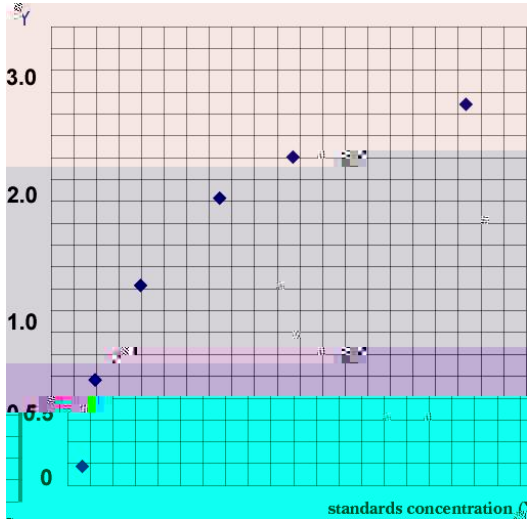
BALP


Y
3.0
2.0
1.0
0.5
0
standards concentration

This BALP ELISA kit is intended Laboratory for Research use only and is not for use in diagnostic or therapeutic procedures.The Stop Solution changes the color from blue to yellow and the intensity of the color is measured at 450 nm using a spectrophotometer. In order to measure the concentration of BALP in the sample, this BALP ELISA Kit includes a set of calibration standards. The calibration standards are assayed at the same time as the samples and allow the operator to produce a standard curve of Optical Density versus BALP concentration. The concentration of BALP in the samples is then determined by comparing the O.D. of the samples to the standard curve.

- Use a serum separator tube and allow samples to clot for 30 minutes before centrifugation for 10 minutes at approximately 3000Ǐg. Remove serum and assay immediately or aliquot and store samples at -20 or -80 .Avoid repeated freeze-thaw cycles

- Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge samples for 30 minutes at 3000Ǐg at 2-8 within 30 minutes of collection. Store samples at -20 or -80 . Avoid repeated freeze-thaw cycles.

Remove particulates by centrifugation and assay immediately or aliquot and store samples at -20 or -80 . Avoid repeated freeze-thaw cycles.

The samples shoule be

1. Standard microplate reader(450nm)
2. Precision pipettes and Disposable pipette tips.
3. 37 incubator

1. Do not substitute reagents from one kit to another. Standard, conjugate and microplates are matched for optimal performance. Use only the reagents supplied by manufacturer.
2. Do not remove microplate from the storage bag until needed. Unused strips should be stored at 2-8ÁC in their pouch with the desiccant provided.
3. Mix all reagents before using.

Remove all kit reagents from refrigerator and allow them to reach room temperature (20-25ÁC)

Name	96 determinations	48 determinations
Microelisa stripplate	12*8strips	12*4strips
Standard	0.3ml*6tubes	0.3ml*6tubes
Sample Diluent	6.0ml	3.0ml
HRP-Conjugate reagent	10.0ml	5.0ml
20X Wash solution	25ml	15ml
Chromogen Solution A	6.0ml	3.0ml
Chromogen Solution B	6.0ml	3.0ml
Stop Solution	6.0ml	3.0ml

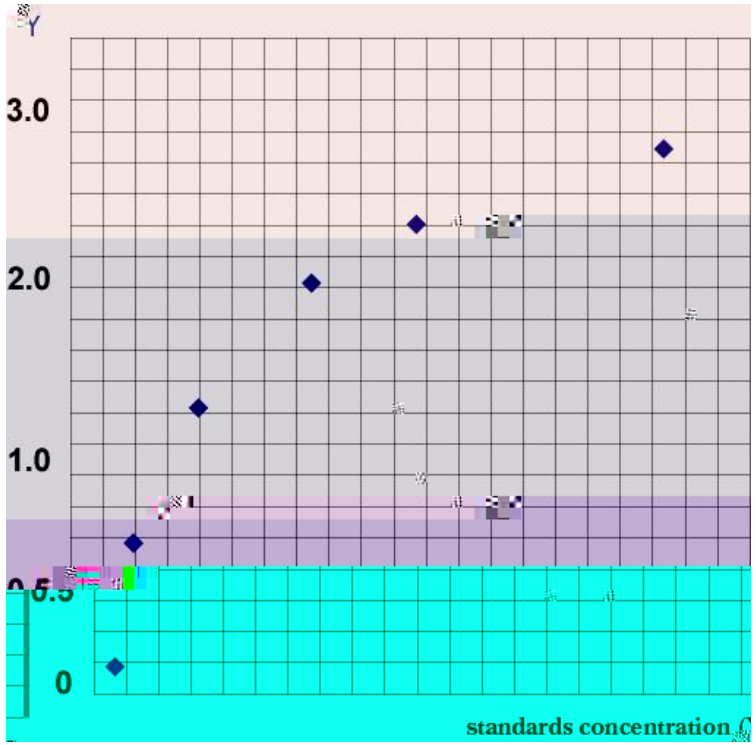
Closure p[illegible] membran[illegible]	2	2
User [illegible]nual	1	1
Sea[illegible]bags	1	1

Standard (S0 Ÿ S5) concentration was followed by:0,0.5,1,2,4,8 ng/ml

20Ťwash solution.Dilute with Distilled or deionized water 1M0.

1. Prepare all reagents before starting assay procedure. It is recommended that all Standards and Samples be added in duplicate to the Microelisa Stripplate.
2. Add standard: Set Standard wells, testing sample wells. Add standard 50εl to standard well.
3. Add Sample: Add testing sample 10εl then add Sample Diluent 40εl to testing [illegible] well; Blank we[illegible]oesnôt [illegible] anyting.
4. Add 100

1. This standard curve is used to determine the amount in an unknown sample. The standard curve is generated by plotting the average O.D. (450 nm) obtained for each of the six standard concentrations on the vertical (Y) axis versus the corresponding concentration on the horizontal (X) axis.



Storage 2-8 .
validity six months.