REFERENCES

- Vanderpump MP; French JM; Appleton D; Tunbridge WM; Kendall-Taylor P. The prevalence of hyperprolactinaemia and association with markers of autoimmune thyroid disease in survivors of the Whickham Survey cohort. Clin Endocrinol (Oxf) 1998; 48(1):39-44.
- Straub RH; Zeuner M; Lock G; Sch olmerich J; Lang B. High prolactin and low dehydroepiandrosterone sulphate serum levels in patients with severe systemic sclerosis. Br J Rheumatol 1997; 36(4):426-32.
- Neidhart M. Elevated serum prolactin or elevated prolactin/cortisol ratio are associated with autoimmune processes in systemic lupus erythematosus and other connective tissue diseases. J Rheumatol 1996; 23(3):476-81.
- Neidhart M Serum levels of interleukin-1 beta, luteinizing hormone, and prolactin correlate with the expression of CD45 isoforms on CD4+ peripheral blood T lymphocytes in healthy women. Ann Hematol 1997; 75(4):155-9.
- Maes M; Mommen K; Hendrickx D; Peeters D; D'Hondt P; Ranjan R; De Meyer F; Scharp'e S. Components of biological variation, including seasonality, in blood concentrations of TSH, TT3, FT4, PRL, cortisol and testosterone in healthy volunteers. Clin Endocrinol (Oxf) 1997; 46(5):587-98.

2016-06-08

Cat#: PR234F (96 Tests)
For Order and Inquiries, please contact
Calbiotech Inc.,
1935 Cordell Ct., El Cajon, CA 92020
Tel (619) 660-6162, Fax (619) 660-6970,
www.calbiotech.com



Prolactin ELISA

Catalog No.: PR234F (96 Tests)

INTENDED USE

For Research Use Only. Not for use in diagnostic procedures.

	96 Tests	
1.	Microwells coated with Streptavidin	12x8x1
2.	Prolactin Standards: 6 vials (ready to use)	0.5 ml
3.	Enzyme Conjugate: 1 bottle (ready to use)	12 ml
4.	TMB Substrate: 1 bottle (ready to use)	12 ml
5.	Stop Solution: 1 bottle (ready to use)	12 ml
6.	20X Wash concentrate: 1 bottle	25 ml

MATERIALS NOT PROVIDED

- Distilled or deionized water
- 2. Precision pipettes
- Disposable pipette tips
- 4. ELISA reader capable of reading absorbance at 450nm
- 5. Absorbance paper or paper towel
- Graph paper

STORAGE AND STABILITY

- Store the kit at 2 8° C.
- 2. Keep microwells sealed in a dry bag with desiccants.
- 3. The reagents are stable until expiration of the kit.
- 4. Do not expose test reagents to heat, sun, or strong light.

WARNINGS AND PRECAUTIONS

- For Research Use Only. Not for use in diagnostic procedures.
- 2. For laboratory use.
- 3. Potential biohazardous materials:

The calibrator and controls contain human source components, which have been tested and found non-reactive for hepatitis B surface antigen as well as HIV antibody with FDA licensed reagents. However, there is no test method that can offer complete assurance that HIV, Hepatitis B virus or other infectious agents are absent. These reagents should be handled at the Biosafety Level 2, as recommended in the Centers for Disease Control/National Institutes of Health manual, "Biosafety in Microbiological and Biomedical Laboratories" 1984.

- Do not pipette by mouth. Do not smoke, eat, or drink in the areas in which specimens or kit reagents are handled.
- The components in this kit are intended for use as an integral unit. The components of different lots should not be mixed.
- 6. It is recommended that standards, control and serum samples be run in duplicate.
- Optimal results will be obtained by strict adherence to this protocol. Accurate and precise pipetting, as well as following the exact time and temperature requirements prescribed are essential. Any deviation from this may yield invalid data.

SPECIMEN COLLECTION HANDLING

- 1. Collect blood specimens and separate the serum immediately.
- Specimens may be stored refrigerated at (2-8° C) for 5 days. If storage time exceeds 5 days, store
 frozen at (-20° C) for up to one month.
- 3. Avoid multiple freeze-thaw cycles.
- 4. Prior to assay, frozen sera should be completely thawed and mixed well.
- 5. Do not use grossly lipemic specimens.

REAGENTS PREPARATION

Prepare 1X Wash buffer by adding the contents of the bottle (25 ml, 20X) to 475 ml of distilled or deionized water. Store at room temperature (20-25°C).

ASSAY PROCEDURE

Prior to assay, allow reagents to stand at room temperature.

Gently mix all reagents before use.

- 1. Place the desired number of coated strips into the holder
- Pipette 25 μl of Prolactin standards, control and patient's sera.
- 3. Add 100 µl of enzyme conjugate to all wells.
- 4. Cover the plate and incubate for 60 minutes at room temperature (20-25°C).
- Remove liquid from all wells. Remove liquid from all wells. Wash wells three times with 300 μl of 1X wash buffer. Blot on absorbance paper or paper towel.
- Add 100 ul of TMB substrate to all wells.
- 7. Incubate for 15 minutes at room temperature.
- 8. Add 50 µl of stop solution to all wells. Shake the plate gently to mix the solution.
- 9. Read absorbance on ELISA Reader at 450 nm within 15 minutes after adding the stopping solution.

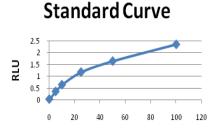
CALCULATION OF RESULTS

The standard curve is constructed as follows:

- Check Prolactin standard values on each standard vial. This value might vary from lot to lot. Make sure you check the value on every kit. See example of the standard attached.
- To construct the standard curve, plot the absorbance for the standards (vertical axis) versus the standard concentrations (horizontal axis) on a linear graph paper. Draw the best curve through the points.
- Read the absorbance for controls and each unknown sample from the curve. Record the value for each control or unknown sample.

Example of a Standard Curve

	OD 450 nm	Conc. ng/mL
Std 1	0.037	0
Std 2	0.363	5
Std 3	0.648	10
Std 4	1.181	25
Std 5	1.647	50
Std 6	2.353	100



Conc.

LIMITATIONS OF THE TEST

1. Do not use sodium azide as preservative. Sodium azide inhibits HRP enzyme activities.