Effective November 1, 2023, refer to <u>CMS</u> <u>Manual 100-02, Chapter</u> <u>16-General Exclusions</u> <u>from Coverage</u> for services included in this policy.



# Name of Blue Advantage Policy: Analysis of Proteomic Patterns in Serum to Identify Cancer

Policy #: 176

Latest Review Date: May 2023

Category: Laboratory

**ARCHIVED EFFECTIVE 11/1/2023** 

# **BACKGROUND:**

**Blue Advantage** medical policy does not conflict with Local Coverage Determinations (LCDs), Local Medical Review Policies (LMRPs) or National Coverage Determinations (NCDs) or with coverage provisions in Medicare manuals, instructions or operational policy letters. In order to be covered by Blue Advantage the service shall be reasonable and necessary under Title XVIII of the Social Security Act, Section 1862(a)(1)(A). The service is considered reasonable and necessary if it is determined that the service is:

- 1. Safe and effective;
- 2. Not experimental or investigational\*;
- 3. Appropriate, including duration and frequency that is considered appropriate for the service, in terms of whether it is:
  - Furnished in accordance with accepted standards of medical practice for the diagnosis or treatment of the patient's condition or to improve the function of a malformed body member;
  - Furnished in a setting appropriate to the patient's medical needs and condition;
  - *Ordered and furnished by qualified personnel;*
  - One that meets, but does not exceed, the patient's medical need; and
  - At least as beneficial as an existing and available medically appropriate alternative.

\*Routine costs of qualifying clinical trial services with dates of service on or after September 19, 2000 which meet the requirements of the Clinical Trials NCD are considered reasonable and necessary by Medicare. Providers should bill **Original Medicare** for covered services that are related to **clinical trials** that meet Medicare requirements (Refer to Medicare National Coverage Determinations Manual, Chapter 1, Section 310 and Medicare Claims Processing Manual Chapter 32, Sections 69.0-69.11).

## **POLICY:**

Blue Advantage will treat analysis of proteomic patterns in serum to identify cancer as a non-covered benefit and as investigational.

Blue Advantage does not approve or deny procedures, services, testing, or equipment for our members. Our decisions concern coverage only. The decision of whether or not to have a certain test, treatment or procedure is one made between the physician and his/her patient. Blue Advantage administers benefits based on the members' contract and medical policies. Physicians should always exercise their best medical judgment in providing the care they feel is most appropriate for their patients. Needed care should not be delayed or refused because of a coverage determination.

# **DESCRIPTION OF PROCEDURE OR SERVICE:**

The analysis of proteomic patterns in serum for early detection of cancer has been proposed. Several of these proteomic tests are being studied, particularly in ovarian and prostate cancer.

Proteomics involves the use of mass spectrometry to study differences in patterns of protein expression. Proteomic patterns are patterns of proteins present within a clinical sample used as a diagnostic "fingerprint" that is proposed to provide an early diagnosis for some types of cancer.

## **KEY POINTS:**

The most recent literature update was performed through May 17, 2023.

## **Summary of Evidence**

While patterns of protein expression have been proposed to yield more biologically relevant and clinically useful information than assays of single proteins, many limitations in the use of proteomics exist. In contrast to genomics, in which amplification techniques like polymerase chain reaction (PCR) allow for the investigation of single cells, no technology is available at the protein level. Other issues between studies have been lack of uniform patient inclusion and exclusion criteria, small patient numbers, absence of standardized sample preparations, and limited analytical reproducibility. Data in the peer-reviewed literature are inadequate to permit scientific conclusions regarding ovarian, prostate, or other malignancies. The utility of this technology is unproven and is considered investigational.

## **Practice Guidelines and Position Statements**

# National Comprehensive Cancer Network (NCCN) Guidelines

NCCN guidelines for the common cancers addressed in this policy do not comment on the use of proteomics.

## **U.S. Preventative Services Task Force**

Not applicable.

## **KEY WORDS:**

Proteomics, ovarian cancer, OvaCheck<sup>TM</sup>, Correlogic Systems, ProstaCheck, MammoCheck, NovellusDX, Vermillion

# APPROVED BY GOVERNING BODIES:

Clinical laboratories may develop and validate tests in-house and market them as a laboratory service; laboratory-developed tests must meet the general regulatory standards of the Clinical Laboratory Improvement Amendments. Laboratories that offer laboratory-developed tests must be licensed by the Clinical Laboratory Improvement Amendments for high-complexity testing. To date, the U.S. Food and Drug administration has chosen not to require any regulatory review of this test.

## **BENEFIT APPLICATION:**

Coverage is subject to member's specific benefits. Group-specific policy will supersede this policy when applicable.

## **CURRENT CODING:**

#### **CPT Codes:**

There is no specific code for this type of testing. One of the following codes might be used to report the test:

83789	Mass spectroscopy and tandem mass spectrometry (MS, MS/MS), analyte not elsewhere specified, quantitative, each specimen
84999	Unlisted chemistry procedure

## **REFERENCES:**

- 1. Bast RC Jr, Brewer M, Zou C, et al. Prevention and early detection of ovarian cancer: Mission impossible? Recent Results Cancer Res 2007; 174: 91-100.
- 2. Belluco C, Petricoin EF, Mammano E, et al. Serum proteomic analysis identifies a highly sensitive and specific discriminatory pattern in stage 1 breast cancer. Ann Surg Oncol, September 2007; 14(9): 2470-2476.
- 3. Conrads TP, Fusaro VA, Ross S et al. High-resolution serum proteomic features for ovarian cancer detection. Endocr Relat Cancer 2004; 11(2):163-78.
- 4. Conrads TP, Veenstra TD. The utility of proteomic patterns for the diagnosis of cancer. Curr Drug Targets Immune Endocr Metabol Disord 2004; 4(1): 41-50.
- 5. Conrads, T., Zhou, M., Petricoin, E., Liotta, L., & Veenstra, T. (2003). Cancer diagnosis using proteomic patterns. Expert Review in Molecular Diagnostics, 3(4), 411-421.

- 6. Cristea IM, Gaskell SJ, Whetton AD. Proteomics techniques and their application to hematology. Blood 2004; 103(10): 3624-34.
- 7. Diamandis EP. Analysis of serum proteomic patterns for early cancer diagnosis: drawing attention to potential problems, J Natl Cancer Inst 2004; 96(5): 353-6.
- 8. Dziadziuszko R and Hirsch FR. Advances in genomic and proteomic studies of non-small-cell lung cancer: clinical and translational research perspective. Clin Lung Cancer, March 2008; 9(2): 78-84.
- 9. Garrisi VM, Abbate I, Quaranta M, et al. SELDI-TOF serum proteomics and breast cancer: Which perspective? Expert Rev Proteomics 2008; 5(6): 779-785.
- 10. IOM (Institute of Medicine). 2011. Clinical Practice Guidelines We Can Trust. Washington, DC: The National Academies Press.
- 11. Leman ES, Schoen RE, Weissfeld JL, et al. Initial analyses of colon cancer-Specific antigen (CCSA)-3 and CCSA-4 as colorectal cancer-Associated serum markers. Cancer Res 2007; 67(12): 5600-5605.
- 12. Li J, Zhuang Z, Okamoto H et.al. Proteomic profiling distinguishes astrocytomas and identifies differential tumor markers. Neurology 2006; 66(5):733-6.
- 13. Lim, L., Looi, M., Zakaria, S., Sagap, I., Rose, I., Chin, S., et al. (2016). Identification of differentially expressed proteins in the serum of colorectal cancer patients using 2D-DIGE proteomics analysis. Pathology Oncology Research, 22, 169-177.
- 14. Lin Y, Dynan WS, Lee JR, et al. The current state of proteomics in GI oncology. Dig Dis Sci, March 2009; 54(3): 431-457.
- 15. Lomnytska M and Souchelnytskyi S. Markers of breast and gynecological malignancies: The clinical approach of proteomics-based studies. Proteomics Clinical Applications, REVIEWS 2007, Vol. 1, Issue 9, pp. 1090-1101.
- 16. McLerran D, Grizzle WE, Feng Z, et al. SELDI-TOF MS whole serum proteomic profiling with IMAC surface does not reliably detect prostate cancer. Clin Chem 2008; 54(1): 53-60.
- 17. Ornstein DK, Rayford W, Fusaro VA et al. Serum proteomic profiling can discriminate prostate cancer from benign prostates in men with total prostate specific antigen levels between 2.5 and 15.0 ng/ml. J Urol 2004; 172(4 pt 1):1302-5.
- 18. Petricoin EF, Ardekani AM, Hit, BE, et al. Use of proteomic patterns in serum to identify ovarian cancer, Lancet 2002; 359(9306): 572-7.
- 19. Shao, S., Neely, B. A., Kao, T. C., Eckhaus, J., Bourgeois, J., Brooks, J., et al. (2016). Proteomic profiling of serial pre-diagnostic serum samples for early detection of colon cancer in the U. S. military. Cancer Epidemiology Biomarkers Prevention, 2016, Epub ahead of print. Abstract retrieved February 2, 2017 from PubMed database.
- 20. Tanase, C., Albulescu, R., Codrici, E., Popescu, D., Mihai, S., Enciu, A., et al. (2015) Circulating biomarker panels for targeted therapy in brain tumors. Future Oncology, 11 (3), 511-524.
- 21. Unwin RD and Whetton AD. How will haematologists use proteomics? Blood Rev, November 2007; 21(6): 315-326.
- 22. Van Gorp T, Cadron I, Vergote I. The utility of proteomics in gynecologic cancers. Curr Opin Obstet Gynecol, 2011 Feb; 23(1):3-7.

23. Xu, Y., Zhuo, J., Duan, Y., Shi, B., Chen, X. Zhang, X., et al., (2014). Construction of protein profile classification model and screening of proteomic signature of acute leukemia. International Journal of Clinical Expert Pathology, 7 (9), 5569-5581.

# **POLICY HISTORY:**

Adopted for Blue Advantage, November 2006

Available for comment December 5, 2006-January 18, 2007

Medical Policy Group, November 2007

Medical Policy Group, March 2009

Medical Policy Group, March 2010

Medical Policy Group, July 2011

Medical Policy Group, July 2012

Medical Policy Group, August 2013

Medical Policy Group, December 2015

Medical Policy Group, August 2016

Medical Policy Group, June 2017

Medical Policy Group, June 2019

Medical Policy Group, May 2021

Medical Policy Group, October 2021: Reviewed by consensus. No new published peer-reviewed literature available that would alter the coverage statement in this policy.

Medical Policy Group, May 2022: Reviewed by consensus. No new published peer-reviewed literature available that would alter the coverage statement in this policy.

Medical Policy Group, May 2023: Reviewed by consensus. No new published peer-reviewed literature available that would alter the coverage statement in this policy.

Medical Policy Group, November 2023: Archived effective 11/1/2023.

This medical policy is not an authorization, certification, explanation of benefits, or a contract. Eligibility and benefits are determined on a case-by-case basis according to the terms of the member's plan in effect as of the date services are rendered. All medical policies are based on (i) research of current medical literature and (ii) review of common medical practices in the treatment and diagnosis of disease as of the date hereof. Physicians and other providers are solely responsible for all aspects of medical care and treatment, including the type, quality, and levels of care and treatment.

This policy is intended to be used for adjudication of claims (including pre-admission certification, pre-determinations, and pre-procedure review) in Blue Cross and Blue Shield's administration of plan contracts.