70-Gene Assay (MammaPrint) **is the ONLY Genomic Assay with FDA Clearance and Category 1 Evidence According to Both National Comprehensive Cancer Network®** (NCCN®) **and American Society of Clinical Oncology** (ASCO®)

AGENDIA®
MAMMAPRINT

National Comprehensive Cancer Network® (NCCN®)

PATIENT GROUP	RECOMMENDATION	CATEGORY OF EVIDENCE
ER/PgR-positive, HER2-negative, **lymph node negative** breast cancer	70-gene assay (MammaPrint) received an NCCN Category 1 Evidence and Consensus recommendation.	1
ER/PgR-positive, HER2-negative, **1-3 lymph node positive** breast cancer	Of the multigene assays considered for lymph node positive breast cancers, only the 70-gene assay (MammaPrint) was given Category 1 endorsement. All other multigene assays for lymph node-positive breast cancer were given a Category 2A recommendation.	1

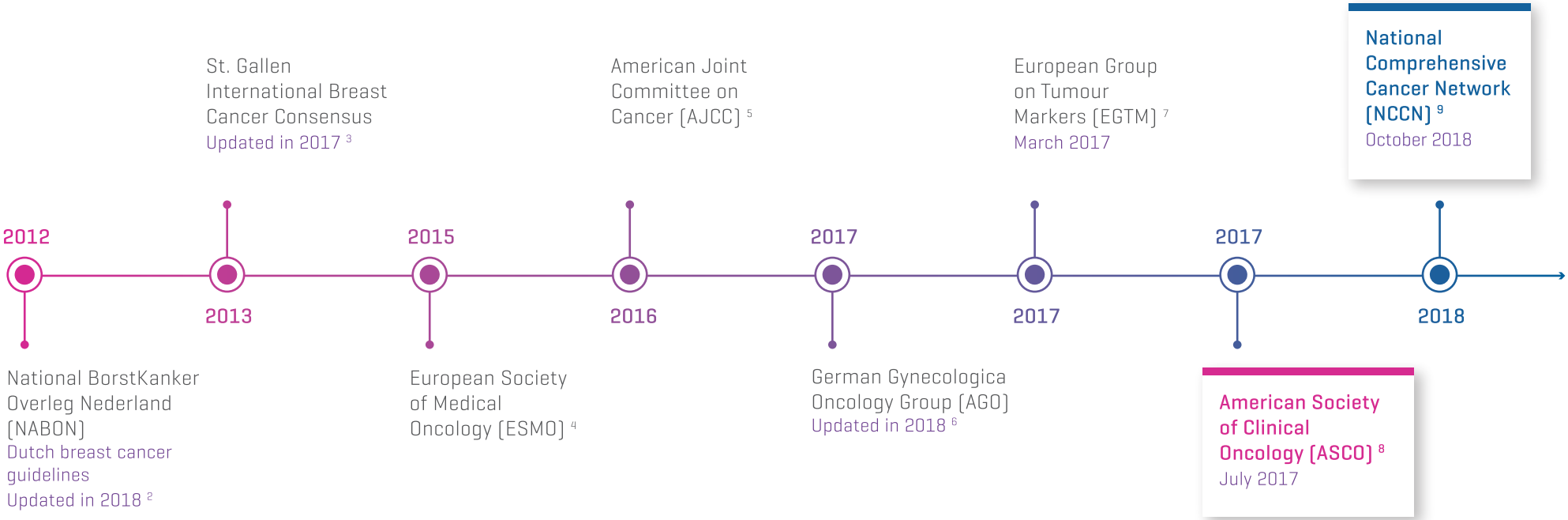
American Society of Clinical Oncology (ASCO®)

PATIENT GROUP	RECOMMENDATION	EVIDENCE
ER/PgR-positive, HER2-negative, **lymph node negative** breast cancer	MammaPrint may be used in patients with high clinical risk per MINDACT categorization to inform decisions on withholding adjuvant systemic chemotherapy due to its ability to identify a good prognosis population with potentially limited chemotherapy benefit.	**TYPE:** Evidence-based **EVIDENCE QUALITY:** High **STRENGTH:** Strong
ER/PgR-positive, HER2-negative, **1-3 lymph node positive** breast cancer	MammaPrint may be used in patients at high clinical risk per MINDACT categorization to inform decisions on withholding adjuvant systemic chemotherapy. However, such patients should be informed that a benefit of chemotherapy cannot be excluded, particularly in patients with greater than one involved lymph node.	**TYPE:** Evidence-based **EVIDENCE QUALITY:** High **STRENGTH:** Moderate

Adapted from Use of Biomarkers to Guide Decisions on Adjuvant Systemic Therapy for Women with Early-Stage Invasive Breast Cancer: American Society of Clinical Oncology Practice Guidelines.

70-Gene Assay (MammaPrint): Recommended in International Breast Cancer Clinical Practice Guidelines

Following the publication of the results of MINDACT [1] in the New England Journal of Medicine in August 2016, the medical societies that reviewed the findings have included or expanded their recommendation of MammaPrint based on this landmark clinical trial.



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