

REF



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### Intended Use

The Aurora Vitamin B12 Assay is a chemiluminescent microparticle immunoassay (CMIA) for the quantitative determination of Vitamin B12 (VB12) in human serum and plasma. The Aurora Vitamin B12 Assay is intended for in vitro diagnostic use.

### Summary

Vitamin B12, also known as cobalamin, is a water-soluble vitamin involved in metabolism. It is derived from a tetrapyrrolic structural framework created by the enzymes deaminase and cosynthetase which transform aminolaevulinic acid via porphobilinogen and hydroxymethylbilane to uroporphyrinogen III. The latter is the first macrocyclic intermediate common to heme, chlorophyll, siroheme and B12 itself. It is required by animals, which use it as a cofactor in DNA synthesis, in both fatty acid and amino acid metabolism. It is important in the normal functioning of the nervous system via its role in the synthesis of myelin, and in the maturation of red blood cells in the bone marrow. Vitamin B12 is an essential cofactor in intermediary metabolism and is required for the biosynthesis of RNA and DNA. Since a deficiency of either vitamin may be the cause of megaloblastic anemia, it is essential to determine the levels of both vitamin B12 and folic acid to establish the etiology of the anemia. The deficiency of Vit.B12 is due to the lack of intrinsic factor (IF) by the stomach and intestinal malabsorption. Vitamin B12 deficiency can potentially cause severe and irreversible damage, especially to the brain and nervous system. At levels only slightly lower than normal, a range of symptoms such as feeling tired and weak, feeling like one may faint, difficulty walking (staggering balance problems), depression, poor memory, poor reflexes, confusion, breathlessness, headaches, and pale skin, feeling abnormal sensations, among others. Vitamin B12 deficiency can also cause symptoms of mania and psychosis. Among other problems, weakened immunity, reduced fertility and interruption of blood circulation in women may occur.

### Test Principles

The Vitamin B12 assay is a quantitative competitive immunoassay to determine the presence of VB12 in human serum and plasma using CMIA technology with flexible assay protocols.

1. Sample, assay auxiliary 2, assay auxiliary 3 are mixed, after incubation, then add assay auxiliary 1 and a conjugate containing acridinium -labeled intrinsic factor, VB12 present in the sample binds to intrinsic factor labeled acridinium, forming a complex.
2. After incubation, biotin-labeled VB12 coated microparticles are added to the reaction mixture and VB12 binds to unoccupied binding sites of the intrinsic factor labeled acridinium.
3. After further incubation and washing, Pre-Trigger and Trigger Solutions are added to the reaction mixture.
4. The resulting chemiluminescent reaction is measured as relative light units (RLUs). There is a relationship between the amount of VB12 in the sample and the RLUs detected by the Optical system. Results are calculated automatically based on the previously established calibration curve.

### Reagents

R1	3.0 mL
R2	4.8 mL
R3	4.5 mL
R4	2.2 mL
R5	2.2 mL
CAL	Contains 3 levels, 1.0 mL each level
CON	Contains 2 levels, 1.0 mL each level
R1:	Microparticles. Biotin-labeled VB12 coated microparticles. Preservative: 0.05% ProClin 300
R2:	Conjugate. Acridinium-labeled intrinsic factor. Preservative: 0.05% ProClin 300.
R3:	Assay auxiliary 1. Phosphate Buffer. Preservative: 0.05% ProClin 300
R4:	Assay auxiliary 2. Protein lysing agent. Preservative: 0.05% ProClin 300
R5:	Assay auxiliary 3. 0.25M Sodium hydroxide
CAL:	Calibrator. Solutions of different

concentrations of VB12. Preservative: 0.05% ProClin 300

**CON:**

Control. Tris buffer solution for quality control of VitaminB12 (VB12)

## Required Materials

- Pre-Trigger Solution: Hydrogen peroxide solution.
- Trigger Solution: Sodium hydroxide solution.
- Wash Buffer: Phosphate buffered saline solution with 0.05% ProClin 300.

## Safety Precautions

- Exercise the normal precautions required for handling all laboratory reagents.
- Disposal of all waste material should be in accordance with local guidelines.
- Wear gloves when handling specimens or reagents.
- Clean and disinfect all spills of specimens or reagents using a suitable disinfectant.
- Trigger solution contains sodium hydroxide (NaOH) and should be avoided contact with eyes.

## Warning (Contains Proclin 300)

Hazardous Component: 0.05% Proclin 300

- Reaction mass of:  
5-chloro-2-methyl-4-isothiazolin [EC no. 247-500-7] and 2-methyl-4-isothiazolin-3-one [EC no. 200-239-6] (3:1)

## Hazard Statement

- H317: May cause an allergic skin reaction.
- H319: Causes serious eye irritation.
- H410: Very toxic to aquatic life with long-lasting effects.

## Precautionary Statement

- P261: Avoid breathing dust/fume/gas/mist/vapors/spray.
- P264: Wash hands thoroughly after handling.
- P272: Contaminated work clothing should not be allowed out of the workplace.
- P280: Wear protective gloves/protective clothing/eye protection/face protection.
- P305+P351+P338: IF IN EYES: Rinse cautiously with water for several minutes. Remove contact

lenses, if present and easy to do. Continue rinsing.

- P337+P313: If eye irritation persists: Get medical advice/attention.
- P333+P313: If skin irritation or rash occurs: Get medical advice/attention.
- P302+P352: IF ON SKIN: Wash with plenty of soap and water.
- P321: Seek immediate care from a doctor.
- P363: Wash contaminated clothing before reuse.
- P273: Avoid release to the environment.
- P391: Collect spillage.
- P501: Dispose of contents/container in a safe way.

## Reagent Handling

- Do not use reagent kits beyond the expiration date.
- Do not pool reagents within a kit or between kits.
- Before loading the reagent kit on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment.
- Septums MUST be used to prevent reagent evaporation and contamination and to ensure reagent integrity. Reliability of assay results cannot be guaranteed if septums are not used according to the instructions in this package insert.
- To avoid contamination, wear clean gloves when placing a septum on an uncapped reagent bottle.
- Once a septum has been placed on an open reagent bottle, do not invert the bottle as this will result in reagent leakage and may compromise assay results.

## Reagent Storage

REAGENT	Storage Temperature	Maximum Storage Time
Unopened	2°C~8°C Do not freeze.	12 months
On board/ Opened	2°C~8°C Do not freeze.	28 days

- Reagents may be stored on or off the Chemiluminescence immunoassay analyzer. If reagents are removed from the analyzer, store them at 2-8°C (with septums and replacement caps) in an upright position. For reagents stored off the system,

it is recommended that they be stored in their original trays and boxes to ensure they remain upright. If the microparticle bottle does not remain upright (with a septum installed) while in refrigerated storage off the system, the reagent kit must be discarded.

### Calibrator & Control Storage

CAL	CON	Storage Temperature	Maximum Storage Time
Unopened		2°C~8°C	12 months
Opened		2°C~8°C	30 days

### Applicable Analyzer

Automatic Chemiluminescence Immunoassay Analyzer (model: Aurora S-01).

### Specimen Types

Verified specimen types to be used with this assay:

Specimen Types	Collection Tubes
Serum	Serum separator tubes (SST)
Plasma	Dipotassium EDTA Tripotassium EDTA Sodium heparin Lithium heparin powder Plasma separator tubes (PST) -lithium heparin gel

- Other specimen collection tube types have not been tested with this assay.
- Liquid anticoagulants may have a dilution effect resulting in lower concentrations for individual patient specimens.
- The instrument does not provide the capability to verify specimen type. It is the responsibility of the operator to verify that the correct specimen types are used in the assay.

### Specimen Conditions

Do not use specimens with the following conditions:

- heat-inactivated
- pooled
- grossly hemolyzed (> 500 mg/dL hemoglobin)
- obvious microbial contamination
- fungal growth
- For accurate results, serum and plasma

specimens should be free of fibrin, red blood cells, and other particulate matter. Serum specimens from patients receiving anticoagulant or thrombolytic therapy may contain fibrin due to incomplete clot formation.

- To prevent cross contamination, use of disposable pipettes or pipette tips is recommended.

### Preparation for Analysis

- Follow the tube manufacturer's processing instructions for collection tubes. Gravity separation is not sufficient for specimen preparation.
- Mix thawed specimens thoroughly by low speed vortexing or by inverting 10 times. Visually inspect the specimens. If layering or stratification is observed, continue mixing until specimens are visibly homogeneous.
- Avoid more than 3 freeze/thaw cycles.
- To ensure consistency in results, specimens must be transferred to a centrifuge tube and centrifuged for a minimum of 30,000 g-minutes before testing if they contain fibrin, red blood cells, or other particulate matter, they were previously frozen.
- Examples of acceptable time and force ranges that meet this criterion are listed in the table below. Centrifugation time using alternate Relative Centrifugal Force values (RCF) can be calculated using the following formula:

Centrifugation Time (Minutes)	RCF (×g)	g × minutes
10	3,000	30,000
15	2,000	30,000
20	1,500	30,000

- Transfer clarified specimen to a sample cup or secondary tube for testing. For centrifuged specimens with a lipid layer, transfer only the clarified specimen and not the lipemic material.
- Inspect all specimens for bubbles. Remove bubbles with an applicator stick before analysis. Use a new applicator stick for each specimen to prevent cross contamination.

### Specimen Storage

Specimen Type	Storage	Maximum
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	Temperature	Storage Time
Serum/Plasma	Room temperature	≤ 8 hours
	2°C~8°C	≤ 7 days
	-20°C	≤ 90 days

- Remove serum or plasma from the clot, red blood cells, or separator gel if stored longer than the maximum room temperature storage time.
- Remove serum or plasma from the clot, red blood cells, or separator gel if stored longer than the maximum 2-8°C storage time and store frozen.
- Frozen specimens must be mixed thoroughly after thawing.
- Use caution in handling patient specimens to prevent cross-contamination.
- Do not exceed the storage limitations listed above.

#### Assay Procedure

- Refer to the system operating instruction or the online help system for detailed information on preparing the system.
- Before loading the Aurora Vitamin B12 Assay on the system for the first time, the microparticle bottle requires mixing to resuspend microparticles that may have settled during shipment. After the first time the microparticles have been loaded, no further mixing is required.
- Invert the microparticle bottle 30 times. Visually inspect the bottle to ensure microparticles are resuspended. If microparticles are still adhered to the bottle, continue to invert the bottle until the microparticles have been completely resuspended. If the microparticles do not resuspend, DO NOT USE. Once the microparticles have been resuspended, place a septum on the bottle.
- Load the reagent kit on the Chemiluminescence immunoassay analyzer.
- Verify that all necessary reagents are present.
- Verify adequate sample volume is present prior to running the test.
- Sample volume for first test: 250 µL
- Sample volume for each additional test from

same sample cup: 34 µL

- The test-specific parameters stored in barcode on the reagent pack are read in. In cases the barcode cannot be read, enter the sequence numbers
- Order calibration, if necessary.
- Prepare Vitamin B12 Calibrators and Controls.
- Mix calibrator(s) and controls by gentle inversion before use.
- Hold bottles vertically and dispense recommended volumes into each respective sample cup.
- Place the calibrators in the calibrator rack in the sample zone.
- Calibration.
- Load samples. For information on loading samples, refer to the Analyzer's Operations Manual.
- Press RUN.
- The Chemiluminescence immunoassay analyzer performs all the functions automatically and calculates the results.

For optimal performance, it is important to perform routine maintenance as described in the Analyzer's Operations Manual. Perform maintenance more frequently when required by laboratory procedures.

#### Sample Dilution Procedures

- Samples with a VB12 value exceeding 2000 pg/mL may be diluted using the Manual Dilution Procedure.

#### Manual Dilution Procedure

Suggested dilution: 1:2

- Add 100 µL of the sample to 100 µL of Stroke physiological saline solution or dilutions.
- The operator must enter the dilution factor in the Patient or Control order screen. The system will use this dilution factor to automatically calculate the concentration of the sample before dilution and report the result.

#### Calibration

- Traceability: This assay has been standardized against the NRCCRM, NIM-RM3603.
- Every VB12 assay kit has a two-dimension code label containing the predefined master curve of the

particular reagent lot.

- Test Calibrators in duplicate. The calibrators should be priority loaded. A replicate of each control level must be tested to evaluate the assay calibration. Ensure that assay control values are within the ranges specified in the respective control package insert.
- Once calibration is accepted and stored, all subsequent samples may be tested without further calibration unless:
  - After 28 days when using the same lot reagent.
  - A reagent kit with a new lot number is used.
  - Controls are out of range.
  - Required by pertinent regulations.
  - Assay may also need to be recalibrated after specified service procedures have been performed or maintenance to critical part or subsystems that might influence the performance of the assay. For detailed information on how to perform an assay calibration, refer to the Analyzer's Operations Manual.
- Calibration Range: 50 pg/mL~1600 pg/mL.

#### Quality Control Procedures

- Order Control, if necessary.
- The recommended control requirement for the VB12 assay is that a single replicate of each control level be tested:
  - Once every 24 hours each day of use
  - After performing calibration
  - After instrument service procedures or maintenance that may affect assay performance have been performed.
  - If the quality control procedures in your laboratory require more frequent use of controls to verify test results, follow your laboratory-specific procedures.
  - Each laboratory should establish control ranges to monitor the acceptable performance of the assay. If a control is out of its specified range, the associated sample results are invalid and the samples must be retested. Recalibration may be indicated.
- These results should be applied to your

laboratory's quality control practices. In addition, the laboratory must ensure that the matrix of the control material is suitable for use in the assay per the assay package insert.

- Unless specified, target values and ranges provided with the commercial control product insert are guidelines only and should not be used for quality control purposes.
- Refer to Clinical and Laboratory Standards Institute (CLSI) Document C24-A3, or other published guidelines for general quality control recommendations.

#### Results

##### Calculation

- The analyzer automatically calculates the concentration of each sample either in pg/mL or pmol/L (selectable).
- Conversion factors:
  - $\text{pg/mL} \times 0.738 = \text{pmol/L}$
  - $\text{pmol/L} \times 1.355 = \text{pg/mL}$

##### Limits

- Results should be used in conjunction with other data; e.g., symptoms, results of other tests, and clinical impressions.
- If the VB12 results are inconsistent with clinical evidence, additional testing is recommended.
- Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may contain human anti-mouse antibodies (HAMA). Such specimens may show either falsely elevated or depressed values when tested with assay kits that employ mouse monoclonal antibodies. Additional information may be required for diagnosis.
- Heterophilic antibodies in human serum can react with reagent immunoglobulins, interfering with in vitro immunoassays. Patients routinely exposed to animals or to animal serum products can be prone to this interference, and anomalous values may be observed. Additional information may be required for diagnosis.
- Rheumatoid factor (RF) in human serum can react with reagent immunoglobulins, interfering

with in vitro immunoassays. Additional information may be required for diagnosis.

- The Vitamin B12 assay is susceptible to interference effects from triglycerides at > 1000 mg/dL.

### Expected Values

- It is recommended that each laboratory establish its own reference range, which may be unique to the population it serves depending upon geographical, season, patient, dietary, or environmental factors. A study was performed based on guidance from Clinical and Laboratory Standards Institute (CLSI) C28-A3c.
- Human serum specimens from apparently healthy individuals were collected the 125 specimens, 63 were female and 62 were male, age between 21 and 90 years.

The population	n	Reference range*
Apparently healthy individuals	125	201 pg/mL~886 pg/mL

\*According to the 2.5th~97.5th percentile.

\*Representative data; results in individual laboratories and in different geographical areas may vary from these data.

### Specific Performance Characteristics

- Assay results obtained in individual laboratories may vary from data presented.

### Limit of Blank (LoB)

- The Limit of Blank and was determined in accordance with the CLSI (Clinical and Laboratory Standards Institute) EP17-A requirements.
- The Limit of Blank is the 95th percentile value from  $n \geq 20$  measurements of analyte-free samples over several independent series. The Limit of Blank corresponds to the concentration below which analyte-free samples are found with a probability of 95 %.
- The observed LoB value was  $\leq 50$  pg/mL.

### Accuracy

#### Intra Assay Variation

- Within run variation was determined by replicate determination (n=10) of two different control sera in one assay. The within assay variability is  $\leq 8.0\%$ .

#### Inter Assay Variation

- Between run variation was determined by replicate measurements (n=10) of two different control sera in 3 different lots. The between assay variability is  $\leq 10.0\%$ .

Intra-Assay, n=10			Inter-Assay, n=10x3		
Sample	Mean (pg/mL)	CV	Sample	Mean (pg/mL)	CV
1	233.07	4.8%	1	231.57	4.9%
2	605.42	3.8%	2	618.86	5.4%

#### Linearity

- The linearity was determined in accordance with the CLSI (Clinical and Laboratory Standards Institute) EP6-A requirements.
- The linearity range was verified by more than 6 concentration levels which encompass or be equal to the minimum and the maximum values of linearity range and duplicate assays (n=3) at all levels.
- The Aurora Vitamin B12 Assay has been demonstrated to be linear from 100 pg/mL to 1600 pg/mL, regression  $\geq 0.99$  and max diff  $\leq 15\%$  in this interval.

#### Interference

- A study was performed based on guidance from CLSI EP7-A2.
- Potentially interfering substances were evaluated to determine whether VB12 concentrations were affected when using the Vitamin B12 assay. Samples containing the potential interferents were prepared at two VB12 concentrations. The samples were assayed, and the VB12 concentrations of the spiked samples were compared to the reference samples.





















Potential Interferent	Interferent Concentration	% Interferent Bias
Bilirubin	20 mg/dL	$\leq 10\%$
Hb	500 mg/dL	$\leq 10\%$

Potential Interferent	Interferent Concentration	% Interferent Bias
Intralipid	1000 mg/dL	≤ 10%
Total protein	10 g/dL	≤ 10%
RF	1000IU/mL	≤ 10%
ANA	400AU/mL	≤ 10%
HAMA	600ng/mL	≤ 10%

## Reference

1. Allen LH, Miller JW, de Groot L, et al. Biomarkers of Nutrition for Development (BOND): Vitamin B-12 Review. *J Nutr.* 2018;148(suppl\_4):1995S-2027S.
2. Miller A, Korem M, Almog R, Galboiz Y. Vitamin B12, demyelination, remyelination and repair in multiple sclerosis. *J Neurol Sci.* 2005 Jun 15;233(1-2):93-7.
3. "Vitamin B12 or folate deficiency anaemia – Symptoms". National Health Service, England. 23 May 2019.
4. Masalha R, Chudakov B, Morad M, Rudoy I, Volkov I, Wirguin I. Cobalamin-responsive psychosis as the sole manifestation of vitamin B12 deficiency. *Isr Med Assoc J.* 2001 Sep;3(9):701-3.
5. Lederer AK, Hannibal L, Hettich M, et al. Vitamin B12 Status Upon Short-Term Intervention with a Vegan Diet-A Randomized Controlled Trial in Healthy Participants. *Nutrients.* 2019;11(11):2815.
6. Obeid R, Heil SG, Verhoeven MMA, van den Heuvel EGHM, de Groot LCPGM, Eussen SJPM. Vitamin B12 Intake from Animal Foods, Biomarkers, and Health Aspects. *Front Nutr.* 2019; 6:93.
7. Owen MD, Baker BC, Scott EM, Forbes K. Interaction between Metformin, Folate and Vitamin B12 and the Potential Impact on Fetal Growth and Long-Term Metabolic Health in Diabetic Pregnancies. *Int J Mol Sci.* 2021;22(11):5759.
8. Watanabe F, Bito T. Vitamin B12 sources and microbial interaction. *Exp Biol Med (Maywood).* 2018 Jan;243(2):148-158.

## Symbols

	Manufacturer
	Date of manufacture
	Use-by date
	Contains sufficient for <n> tests
	Consult instructions for use
	Biological risks
	Temperature limit
	CE Marking
	EU Representative
	In Vitro diagnostic medical device
	Catalogue Number
	Batch code
	Reagent
	Microparticles
	Conjugate
	Assay auxiliary 1
	Assay auxiliary 2
	Assay auxiliary 3
	Calibrator
	Control

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Release Date: ... .. Date of Manufacture: ... ..