

AIA-PACK Series
Tumor Markers



Meet the Challenges of Oncology Testing

Cancer incidence and mortality rates are rapidly growing worldwide. In 2018, the World Health Organization estimated that 9.6 million people worldwide would die from cancer - nearly 1 in 6 global deaths - leading to a significant economic burden. Between 30-50% of cancer deaths are preventable. Early diagnosis, screening and monitoring increase the likelihood of survival and lower morbidity rates.

Tosoh is committed to delivering the highest standard of quality, precision and accuracy in clinical diagnostics to aid in the fight against cancer. Our extensive tumor marker menu offered in a unique dry reagent test cup technology combined with our automated immunoanalyzers provide a comprehensive solution for oncology testing.

TUMOR MARKERS AVAILABLE

Total PSA

27.29

CA 125

CA 19-9

AFP

CEA

TOSOH BIOSCIENCE

www.tosohbioscience.us



Features and Benefits

Tumor Marker	Disease	Indication	Assay Principle	Distinguishing Feature	Benefit
Prostate Specific Antigen (PSA)	Prostate	Screening, diagnosis and monitoring of prostate cancer	Hybritech® Equimolar Response Total PSA sandwich immunoassay	Equimolar assays that shows better diagnostic sensitivity and specificity compared to skewed assays ¹	<ul style="list-style-type: none"> ■ Accurate assay with better discrimination between benign prostatic hyperplasia and prostate cancer² ■ Avoid unnecessary biopsies³
Cancer Antigen 27.29	Breast	Clinical management of breast cancer	Sandwich assay with unique capture and tracer antibodies	Assay targets both epitopes for CA27.29 (B27.29) and CA15-3 (DF3)	Monitoring of breast cancer recurrence, staging and response to therapy with single assay ⁴

Universal Reagent

Tosoh's AIA-PACK test cup format works with every Tosoh automated immunoassay system allowing for a seamless transition from one system to the other, ensuring consistent results in an efficient and economical process.

Save Time and Money

Tosoh's AIA-PACK test cups are single, unitized cups that use a dry reagent format that ensures calibration stability of up to 90 days.

No Contamination, More Traceability

Because there is no transfer of reagents the risk of contamination is eliminated. The unique bichromatic fluorescence kinetic measurement ensures a high analytical and functional sensitivity for all assays. AIA-PACK test cups and trays are labelled with the assay code and lot number for automated scheduling and inventory.

Choose Wisely. Choose Tosoh.



Accurate & precise



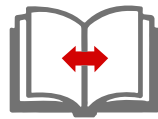
Specific & sensitive



Correlation with gold standard methods



Biotin-free immunoassays



Extensive test menu



Up to 90 day calibration stability



10 min incubation times



Automated workflows

Available for Labs and Workloads of All Sizes



AIA-360



AIA-900
Benchtop



AIA-900
with 9 Tray Sorter
Also available with 19 tray sorter option and as Loader model



AIA-2000

Tosoh products are for Prescription use only as In-Vitro Diagnostics



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References

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2. Andrew W. Roddam, et al. Assessing the Clinical Impact of Prostate-Specific Antigen Assay Variability and Nonequimolarity: A Simulation Study Based on the Population of the United Kingdom Clinical Chemistry. Vol. 50, Issue 6 June, 2004
3. Onur R, et al. Increased discrimination between benign prostatic hyperplasia and prostate cancer with equimolar total prostate specific antigen measurement. World J Urol 2003;21:43-47.
4. Klee, G., et al. MUC1 Gene-Derived Glycoprotein Assays for Monitoring Breast Cancer (CA15-3, CA 27.29, BR). Arch Pathol Lab Med -Vol 128, October 2004.