

in intrinsic spatial resolution does, in fact, interfere with lesion detection.

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Analytical Performance of the ARIA II automatic system for TSH Assay

The widespread use of radioimmunological techniques in clinical practice has prompted numerous attempts to automate radioimmunoassay (RIA) partially or completely. The evaluation of the performances of these automatic systems has to be done not only in terms of practicability and throughput, but also and chiefly in terms of analytical reliability.

The fully automatic system ARIA II* has attained some popularity in laboratories, mainly for assaying T_3 , T_4 , and TSH. The performances of this system for T_3 and T_4 measurements have already been evaluated, either from results produced in a single laboratory (1-3) or from data gained in interlaboratory survey (4).

We report here estimates of the accuracy and precision of the TSH determinations carried out using ARIA II. This evaluation is based on data collected from a national external quality-control survey (EQCS) (5,6), which involved about 150 laboratories, nine of them being ARIA II users. The analysis was performed on the results of 51 EQCS samples sent in 11 monthly dispatches from December, 1981, to May, 1983; the majority of these samples (36) were unidentified replicates for the estimation of the between-laboratory, between-batch precision.

The precision was computed subdividing the results into two concentration ranges; the precision (CV) achieved by ARIA II users was 28.6% CV% (for samples with concentrations in the range 3-5 $\mu\text{IU/ml}$) and 17.2 CV% (range 5-20 $\mu\text{IU/ml}$). For comparison, the precisions of the other five most popular kits used in the survey, turned out as follows (respectively for the low and the high samples): Corning Immophase (7 labs) CV = 13.4 and 10.9 CV%, Cis-Sorin (9 labs) 18.1 and 12.6 CV%, Byk-Mal-linckrodt (36 labs) 18.0 and 17.5 CV%, Biodata-Serono (27 labs) 28.7 and 20.6 CV%, Diagnostic Product Corp. (15 labs) CV = 34.7 and 18.7 CV%.

The accuracy of ARIA II was estimated with respect to the median (after rejection of outliers) of all results reported by participants in the survey. The results, shown in Fig. 1, indicate that ARIA II system consistently underestimates TSH concentrations. This negative bias was confirmed by the results of three recovery experiments (see Table 1) carried out by sending to the participants

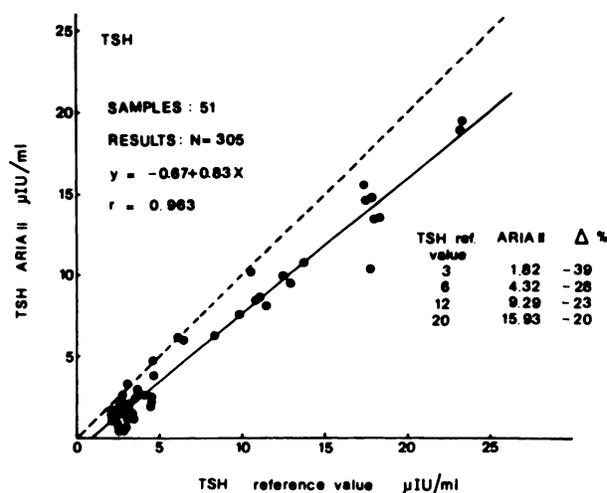


FIG. 1. Regression analysis of 305 results from laboratories using ARIA II system, against respective consensus medians taken as reference values. Closed circles represent mean values of ARIA II results found in each EQCS sample; regression and identity are shown as full and dashed lines respectively. Inset table reports mean readings by ARIA II users (computed from regression line) corresponding to four TSH levels, together with % deviations.

low-concentration samples spiked with known amounts of TSH standard (First IRP WHO 68/38 supplied by NIBSC, Holly Hill, Hampstead, London, UK).

We conclude that the TSH assays of the ARIA II system are not as good as those found for T_3 and T_4 —in fact, ARIA II measurements of TSH are clearly inaccurate and, in addition, do not display better precision than that achieved by the nonautomated methods or kits.

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FOOTNOTE

* Becton-Dickinson Immunodiagnosics, 180 West 2950 South, Salt Lake City, UT 84115, USA.

TABLE 1. TSH RECOVERY OF ARIA II

Sample No.	Median of ARIA II results ($\mu\text{IU/ml}$)	TSH diff.	TSH 68/38 added	Recovery %
C092	18.90			
C090	1.70	17.20	22.5	76.4
C107	6.25			
C105	1.30	4.95	6.0	82.5
C112	13.45			
C109	2.70	10.75	15.0	71.7

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Announcement from the Editor's Office

On January 1, 1985, Thomas P. Haynie, M.D. will assume the Editorship of *The Journal of Nuclear Medicine*. To ensure a smooth transition of duties, it is necessary that Dr. Haynie initiate manuscript processing in September, 1984. Beginning September 15, 1984, all manuscripts, original submissions, and revisions, should be forwarded to:

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