

BASIC AND CLINICAL EVALUATION FOR PTH RADIOIMMUNO-
ASSAY KIT EIKEN

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Radioimmunoassay of parathyroid hormone(PTH) is
thought to be one of the most difficult assays be-
cause of the heterogeneity of PTH and the lower sen-
sitivity of the method. Recently, a sensitive and
high specific radioimmunoassay procedure for PTH
was developed. We checked the PTH kit (EIKEN) ba-
sically, and determined serum PTH levels in normal
subjects and various parathyroid diseases. The re-
sults were as follows.

(1) Standard curve showed dose-responses from 0.1
to 20.0ng/ml.

(2) The cross-reactivities to other peptides,
ACTH, LH, FSH, GH, CT, PTH 1-34, TSH and ADH were
not observed.

(3) The dilution curves of serum PTH of primary
hyperparathyroidism and chronic renal failure were
parallel to this standard curve.

(4) The coefficients of variation for intraassay
were 4.17% and for interassay were 9.0%. Correla-
tion between this assay and our own method revealed
good, $r=0.86$ ($n=100$).

(5) The concentrations of serum PTH were as follows:
normal subjects, 0.43 ± 0.66 (M \pm S.D.) ng/ml ($n=20$),
patients with primary hyperparathyroidism, $4.75\pm$
 2.18 ng/ml ($n=13$); chronic renal failure, 4.43 ± 0.21
ng/ml ($n=35$); hypercalcemia, 0.54 ± 0.74 ng/ml ($n=5$);
idiopathic hypoparathyroidism, 0.15 ± 0.22 ng/ml ($n=$
 16); pseudo-hypoparathyroidism, 0.83 ± 0.91 ng/ml ($n=$
 3); osteomalacia, 0.75 ± 0.87 ng/ml ($n=18$).

(6) Serum PTH levels in the most of the patients
with primary hyperparathyroidism were not suppres-
sed during Ca infusion for 4 hours.

A RADIOIMMUNOASSAY FOR PARATHYROID HORMONE
BY THE USE OF BOVINE 1-34 PTH ANTIBODY

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A sensitive and specific radioimmunoassay
for parathyroid hormone(PTH) was devised by
the use of rabbit antiserum to synthetic
bovine 1-34 PTH(Beckman).

The PTH antibody prepared by immunization
with the synthetic bovine 1-34 PTH, conjug-
ated with glutaraldehyde to BSA, showed a
high titer of about 1:20,000.

A preparation of synthetic bovine 1-34 PTH
was radioiodinated by the modified method
of Hunter & Greenwood, that is, chromamine
T procedure described in this report.

125 I-bPTH was purified by means of adsorp-
tion on Quso G-32 powder(Sigma).

Antibody-bound 125 I-bPTH(B) and antibody
free 125 I-bPTH(F) were separated by adsorp-
tion on dextran-coated charcoal (D-C-C).

The minimal detectable quantity of PTH by
this assay system was 100 pg/ml in plasma.

Synthetic h.ACTH, h.calcitonin, h.GH, TSH, so-
matostatin, LH-RH, glucagon, insulin, neuroten-
sin and FSH showed not significant cross re-
action in the range of 100 pg/ml-10 ng/ml.

PTH concentrations in the plasma of health-
y subjects were not detectable-290 pg/ml.

PTH concentrations in the plasma of patien-
ts with primary hyperparathyroidism were
190-510 pg/ml (342.0 ± 90.6 pg/ml, M \pm sD).

PTH concentrations in the plasma of patie-
nts with renal insufficiency were 100-340
pg/ml.

The values for the patients with primary
hyperparathyroidism overlapped with those
for the healthy subjects.