ORISE Beryllium Lab Guide to Results

display a normal response to beryllium sulfate. A normal test is indicated when all the Day 5 and Day 7 Beryllium Stimulation Index (SI) values are below the Normal Beryllium SI Reference Range. The result of the Beryllium Lymphocyte Proliferation test is ABNORMAL. The cells display an abnormal response to beryllium sulfate. An abnormal test is indicated when two or more Beryllium Stimulation Index (SI) values are above the Normal Beryllium SI Reference Range. BORDERLINE The result of the Beryllium Lymphocyte Proliferation Test is BORDERLINE. The blood cells display an abnormal response in one beryllium stimulation Index (SI) value is above the Normal Beryllium SI Reference Range. The result of the Beryllium Lymphocyte Proliferation test is UNINTERPRETABLE due to unacceptably high variability of cell growth (CV) in one or more data sets. Elevated CV(s) in control or test wells invalidates the corresponding Stimulation Index (SI) values. A repeat BeLPT is recommended. The result of the Beryllium Lymphocyte Proliferation test is UNINTERPRETABLE due to increased cell killing in test wells. A repeat BeLPT in a secondary serum is recommended. PG UNINTERPRETABLE OPE UNINTERPRETABLE OPE VININTERPRETABLE O		
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PM UNINTERPRETABLE Poor Mitogen	The result of the Beryllium Lymphocyte Proliferation test is UNINTERPRETABLE due to poor proliferative response of cells to positive growth controls (mitogens). A repeat BeLPT is recommended.
UNACCEPTABLE sample Low Cell Recovery	Low Cell Recovery: The sample volume was adequate; however, the number of lymphocytes recovered was not sufficient to set up a Beryllium Lymphocyte Proliferation Test. Please send a minimum of 40 ml of sample if repeated.
UNACCEPTABLE sample > 48 Hours	Greater Than 48 Hours: The sample received by the ORISE Beryllium Laboratory was greater than 48 hours old due to shipping delay or wrong draw date. ORISE recommends that specimens be less than 30 hours old at time of processing or results could be compromised. A repeat sample is recommended to be drawn and shipped on Tuesday, Wednesday, or Thursday.
UNACCEPTABLE sample 48 Hours	Receipt at 48 Hours: The sample received by the ORISE Beryllium Laboratory was 48 hours old due to shipping delay or wrong draw date. Specimen age compromised sample viability and quality is not sufficient to set up a Beryllium Lymphocyte Proliferation Test. ORISE recommends that specimens be less than 30 hours old at time of processing or results could be compromised. A repeat sample is recommended to be drawn and shipped on Tuesday, Wednesday, or Thursday.
UNACCEPTABLE sample Quantity Not Sufficient	QNS (Quantity Not Sufficient): Less than 30 ml of sample received. Please send 30 ml of sample at a minimum if repeated.
UNACCEPTABLE sample Unresolved Labeling Errors	Sample Unacceptable: Labeling issues on the tubes or requisition prevented processing. Please resolve and recollect sample. Examples of unresolved labeling errors include: Unlabeled tubes; wrong patient
	collected; incorrect identifiers; and/or other errors preventing confirmation of patient identity. This is the only entry that does not generate a Doctor's report from the Web App.
UNACCEPTABLE sample Other	Sample Unacceptable: XXX
	Note: The laboratory will insert the reason for unacceptable status and/or other information as needed.

Chronic beryllium disease (CBD), a chronic disorder mainly affecting the lungs, is found in a small percentage of persons exposed to beryllium dust. In these individuals, exposure to beryllium elicits a delayed-type hypersensitivity resulting in granulomas. The blood beryllium lymphocyte proliferation test (BeLPT) is an in-vitro test, which measures the response of an individual's lymphocytes to the presence of beryllium sulfate. An abnormal BeLPT indicates sensitivity to beryllium but alone is not diagnostic for chronic beryllium disease. However, individuals that are sensitive to beryllium are at a greater risk for developing CBD. The present lymphocyte proliferation test is the best non-invasive screening method available to detect individuals at risk.

