# Analytical evaluation of chemiluminescence in alpha - fetoprotein estimation.

Thesis

Submitted for partial fulfillment for Master degree in Clinical pathology.

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حجق الله العظيم

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#### ABBREVIATION LIST

AE : Acridinium ester

AFAFp: Amniotic fluid alpha fetopriein

AFp: Alpha fetoprotein

ALP: Alkaline phosphatase

AMPPD: 3-(2'-spiroadamantane) - 4 - methoxy - 4 - (3" phosphoryloxy)

phenyl - 1, 2 - dioxetane

ATP: Adenosine triphophosphate

CEA: Carcinoembryonic antigen

CIEP: Counter immunoelectrophoresis

CLLA: chemiluminescence immunoassav

CNR: Concanavalin A non-reactive

Con A: Concanavalin A

CT: Computerized tomography

CV: Coefficient of variation

DDL4: Double diffusion in agar

DELFLA: Delayed enhanced lanthanide fluorescent immunoassay

DMAE: Dimethylated acridinium ester

EL4: Enzyme immunoassay

ELISA: Enzyme linked immunosorbent assay

FDA: Food and Drug A im \*\*t sin

FLAEC: Flow injection analyzer electrochemical detection system

G0: Glucose oxidase

G: : One step glutaraldehyde

HCC: Hepatocellular carcinoma (primary hepatic malignancy)

ICMA: Immunochemiluminometric assav

IEP: Immunoelectrophoresis

IRMA: Immunoradiometric assav

M-AFP: Monoclonal Alpha fetoprotein

M-ICMA: Monoclonal Immunochemiluminometric assay

M-ab: Monoclonal antibody

MSAFP: Maternal serum Alpha fetoprotein

NTD: Neural tube defect

ONTDS: Open Neural tube defect

P-AFP: Polyclonal Alpha fetoprotein

P-ICMA: polyclonal Immunochemiluminometric assay

P : Probability

PNPP: Paranitrophenyl phosphate

 $RAI(^{125}I)$ : Radioactive iodine ( $^{125}I$ )

RIA: Radioimmunoassay

RLU: Relative light unit

SB: Saccharides and Schiff base

U/S: Ultrasonography

#### AMINO ACID ABBREVIATION

A Sp: Aspartic acid

Thr: Threonine

Ser : Serine

Glu: Glutamic acid

Pro: Proline

Gly: Glycine

Ala: Alanine

Cys: Cysteine

Val: Valine

Met: Methionine

Ile: Isoleucine

Leu: Ieucine

Tyr: Tyrosine

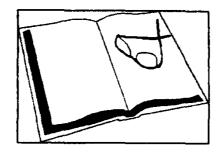
Phe: Phenyl alanine

Lys: Lysine

His: Histidine

Arg: Arginine

Trp: Tryptophan



# INTRODUCTION AND AIM OF THE WORK

### INTRODUCTION AND AIM OF THE WORK

#### INTRODUCTION

Measurement of serum alpha fetoprotein (AFp) plays a very important role in diagnosis, predicting prognosis and offering follow- up programs for many serious diseases, either malignant or not. The development of immunoassay techniques during the last three decades have permitted a sensitive quantification of AFp.

Problems of performance and stability are common to all immunoassays, which utilize  $I^{\omega}$  since it is a high energy emitter with a half life of only sixty days. Several disadvantages are encountered in the technique, including counting time (Several minutes). Inconsistency in separation of the free and bound complexes and radiation hazards.

Therefore non-isotopic immunoassay techniques have been introduced including Fluorescent immunoassay and ELISA which solved some of the previously mentioned RIA limitations. However new problems arouse with the use of the ELISA technique. The most critical factors for reaction were the time and temperature accuracy.

Introduction and Aim of the Work

Recently chemiluminescent compounds are introduced as labels for immunoassay. These compounds have several advantages over RIA and even ELISA techniques. These include very short counting time (Seconds), long reagent stability, safety, temperature independent reactions, no centrifugation and incubation could be done at room temperature

#### AIM OF THE WORK

Evaluation of the chemiluminescence technique as regards reliability characteristics as precision, sensitivity, specificity and accuracy as well the practicability characteristics.

Introduction and Aim of the Work



# Review of Literature