BioCLIA Autoimmune Reagent Kit, DGP IgG

(Chemiluminescent Microparticle Immunoassay)

Kit size	Cat. No.	GTIN Code
50 Tests/kit	MY00141	06924030402563
100 Tests/kit	MY00192	06924030403072

INTENDED USE

The BioCLIA Autoimmune Reagent Kit, DGP IgG assay is intended for the in vitro semi-quantitative measurement of IgG antibodies directed to deamidated gliadin peptide in human serum and plasma as an aid in the diagnosis of celiac disease (CD). It is an *in vitro* diagnostic medical device intended for laboratory professional use.

SUMMARY AND EXPLANATION OF THE TEST

Anti-deamidated gliadin peptide antibodies (anti-DGP antibodies) are produced in response to gliadin, a prolamin found in wheat. DGP IgA is found in about 80% of patients with coeliac disease.[1, 2] It is directed against the alpha/beta and gamma (α, β, γ) gliadins. [3] It is also found in a number of patients who are not enteropathic. Some of these patients may have neuropathies that respond favorably to a gluten elimination diet. This is referred to as gluten-sensitive idiopathic neuropathy.[4] Clinically these antibodies and IgG antibodies to gliadin are abbreviated as AGA. The IgG antibody is similar to AGA IgA, but is found at higher levels in patients with the IgA-less phenotype. It is also associated with celiac disease and non-celiac gluten sensitivity. [5-7]

AGA was one of the first serological markers for coeliac disease. Problematic with AGA is the typical sensitivity and specificity was about 85%. Gliadin peptides which are synthesized as the deamidated form have much higher sensitivity and specificity, creating 2 serological tests for CD that approach biopsy diagnostic in performance. [8]

PRINCIPLES OF THE PROCEDURE

BioCLIA Autoimmune Reagent Kit is a two-step immunoassay. The antigen of interest is coated onto magnetic microparticles. Specific IgG in the patient's serum/plasma reacts with these to form an IgG-antigen complex. After incubation, a washing step removes the free and non-specifically bound molecules. Subsequently enzyme labeled anti-human IgG antibodies (conjugate) is added and this binds to the IgG-antigen complex. After further incubation a second washing step removes the unbound conjugate. Then addition of substrate results in the emission of light and the relative light unit (RLU) intensity is measured. The relative light unit (RLU) intensity is proportional to the amount of antigen specific IgG present in the sample.

The BioCLIA Autoimmune Reagent Kit utilizes a predefined lot specific Master Curve which is uploaded into the instrument via the barcode provided in the kit. Based on the results of running two calibrators, the instrument specific Working Calibration Curve is generated and is used to calculate the concentration from the RLU obtained for each patient.

WARNINGS AND PRECAUTIONS

- For professional *in vitro* diagnostic use only.
- Used on BioCLIA 6500 and BioCLIA 500 instruments only.
- Do not use reagents beyond the expiration dates.
- The kit contains human sourced materials. All recommended precautions for the handling of blood derivatives should be taken. Please refer to the existing laboratory safety regulations and good laboratory practice.
- Liquid waste and solid waste taken from BioCLIA 6500 and BioCLIA 500 should also be handled in accordance with the National or Local legislation.
- Once opened, the reagent cartridge must be stored in the instrument's reagent carousel. For the first placement of reagent into the instrument, please take care to avoid spilling the reagents.
- Spilled reagents should be cleaned up immediately. Comply with all National and local environmental regulations when disposing of wastes.

  Improper cleaning or rinsing of the instrument may lead to chemical contamination of the reagents. Residues from common laboratory chemicals such as formalin, bleach, ethanol, or detergent can cause interference in the assay. Be sure to follow the recommended cleaning procedure as outlined in the BioCLIA 6500 and BioCLIA 500 User's Manual.

Precautions

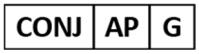
- The assay contains ProClin 300 0.0015%~3% as preservative and may cause an allergic skin reaction by skin contact. Avoid contact with skin. Wear protective gloves, protective clothing and protective glasses.
- The assay contains TritonX-100 0.0015%~0.5% as surfactant and may cause an allergic skin reaction by skin contact. Avoid contact with skin. Wear protective gloves, protective clothing and protective glasses.
- Any serious incident that has occurred in relation to the device shall be reported to the manufacturer and the competent authority of the Member State in which the user and/or the patient is established.

MATERIALS SUPPLIED

- **Buffer** 0.01M PBS (pH7.4)

Preservatives: 0.0015%＜ProClin 300 < 3%.
Surfactant: 0.0015%＜Triton X-100 < 0.5%.

- **Conjugate** Alkaline phosphatase (AP) labeled anti-human IgG



antibodies in 0.05 M MES (pH6.0) Buffer with stabilizer.
Preservatives: 0.0015%＜ProClin 300 < 3%.

• **Microparticle** Antigen coated on microparticles in 0.01M PBS (pH 7.4).

M | ANTIGEN

Preservatives: 0.0015%＜ProClin 300 < 3%.

Components	50 Tests/Kit	100 Tests/Kit
Buffer	1 x 2.5 mL	1 x 5 mL
Conjugate	1 x 6.75 mL	1 x 13.5 mL
Microparticle	1 x 2.5 mL	1 x 5 mL

ADDITIONAL MATERIALS SUPPLIED SEPARATELY

Product	CATALOGUE No.
BioCLIA Autoimmune Calibrator Set, DGP IgG	MY00243 (2 x 1 mL) MY00294 (4 x 1 mL)
BioCLIA Autoimmune Control Set, DGP IgG	MY00345 (2 x 1 mL) MY00396 (4 x 1 mL)
BioCLIA Sample Diluent I	MY00965
BioCLIA System Wash Buffer	MY00404
BioCLIA System Substrate	MY00405
BioCLIA 6500	MA00243
BioCLIA 500	MA00502
BioCLIA Cuvettes	MA00244 (2000 pcs/bag) MA00549 (65 pcs/box)
BioCLIA Silicone gasket (Small)	MV00195
BioCLIA Silicone gasket (Large)	MV00196

MATERIALS REQUIRED

- Distilled or deionized Water

STORAGE AND STABILITY

- Store the kit at 2-8 °C.
- The shelf life of the unopened kit is 12 months from day of production.
- Opened reagents or onboard reagents may be used for 28 successive days. The software of the BioCLIA instruments monitors the onboard (in-use) expiration of the reagent cartridge. The system will not allow use of a reagent which has expired.

SPECIMEN COLLECTION, STORAGE AND HANDLING

- Serum from venous can be used.
- Collect blood specimens using standard procedures.
- Test serum should be clear and free from hemolysis.
- Cloudy samples should be clarified by centrifugation at 5000 rpm for 5 minutes before use. For samples with the presence of fibrin, ensure that complete clot formation has taken place prior to centrifugation of samples. Some samples, particularly those from patients receiving anticoagulant therapy, may require increased clotting time.
- Specimens may be refrigerated at 2-8 °C for up to seven days or stored at -20°C up to six months.
- Specimens may be kept onboard on BioCLIA instruments under room temperature (18-25°C) for up to 2 hours.
- Avoid repeated freezing and thawing.

ASSAY PROCEDURE

Detailed information about operating the BioCLIA instruments can be taken from the Instrument User's Manual.

Note that, it is important to perform all routine maintenance procedures for optimal performance.

Sample Dilution

The specimens are diluted with BioCLIA Sample Diluent I before testing (dilution ratio 1:19) by the BioCLIA instruments automatically.

Assay Calibration

The BioCLIA Autoimmune Reagent Kit utilizes a predefined lot specific Master Curve which is uploaded into the instrument via the barcode provided in the kit. Based on the results of running two calibrators, the instrument specific Working Calibration Curve is generated and is used to calculate the concentration from the RLU obtained for each patient.

For each new lot of reagent, please calibrate prior to the first time use, and every 28 days thereafter. The software will not allow the lot to be used if the above requirements are not meet.

Control

The control procedure should be done before running the specimens each day. Users also can adjust the control procedure period according to their own lab frequency.
Each Laboratory should establish its own reference ranges.

Programming and Running samples

1. Put the kit into any empty position of the reagent chamber of the BioCLIA instruments. Details of the kit can be uploaded into the instrument system through the scanning of reagent barcode and can also be set manually.
2. The information of calibrator / quality control is identified by scanning the calibrator / control barcodes, and the position of calibrator / quality control is assigned in the instrument system.
3. The sample to be tested is placed on the instrument sample rack chamber, and the corresponding test information is edited through the instrument supporting software.
4. Start the operation procedure, and all calibrator / quality control / sample processing steps will be automatically executed.

CALCULATION OF RESULTS

Calculation and interpretation of results will be performed automatically by software on BioCLIA instruments.

RESULT INTERPRETATION

Specimens with concentration <20 RU/mL, should be interpreted as negative;
Specimens with concentration ≥20 RU/mL, should be interpreted as positive.

The test results only reflect the amount of antigen specific IgG present in the sample and should be diagnosed in conjunction with other laboratory and clinical findings.

CUT-OFF VALUE DETERMINATION

120 clinical samples, including 30 positive sera, 30 negative sera, 30 positive plasmas and 30 negative plasmas were collected and evaluated. Results were analyzed using the receiver-operating characteristic curve (ROC) and the cut-off value was determined at 20 RU/mL.

PERFORMANCE CHARACTERISTICS

ACCURACY / SPIKED RECOVERY

The accuracy/spiked recovery was determined by analyzing samples spiked with known amounts of antibody into sample matrix. Specific antibody positive samples (low 100 RU/mL, mid 200 RU/mL, high 300RU/mL) were spiked into two matrixes (50 and 100 RU/mL) separately at the volume ratio of 1:9, making totally 6 spiked samples and each sample was tested in triplicate. The spiked recovery for the concentration of autoantibodies to specific antigen was calculated. *

	Matrix 50 RU/mL			Matrix 100 RU/mL		
Spiked Conc.	Obs	Exp.	Obs/Exp	Obs	Exp.	Obs/Exp
Neat	47.57			102.39		
100 RU/mL	50.27	52.81	95.20%	100.98	102.15	98.90%
200 RU/mL	63.62	62.81	101.30%	111.23	112.15	99.20%
300 RU/mL	71.22	72.81	97.80%	122.21	122.15	100.00%

*Representative data; results in individual laboratories may vary from these data.

TRACEABILITY

This assay system is calibrated in relative arbitrary units since no international reference preparation is available for this assay. The reported values were determined with multiple runs on the BioCLIA 6500 and BioCLIA 500 using specific reagents against an in-house standard.

PRECISION

A study based on guidance from (NCCLS) document EP-A[18] was performed.

Intra-assay precision: Four samples (negative, low, moderate, and high) were taken and tested with 10 replicates for each in a single run. Coefficient of variation (CV) was calculated for each of four samples. The results for intra-assay precision are shown in the table below.

Inter-assay precision: Four samples (negative, low, moderate, and high) were taken and tested with 4 replicates in a single run, two runs per day for 10 days. Coefficient of variation (CV) was calculated for each of four samples. The results for inter-assay precision are shown in the table below. *

Intra-assay precision: CV < 10%

Intra-Assay	Sample1	Sample2	Sample3	Sample4
Mean(RU/mL)	10.77	20.19	99.79	359.96
CV	4.56%	4.62%	4.71%	4.28%

Inter-assay precision: CV< 15%

Inter-Assay	Sample1	Sample2	Sample3	Sample4
Mean(RU/mL)	10.41	19.39	99.25	357.09
CV	6.00%	5.71%	5.79%	4.30%

*Representative data; results in individual laboratories may vary from these data.

LIMIT OF BLANK / DETECTION (LOB/LOD)

LOB/LOD was determined according to CLSI EP17-A guideline.
The assay is designed to have LoB/LoD of ≤ 0.5 RU/mL.

LINEARITY

The linear range of the assay is 2-400 RU/mL.
The linear range was determined by serially diluting a sample containing high levels of antigen specific IgG with a negative sample and covering the entire assay linear range according to the scheme in CLSI EP6-A. The expected value was plotted against the observed value, and linear regression analysis was performed to get slope, intercept and coefficient of correlation (r) values. The results are summarized in the table below*:

Slope	Intercept	r
0.98	+1.34	0.99

*Representative data; results in individual laboratories may vary from these data.

INTERFERENCE

No interference has been observed with bilirubin, hemoglobin, triglycerides, rheumatoid factor (RF), human anti-mouse antibody (HAMA) at the levels indicated below.

- Bilirubin ≤ 40 mg/dL;
- Hemoglobin ≤ 150 mg/dL;
- Triglycerides ≤ 1,000 mg/dL;
- Rheumatoid factor (RF) ≤ 1,000 IU/mL;
- Human anti-mouse antibody (HAMA) ≤ 2,000 ng/mL.

METHOD COMPARISON

Method comparison was implemented by comparing clinical sample results of the assay to the results of predicated assay. The results are shown in the table below. *

Clinical Sample		BioCLIA Autoimmune Reagent Kit, DGP IgG		
		-	+	Total
Predicated Method	-	48	2	50
	+	4	46	50
	Total	52	48	100

Sensitivity	92.0%
Specificity	96.0%
Total agreement	94.0%

LIMITATIONS

- The effectiveness of this kit is only confirmed for human serum/plasma, the applicability of the other kinds of samples is not verified.
- Reliable and reproducible results will be obtained when the assay procedure is carried out in accordance with the instructions and with adherence to good laboratory practice.
- Clinical diagnosis should not be made on the findings of a single test result, but should be interpreted with all clinical and laboratory findings.

SYMBOLS

REF	Catalog Number		Use-by date
IVD	In Vitro diagnostic medical device	LOT	Lot Number
+8°C +2°C	Store between +2°C and +8°C		Consult Instruction for Use
	Manufacturer	EC REP	Authorized Representative in the European Community
CE	CE Marking		Contains Sufficient for<n>Tests
	Biological Risk		GHS07 Warning

BUFFER	Buffer
CONJ AP G	AP labeled anti-human IgG antibody
M ANTIGEN	Antigen coated microparticles

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TECHNICAL ASSISTANCE

For technical assistance, contact your National Distributor.

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