

REF 57838, 57839, 57840



Stat Profile Prime Plus® Blood Gas, CO-Oximeter, Chemistry Controls Auto-Cartridge

Cartucho automático para controles de química, gases en sangre y cooximetría Stat Profile Prime Plus®, Cartouche automatique de contrôles de gaz du sang/CO-oxymètre, chimie Stat Profile Prime Plus®, Stat Profile Prime Plus® Auto-Kassette für Blutgas, CO-Oximeter- und Blutochemiekontrollen, Stat Profile Prime Plus® Αυτόματος φυσίγγιο ελεγχέου χημείας αερίων αίματος, CO-Οξόμετρο, Cartuccia per controlli automatici chimici per gas ematici/CO-ossimetria Stat Profile Prime Plus®, Cartucho automático de controles de química, de CO-oxímetro e de gás no sangue Stat Profile Prime Plus®, Stat Profile Prime Plus® көрөгчтө CO-oximéter, Stat Profile Prime Plus® Кан Газ, CO Оксиметр, Kimya Kontrolleri Otomatik Kartuşu, kemiai kontrollok automatikus patron, Stat Profile Prime Plus® מוסך בדם לנורמליזציה אוטומטית לנו בדם, Stat Profile Prime Plus® חוקימת, Stat Profile Prime Plus® 血液ガス, CO オキシメーター, 生化学検査用コントロール自動カートリッジ, Stat Profile Prime Plus® 혈액 가스, CO-산소 농도계, 화학 조절제 자동 카트리지, Stat Profile Prime Plus® 血气, CO-一氧化碳-血氧仪, 化学对照液自动试剂盒

LOT 24355031

CONTROL 1 2 3 4 5

2026-06-11

EN

Product Description
Accepts quality control material for monitoring the performance of pH, PCO₂, PO₂, SO₂, hematoct (Hct), fetal hemoglobin (HbF), total hemoglobin (HbT), total bilirubin (TbI), albumin (Alb), cyanmethemoglobin (CyHb), carboxyhemoglobin (COHb), methemoglobin (MetHb), and deoxyhemoglobin (HbD) in levels 1, 2 and 3 as well as Na⁺, K⁺, Cl⁻, Ca, Mg, Glucose, and Lactate in Level 4 and 5. For use with Stat Profile Prime Plus Analyzers ONLY.

Intended Use
Intended for in vitro diagnostic use by healthcare professionals for monitoring the performance of the Stat Profile Prime Plus Analyzers.

Methodology
Refer to Stat Profile Prime Plus Analyzer Instructions For Use Manual for Methodology and Principles.

Composition
Controls Levels 1, 2 and 3 are known bicarbonate solutions containing dye, salts and preservatives. Each level has a known pH and is equilibrated to a known O₂, CO₂ and N₂ value. Controls Level 4 and 5 are buffered solutions containing known concentrations of Na⁺, K⁺, Cl⁻, Ca, Mg, Glucose, Lactate and preservatives. Each pouch contains a minimum of 100 mL. Controls contain no constituents of human origin, however good laboratory practices should be followed during handling of these materials. (REF: NCCLS DOCUMENT M29-T2)

Warnings and Cautions:
DO NOT FREEZE. Mix the cartridge by gently inverting for several seconds. DO NOT SHAKE CARTRIDGE. Refer to the Stat Profile Prime Plus Analyzer Instructions for Use Manual for complete information.

Storage
Store at 2-8°C (37-46°F). DO NOT FREEZE.

Directions for use
Ensure controls are at room temperature prior to installation. Mix Cartridge well by gently inverting for 1 minute. Verify that the lot number on the Expected Ranges Table corresponds to the Lot Number on the cartridge. Refer to Stat Profile Prime Plus Analyzer Instructions for Use Manual for complete directions.

Limitations
PO₂ values vary inversely with temperature (approximately 1%/°C). Therefore, it is critical to follow the temperature guidelines described in "Directions for Use." The Expected Range values are specific for instruments and controls manufactured by Nova Biomedical. Once installed, each Stat Profile Prime Plus Cartridge may be used for a maximum of 35 days from the initial installation date on the system at which time the system will indicate the cartridge is invalid. Each cartridge may be inserted and removed from the analyzer a maximum of 10 times.

Traceability of Standards
Total Hemoglobin (HbT) and Methemoglobin (MetHb) are traceable by using Cyanmethemoglobin Method. Carboxyhemoglobin (COHb) and Deoxyhemoglobin (HbD) are traceable using Spectrophotometry. Analytes are traced to NIST Standard Reference Materials.

Reference Intervals
Concentrations are formulated at normal and abnormal expected values in patient blood. The expected clinical range of these values in patient blood is referenced in Tietz, NW ed. 1985 Textbook of Clinical Chemistry, W.B. Saunders Co. Users may wish to determine Mean Values and Expected Ranges in their own laboratory.

Expected Ranges
The expected range for each parameter was determined at Nova Biomedical using replicate determinations on Nova analyzers. The expected range indicates the maximum deviation from the Mean Value that may be expected under differing laboratory conditions for instruments operating within specifications. Refer to Expected Ranges Table.

*Not available in the USA or for Point-of-Care/Non-Patient Testing only.

NCCLS Document M29-T2
How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

Expected Ranges, Rangos esperados, Plages attendues, Erwartungsbereiche, Αναμενόμενο εύρος, Intervalli previsti, Intervalos previstos, Beklenen Aralıklar, Várt tartományok, הצפויים הטווחים, 予測範囲, 예상 범위, 预期范围值

		CONTROL 1		CONTROL 2		CONTROL 3		CONTROL 4		CONTROL 5			
		min	max	min	max	min	max	min	max	min	max		
pH		7.183	7.213	7.243	7.384	7.414	7.444	7.611	7.641	7.671			
H ⁺	nmol/L	66	61	57	41	39	36	24	23	21			
PCO ₂	mmHg	51.0	58.0	65.0	36.0	41.0	46.0	16.2	20.2	24.2			
PO ₂	kPa	6.8	7.7	8.6	4.8	5.5	6.1	2.2	2.7	3.2			
PO ₂	mmHg	51.8	61.8	71.8	36.0	41.0	46.0	128.9	143.9	158.9			
PO ₂	kPa	6.9	8.2	9.5	4.8	5.5	6.1	17.1	19.1	21.1			
SO ₂	%	47	50	53	77	80	83	88	91	94			
Hct	%	56	59	62	36	39	42	22	25	28			
Na ⁺	mmol/L							139.3	143.3	147.3	114.2	115.2	119.2
K ⁺	mmol/L							3.73	3.98	4.23	5.89	6.19	6.49
Cl ⁻	mmol/L							123.4	127.9	132.4	93.4	97.9	102.4
iCa	mmol/L							1.00	1.08	1.16	1.36	1.48	1.60
iCa	mg/dL							4.0	4.3	4.6	5.5	5.9	6.4
iMg	mmol/L							0.53	0.60	0.67	1.01	1.16	1.31
iMg	mg/dL							1.3	1.5	1.6	2.5	2.8	3.2
Glucose	mg/dL							73	81	89	245	270	295
Glucose	mmol/L							4.1	4.5	4.9	13.6	15.0	16.4
Lac	mmol/L							1.7	2.0	2.3	6.2	6.9	7.6
Lac	mg/dL							15.1	17.8	20.5	55.2	61.5	67.7
HbF*	%	79.0	87.0	95.0	44.6	59.6	74.6	22.8	27.8	32.8			
TbI	g/dL	18.9	20.7	22.5	13.2	14.7	16.2	6.1	7.1	8.1			
iHb	g/L	189	207	225	132	147	162	61	71	81			
iHb	mmol/L	11.7	12.9	14.0	8.2	9.1	10.1	3.8	4.4	5.0			
O ₂ Hb	%	19.3	21.8	24.3	45.7	49.7	53.7	76.2	81.2	86.2			
COHb	%	24.7	28.7	32.7	16.1	20.1	24.1	2.0	6.0	10.0			
MetHb	%	24.3	27.3	30.3	14.9	17.9	20.9	2.4	5.4	8.4			
HbH	%	18.2	22.2	26.2	8.3	12.3	16.3	3.5	7.5	11.5			
TbI*	mg/dL	17.9	21.9	25.9	9.3	11.3	13.3	5.5	5.9	6.3			
TbI*	µmol/L	306.1	374.5	442.9	159.0	193.2	227.4	94.1	100.9	107.7			
TbI*	mmol/L	179.0	219.0	259.0	93.0	113.0	133.0	55.0	59.0	63.0			

ES

Descripción del producto
Material usado de control de calidad para supervisar el desempeño de pH, PCO₂, PO₂, SO₂, hematocrito (Hct), hemoglobina fetal (HbF), hemoglobina total (HbT), bilirubina total (TbI), albúmina total (Alb), oxihemoglobina (O₂Hb), carboxihemoglobina (COHb), metahemoglobina (MetHb) y deoxihemoglobina (HbD) en niveles 1, 2 y 3, además de Na⁺, K⁺, Cl⁻, Ca, Mg, glucosa y lactato en niveles 4 y 5. Para uso ÚNICAMENTE con los analizadores Stat Profile Prime Plus.

Usos indicados
Destinado al uso diagnóstico in vitro por parte de profesionales de la salud para supervisar el desempeño de los analizadores Stat Profile Prime Plus.

Metodología
Para conocer la metodología y los principios de prueba, consulte el Manual de instrucciones de uso del analizador Stat Profile Prime Plus.

Composición
Los controles de nivel 1, 2 y 3 son soluciones tamponadas de bicarbonato que contienen tintura, sales y conservantes. Cada nivel tiene un pH conocido e está equilibrado a un valor conocido de O₂, CO₂ y N₂. Los controles de nivel 4 y 5 son soluciones tamponadas que contienen concentraciones conocidas de Na⁺, K⁺, Cl⁻, Ca, Mg, glucosa, lactato y conservantes. Cada envase contiene 100 mL, como mínimo. Los controles no contienen ninguna sustancia de origen humano. Sin embargo, se deben cumplir las buenas prácticas de laboratorio al manipular estos materiales. (REF: NCCLS DOCUMENT M29-T2)

Advertencias y precauciones:
NO CONGELAR. Mezcle el cartucho invertiéndolo suavemente durante unos segundos. NO AGITAR EL CARTUCHO. Para conocer la información completa, consulte el Manual de instrucciones de uso del analizador Stat Profile Prime Plus.

Almacenamiento
Conserve a 2-8°C (37-46°F). NO CONGELAR.

Modo de empleo
Asegúrese de que los controles estén a temperatura ambiente antes de usar. Mezcle bien el cartucho invertiéndolo suavemente durante 1 minuto. Verifique que el número de lote que figura en la tabla de rangos esperados coincida con el número de lote impreso en el cartucho. Para conocer las instrucciones completas, consulte el Manual de instrucciones de uso del analizador Stat Profile Prime Plus.

Limitaciones
Los valores de PO₂ varían en proporción inversa a la temperatura (aproximadamente 1%/°C). Por lo tanto, es esencial seguir las normas de temperatura que se describen en la sección "Instrucciones de uso". Los valores de rangos esperados son específicos para los instrumentos y controles fabricados por Nova Biomedical. Una vez instalado, cada cartucho Stat Profile Prime Plus se puede utilizar por un máximo de 35 días a partir de la fecha de instalación en el sistema. Pasado ese lapso, el sistema indicará que el cartucho no es válido. Cada cartucho puede ser insertado y retirado del analizador hasta 6 veces como máximo.

Cumplimiento de normas
La hemoglobina total (HbT) y la metahemoglobina (MetHb) son trazables al método de la Cianmetahemoglobina. La carboxihemoglobina (COHb) y la deoxihemoglobina (HbD) son trazables a la técnica espectrofotométrica. Analitos trazables a los materiales de referencia estándar del NIST.

Intervalos de referencia
Las concentraciones están formuladas como valores esperados normales y anormales en el suero del paciente. Se puede consultar el rango clínico esperado de estos valores en Tietz, NW ed. 1985 Textbook of Clinical Chemistry, W.B. Saunders Co. Es posible que los usuarios deseen determinar valores medios y rangos esperados en su propio laboratorio.

Rangos esperados
El rango esperado para cada parámetro ha sido determinado en Nova Biomedical usando determinaciones replicadas en analizadores Nova. El rango esperado indica las desviaciones máximas del valor medio que pueden esperarse bajo condiciones de laboratorio óptimas para instrumentos que funcionan dentro de las especificaciones. Consulte la tabla de rangos esperados.

*No disponible en EE. UU. o para uso en pruebas en punto de atención del paciente.

NCCLS Document M29-T2
How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

How to Define and Determine Reference Intervals in the clinical laboratory, approved guideline-second edition, NCCLS C28-A2, Volume 20, Number 13.

