contains 1mL, ready to use reagent. Control 2 contains anti-RF IgG antibodies in MOPS (PH6.0-7.0) buffer (High).

RF-G	Control	H

Preservatives: 5-Bromo-5-Nitro-1, 3-Dioxane (BND) < 1%.

Target value and acceptable range for the controls are indicated on control information sheet in each kit.

Warnings and Precautions

The human derived material in this product was tested by FDA approved methods and found nonreactive for Hepatitis B Surface Antigen (HBsAg), Anti-HCV and HIV 1/2 antibodies. Handle as if potentially infectious. [9] Avoid contacting with skin and eyes. Do not empty into drains. Wear suitable protective clothing.

Precautions:

 Human serum is added in the controls.

- The product is for *in vitro* diagnostic use only.
- Do not use any controls beyond their expiration dates. Do not mix controls from different lots unless specified.
- Instructions must be carefully followed for using and storing of controls. Any modification in procedure may interfere with the results. Controls and contaminated vials must be handled strictly following safety guidelines or rules of biological hazards to ensure the users' and environmental safety.
- Controls contain chemical and biological components. Avoid ingesting or splashing onto skin and mucous membrane. If direct contact with controls happens, rinse the contact surface with plenty of water immediately and see a doctor if necessary.

Storage Conditions

The kit is stable until the expiration date, if it is stored and handled as directed. Routine store the kit in refrigerator (2-8°C). Once a control tube is opened, it is good for a total of 15 times, no more than 2 hours per time when kept uncapped, onboard the instrument, after which the reagent must be discarded. Three freeze-thaw cycles before testing has no effect on the kit reagents.

Assay Procedure

Note that for obtaining performance, it is important to perform all routine maintenance procedures as defined in the BioCLIA® 1200 and BioCLIA® 6500 User Manual.

See the BioCLIA® 1200 and BioCLIA® 6500 User Manual for preparation, setup, dilutions, adjustment, assay and quality control procedures.

The control procedure can be done before running the specimens each day. Users also can adjust the control procedure period according to their own lab frequency.

Limitations

This product is designed as controls for monitoring the performance of the BioCLIA RF IgG. These controls are subjected to the limitations of the assay system. Deviations may indicate possible problems with one or more components in the test system.

References

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6. Visser H. Early diagnosis of rheumatoid arthritis. Best Practice & Research Clinical Rheumatology 2005;19:55-72.

7. Wilson D. Rheumatoid factors in patients with rheumatoid arthritis. Canadian Family Physician Médecin De Famille Canadien 2006;52:180-1.

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9. Richmond JY, Mckinney RW. Biosafety in microbiological and biomedical laboratories: U.S.GPO. 1999.

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EC REP

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Technical Assistance

For technical assistance, contact your National Distributor.

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