

Patient Name

Peterson, Janette

Date of Birth

22-DEC-1975

EpiSwitch CiRT Response Profile

Your Response Level

Low Probability

High Probability

Patient Information

Gender Female

Medical Record # c14456d0

Report # CiRT0254

Physician Release

Physician Name Christopher Smith, MD, MS

Facility Name XXX-XXX-XXX-XXX

Address [Address line]  
[City], [State] [Zip]

Phone 811-555-4210

Account Ref FL

Specimen Information

Report Date 28-FEB-2021

Receipt Date 28-FEB-2021

Collection Date 20-FEB-2021

Specimen Type WB EDTA K3

Specimen ID D000097422

The **EpiSwitch Checkpoint inhibitor Response Test (CiRT)** result for this specimen is in the **High Likelihood of Response** range to an Immune Checkpoint Inhibitor (ICI) therapy. Individuals in the group have a greater chance of benefiting from an ICI agent. This result should be considered along with other clinical features for interpretation by a licensed medical professional.

EpiSwitch Checkpoint inhibitor Response Test Description

**Intended Use:** Checkpoint inhibitor Response Test (CiRT) is a blood test that evaluates DNA conformational structures in the immune cells to assess the likelihood of response to immune checkpoint inhibitor (ICI) therapy targeting PD-L1 (Atezolizumab, Avelumab, Duralumab) and PD-1 (Pembrolizumab). The results are not specific to ICI agent or to type of tumor. The test is intended to provide supplemental information to cancer treatment professionals on the overall clinical picture. It should not be used as the sole data point in treatment decisions.

**EpiSwitch CiRT Response Test** is a quantitative laboratory developed blood test to assess the likelihood of response to an ICI therapy. The test evaluates eight (8) DNA regulatory (epigenetic) markers called chromatin-conformational signatures (CCS) and stratifies patients based on their probability of responding to an ICI. The test may be run at any stage of treatment, including post-initiation of an ICI.

References

- Hunter, E., Salter, M., et al., (2023). Development and Validation of Blood-Based Predictive Biomarkers for Response to PD-1/PD-L1 Checkpoint Inhibitors: Evidence of a Universal Systemic Core of 3D Immunogenetic Profiling across Multiple Oncological Indications. *Cancers*. 2023; 15(10):2696. <https://doi.org/10.3390/cancers15102696>.
- Hunter, E., Dizfouli, M., et al., (2021). Development and validation of blood-based predictive biomarkers for response to PD-(L)-1 checkpoint inhibitors: evidence of a universal systemic core of 3D immunogenetic profiling across multiple oncological indications. *MedRxiv*. <https://doi.org/10.1101/2021.12.21.21268094>.
- Shah, P., Hunter, E., et al., (2019). Development and validation of baseline predictive biomarkers for response to avelumab in second-line (2L) non-small cell lung cancer (NSCLC) using EpiSwitch epigenetic profiling. *SITC, J. Immunotherapy Cancer* 7(282) P142. <https://dx.doi.org/10.1186/s40425-019-0763-1>. Co- published by authors from Oxford BioDynamics, EMD Serono, Pfizer, and the Mayo Clinic.
- Shah, P., Hunter, E., et al., (2019). Development and validation of baseline predictive biomarkers for response to immuno-checkpoint treatments in the context of multi-line and multi-therapy cohorts using EpiSwitch epigenetic profiling. *SITC, J. Immunotherapy Cancer* 7(282) P143. <https://dx.doi.org/10.1186/s40425-019-0763-1>. Co- published by authors from Oxford BioDynamics, EMD Serono, Pfizer, and the Mayo Clinic.

**Disclaimer:** The EpiSwitch CiRT Response Test is a laboratory developed test (LDT). It has not been reviewed or cleared by the US Food and Drug Administration. The laboratory is certified under the Clinical Laboratory Improvement Amendments (CLIA) to perform high-complexity clinical testing. Decisions regarding patient care and treatment should not be solely based on a single test such as this test, rather, on the independent medical judgment of the treating physician taking into consideration all available information concerning the patient's conditions, including other clinical tests, in accordance with the standard of care in each healthcare setting.

This test was performed at **NEXT Bio Research Services, LLC, 11601 Ironbridge Rd, Chester, VA 23831 – CLIA #49D2104154**. For questions about the report, email [CiRT.TEST@myOBDX.com](mailto:CiRT.TEST@myOBDX.com) or call **1.888.236.8896**.

Medical Director Signature

Date

Patient Name

Peterson, Janette

Date of Birth

22-DEC-1975

EpiSwitch CiRT Response Profile

Your Response Level

Low Probability

High Probability

The **EpiSwitch Checkpoint inhibitor Response Test (CiRT)** result for this specimen is in the **Low Likelihood of Response** range to an Immune Checkpoint Inhibitor (ICI) therapy. This result should be considered along with other clinical features for interpretation by a licensed medical professional.

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Gender Female  
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Report # CiRT0254

Physician Release

Physician Name Christopher Smith, MD, MS  
Facility Name XXX-XXX-XXX-XXX  
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