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شبكة المعلومات الحامعية

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شبكة العلومات الحامعية



شبكة المعلومات الجامعية التوثيق الالكتروني والميكروفيلم





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جامعة عين شمس

التوثيق الإلكتروني والميكروفيلم

قسو

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بالرسالة صفحات لم ترد بالأصل



BLOOD LEAD LEVEL IN PRETERM AND FULL-TERM DELIVERING WOMEN AND THEIR NEONATES

Thesis

SUBMITTED FOR PARTIAL FULFILLMENT OF M.Sc. DEGREE
IN CLINICAL BIOCHEMISTRY

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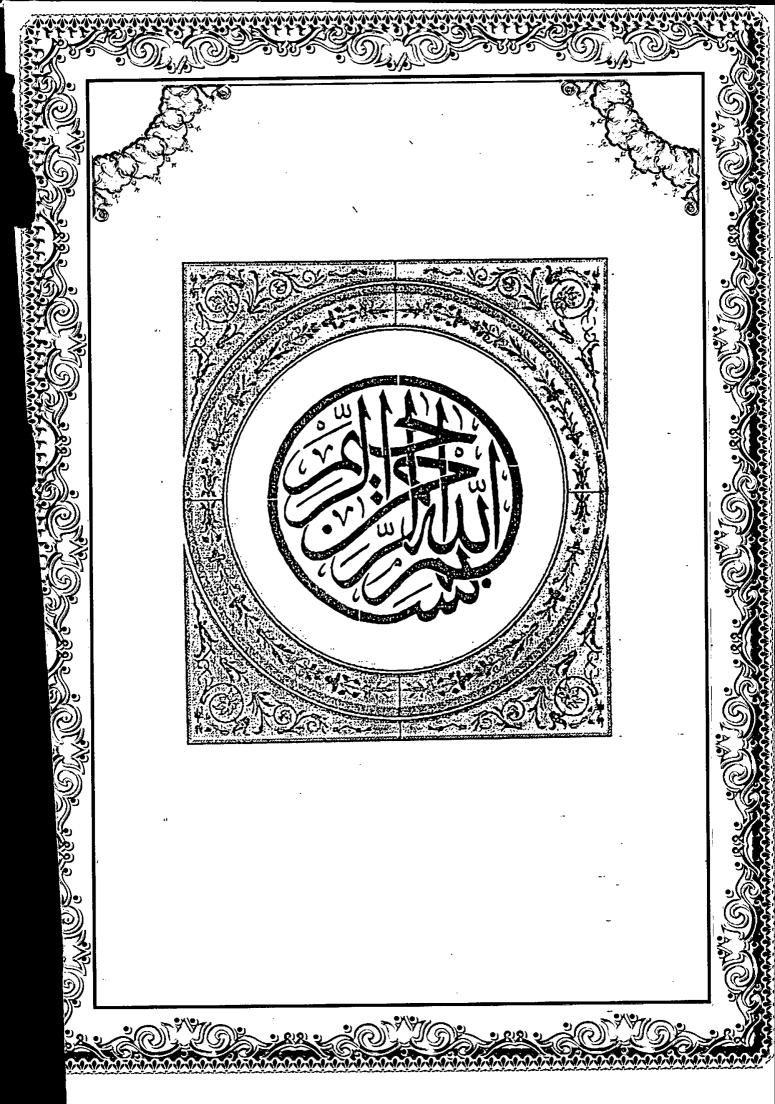
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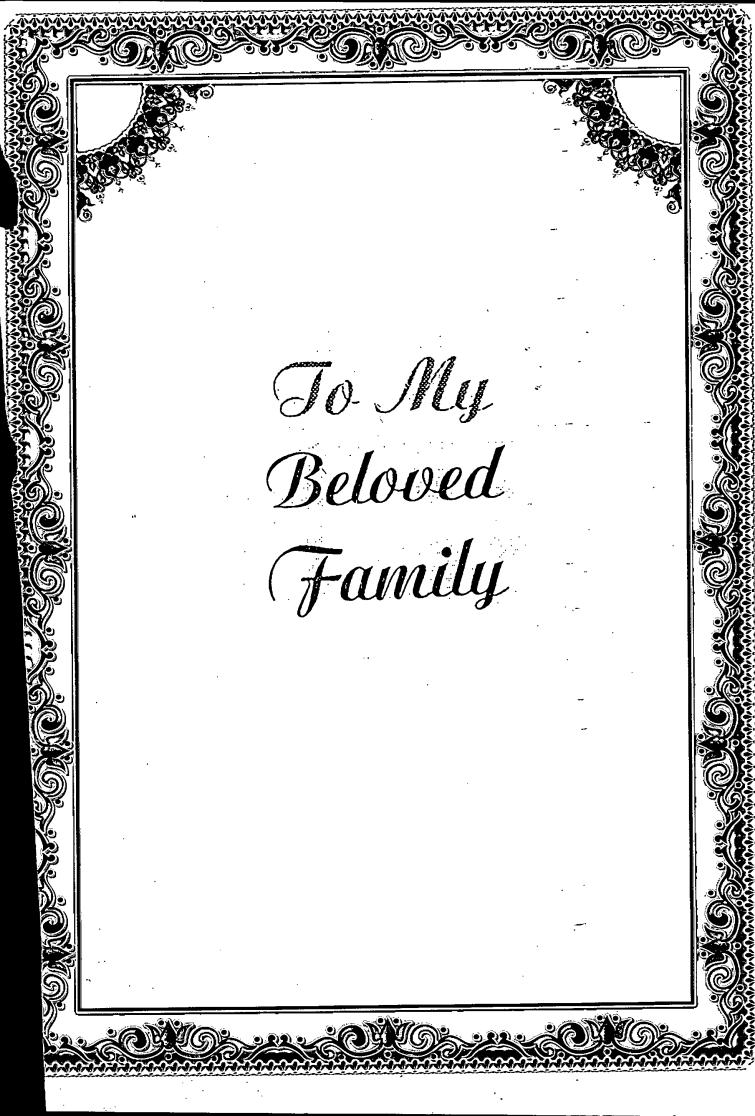
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PVC Polyvinyl chloride

RBCs Red blood cells

RLUS Relative light units.

RNA Ribonucleic acid

SGA Small gestational age

TEL Tetraethyl lead

TIBC Total iron binding capacity

TML Tetramethyl lead

TRH Thyrotropin-releasing hormone

TSH Thyroid stimulating hormone

U ALAD urinary δ-aminolevulinic acid dehydratase

UCP Urinary coproporphyrin

WBCs White blood cells

WHO World Health Organization

X-RF X-ray film

ZPP Zinc protoporphyrin

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INTRODUCTION

Lead is the fifth most abundant metal in the world, after iron, copper, zinc, and aluminum. High blood lead levels can affect survival and development of the fetus and the infant. Sterility, abortion, stillbirth, premature delivery and adverse pregnancy outcomes such as low birth weight, failure to thrive and impaired mental development of infants were recognized at low lead exposure levels. Lead can readily cross the placenta during the entire pregnancy and fetal levels ultimately equilibrate with maternal levels (Rom, 1976 and Mirghani, 1997).

Mayer et al. (1986) consistently identified a link between maternal blood level of $100 - 150 \,\mu\text{g/L}$ and disturbance in early infant and neurological behavior performance. They concluded that the previously accepted safe maternal blood level of $250 \,\mu\text{g/L}$ is too high, and that adverse reproductive and neurobehavior effects may occur at blood lead levels commonly found in the population of many nations today.

McMichael et al. (1986) showed that preterm deliveries were significantly related to maternal blood lead levels at