

**USE OF RECOMBINANT HUMAN PROLACTIN FOR
RADIOIODINATION AND STANDARD PREPARATION
IN RADIOIMMUNOASSAY**

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Recombinant human prolactin (hPRL) with a 6His affinity tag and a cleavage site for factor Xa was expressed in *Escherichia coli* (E.coli) using a high-level expression vector and obtaining the hormone in the periplasmic space of the bacteria. 6His-11e-Glu-Gly-Arg-hPRL after purification and characterization was used for the preparation of RIA reagents. Standard curves (logit %B/Bo x log dose) had a good parallelism when recombinant (rec) 6His-11e-Glu-Gly-Arg-hPRL and a pituitary (pit) hPRL were used as reference preparations in a RIA system, with ^{125}I pit hPRL as a tracer. The slopes at the standard curves were 0.935 for pit hPRL and 0.971 for rec hPRL with a correlation coefficient of 0.990 and 0.996 respectively. When using ^{125}I rec hPRL as a tracer, the slopes obtained were 1.1131 for pit hPRL and 1.0874 for rec hPRL and the respective correlation coefficients were 0.999 and 0.994.

Unknown samples (n = 11) at various concentrations (from 5 to 300 ng/mL) were determined using a pituitary reference preparation to study the correlation between the values obtained with the use of the tracers ^{125}I -6His-11e-Glu-Gly-Arg-hPRL and ^{125}I -pit hPRL prepared by the Chloramine T iodination method. In the correlation curve practically no bias was introduced (slope = 0.993) while the correlation coefficient was $r = 0.997$.

These preliminary data suggest that this recombinant hPRL preparation is suitable for the preparation of tracer and for use as a standard preparation in RIA assay.

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