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RAT GROWTH HORMONE ENZYME IMMUNOASSAY KIT

catalogue # A05104

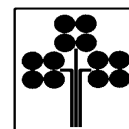
96 wells

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U.S. patent # 50 47 330
European patent # 89 139 552

**THE RAT GROWTH HORMONE
ENZYME IMMUNOASSAY HAS
BEEN DEVELOPED AND
VALIDATED BY SPI-BIO.**

*For research laboratory use only.
Not for human diagnostic use.*



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RAT GROWTH HORMONE EIA KIT

96 wells

Storage: -20°C

Expiry date: stated on the package

This kit contains:

- ☞ A covered 96 well plate, pre-coated with rabbit polyclonal anti-goat immunoglobulin antibodies, ready to use after thawing
- ☞ One vial of Rat growth hormone tracer, lyophilised
- ☞ Two vials of Rat growth hormone standard, lyophilised
- ☞ One vial of Rat growth hormone antiserum, lyophilised
- ☞ One vial of EIA buffer, lyophilised
- ☞ One vial of concentrated Wash buffer, liquid
- ☞ One vial of tween 20, liquid
- ☞ Two vials of Quality Control sample, lyophilised
- ☞ Two vials of Ellman's reagent, lyophilised
- ☞ One instruction booklet
- ☞ One template sheet
- ☞ One well cover sheet

Each kit contains sufficient reagents for 96 wells. This allows for the construction of one standard curve in duplicate and the assay of 33 samples in duplicate.

PRECAUTIONS FOR USE

Users are recommended to read all instructions for use before starting work.

Each time a new pipet tip is used, aspirate a sample or reagent and dispense it back into the same vessel. Repeat this operation two or three times before distribution.

For research laboratory use only.

Not for human diagnostic use.

Do not pipet liquids by mouth.

Do not use kit components beyond the expiration date.

Do not eat, drink or smoke in area in which kit reagents are handled.

Avoid splashing.

The QC samples provided in this kit have been prepared by diluting rat plasma (Sprague Dawley rat) in EIA buffer. A sanitary control has been completed on Sprague Dawley rats following the Felasa Health Monitoring Recommendations. However, handle the CQ samples as a possible source of infection.

The total amount of reagents contains less than 100 µg of sodium azide. Flush the drains thoroughly to prevent the production of explosive metal azides.

PRINCIPLE OF THE ASSAY

This Enzyme Immunoassay (EIA) is based on the competition between unlabelled rat growth hormone (rGH) and acetylcholinesterase (AChE) linked to rat growth hormone (tracer) for limited specific goat anti-rat GH antiserum sites.

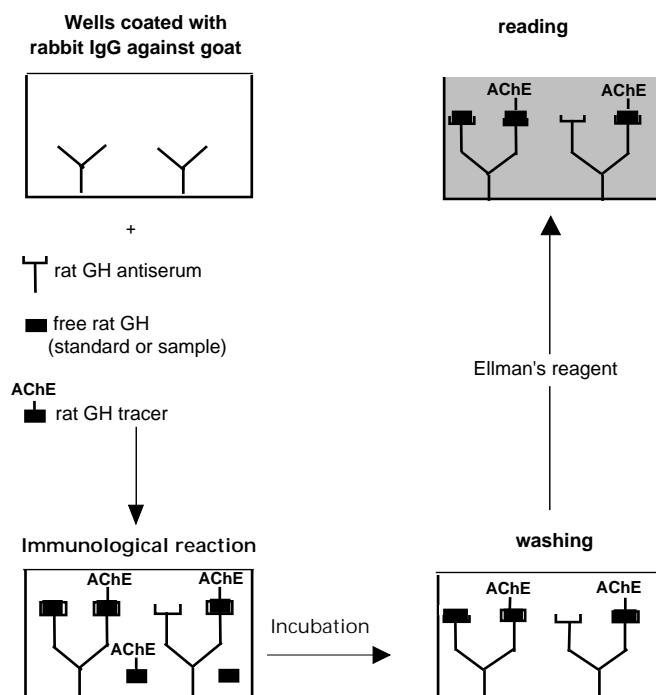
The complex goat antiserum-rat GH (free GH or tracer) binds to the rabbit polyclonal antibody anti-goat immunoglobulin antibodies that are attached to the well.

The plate is then washed and Ellman's Reagent (enzymatic substrate for AChE and chromogen) is added to the wells.

The AChE tracer acts on the Ellman's Reagent to form a yellow compound.

The intensity of the colour, which is determined by spectrophotometry, is proportional to the amount of tracer bound to the well and is inversely proportional to the amount of free rat GH present in the well during the immunological incubation.

The principle of the assay is summarised below:



MATERIALS AND EQUIPMENT REQUIRED

In addition to standard laboratory equipment, the following material is required:

- ☞ Precision micropipettes (20 to 1000 μ L)
- ☞ Spectrophotometer plate reader (405 or 414 nm filter)
- ☞ Microplate washer (or washbottles)
- ☞ Microplate shaker
- ☞ Distilled or deionized water
- ☞ Polypropylene tubes



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SAMPLE COLLECTION & PREPARATION

This assay may be used to measure rat GH in cell culture media or in plasma samples (see Bibliography hereafter for further information).

Blood samples may be collected in tubes containing EDTA, lithium heparin, potassium oxalate or sodium citrate. The samples are centrifuged at 1 600 g for 20 minutes. Plasma are collected and kept at -20°C until assay.

No prior extraction procedure is necessary to measure GH in plasma samples and cell culture media.

REAGENT PREPARATION

The coated plates and reagents are provided ready to use.

☞ EIA buffer

Reconstitute one vial with 50 mL of distilled or deionized water. Use a magnetic stirrer to mix the contents. Stability at 4°C: 1 month.

☞ Rat GH standard (calibrated against the reference preparation NIDDK standard rGH-RP2)

Reconstitute the vial with 1 mL of distilled or deionized water. Allow it to stand 5 minutes until completely dissolved and then mix thoroughly by gentle inversion. The concentration of the first standard is 40 ng/mL. Prepare seven propylene tubes (for the seven other standards) and add 500 µL of EIA buffer into each tube. Add 500 µL of the first tube (containing the first standard) to the second tube. Continue this procedure for the other tubes. Thus, standard concentrations are: 40 (S1), 20 (S2), 10 (S3), 5 (S4), 2.5 (S5), 1.25 (S6), 0.63 (S7) and 0.31 ng/mL (S8), respectively. Stability at 4°C: 1 week.

☞ Quality control (on day 1)

Reconstitute one vial with 1 mL of distilled or deionized water. Allow it to stand 5 minutes until completely dissolved and then mix thoroughly by gentle inversion. Stability at 4°C: 1 week.

☞ Rat GH antiserum (on day 1)

Reconstitute one vial with 5 mL of EIA buffer. Allow it to stand 5 minutes until completely dissolved and then mix thoroughly by gentle inversion. Stability at 4°C: 1 week.

☞ Wash buffer (on day 1)

Dilute 1 mL of the concentrated wash buffer to 400 ml with distilled or deionized water. Add 200 µL of tween 20 (Use a magnetic stirrer to mix the contents). Stability at 4°C: 1 week.

☞ Rat GH-AChE tracer (on day 2)

Reconstitute one vial with 5 mL of EIA buffer. Allow it to stand 5 minutes until completely dissolved and then mix thoroughly by gentle inversion. Stability at 4°C: 1 month.

☞ Ellman's Reagent (on day 3)

Five minutes before use, reconstitute with 50 mL of distilled or deionized water. The tube contents should be thoroughly mixed. Stability at 4°C and in the dark: 4 days.

ASSAY PROCEDURE

It is recommended to perform the assays in duplicate and to follow the instructions hereafter.

PLATE PREPARATION

Prepare the wash buffer as indicated in the reagent preparation section. Open the plate packet and select the sufficient strips for your assay and place the unused strips back in the packet (stored at 4°C). Rinse each well five times with the wash buffer (300 µL/well).

Just before distributing reagents and samples, remove the buffer from the wells by inverting the plate and shaking out the last drops.



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INCUBATING THE PLATE (SECOND STEP)

Cover the plate with a plastic film and incubate for 20 hours at room temperature.

DEVELOPING AND READING THE PLATE

Reconstitute the wash buffer and Ellman's Reagent as indicated in reagent preparation section. Empty the plate by turning over and shaking. Then, wash each well five times with the wash buffer (300 μ L/well). Dispense 200 μ L of Ellman's Reagent to the 96 wells. Incubate in the dark (plate covered with an aluminium sheet) at room temperature. Optimal development is obtained using an orbital shaker. The plate should be read between 405 and 414 nm (yellow colour) when the Maximum Binding (B_0) wells reach an absorbance of 0.2-0.8 unit.

Enzyme Immunoassay Protocol (Volumes are in μ L)					
	Blank	Non specific binding	Maximum binding	Standard	Sample
Buffer	-	100	50	-	-
Standard	-	-	-	50	-
Sample	-	-	-	-	50
Antiserum	-	-	50	50	50
Tracer	-	50	50	50	50
Ellman's reagent	200	200	200	200	200
Cover the plate, incubate at 20°C for 20h Cover the plate, incubate at 20°C for 20h Wash the plate 5 times Incubate the plate with an orbital shaker in the dark at room temperature Read the plate between 405 and 414 nm					

DATA ANALYSIS

Make sure that your plate reader has subtracted the absorbance readings of the blank well (absorbance of Ellman's reagent) from the absorbance readings of the rest of the plate. If not, do it now.

- ↪ Calculate the average absorbance for each NSB, B_0 , standards and samples.
- ↪ Calculate the B/ B_0 (%) for each standard and sample: (average absorbance of standards or sample - average absorbance of NSB) divided by (average absorbance of B_0 - average absorbance of NSB) & multiplied by 100.
- ↪ Using a semi-log graph paper, plot the B/ B_0 (%) for each standard point (y axis) versus the concentration (x axis). Draw a best-fit line through the points.
- ↪ To determine the concentration of your samples, find the B/ B_0 (%) value on the y axis. Read the corresponding value on the x axis which is the concentration of your unknown sample. Samples with a concentration greater than 40 ng/mL should be re-assayed after dilution in EIA buffer.
- ↪ Most plate readers are supplied with curve-fitting software capable of graphing this type of data (logit/log or 4-parameter). If you have this type of software, we recommend using it. Refer to it for further information.



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TYPICAL DATA

EXAMPLE DATA

The following data are for demonstration purpose only. Your data may be different and still correct. These data were obtained using all reagents as supplied in this kit under the following conditions: 2 hours developing at 20°C, reading at 414 nm. A logit/log curve fitting was used to determine the concentrations.

	mAU	B/Bo (%)
NSB	3	
Bo	522	100
Standard 40 ng/mL	56	10.2
Standard 20 ng/mL	97	18.1
Standard 10 ng/mL	139	26.2
Standard 5 ng/mL	195	37.0
Standard 2.5 ng/mL	259	49.3
Standard 1.25 ng/mL	339	64.7
Standard 0.63 ng/mL	405	77.5
Standard 0.31 ng/mL	441	84.4
QC	241	

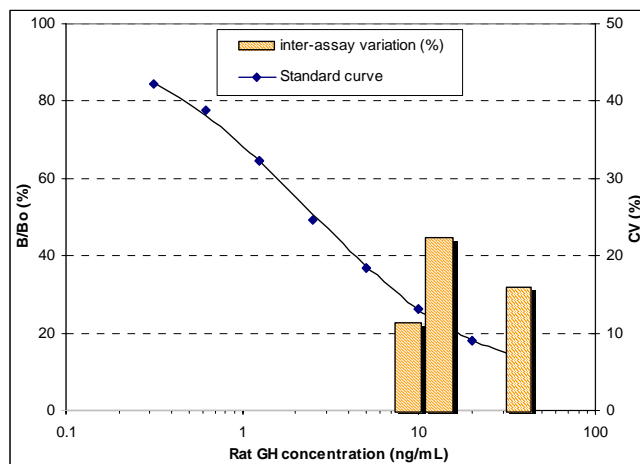
ACCEPTABLE RANGE

☞ Bo absorbance: > 200 mAU in the conditions indicated above.

☞ Ratio NSB absorbance / Bo absorbance: < 0.1.

☞ 50% B/Bo (%): 2.9 to 4.3 ng/mL (mean: 3.6 ng/mL).

RAT GH STANDARD CURVE



ASSAY VALIDATION AND CHARACTERISTICS

The Enzyme Immunoassay of rat GH has been validated by Ezan *et al.* for its use in rat plasma and cell culture media (see Bibliography hereafter). Its main characteristics are the following:

☞ Cross-reactivity:

Compound	%	Compound	%
Mouse GH	91 %	Human GH	<0.1 %
Rat prolactin	<1 %	Rat TSH	<0.1 %
Rat FSH	<0.1 %	Rat LH	<0.1 %

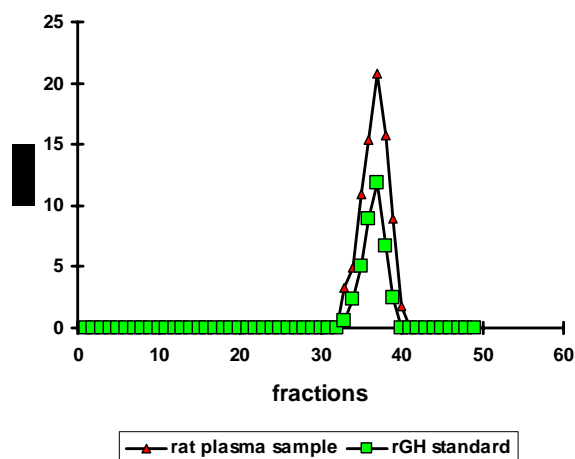
☞ The limit of detection calculated as the concentration producing 15 % displacement of initial tracer: 0.5 ng/mL

☞ Intra- and inter-assay coefficients of variation are 4 % (n=24) and 14 % (n=9).

☞ Accuracy in a hypophysectomised rat plasma sample:

GH added	GH measured	Recovery
0 ng/mL	0 ng/mL	-
0.6 ng/mL	0.5 ng/mL	83 %
1.2 ng/mL	1.3 ng/mL	108 %
2.5 ng/mL	2.5 ng/mL	100 %
5 ng/mL	5 ng/mL	100 %
10 ng/mL	10.1 ng/mL	101 %
20 ng/mL	24.4 ng/mL	122 %

☞ Specificity: comparison of HPLC profiles of rGH standard and a plasma sample.



ASSAY TROUBLE SHOOTING

- ☞ Bo value is too low: incubation in wrong conditions (time or temperature) or reading time too short or Rat GH-AChE tracer, Rat GH antiserum or Ellman's reagent have not been dispensed.
- ☞ NSB value too high: contamination of NSB wells with Rat GH antiserum or inefficient washing.
- ☞ High dispersion of duplicates: poor pipetting technique or irregular plate washing.
- ☞ IC₅₀ or QC concentrations not within the expected range: wrong preparation of standards.
- ☞ Analyses of two dilutions of a biological sample do not agree: Interfering substances are present. Sample must be purified prior to EIA analysis (excepting plasma samples).

These are a few examples of trouble shooting that may occur. If you need further explanation, SPI-BIO will be happy to answer any questions or information about this assay. Please feel free to contact our technical support staff by letter, phone (33 (0)1 39 30 62 60), fax (33 (0)1 39 30 62 99) or E-mail (spibio@spibio.com), and be sure to indicate the lot number of the kit (see outside of the box).

SPI-BIO offers a training workshop in EIA practice & theory. This workshop is given twice a year. For further information, please contact our Customer Relation Representative (33 (0)1 39 30 62 60).

BIBLIOGRAPHY

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