

An-Najah National University
Faculty of Graduate Studies

**EVALUATION OF DAILY CHANGES IN AIR POLLUTANTS
LEVELS ON THE INDUCTION OF RESPIRATORY
SYMPTOMS IN SCHOOL CHILDREN:
THE Y'ABED CHARCOAL PRODUCTION BASIN**

By

Handwritten signature and date: 27/12/01

YASER OMAR HAFETH KAYED

Supervisor

PROF. DR. MOHAMMED S. ALI-SHTAYEH

Co-supervisor

DR. MUTASIM BABA

Submitted in Partial Fulfillment of the Requirements for the Degree of Master of Environmental Sciences in, Faculty of Graduate Studies, at An-Najah National University, Nablus, Palestine.


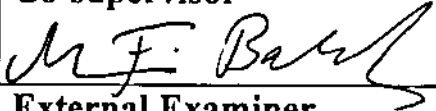
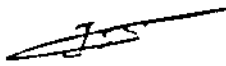
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This Thesis was defended successfully on 1/7/2001 and approved on By:

Committee members	Signature
1. Prof. Dr. Mohammed S. Ali-Shtayeh Prof. of Biological Sciences	Supervisor 
2. Dr. Mutasim Baba Electrical Engineering	Co-supervisor 
3. Assist. Prof. Dr. Issam A. Al-Khatib Environmental Engineer	External Examiner 

*TO
MY DEAR FATHER AND MOTHER FOR
THEIR SUPPORT, TO MY WIFE JUMANA
AND TO MY BROTHERS AND My FRIEND
WITH LOVE AND RESPECT*

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TABLE OF CONTENT

Subject	Page
Dedication	II
Acknowledgment	III
Table of Contents	IV
List of Tables	VI
List of Figures	VII
Abstract	X
CHAPTER ONE: INTRODUCTION	
1.1 Air pollution	1
1.2 Air pollution in the West Bank	3
1.3 Charcoal production	5
1.4 Health hazards during charcoal production	8
1.4.1 Particulate	8
1.4.1.1 Particles measuring	8
1.4.2 Gases	10
1.4.2.1 CO	10
1.4.2.2 CO ₂	11
1.4.2.3 NO ₂	12
1.4.2.4 SO ₂	14
1.4.2.5 H ₂ S	15
1.4.2.6 Methane	16
1.5 Charcoal production in Palestine	16
1.5.1 Kilns in Y'abed	19
1.6 Objectives	28
CHAPTER TWO: MATERIALS AND METHODS	
2.1 Study area : The Y'abed charcoal production basin	29
2.2 Study sites	30
2.3 Environmental monitoring system	31
2.4 Experimental design	36
2.5 Data collection	37
2.5.1 Statistical study	38
2.5.2 Graphics	38
2.6 Questionnaire	38
CHAPTER THREE: RESULTS	
3.1 Air pollutant levels in the Y'abed area	41
3.1.1 CO monitoring	41
3.1.2 NO ₂ monitoring	47
3.1.3 SO ₂ monitoring	59
3.1.4 CO ₂ monitoring	62
3.1.5 H ₂ S monitoring	63

LIST OF TABLES

Table	Page
A.1 The concentration of CO, H ₂ S, NO ₂ and CO ₂ in relation to RH and temperature. during May 2-11,1999	85
A.2 The 8-hour concentration of CO, H ₂ S, NO ₂ and SO ₂ gas and the 8-hour reading of RH, Air temperature and wind speed during the period of May 31-June 9,1998	86
A.3 The 8-hour concentration of CO, H ₂ S, NO ₂ , SO ₂ and CO ₂ in addition to RH, temperature and wind speed. during the period of September 27-29,1998	87
A.4 The concentration of CO and H ₂ S gasses and temperature. During the period of June 16-21,1999	88
A.5 The concentration of CO, NO ₂ and SO ₂ gasses and RH and wind speed during the period of July 2-5,1999.	89
A.6 The 8-hour concentration of H ₂ S, NO ₂ , SO ₂ and CO ₂ gases and RH, and temperature. during the period of March 19 -April 22,1999	90
A.7 The 8-hour average concentration of H ₂ S, NO ₂ , and the 8-hour average reading of RH, temperature ,and wind speed during the period of May 28-30,1998	93
A.8 The 8-hour concentration of H ₂ S, NO ₂ , SO ₂ and CO ₂ gases and RH, temperature wind speed and wind direction. during the period of Aug. 17-24,1998	94
A.9 The 8-hour concentration of H ₂ S, NO ₂ , SO ₂ and CO ₂ gases and RH, temperature and wind speed. during the period of September 1-8,1998	95
A.10 The 8-hour concentration of H ₂ S, NO ₂ , SO ₂ and CO ₂ gases and RH, and temperature. during the period of April 22-29,1999	96
A.11 The 8-hour concentration of H ₂ S, NO ₂ , and SO ₂ gases and RH, and temperature. during the period of May 12-19,1999	97
A.12 The concentration of H ₂ S, NO ₂ and SO ₂ gasses and wind speed and temp. during Sep.28-Oct.5,1999	98
A.13 The 8-hour concentration of CO, H ₂ S, NO ₂ and SO ₂ gasses in relation to RH, temperature, and wind direction and wind speed. During the period of May 19-20,1999	99

LIST OF FIGURES

Figure	Page
3.1.A CO ₂ and CO concentration in relation to temperature during the period of May 2-11,1999	42
3.1.B. H ₂ S and SO ₂ concentration in relation to RH during the period of May 2-11,1999	42
3.1.C H ₂ S and SO ₂ concentration in relation to temperature during the period of May 2-11,1999	42
3.1.D.CO and NO ₂ concentration in relation to RH during the period of May 2-11,1999	42
3.1.E. NO ₂ concentration in relation to temperature during the period of May 2-11,1999	42
3.1.F. CO ₂ concentration in relation to temperature during the period of May 2-11,1999	42
3.2.A. H ₂ S and SO ₂ concentration in relation to RH during the period of May 31-June 9,1998	44
3.2.B H ₂ S and SO ₂ concentration in relation to wind speed during the period of May 31-June 9,1998	44
3.2.C CO and NO ₂ concentration in relation to RH during the period of May 31-June 9,1998	44
3.2.D. CO and NO ₂ concentration in relation to wind speed during the period of May 31-June 9,1998	44
3.3.A Concentration of CO gas in relation to temperature during the period of June 16-21,1999	45
3.3.B. Concentration of H ₂ S gas in relation to temperature during the period of June 16-21,1999	45
3.4.A. Concentration of CO in relation to RH during August 2-5, 1998.	46
3.4.B. Concentration of SO ₂ in relation to RH during August 2-5, 1998.	46
3.5.A. concentration of NO ₂ in relation to temperature during the period of March 19-April 22, 1999	48

3.5.B concentration of NO ₂ in relation to RH during the period of March 19-April 22, 1999	48
3.5.C. concentration of CO ₂ in relation to temperature during the period of March 19-April 22, 1999	49
3.5.D. concentration of CO ₂ in relation to RH during the period of March 19-April 22, 1999	49
3.5.E. concentration of H ₂ S and SO ₂ in relation to temperature during the period of March 19-April 22, 1999	50
3.5.F. concentration of H ₂ S and SO ₂ in relation to RH during the period of March 19-April 22, 1999	50
3.6.A CO and NO ₂ concentration in related to RH during May 28-30,1998	52
3.6.B H ₂ S and SO ₂ concentration in related to temperature during May 28-30,1998	52
3.6.C H ₂ S and SO ₂ concentration in related to RH during May 28-30,1998	52
3.6.D NO ₂ concentration in related to temperature during May 28-30,1998	52
3.7.A H ₂ S and SO ₂ concentration in relation to temperature during the period of Aug. 17-24,1998	53
3.7.B H ₂ S and SO ₂ concentration in relation to RH during the period of Aug. 17-24,1998	53
3.7.C NO ₂ and CO ₂ concentration in relation to temperature during the period of Aug. 17-24,1998	53
3.7.D NO ₂ and CO ₂ concentration in relation to RH during the period of Aug. 17-24,1998	53
3.8.A. H ₂ S and SO ₂ concentration in relation to temperature during the period of Sep. 1-8,1998	55
3.8.B. H ₂ S and SO ₂ concentration in relation to humidity during the period of Sep. 1-8,1998	55
3.8.C. NO ₂ concentration in relation to temperature during the period of Sep. 1-8,1998	55
3.8.D. NO ₂ concentration in relation to RH during the period of Sep. 1-8,1998	55

3.8.E. CO ₂ and CO concentration in relation to temperature during the period of Sep. 1-8,1998	55
3.8.F CO ₂ and CO concentration in relation to RH during the period of Sep. 1-8,1998	55
3.9.A H ₂ S,SO ₂ and NO ₂ concentration in relation to temperature during the period of April 22-29,1999	56
3.9.B H ₂ S,SO ₂ and NO ₂ concentration in relation to RH during the period of April 22-29,1999	56
3.10.A H ₂ S,SO ₂ and NO ₂ concentration in relation to RH during the period of May 12-19,1999	57
3.10.B H ₂ S,SO ₂ and NO ₂ concentration in relation to temperature during the period of May 12-19,1999	57
3.11.A H ₂ S and SO ₂ concentration in relation to humidity during the period of September 28-October 5,1999	58
3.11.B H ₂ S and SO ₂ concentration in relation to temperature during the period of September 28-October 5,1999	58
3.11.C NO ₂ concentration in relation to humidity during the period of September 28- October 5,1999	58
3.11.D.NO ₂ concentration in relation to temp. during the period of September 28- October 5,1999	58

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ABSTRACT

The effect of charcoal production in Ya'bad area was studied by evaluation of gasses emitted from the process of carbonization. The study was carried out over 14-month period (May 1998 – August 1999). An ELE 8000/EMS 1417 Environmental Data Logger was used to measure the concentrations of CO, CO₂, NO₂, SO₂ and H₂S in three different locations. These locations are: site A located at about 180 meter to the north east of the kilns, site B about 70 meter forward to the smoke comes out of the kilns in Ya'bad and site C in the site of the kilns. Other physical factors; wind speed, wind direction, temperature and relative humidity were measured by the Logger. The effect of gas production on respiratory symptoms on Ya'bad inhabitants was studied by a 14 questions questionnaire. Primary school children from

the fifth grades living in four communities (supposed to be polluted), and in three communities (low polluted) answered the questionnaire. Our study has demonstrated that the concentrations of the gases measured were inversely correlated with distance from charcoal kilns, with the highest levels being encountered in site C (CO average concentration was 100.57ppm, NO₂ 11.7 ppm and SO₂ 1.75 ppm). In site A gases levels were very low (CO average concentration was 0.19ppm, NO₂ 0.29 ppm, SO₂ 0.26ppm, CO₂ 1.6%, and H₂S 0.36ppm). In site B, the levels of gases were also lower than that of site C (CO average concentration was 37.9ppm, NO₂ 14.9 ppm, SO₂ 14.7 ppm, CO₂ 1.98%, and H₂S 12.2 ppm). In sites C and B the gases levels exceeded the maximum standards limits and in site A the levels did not exceed the standards. Statistical analysis using ANOVA and F-tests ($p < 0.05$) showed significant difference in gases levels in the three locations. Our study showed also a considerable effect of charcoal production was significant (using ANOVA and t-tests) between school children close to the kilns and school children far from the kilns. Moreover, there was significant difference in respiratory symptoms between male and female students in schools closer to the kilns.

CHAPTER ONE

INTRODUCTION

Chapter 1

Introduction

1.1 Air Pollution

Air pollution is defined as substances in the atmosphere that have harmful effects on breathing (Nebel and Wright, 1993). The main sources of air pollution are energy production, transportation and industry.

One of the most problematic air pollutants in Israel as confirmed by air quality monitoring carried out, is sulfur dioxide (Gabbay, 1994). In 1993, over 11.5 million tons of crude oil and 5.8 million tons of coal were imported in to Israel (Gabbay, 1994).

In a study in Jordan there was an elevation in SO_2 and CO during the cold periods and exceeds the standard (Environmental Research Center, 1990).

Industrial air pollution is made up largely of smog and sulphur dioxide from the burning of coal and generally occurs in winter. Exposure to carbon monoxide at high concentration some times encountered in heavy traffic can raise carboxyhemoglobin

concentration in the blood of normal subjects, and can limit exercises tolerance in-patients with angina.

A small consistent association between standardized mortality and concentration of particulates in the environment particularly those below 2.5 micro meter in comparisons was found between similar cities with differing levels of pollution in the U.S.A. (Greef, 1995).

A significant association between daily SO₂ levels originated from a coal-fueled power plant in France and prevalence of upper and lower respiratory symptoms, in children was demonstrated (Charpin et al., 1988).

The evaluation of adverse health effects of exposure to air pollutants, even at relatively low concentrations, had become recently a major concern in many parts of the world. Significantly higher prevalence of respiratory symptoms in children were reported to be induced by moderate daily changes in SO₂ concentration in coal-basin area in France (Charpin et al., 1988).

Effects of air pollution have been mainly studied through examination of pulmonary function tests, whereas the association between air pollution and respiratory symptoms has been studied less extensively in sensitive subjects, e.g., children and asthmatic or bronchitic patients (Charpin et al., 1988).

1.2 Air pollution in the West Bank

There are no data available on air pollution in the West Bank due to the lack of air quality monitoring stations or programs so far. Quantification of air pollution has been carried out based on the estimation of possible air pollution contributing factors such as industrial activities, transportation, energy consumption, open burning of solid waste, and transboundary air pollutants. Although West Bank houses have little heavy industry, it suffers from substantial air pollution especially in the main urban areas and vicinities. More than 90% of the pollution in the West Bank comes from: human activities; increase in population; expanding industrial activity; and transboundary air pollutants. These are probably the key factors of air quality

deterioration in the West Bank. The lack of preventive legislation and codes and regulations to protect the environment; leads to an increase in human activities that affect air pollution. More over, climate and topography of the West Bank play a crucial role in transmitting air pollutants from one place to another (Applied Research Institute Jerusalem ARIJ, 1996).

The most expected common air pollutants in the West Bank are sulfur dioxide, suspended particulate matter, nitrogen dioxide, carbon monoxide, carbon dioxide and lead. Industrial activities with regard to air pollution are almost unregulated and their impact is heightened by the close proximity of the industrial facilities to populated areas. Quarries and stone cutting factories emit huge amounts of dust to the air (Shtayeh and Hamad, 1995). Charcoal production mainly found in Jenin area is a major contributor to air pollution. Considerable amounts of particulate, carbon monoxide, nitrogen oxides and VOC (volatile organic compounds) were expect to be emitted from charcoal production (ARIJ, 1996; Harris, 1978).

1.3 Charcoal production

Charcoal is an energy source in some areas of the world. It is considered as a clean energy source in comparison to coal and petroleum (Ballester et al., 1996). Charcoal by itself is a nontoxic material. However, it can be used to treat some digestive system problems (Baba, 1995). However, gases that are produced as a result of converting wood to charcoal may cause some adverse effects to humans, animals, plants and other organisms health. It is also a serious cause of air pollution (Charpin et al., 1988).

The process of charcoal production is known as “carbonization”. This process occurs in kilns and consists essentially of burning some wood charges to carbonize the rest in the presence of limited controlled quantities of air (Harris, 1978).

The process may require from 5 to 15 days of continuous vigilance by the workers with burning materials like hay and dry branches until actual carbonization happened (Baba, 1995).

The temperature of wood is raised inside the kiln up to 100-120 °C so that all of the free moisture will be driven off as steam, until wood become absolutely dry. Then the temperature increased to 170 °C at which wood will ignite in the presence of air. Decomposition of wood starts at 270 °C and the temperature continues to rise and may reach 400 °C or more until carbonization process is completed. This process results in: charcoal that contains 75-80% of fixed carbon, some ash and 20-25% volatile matter including tars and gases such as methane, carbon monoxide, carbon dioxide and others (Harris, 1978).

Production of charcoal using old kilns involves internal combustion of part of wood charged. These old kilns consume approximately twice as much wood per ton of charcoal as modern methods. In the ancient time charcoal was produced in earth pits or caves involving little or no capital. The process of charcoal burning requires days of continuous vigilance by the charcoal burner whose occupation was nevertheless regarded as

lowly one, although it required a considerable skill (Harris, 1978).

After the industrial revolution more sophisticated kilns were developed using brick, concrete or even steel independently or in combination but with no essential change in the practices, The best results were given by the tranchant kiln which was made of steel and was portable which consists of two cylindrical sections and a lid, other kilns which are operated widely and successfully in Brazil, and especially in the state of Minas Gerais are internally heated fixed , batch type , the important iron and steel companies operate several thousand of them .

They are circular , with a domed roof and are built of ordinary fire bricks , the circular wall is totally in contact with the outside air , this type of kiln is referred as “Beehive Brick kiln, this type has many advantages such as the gases pass through the wood charge, the heat contained in the gases is partially used in the processes of wood drying and carbonization. Good yield of charcoal and low cost (Harris, 1978).

1.4 Health hazards during charcoal production

1.4.1 Particulates

During the process of charcoal production (wood carbonization), different suspended particles are emitted into the air breathed by the workers and the people living in the region surrounding the production area. In addition to that, these particulates affect the trees, buildings and transportation. These particles include pure CO and dust. Also as a result of carbonization and after the process is completed, a huge amount of ash is produced. When the burning wood is cooled and washed with water, ash is mixed with the ground soil causing an increase in acidity of the soil (Baba, 1995; Schwartz, 1993).

1.4.1.1 Particles measuring

Particulate air pollution has been traditionally measured by drawing air through a filter paper and measuring the density of the black stain on the filter. This method is called black smoke method (Seaton et al., 1995).

Respirable particles, typically with a 4.5 micro meter aerodynamic diameter (50% cut-off point) are collected by the black smoke method and its variation; some particles up to 7-9 micrometer are also collected. Methods to measure total suspended particulate (by high volume sampler) have been used extensively in the USA. There are problems with this method, however in that the size range of particles sampled extends well beyond those particles that are able to penetrate the upper respiratory tract and in arid regions the method is liable to sample wind – entrained dust of non combustive origin, and this problem has been recognized by US EPA who recommended that particulate matter of less than 10 micro meter aerodynamic diameter (PM10) be measured, as a better indicator of health – related particles (WHO,1987).

The measurements of black smoke or PM10 are both expressed in micro gram per meter cubed, but they don not measure quite the same thing (Seaton et al., 1995).

1.4.2 Gases

1.4.2.1 CO

Carbon monoxide is an invisible gas that is highly poisonous (Nebel and Wright, 1993). It is a colorless and odorless gas, very stable and has a life time of 2 to 4 months in the atmosphere (Wark and Warner, 1981). This gas comes from pilot lights, unwanted kerosene, gas space heaters, gasoline engines, wood burning and mostly from carbonization (Samet et al., 1987).

Carbon monoxide interferes with oxygen transport by avidly binding to hemoglobin to form carboxyhemoglobin and by shifting the oxyhemoglobin dissociation curve to the left, although it binds to myoglobin, but the physiologic significance of the formation of CO-myoglobin has not been established.

Carboxyhemoglobin reduce oxygen delivery to tissues as dose the hypoxia of altitude, the tissues with highly oxygen needs are myocardium, brain, and exercising muscle, are most affected by the formation of carboxyhemoglobin (Samet et al., 1987).

The relevant standard of CO is 35 ppm EPA 1-hour, 9 ppm EPA 8-hour. The risks of CO poisoning are greater for fetuses, infants, pregnant women and persons with underlying cardiovascular or pulmonary disease.

High concentrations of carbon monoxide can cause physiological and pathological changes and ultimately death. Carbon monoxide is a poisonous inhalant that deprives the body tissues of necessary oxygen (Wark and Warner, 1981).

Hypoxia caused by carbon monoxide leads to deficient function in sensitive organs and tissues like the brain, heart, the inner wall of blood vessels (WHO, 1987).

1.4.2.2 CO₂

Carbon dioxide CO₂ is a colorless gas without smell and has a role in the carbon cycle and is one of the greenhouse gases (Shtayeh and Hamad, 1995). It is emitted from all fossil fuels products. Coal generates 1.8 times as much CO₂ as does natural gas; large increases in atmospheric concentration of CO₂ could produce significant climatic changes mainly through the greenhouse effect. Further more if the rate of fossil fuel and coal

were to continue increasing then the amounts of CO₂ could have severe impacts by the year 2025(The committee on Health and Environmental Effects of increased Coal Utilization, 1980).

1.4.2.3 NO₂

NO₂ has a stinging , suffocating odour and is composed of nitrogen–oxygen compounds (gases) that are converted to nitric acid in the atmosphere and they are a major source of acid deposition (Nebel and Wright, 1993). It plays an important role in formation of troposphere O₃ .NO₂ is often found at higher level indoors than outdoor, levels currently encountered in outdoor and indoors air, have been difficult to characterize.

Most studies of the relationship between residential exposure to NO₂ and health have focused on respiratory symptoms, illness and on level of pulmonary function. Because of its solubility, nitrogen dioxide penetrates into the lung periphery and about 60% of it is deposited there (Bascom, 1996).

NO₂ may damage the lung directly through its oxidant properties or indirectly by increasing susceptibility to respiratory infection (Bascom, 1996).

NO₂ can react with moisture present in the atmosphere to form nitric acid, which can cause considerable corrosion of metal surfaces. It absorbs visible light and at a concentration of 0.25 ppm will cause appreciable reduction in visibility. At a concentration of 0.5 ppm for a period of 10 to 12 days it suppresses growth of some plants. At a concentration found in the atmosphere it is only potentially irritating and potentially related to chronic pulmonary fibroses and some increase in bronchitis in children.

Nitrogen dioxide may trigger biochemical changes at relatively low concentration beginning after a 30-minute exposure to about 0.2 ppm. Long exposure leads to emphysema-like structural changes characterized by thickening of the alveolar capillary membrane. It has effect on the pulmonary function of normal, bronchitis and asthmatic humans (WHO, 1987).

1.4.2.4 SO₂

Sulfur dioxide is a poisonous gas to both plants and animals. It converts to sulfuric acid in the atmosphere and causes acid deposition (Nebeland Wright,1993). SO₂ is one of the most widespread pollutants in the world. It is emitted during the composition of most fuels (Khan and Gibbs, 1996). Domestic fires can produce emissions containing sulfur dioxide (WHO, 1987).

Sulfur dioxide is a nonflammable, nonexclusive, colorless gas that causes a taste sensation at concentrations from 0.3 to 1.0 ppm in air, at concentrations above 3.0 ppm the gas has pungent, irritating odor (Wark and Warner, 1981).

Exposure to sulfur dioxide is sensitive to asthmatics after vigorous exercise. Many of them show excess bronchoconstriction after only a few minutes. Studies have indicated that the bronchoconstrictive effect of SO₂ on asthmatics can be mitigated by anticholenergic drugs beta-adrenergic drugs (Linn et al., 1988).

Broncho constriction may be assessed in terms of slight increase in airway resistance ,most individuals will show are response to SO₂ at concentration of 5.0ppm and above, certain sensitive individuals show slight effects at 1to 2 ppm(Wark and Warner, 1981).

1.4.2.5 H₂S

It is a colorless gas, soluble in various liquids including water and alcohol. It can form under conditions of deficient oxygen in the presence of organic material and sulfate. Human activities can release naturally occurring hydrogen sulfide into ambient air. In industry, hydrogen sulfide could be formed whenever elemental sulfur or sulfur containing compounds come into contacts with organic materials at high temperatures.

In acute form, hydrogen sulfide intoxication is mainly the result of action on the nervous system, at concentration of 15 mg /m³ and above. It causes conjunctiva irritation, it affects the sensory nerves in conjunctivae. If respiration can be maintained, the

Prognosis in a case of a cute hydrogen sulfide intoxication, even a severe one, it cause eye irritation (WHO, 1987).

1.4.2.6 Methane

Methane emissions from agricultural activities in particular the report must include an inventory of activities in rice farming and livestock production and the intentional burning of agricultural waste grassland wood and forests (Ballester et al., 1996).

Methane is produced of microbial fermentation of organic matter in anaerobic environment. The percent of methane in the atmosphere is about 0.0002%. Increase in methane and other greenhouse gases are anticipated to affect global climate dynamics, increasing surface temperatures and altering the distribution of precipitation (Abu Omar and Shtayeh, 1995).

1.5 Charcoal production in Palestine

Charcoal continues to be used in different quantities as domestic fuel in the West Bank and Gaza. It is mainly used for cooking and heating in urban and rural areas. Charcoal was used

in large quantities thirty years ago but it has faced competition from kerosene and liquefied gas. Until now there are no official statistics on the amounts of charcoal produced or consumed in the Palestinian areas, but it is estimated that 18,000 – 20,000 Tons of charcoal is produced annually less than 30% of this quantity is consumed locally (Baba, 1995).

Actually, more than 80% of charcoal produced locally in the West Bank and Gaza is exported to Israel. It is estimated that 18,000 tons of charcoal is produced in the Jenin area and around 2000 tons are produced in the Gaza strip, charcoal is used for cooking (barbecue), home heating, tobacco smocking (babbles) and for small industries. The quality of the produced charcoal is relatively low, and it is consumed locally without any type of processing, the studies shown that the carbonizing process of wood is not complete in more than 85% of the cases. The result of this is higher percentage of carbon monoxide production when burning the charcoal. Using such energy source for home heating in winter is a great danger to human health (Baba, 1995).

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Charcoal production techniques used locally, are extremely poor, since much of the energy of wood is lost in the production process of charcoal. Thus the use of charcoal as a viable cooking and heating fuel can be developed and expanded, but this should be on the basis of technology and management that lead to low priced, reliable and sustainable supply. Present methods of cutting the trees and burning the wood in kilns are predatory and waste of energy and can not be sustained beyond a low threshold. More efficient pit kilns would improve the economics of charcoal but not the long term output or validity of the forests (Baba, 1995).

If charcoal, as indigenous sources of energy, is to be reliably supplied to customers, it must come from plantations that are regularly cut and replanted. Energy plantations have generally been promoted for land unfit for agriculture but the availability of such land varies from region to region (Baba, 1995).

In Palestine, old-fashion kilns are used for the production of charcoal. Most of these kilns are located in Y'abad Area near Jinin.

The process of charcoal production used in Palestine is as follows; (Figures 1.1,1.2,1.3), wood pieces are organized together to form a coned shape. A small hole is left at the top of the cone to put the small dry branches and hay to be used as fuel to start the fire. Next, all sides of the cone excluding the top are covered with soil and sand so that smoke can not escape from the cone. After starting the fire by burning the hay and small dry branches at the top the whole cone is closed and covered completely with soil. After that from time to time, the top of the cone is opened to add more wood and hay to keep the fire on inside the cone.

1.5.1 kilns in Y'abed

The charcoal Kilns are mostly located in Y'abed and the surrounding villages see (Figure 1.4). A population of about 35,000 people lives in Y'abed area (Y'abed, Zabadah, Dhahr Al 'abed, Barta'a, Tourah and Nazlet Zeid). Many of these people



Figure 1. 1 The first step in charcoal production: wood is arranged in a cone shape heap.



Figure 1. 2 The second step in charcoal production:
covering wood with a layer of hay (7-10cm) thick

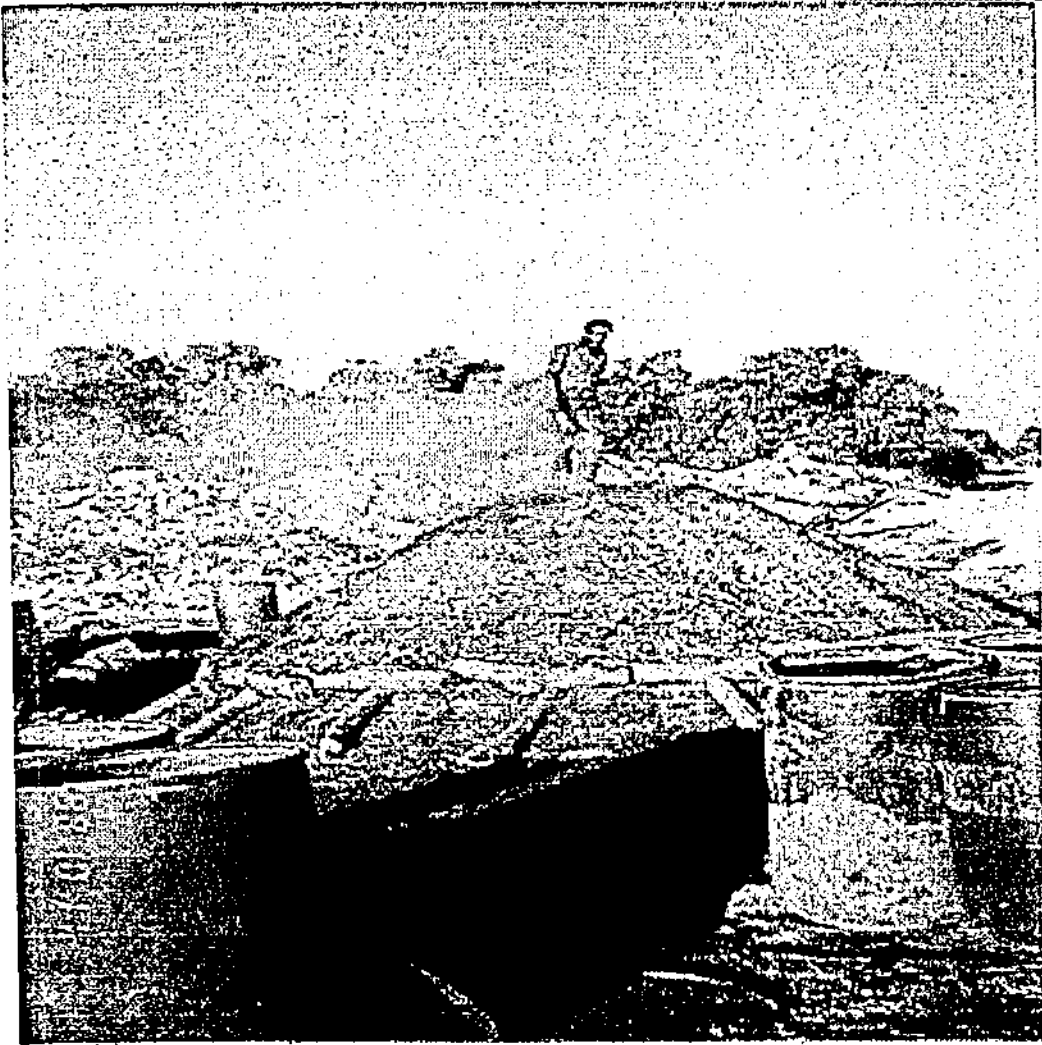


Figure 1.3
Charcoal kiln covered with sand and the worker on the top to see the fire.



Figure 1. 4 charcoal kiln close to houses in Ya'bad

depend, in their living, on olive and tobacco farming, and on charcoal production.

More than 75,000 tons of wood is used annually for charcoal production resulting in about 18,000 tons of charcoal the majority of which is exported to Israel. About 600 workers supporting about one third of Y'abed. The charcoal kilns are located in the agricultural land and not far from the living area, and some of them are very close to houses (Figure1.5), and closer to the main road (Figures 1.6, 1.7).



Figure 1.6 man work in a charcoal kiln



Figure 1.7 Charcoal kilns located near the main road in the vicinity of Y,abed village

1.6 Objectives

This study was aimed at:

1. Providing data on air pollution levels resulted from charcoal kilns in Ya'bed area.
2. Studying the relationship between air pollutants and climate (e.g., temperature, humidity, wind speed and direction).
3. Evaluating the adverse health effects of exposure to air pollutants resulting from charcoal production in Ya'bed area.

CHAPTER TWO

MATERIALS AND

METHODS

Chapter 2

Materials and Methods

2.1 Study area: The Y'abed charcoal production basin

The charcoal kilns are located around Y'abed and surrounding villages. About 200 locations for charcoal production existed in the year 1996.

Y'abed, located about 17-Km south west of Jenin, comprises an area of approximately 35 Km², with a population of 18000 inhabitants. Several other villages with a total population of about 17,000 have charcoal kilns. These villages are: Zabada, Theher Al-abed, Bartaah, Tourah and Nazlet Zaed. Charcoal kilns have been operating there for long time, and many people of the area work in the kilns.

In the year of 1995 more than 75,000 tons of wood were used annually for charcoal production resulting in about 18,000 tons of charcoal the majority of which was exported to Israel. About 600 workers supporting about one third of Yabed. Actually, more than 80% of charcoal produced locally in the

West Bank and Gaza were exported to Israel. It was estimated that 18,000 tons of charcoal were produced in the Jenin area and less than 30% of this quantity is consumed locally (Baba,1995).

2.2 Study sites

Three sites expected to have different air pollution levels that been, in Y'abed town.

Site A:

Located at about 180m to north east of the charcoal production area near Y'abed town on a house in side the town, and expected to have moderate or low air pollution level.

The site was studied in the periods of March 19-April 22,1999 and May 2-11,1999.

Site B:

Which was on a house near the charcoal, kilns near Y'abed town. About 70m from the charcoal kilns and for ward to the smoke comes out of the kilns, we expect this place to be polluted so we took readings in the year 1998 and 1999.

The site was studied in the periods of May 28- June 9-1998, August 17-24-1998, September 1-8-1998, April 22-29-1999, May 12-20-1999, and June 16-21-1999.

Site C:

Was closer to the charcoal kilns near Y'abed town in the field to see the effect on the workers, and we expect that the concentration could be high or heavy polluted, and that was during the period of July 2-5 -1999.

The charcoal kilns are located in an agricultural land mainly planted with olives, many of them are not far from the living area . Soil in the area is a muddy soil.

2.3 Environmental monitoring system

An ELE 8000/EMS 1417 Environmental automatic pollution monitoring station was used throughout this sort. The system is a computerized station that use a computer program called (DEMS) from which the data logger can be set up and programmed.

Dialog 900/EMS Management Software has been developed in conjunction with the MM900/950 series of data loggers and runs

on IBM PC or compatible. The software provides the means to set up and operate the data loggers. Once an experiment has been defined, and the relevant sensors connected the logger can be quickly and easily configured. When the logger has been configured it will run independently from the computer until data has to be collected.

Dialog is a menu driven program with pull down menus, which lead logically through the normal operational sequences. The user interface is intuitive, with context sensitive help available. Dialog is supplied with a database containing the configuration data for standard sensors.

All information about a logging task can be saved in a configuration file, separate configuration files can be created for each experiment, and can be retrieved when required. Dialog can be set up to automatically retrieve a predefined configuration file when the program is first run.

Dialog downloads data from the MM900 via the RS232 serial link and saves it to a data file. The downloading procedure does not interfere with the operation of the logger. Dialog can be

configured to start the download from the end of the last downloaded record in the logger's memory and to append the information to the data file, the data in the files can subsequently be viewed or converted to different formats for export to other programs like Lotus 123, or can be output to a printer.

The data logger depends on a chemical and physical sensors. The chemical sensors are very sensitive to the gases, each sensor is sensitive to one gas, and the sensors are:

Carbon monoxide gas sensor: This sensor uses an electrode, a reference electrode and a counter electrode, the gas diffusing into the sensor reacts with special catalyzed sensing electrode to produce electrons. A built-in circuit amplifies the signal into a millivolt output. Calibration can be checked by exposing the sensor to a known gas construction or by using a cylinder of span gas.

Carbon dioxide sensor: for the estimation of CO₂ gas concentration, the concentration is in (%).

Nitrogen dioxide sensor: This sensor uses an electrode, a reference electrode and a counter electrode, the gas diffusing

into the sensor reacts with special catalyzed sensing electrode to produce electrons. A built-in circuit amplifies the signal into a millivolt output. Calibration can be checked by exposing the sensor to a known gas construction or by using a cylinder of span gas.

The concentration unit is ppm, the reading was recorded on a 10-min. basis.

Sulfur dioxide sensor: This sensor uses an electrode, a reference electrode and a counter electrode, the gas diffusing into the sensor reacts with special catalyzed sensing electrode to produce electrons. A built-in circuit amplifies the signal into a millivolt output. Calibration can be checked by exposing the sensor to a known gas construction or by using a cylinder of span gas.

The concentration unit is ppm, the reading was recorded on a 10-min. basis.

Hydrogen Sulfide sensor, This sensor uses an electrode, a reference electrode and a counter electrode, the gas diffusing into the sensor reacts with special catalyzed sensing electrode to

produce electrons. A built-in circuit amplifies the signal into a millivolt output. Calibration can be checked by exposing the sensor to a known gas construction or by using a cylinder of span gas.

The other sensors are:

Wind speed sensor: this sensor measure wind run and average wind speed.

Measurement method: The wind causes the cups to rotate.

A magnet turns with the rotor spindle and the resulting varying field causes two mercury wetted reed switches to make and break contact each revolution. The unit is m/s.

Relative Humidity sensor: This sensor measures the relative humidity of air.

The measurement method: The water is absorbed according to the relative humidity of the air. This causes a change in the capacitance of the sensor, which the electronics convert, into a linear voltage.

Wind direction sensor: for determining the wind direction.

A wire wound potentiometer system is used to indicate the wind direction. Increased wind angles produce larger resistance's, which can be measured, the sensor gives a smooth rotation through 360 degree.

2.4 Experimental design

The study was carried out over a 14-month period (May - 1998 to August -1999). Monitoring periods are shown in table 2.1 and 2.2

Table 2.1: Air pollution monitoring periods during the year 1998

	May 28-31	June 1-11	July 1-31	August 17-24	September 1-8
CO		++			
CO ₂				++	++
NO ₂	++	++		++	++
SO ₂	++	++		++	++
H ₂ S	++	++		++	++

Table 2.2: Air pollution monitoring periods during the year 1999

	March 19-31	April 1-30	May 2-20	June 16-21	July 2-5
CO			++	++	++
CO ₂	++	++	++		
NO ₂	++	++	++		++
SO ₂	++	++	++		++
H ₂ S	++	++	++	++	

2.5 Data collection

The data were collected every three days and some times every one-week by a computer.

In our study the gases collected were CO, CO₂, NO₂, H₂S and SO₂. Some environmental factors were studied such as humidity, temperature, wind direction and wind speed

2.5.1 Statistical study

The data were statistically done for the averages every 8-hour, because the national standards concern with the 8-hour standard. The maximum and minimum concentrations were found for each period. F-test was done for each gas to see if there was significant difference between the location then statistical test done between each two locations to see the difference between which location and toward which location.

2.5.2. Graphics

Graphs were done for the average data. In the graphs the concentration of gases were put on the main axis and on the secondary axis we put the environmental factors to see if theirs a relation between the concentration of the deferent gases and the environmental factors. We took two axes because of the difference in the unit.

2.6 Questionnaire

The study was including a questionnaire for respiratory symptoms, which was reported at the morning by the help of the

teacher. The questionnaire contains 14 questions and information about the place and the sex, the questions were divided into two parts some of the questions, at the morning had you 1- cough 2- eye irritation 3- a runny nose 4-any sneezing 5- sore throat 6-wheezing in the chest 7- fever (Appendix.B.1).

The symptoms such as eye irritation, runny nose, and sneezing have been considered together under the heading E.N.T symptoms and morning cough and wheezing in the chest appear as pulmonary symptoms.

The questionnaire was answered by primary school children from the fifth grades living in four communities supposed to be polluted (Ya'bed, Zabdeh, Kferet , Nazlet Zeid) and in three low polluted communities (Anin , Arابه, Am alrehan). In each community all children attending the fifth grade were enrolled in the study.

Children were chosen because they do not smoke, not exposed to occupational hazards and spend the day in homogeneous air environment. The children will give indication on the people living in these areas.

Aims of the questionnaire:

To give us indications about respiratory and pulmonary symptoms in different villages in Jenin area.

To see if there was relation ship between air pollution from the charcoal production and respiratory and pulmonary symptoms through the statistical analysis of questionnaire in the area expected to be polluted and the villages expected to be non polluted.

To see if there was relation between the data obtained from the environmental station and the questioner.

The effect of the place and sex on the results.

Statistical analysis was done for the data obtained from the questionnaire between all the villages to see the effect of place and between the polluted and non-polluted villages.

CHAPTER THREE

RESULTS

Chapter 3

Results

3.1 Air pollutant levels in the Y'abed area

Means of concentrations of CO, CO₂, NO₂, SO₂ and H₂S in the air determined at 8-h intervals in relation to relative humidity, temperature and wind speed are presented in Tables A.1-A.13 and Figures 3.1-3.11.

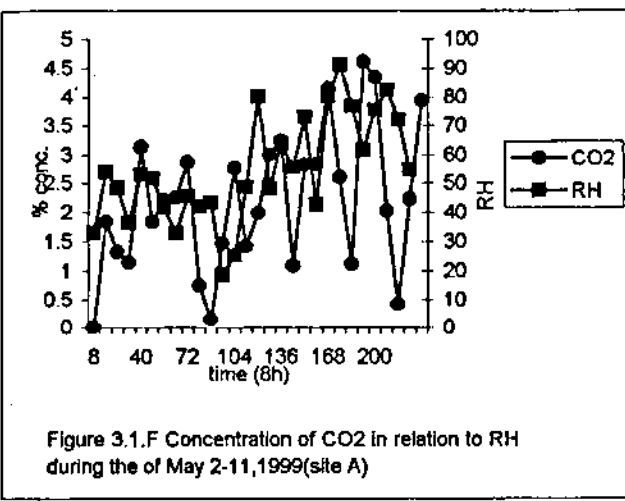
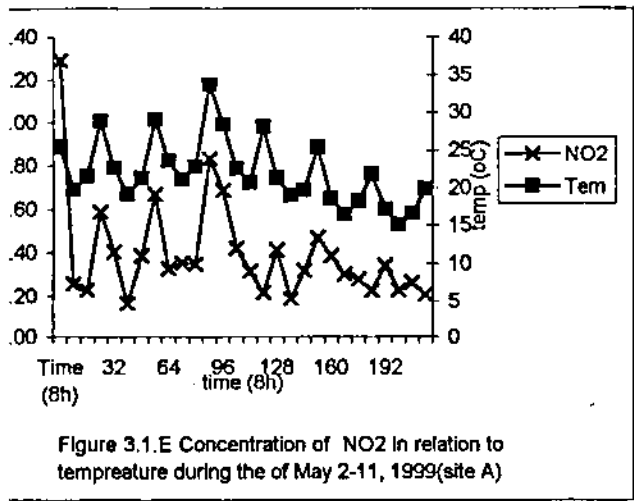
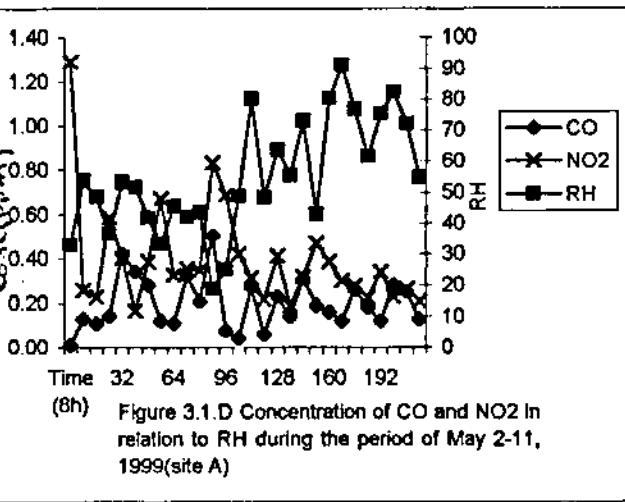
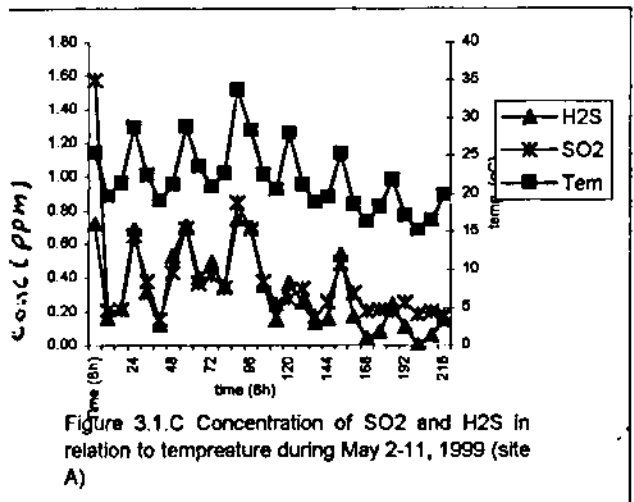
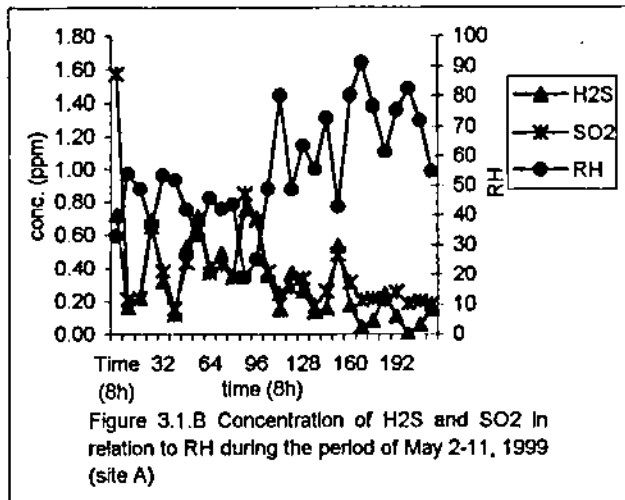
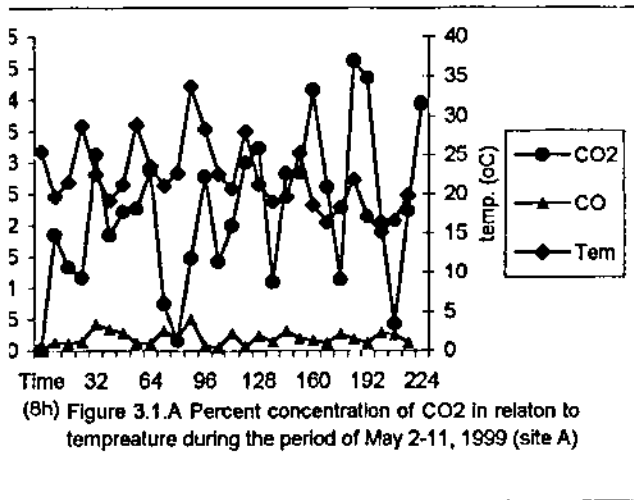
3.1.1 CO monitoring

The 8-hour U.S.A. maximum is 9ppm and the Canadian standard is 13ppm.

Sampling sites (Locations A, B, C) differed significantly at ($p < 0.05$) in CO concentration with location C showing higher CO levels being closer to charcoal kilns.

Location A

During the period of May 2-11,1999 the concentration of CO ranged between 0.01- 0.5 ppm; with an average of 0.19 ppm. In this period there was an increase in the concentration of CO as temperature increased. The correlation with RH was not clear (Figure 3.1.A,D and Table A.1).



Location B

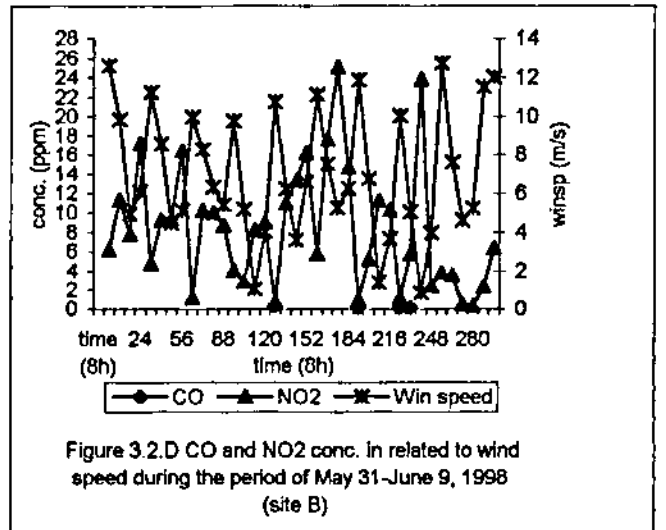
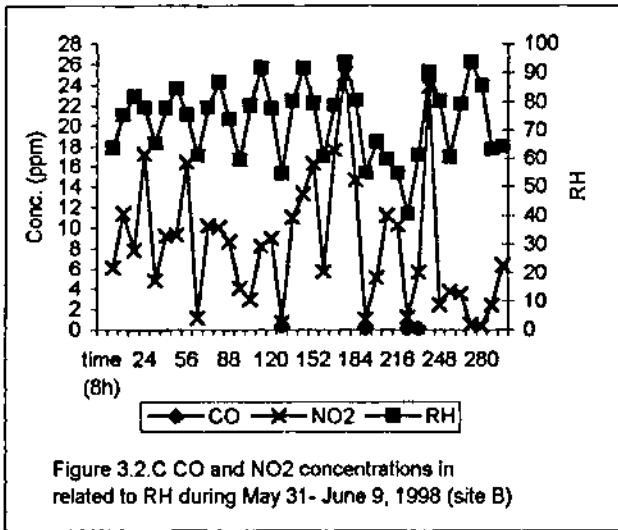
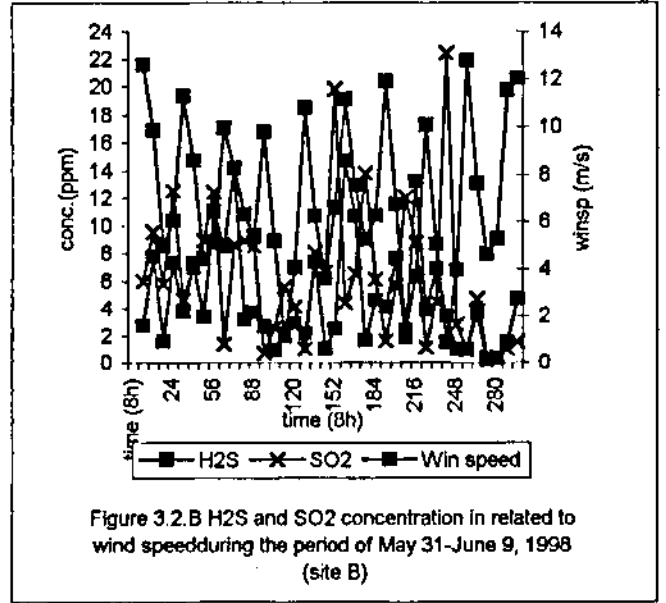
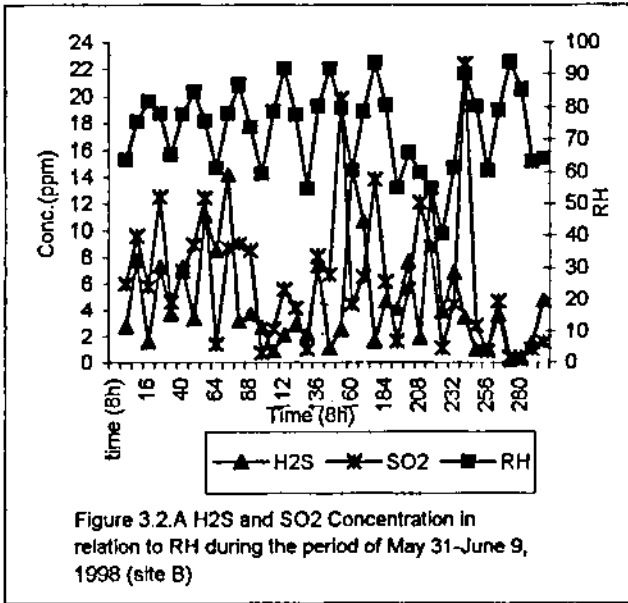
During the period of May31-June9, 1998 CO concentration ranged between 2.12-15.47 ppm; and the average was 10.2-ppm concentration increased with increasing temperatures (Figure 3.2.C, D and Table A.2).

During the period of September 27-29,1998 the concentration was ranged between 9.8-13.8 ppm; with an average of 11.55 ppm (Table A.3).

During June 16-21, 1999 the concentration of CO ranged between 31.65- 50.48 ppm; and the average was 37.9 ppm (Figure 3.3.A, Table A.4).

Location C

During the period of July 2-5,1999 the concentration of CO ranged between (20.86-167.75ppm) and the average was 100.57ppm. The concentration decreased with increasing in relative humidity (Figure 3.4.A and Table A.5).



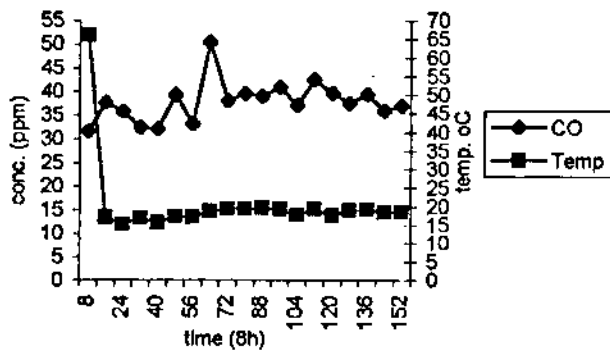


Figure 3.3.A Concentration of CO gas in relation to Temperature during the period of June 16-21, 1999.(site B)

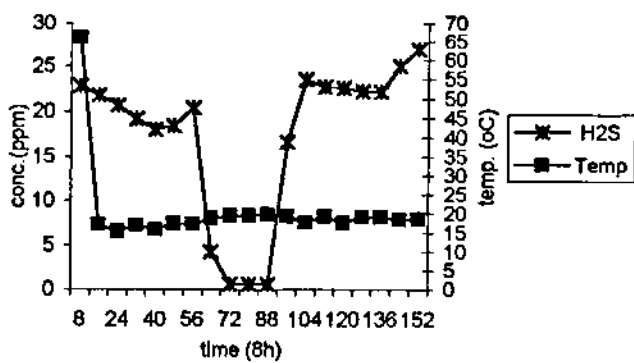
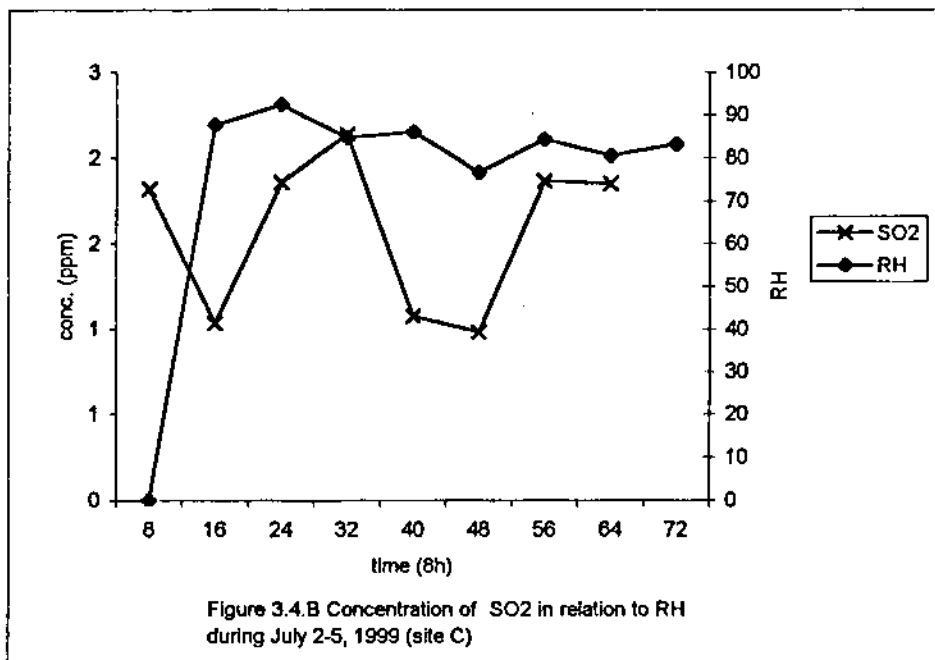
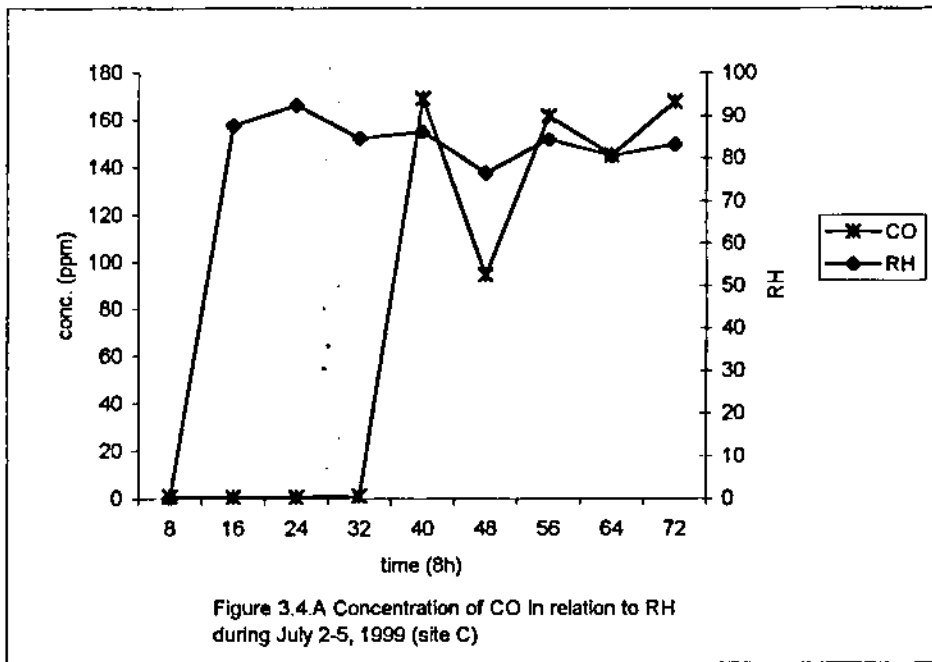


Figure 3.3.B Concentration of H2S in relation to temperature during June 16-21, 1999(site B)



3.2.2 NO₂ monitoring

Sampling sites (locations A,B,C) differed significantly at ($p < 0.05$) in NO₂ concentration with location C showing higher NO₂ level being closer to charcoal kilns.

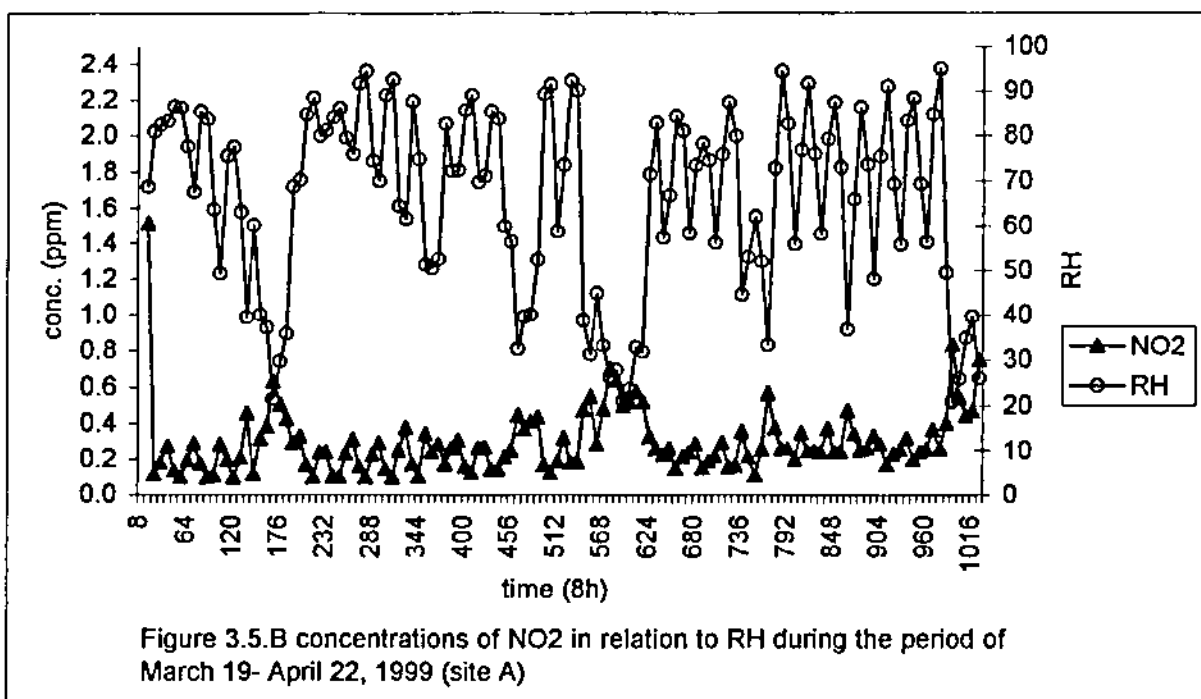
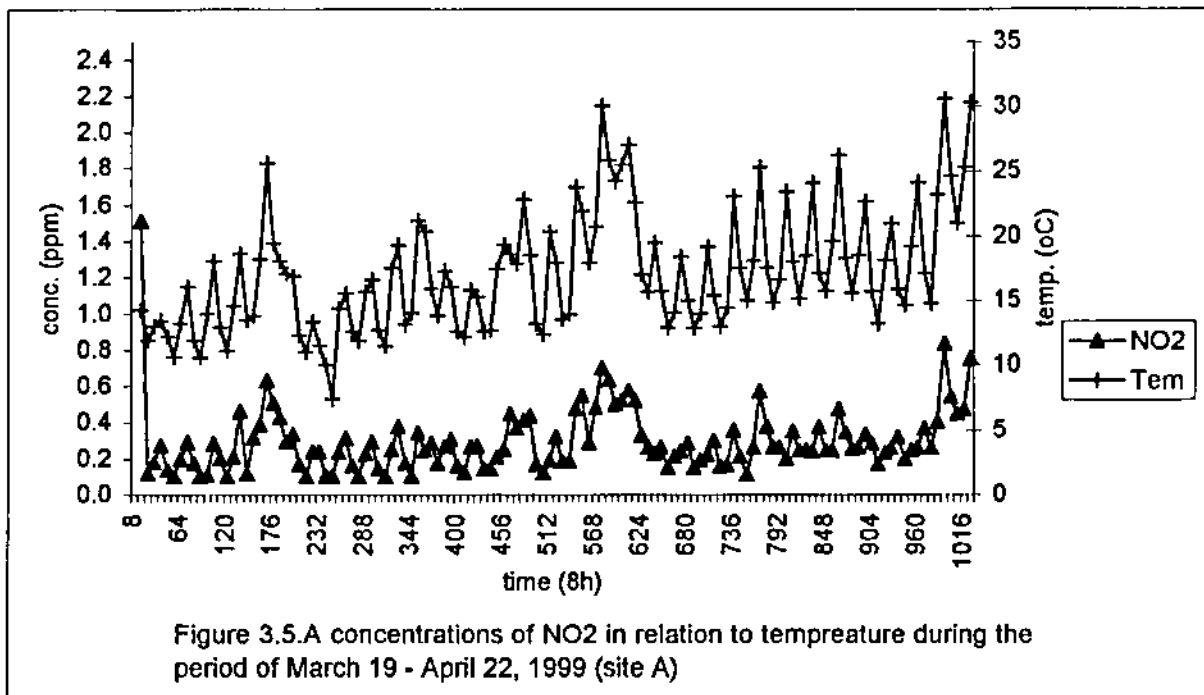
Location A

During the period of March 19-April 22, 1999 the concentration of NO₂ ranged between 0.1- 1.51 ppm; with an average of 0.29 ppm. In most of the reading as the temperature increased the concentration increased, correlation with humidity was not clear (Figure 3.5.A,B, Table A.6).

During the period of May 2-11,1999 the concentration of NO₂ lies between 0.16-1.29ppm; and the average was 0.40 ppm, there was no large change in the concentration during this period. The concentration of NO₂ was increased as the temperature increase, the correlation with humidity increased as the humidity decreased (Figure 3.1.D,E, Table A.1).

Location B

During the period of May 28-30,1998 the concentration of NO₂ ranged between 1-15.29 ppm; the average was 8.57ppm.



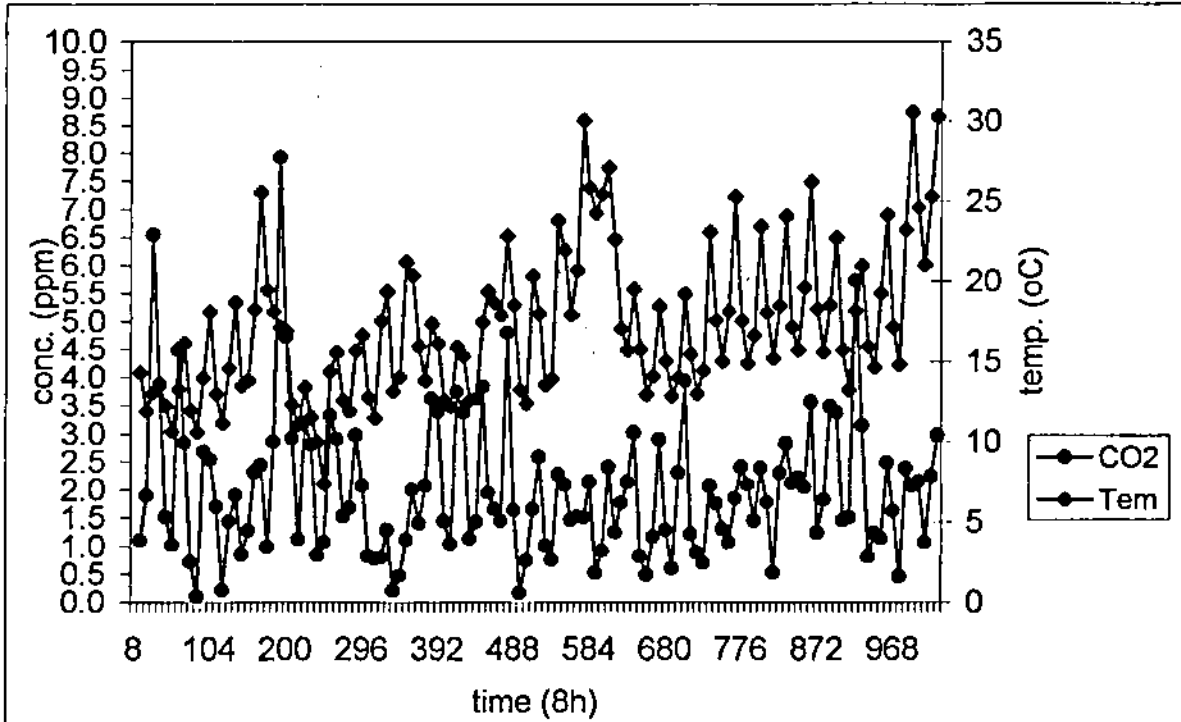


Figure 3.5.C concentrations of CO₂ in relation to temperature during the period of March 19 - April 22, 1999 (site A)

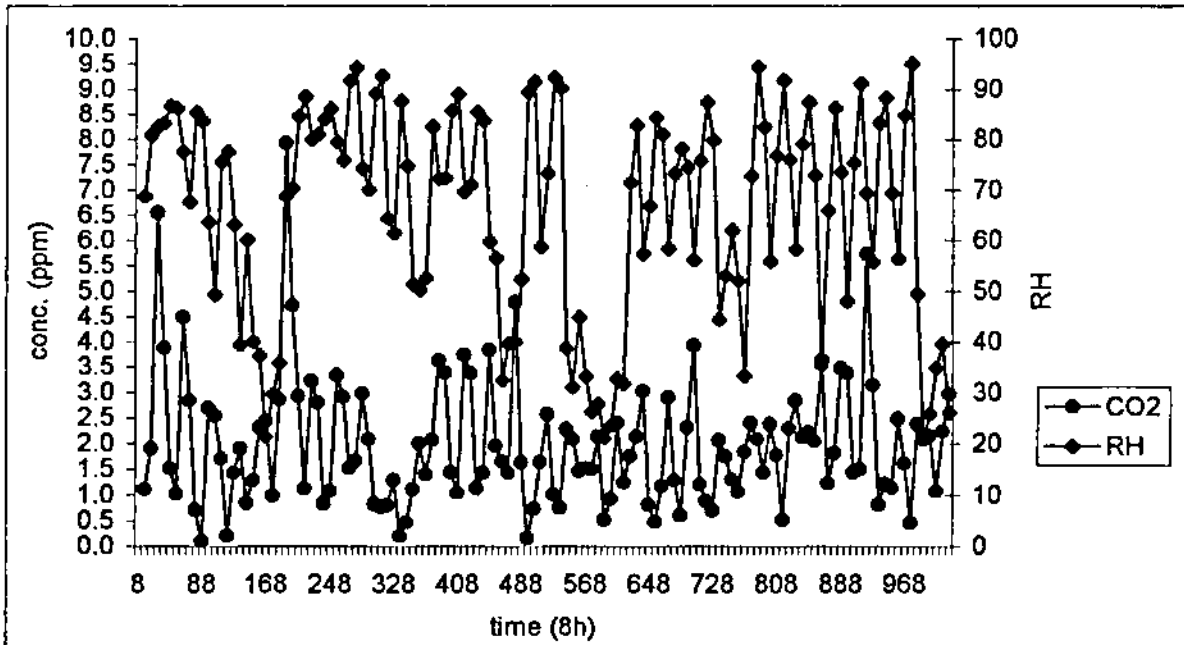
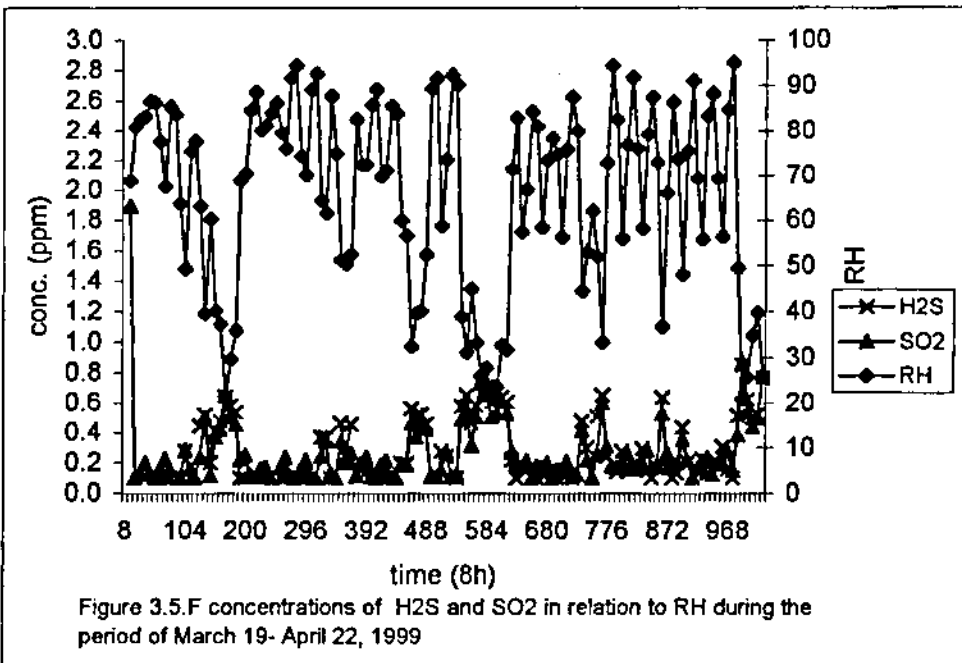
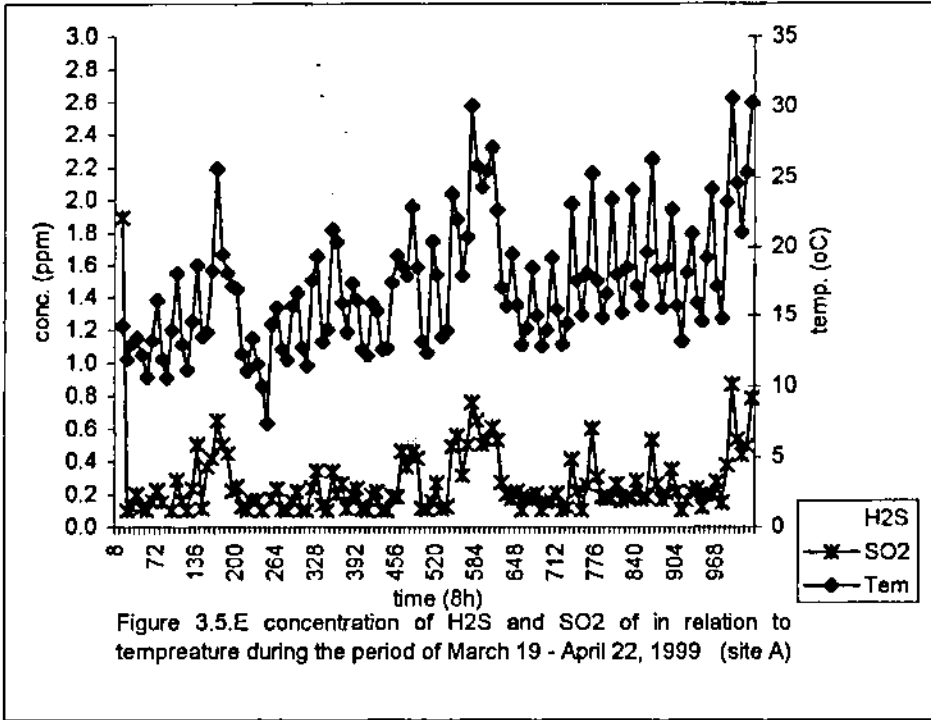


Figure 3.5.D concentrations of CO₂ in relation to RH during the period of March 19- April 22, 1999 (site A)



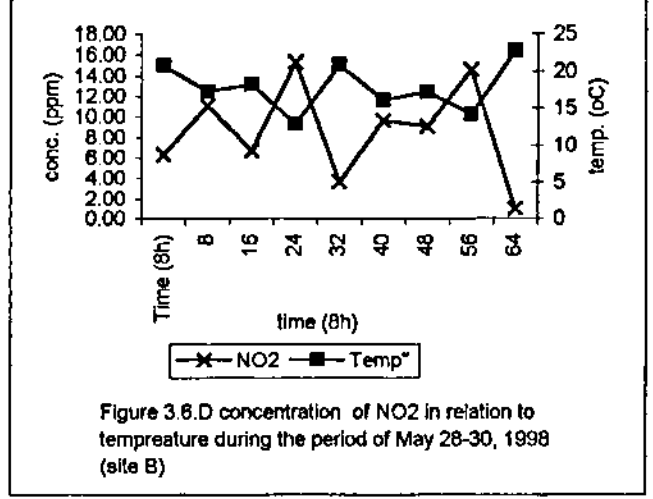
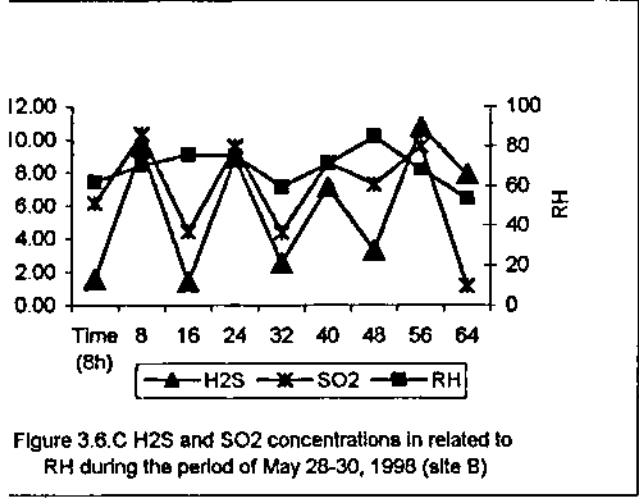
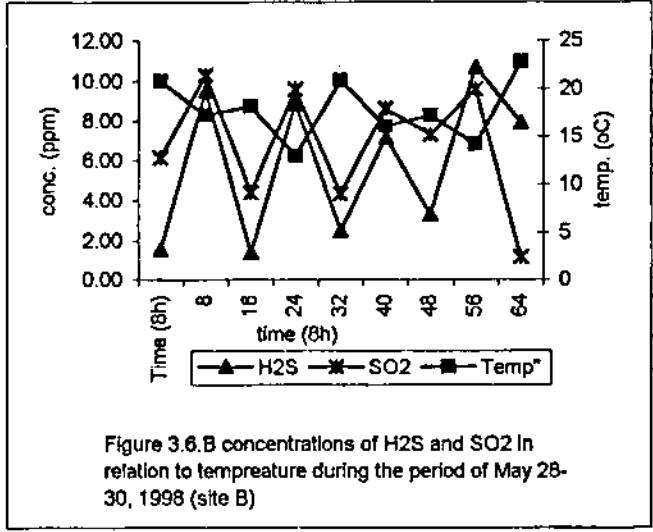
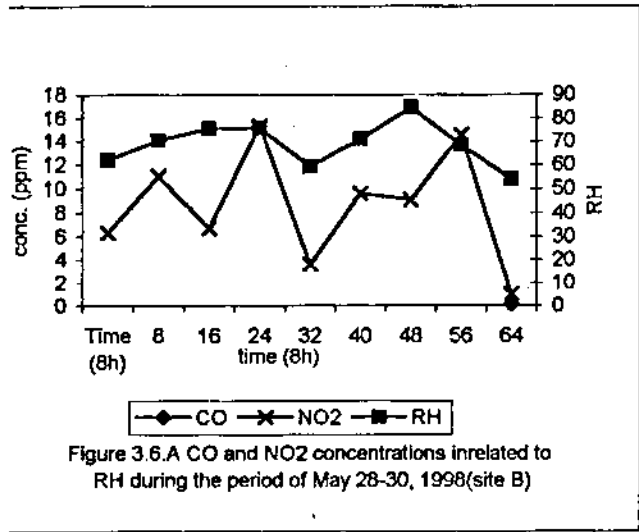
In this period when the humidity increased the concentration of NO₂ increased, (Figure 3.6.A,D , Table A.7).

During the period of May 31-June 9, 1998 the concentration of NO₂ ranged between 0.34-25.03 ppm and the average was 8.37 ppm .The relation with relative humidity the concentration of NO₂ increased as the humidity increased, according to wind speed the concentration decreased as the wind speed increased (Figure 3.2.C,D and Table A.2).

During the period of September 27-29,1998 the concentration ranged between 5.70-12.76 ppm with an average of 8.99-ppm (Table A.3).

During the period of August 17-24,1998 NO₂ concentration ranged between 2.6-28.5 ppm; the average was 14.9 ppm . The relation to relative humidity as the humidity increased the concentration of NO₂ increased (Figure 3.7.C,D and Table A.8) .

During the period of September 1-8, 1998 the concentration of NO₂ was ranged between 3.42-29.09 ppm; with an average of 15.66 ppm. According to humidity as the



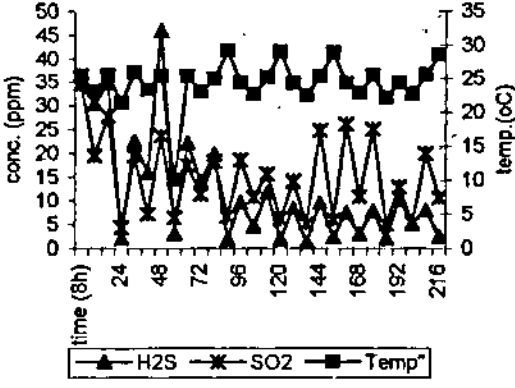


Figure 3.7.A concentration of H2S and SO2 in relation to temperature during the period of Aug. 17-24, 1998, (site B)

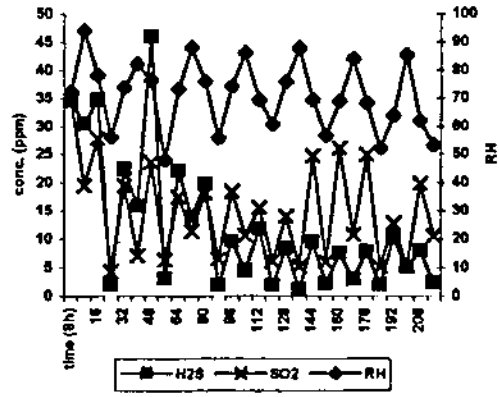


Figure 3.7.B concentrations of H2S and SO2 in relation RH during the period Aug. 17-24, 1998 (site B)

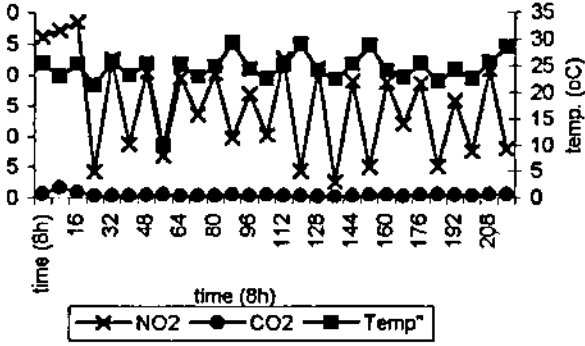


Figure 3.7.C concentration of CO2 and NO2 in relation to temperature during the period of Aug 17-24, 1998 (site B)

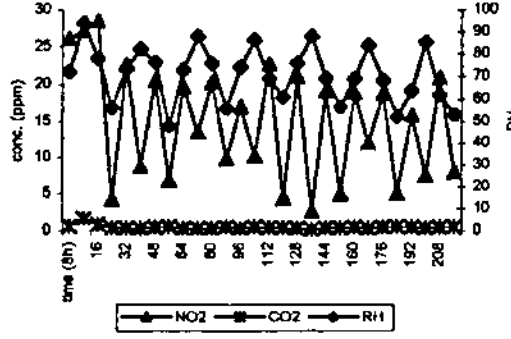


Figure 3.7.D concentrations of CO2 and NO2 in relation to RH during the period of Aug. 17-24, 1998 (site B)

humidity increased the concentration of NO₂ increased, and as temperature increased the concentration of NO₂ increased, there was correlation with wind speed as wind speed increased the concentration decreased (Figure 3.8.C,D and Table A.9) .

During the period of April 22-29,1999, the concentration of NO₂ ranged between 0.23-50.04ppm; the average was 14.58 ppm, (Figure 3.9.A,B and Table A.10).

During the period of May 12- 19,1999, the concentration of NO₂ was low during the first 5days, on the sex day there was an increased in the concentration of NO₂ the concentration ranged between 0.10-25.16 ppm; the average was 6.7 ppm (Figure 3.10.A,B and Table A.11).

During the period of September 28-October 5, 1999 the concentration ranged between 0.1-22.3 ppm; and average of 12.34 ppm.(Figure 3.11.C,D and Table A.12).

Location C

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During the period of July 2-5,1999 the concentration of NO₂ ranged between 7.10-15.7 ppm the average concentration was 11.7 ppm. (Table A.5).

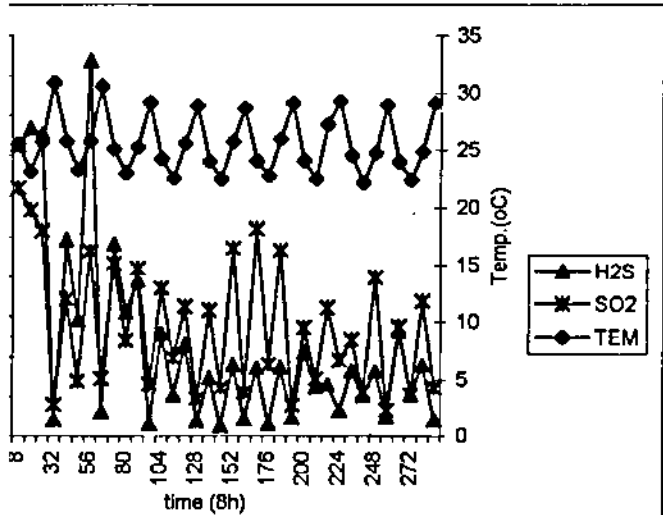


Figure 3.8.A H₂S and SO₂ concentrations in relation to temp. during the period of Sept 1-8, 1998(site B)

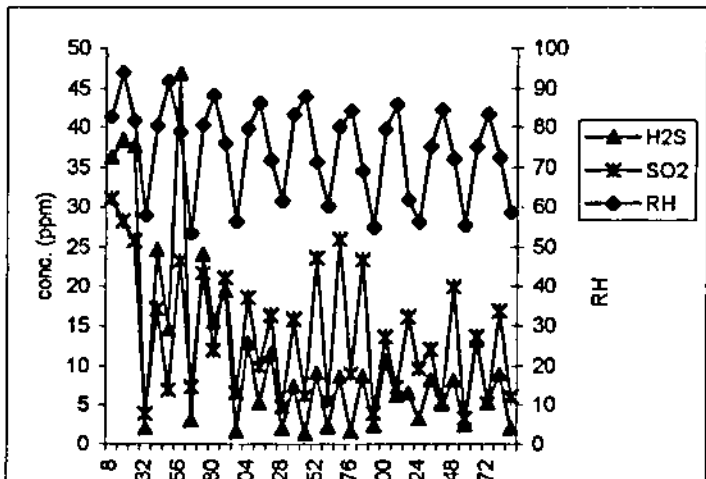


Figure 3.8.B CO₂, CO, H₂S, SO₂ and NO₂ concentrations in relation to RH during the period of Sept 1-8, 1998(site B)

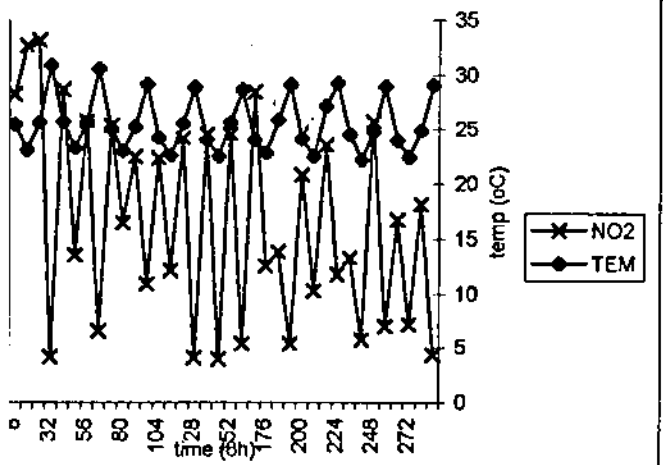


Figure 3.8.C NO₂ concentrations in relation to temperature during the period of Sept 1-8, 1998(site B)

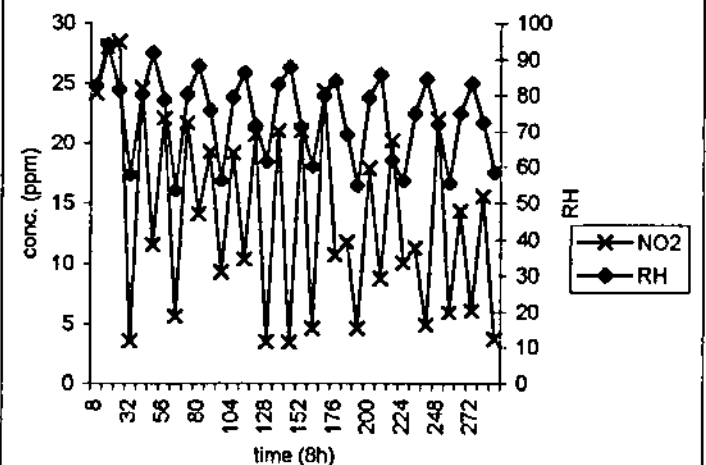
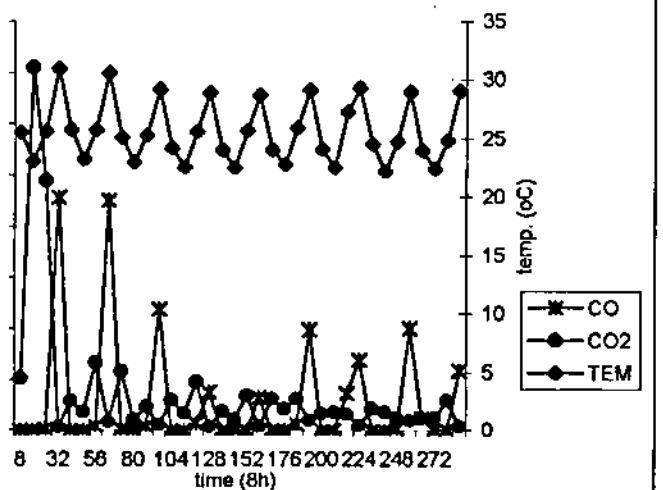


Figure 3.8.D NO₂ concentrations in relation to RH during the period of Sept 1-8, 1998 (site B)



3.8.E CO₂ and CO concentrations in relation to temp. during the period of Sept 1-8, 1998(site B)

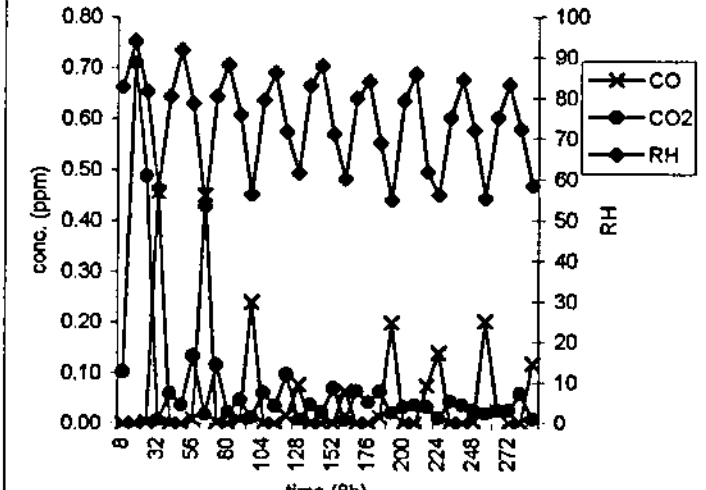
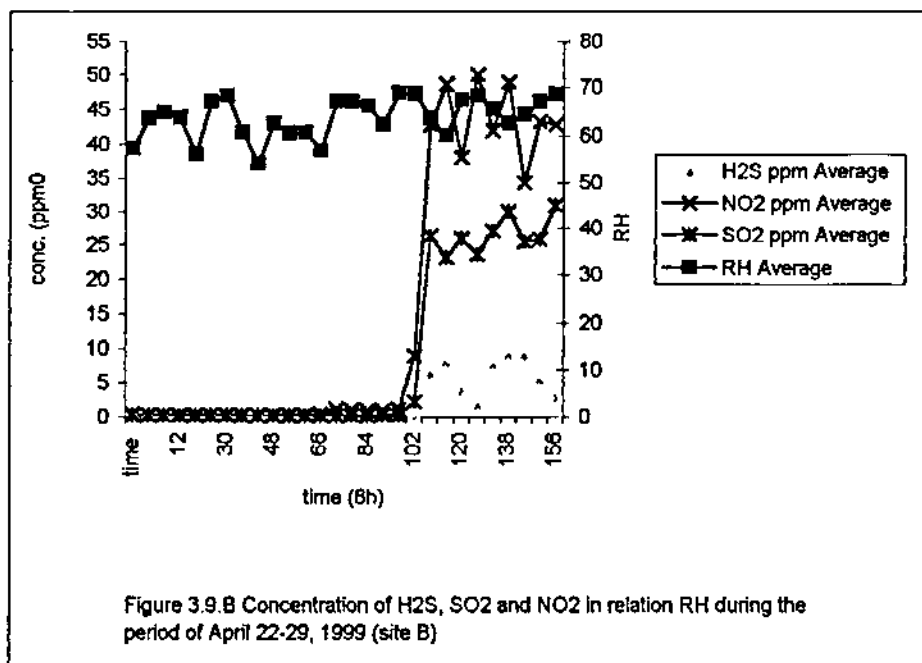
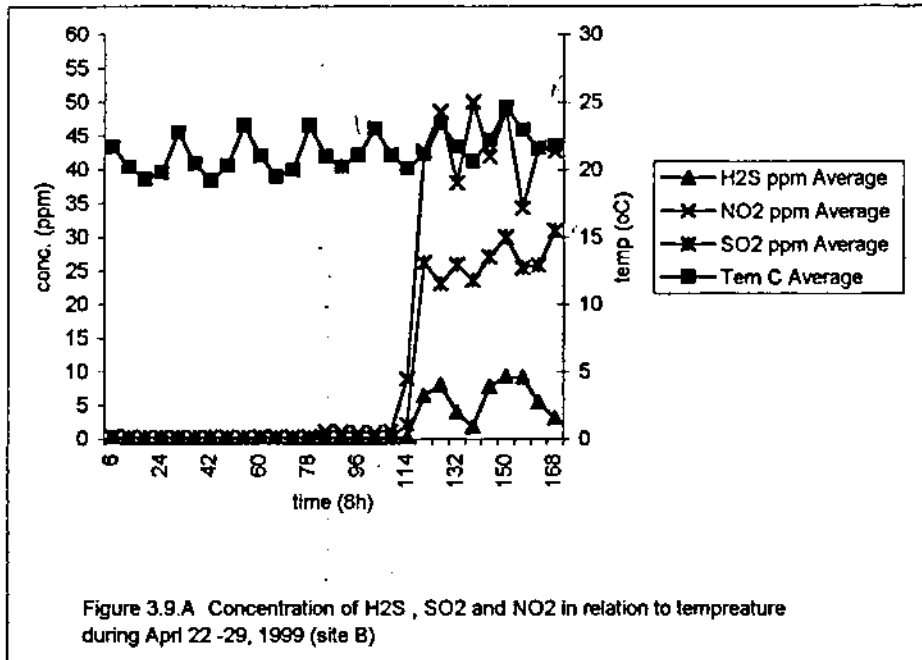
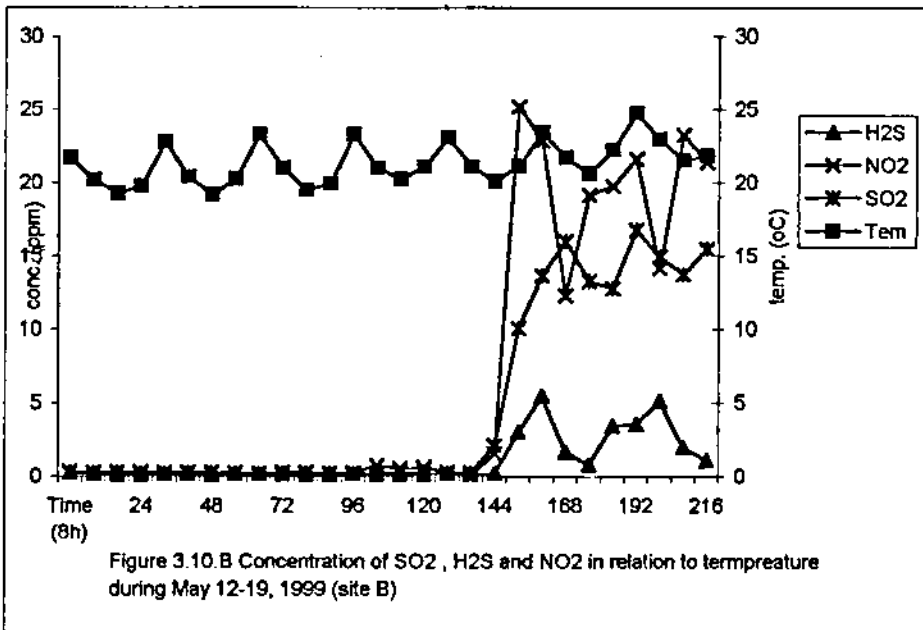
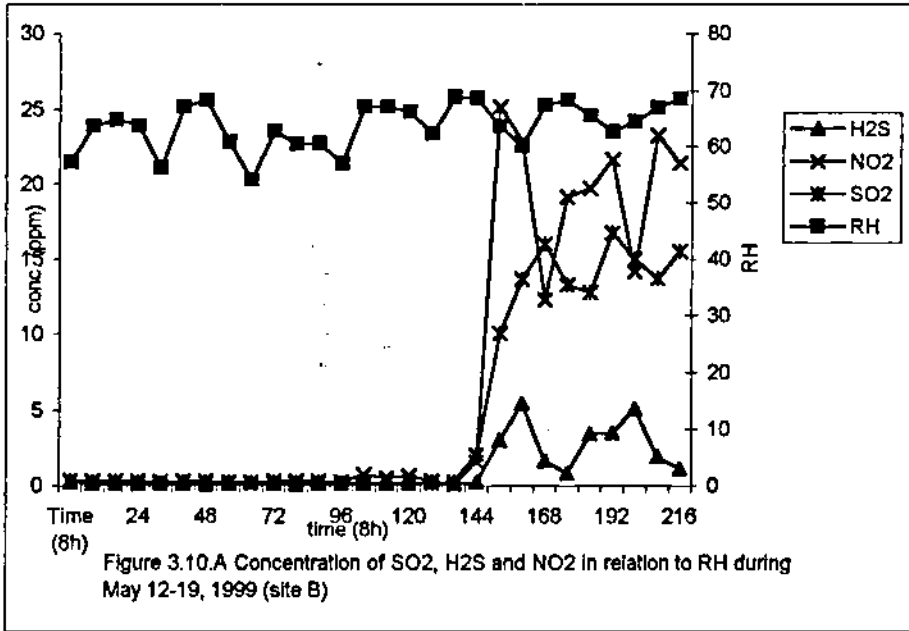


Figure 3.8.F CO₂ and CO concentrations in relation to RH during the period of Sept 1-8, 1998 (site B)





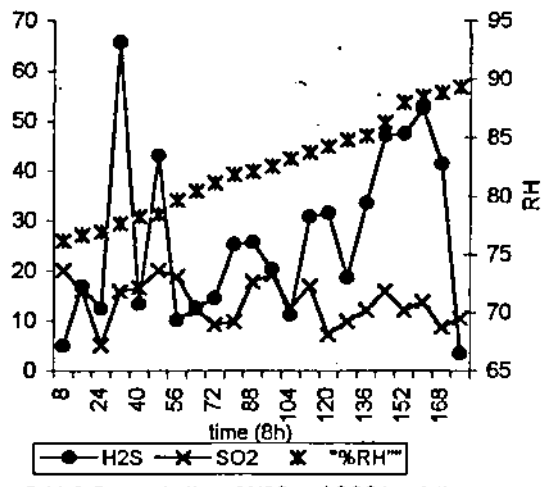


Figure 3.11.A Concentration of H₂S and SO₂ in relation to humidity during the period of September 28-October 9, 1999 (site B)

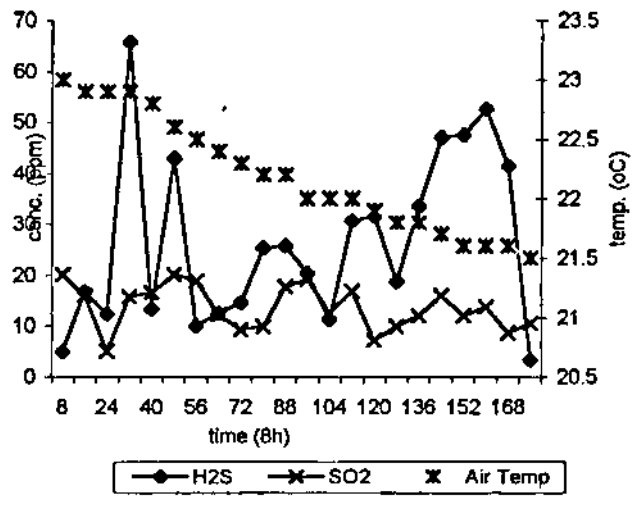


Figure 3.11.B Concentration of H₂S and SO₂ in relation to temperature during September 28- October 5, 1999 (site B)

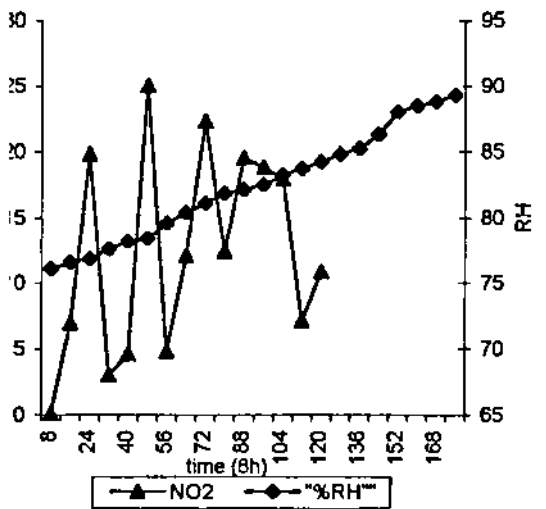


Figure 3.11.C Concentration of NO₂ in relation to RH during the period of September 29 - October 5, 1999 (site B)

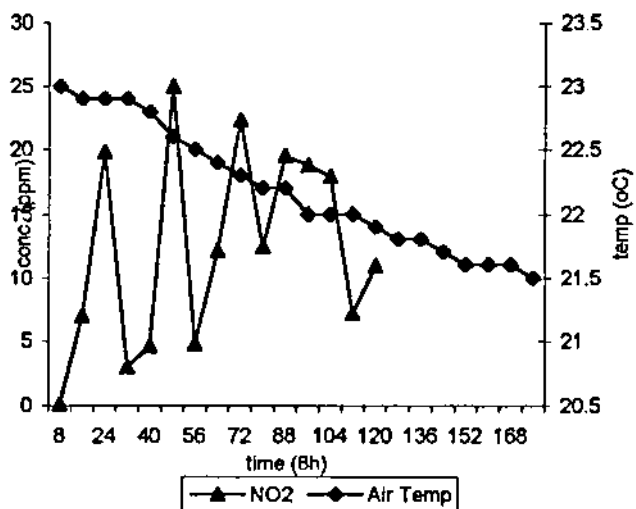


Figure 3.11.D Concentration of NO₂ in relation to air temperature during the period of September 28 - October 5, 1999 (site B)

3.2.3 SO₂ monitoring

The British allowable maximum is 2 ppm , and the American standard is 5 ppm .WHO for quality of air SO₂ must be 0.134 ppm.

Sampling sites (Locations A, B, C) differed significantly at ($p < 0.05$) in SO₂ concentration with location B showing higher SO₂ levels and location A showing lower SO₂ levels.

Location A

During the period of March 19-April 22, 1999 SO₂ concentration was ranged between 0.1- 1.89 ppm with an average of 0.26 ppm. In most of the reading as the temperature increased the concentration increased, the correlation with humidity was not clear (Figure 3.5.E,F, and Table A.6).

During the period of May 2-11,1999 the concentration of SO₂ lies between 0.15-1.57 ppm; the average concentration was 0.38 ppm. The concentration increased as the temperature increased, correlation with humidity the concentration increased as humidity decreased (Figure 3.1.B,C and Table A1).

Location B

During the period of May 28-30,1998 the concentration of SO₂ was arranged between 1.16-10.29 ppm; the average was 6.81 ppm, In this period as the humidity increased the concentration of SO₂ decreased, (Figure 3.6.B,C and Table A.7).

During the period of May 31-June 9, 1998 the concentration of SO₂ ranged 0.37- 22.4 ppm with an average of 6.34 ppm, according to wind speed the concentration decreased as the wind speed increased (Figure 3.2.A,B and Table A.2).

During the period of September 27-29,1998 the concentration of SO₂ ranged between 7.73-23.37 ppm; the average was 12.3 ppm. (Table A.3).

During the period of August 17-24,1998, the concentration ranged between 4.37- 34.8 ppm; the average was 14.7 ppm. In relation to relative humidity as the humidity increase the concentration of SO₂ increased (Figure 3.7.A,B and Table A.8).

During the period of September 1-8, 1998 the concentration of SO₂ was ranged between 3.9-37.98 ppm with an average of 15.18 ppm. According to relative humidity as the humidity

increased the concentration of SO₂ increased, according to temperature as the temperature increased the concentration of SO₂ increased, in correlation to wind speed as wind speed increased the concentration decreased (Figure 3.8.A,B , Table A.9) .

During the period of April 22-29,1999, the concentration of SO₂ ranged between 0.18-30.92 ppm; the average was 8.73 ppm. SO₂ concentration stay low during the first four days in which the concentration don't exceed 1 ppm after that the concentration increased, the temperature ranged between 17- 25 °C. (Figure 3.9.A, B and Table A.10).

During the period of May 12- 19,1999, the concentration SO₂ was ranged between 0.18-16.2 ppm; the average was 4.73 ppm. The concentration was low during the first 5days, on the sex day there was an increased in the concentration of SO₂ (Figure 3.10.A,B and Table A.11).

During the period of September 28-October 5, 1999 the concentration ranged between 5.1-20.1 ppm; the average was 13.7 ppm.(Figure 3.11.A,Band Table A12).

Location C

During the period of July 2-5,1999 the concentration of SO₂ ranged between 1.03-2.2 ppm; with an average of 1.75 ppm. (Figure 3.4.B and Table A.5) .

3.2.4 CO₂ monitoring

The concentration of CO₂ in the air is (0.0325%).

Sampling sites (Locations A, B) different significantly at ($p < 0.05$) in CO₂ concentration with location A showing higher CO₂ levels.

Location A

The concentration of CO₂ was between 0.1-4.79% with an average of 1.98% during the period of March 19-April 22, 1999, in correlation to temperature there was no constant relation, According to the humidity there was a differentiation according to the humidity changes (Figure 3.5.C,D and Table A.6).

During the period of May 2-11,1999 the concentration of CO₂ was between 0.16-4.61%; the average was 2.27%. In most the reading as the temperature increased the concentration increased (Figure 3.1.A,F and Table A.1).

Location B.

During the period of September 27-29,1998 the concentration of CO₂ was between 0.01-0.28% with an average of 0.15% (Table A.3).

During the period of August 17-24,1998 the concentration of CO₂ was between 0.18-1.66%; the average was 0.43%. The relation with temperature and humidity was not clear (Figure 3.8.C,D and Table A.8) .

During the period of September 1-8,1998 the concentration was arranged between 0.03- 1.69% with an average of 0.19% (Figure 3.8.E,F and Table A.9) .

During the period of April 22-29,1999 the concentration of CO₂ was ranged between 0.35- 4.10%; the average was 1.6% (Table A.10).

3.2.5 H₂S monitoring

The USA maximum is 10 ppm and the Canadian maximum concentration is 10-ppm .The National ambient air quality standard for H₂S is 0.07 ppm.

Sampling sites (Locations A, B) differed significantly at ($p < 0.05$) in H_2S concentration with location B showing higher H_2S levels being not far from charcoal kilns.

Location A

During the period of March 19-April 22, 1999 H_2S concentration was between 0.1- 5.43 ppm; with an average of 0.36 ppm. In about 90% of the reading as the temperature increased the concentration increased, the correlation with humidity was not clear (Figure 3.5.E, F and Table A.6).

During the period of May 2-11, 1999 the concentration of H_2S was ranged between 0.06-0.75 ppm; the average was 0.32 ppm. In this period as the temperature increased the concentration of H_2S increased, in correlation to humidity as the humidity decreased the concentration increased (Figure 3.1.B,C and Table A.1).

Location B

During the period of May 28-30, 1998 the concentration of H_2S was ranged between 1.36-10.7 ppm; the average was 5.9 ppm. In this period as the humidity increased the concentration

of H₂S increased , in relation to temperature the concentration decreased as the temperature increased (Figure 3.6.B,C and Table A.7).

During the period of May 31-June 9, 1998 the concentration of H₂S was ranged between 0.18-14.15 ppm; the average was 4.74 ppm. In related to relative humidity the concentration of H₂S increased as the humidity decreased, according to wind speed the concentration decreased as the wind speed increased (Figure 3.2.A,B and Table A.2).

During the period of September 27-29,1998 the concentration of H₂S ranged between 10.44-12.22 ppm. (Table A.3).

During the period of August 17-24,1998, the concentration of H₂S ranged between 1.23 - 45.94 ppm; the average concentration was 12.2 ppm. The relation with temperature was not clear, in related to relative humidity as the humidity increased the concentration of H₂S increased (Figure 3.7.A,B, Table A.8).

During the period of September 1-8, 1998 the concentration of H₂S was ranged between 1.55-46.83 ppm; the

average was 13.06 ppm. According to relative humidity as the humidity increased the concentration of H₂S increased, in related to temperature as temperature increased the concentration of H₂S increased and when the temperature decreased the concentration decreased, during this period as wind speed increased the concentration decreased (Figure 3.8.A,B and Table A.9) .

During the period of April 22-29,1999, the concentration of H₂S ranged between 0.1-9.19 ppm; with an average of 2.05 ppm. H₂S concentration stay low during the first four days in which the concentration don't exceed 1 ppm after that the concentration increased, (Figure 3.9.A,B and Table A.10).

During the period of May 12- 19,1999. The concentration H₂S was low during the first 5 days, the concentration of H₂S ranged between 0.1- 5.99 ppm; (Figure 3.10.A,B, Table A.11).

During the period of June 16-21,1999 the concentration of H₂S ranged between 0.54-26.97 ppm; the average was 17.32 ppm (Figure 3.3.B and Table A.4).

During the period of September 28-October 5, 1999 the concentration ranged 3.4-65.7 ppm; the average was 26.47 ppm (Figure 3.11.A,B and Table A.12).

3.3 Effect of Charcoal Production on School Children:

The effect of charcoal production on school children was obvious in areas where the kilns are close to the schools. Statistical analysis of the questionnaire showed that some children were suffering from respiratory tract diseases (asthma, breathing problems or difficulty and pulmonary symptoms). Using t- and F- tests (at $P < 0.05$), it was shown that there was a significant difference of charcoal production on children between schools that are close to the kilns and those that are far away from the kilns (Tables C.1.A, C.1.B, C.2.A, C.2.B, C.3 and C.4).

The questionnaire also showed that there was significant difference between male and female students in schools of the polluted areas (Table C.5) shows the effect on sex as revealed by statistical analysis using F-test at $P < 0.05$. It is clear from the

table that males are more affected by charcoal production in the above-mentioned areas.

CHAPTER FOUR

DISCUSSION

Chapter 4 Discussion

4.1 CO

The results have demonstrated that CO air pollution level was inversely correlated with distance from charcoal kilns with the highest levels being encountered in site C. In sites A and B; CO levels were very much lower than that in site C. In location C the concentration of CO (100.7 ppm) exceeded the maximum standard of USA and WHO (9ppm). In location A, the concentration (0.5-ppm) did not exceed the standards. In location B the concentration (50.48 ppm) exceeded the standards but it did not reach the levels of location C. In some periods the concentration in location B was close to the maximum and in other periods it was very high but don not reach that in location C.

The above results indicate that CO gas is emitted from charcoal klins. Since the environmental station had recorded the concentrations of CO in places that are close to the kilns. Iris 1975 mentioned that CO gas is one of the gases that produced by carbonizing process of wood.

School children living in areas with higher air pollution levels suffered significantly more frequent respiratory disease symptoms than those living in non-polluted areas. High concentration of carbon monoxide has been reported to cause certain types of blood poisoning and respiratory diseases (Ellegard 1997). Statistical analysis (using ANOVA F and T tests at $P < 0.05$) of the questionnaire showed that there was a significant effect of CO pollution from the kilns.

The relation between temperature and CO concentrations varies between locations. It was not clear in location C in most of the readings. In locations A and B, it was noticed that CO concentration increases as the temperature increases. These two locations are far from the kilns and it can be said that: as the temperature increases the wind weight decreases and it rises up carrying the CO.

4.2 NO₂

The results indicate that there was an increase in the concentration of NO₂, as the station becomes closer to the charcoal kilns so location C contains the highest concentration

The 8-hour concentration of NO_2 gas in location A was elevated but less than the standard. The maximum concentration does not exceeds.51ppm

In location B the concentration of NO_2 was high and exceeds the maximum standard in all the periods. The most elevation was during the period of April 22-29,1999 in which the maximum concentration reaches 50.04 ppm, which indicate that there was NO_2 pollution in that area.

In location C there was an indication that the concentration of NO_2 was high which gives indication that most of the NO_2 comes out from charcoal kilns. Its known that NO_2 formed by the oxidation of both the nitrogen in the air and particles, which are the result of, unburned materials. (The Committee on Health and environmental Effects, 1980).

NO_2 concentration was high in place B and C exceeds all the maximum standards. The statistics shoes significant different in the three locations with location C showing the highest concentration.

NO_2 is known to show more spatial heterogeneity than the other pollutants and hence the central monitoring site may be a poorer

proxy for exposure, NO_2 was associated with reduction in pulmonary function.

NO_2 to nitrate ion, this can occur up to hundreds of miles away from the coal burns plant. (The Committee on Health and environmental Effects, 1980).

The USA maximum for NO_2 is 5ppm and the British maximum is 3 ppm,

4.4 CO_2

At sea level the air ordinary contains about 0.03% CO_2 , To be safe a level of no more than 1% should be maintained.

In our study the concentration of CO_2 was in location A the concentration of CO_2 reaches a maximum of 4.79%, in this location there was an increasing in CO_2 concentration in some parts of the period this may comes from more than one sours the charcoal production and Combustion of all fossil fuels produces emissions of CO_2 , for fossil fuels.

The Committee on Health and environmental Effects ,1980 mentioned that identical energy outputs coal generates 1.8 times as much CO_2 as dose natural gas and 1.2 as much as fuel oil.

.In location B CO₂ concentration reaches 4.10% which is high and gives indication that CO₂ comes out from charcoal kilns.

4.5 SO₂

In this study there was moderate changes in SO₂ levels between the different locations

The maximum concentration of SO₂ in location A was 1.89ppm this less than the maximum standard but more than the WHO quality of air.

In location B the maximum concentration was 37.98ppm and this was during the period of September 1-8,1998 which is very dangerous and exceeds all the maximum standard, in this location the concentration of SO₂ was high during all the periods ,this indicate that SO₂ gas may come out from the carbonization process increase of SO₂ have effects on the people and the building. The workers are the most imposed to the gasses come out from the charcoal kilns, SO₂ has an effect on the teeth it cause teeth damage and this was seen in the mouth of most of the workers. Inhalation studies performed on human volunteers under controlled short-term exposure conditions show that Sulfur Dioxide alone at a concentration of 0.75ppm slightly affects

the respiratory function (Saric, et al., 1978). Timonen and Pekanen, 1997 mentioned that among the children the increase in SO₂ levels was associated with an increase in the incidence of upper respiratory symptoms. Asthmatics tend to be atypically sensitive to Sulfur dioxide exposure, Bronchodilator medication block the brunt-constrictive effects of SO₂ and provide a quantitative estimate of degree of protection that can be expected in typical mild asthmatics (Linn et al., 1988).

The British allowable maximum is 2ppm, and the American standard is 5ppm .WHO for quality of air SO₂ must be 0.134ppm

4.6 H₂S

The maximum U.S.A. standard 10PPM, and the Canadian standard is 10PPM, the national ambient air quality standard is 0.07ppm. In location A maximum concentration of H₂S was 5.43 during one period and this less than the standard. In location B there was an increasing in the concentration of H₂S it reaches 46.83ppm during the period of Septemper1-8, 1998 its very high concentration.

In our study the 8-hour average concentration was high in some period and low in other.

In our study we see that when the temperature increases the average concentration increases and this may be related to the place of the environmental station on the houses which is high so when the temperature is high the gasses goes up.

When the humidity increases the gas concentration decreases ship was not clear between RH. and the concentration of the gases when the RH increase the average concentration of most of the gasses decreases .

The wind direction has an effect on the pollution of the gases we study. When the wind direction is toward the houses and toward the station the concentration of the gases were high. And when the wind direction is to the opposite side the concentration was less, and when we asked the people in the village of Yabad they said; that when the wind is toward the Village the smell was strong.

4.1 Effect of Charcoal Production on School Children

Our study proved that a considerable effect of charcoal production was significant between children of schools close to the Kilns. Those children had defects in respiratory system, and some disease like asthma was severely common between those children. In

the results of the questionnaire there was health problems for the people living near the charcoal kilns. The questions give the indication on the upper respiratory symptoms like asthma, and other pulmonary symptoms problems which comes from air pollution or gas pollution like eye irritation at the morning and cough. In the present study males were more affected by air pollution than females. This may be attribute study to longer exposure time for males than females the more affected was the male more than the female and this because the male or boys go out of the house but the girl stay in the house more than the boys.

The association between respiratory symptoms and air pollution, association between respiratory health and air pollution from coal combustion have been observed elsewhere, in across-sectional study of three areas of Beijing China, heating with coal was associated with reduce lung function in non smoking adults (Archives of Environmental Health). Schwartz, 1993 mentioned that acute respiratory symptoms or illness have also been associated with particulate air pollution.

Ellegard, 1997 mentioned that eye irritation has been used as an indicator of air pollution in developing country.

For lung function and disease incidence studies children are a particularly suitable population group because of their sensitive respiratory system. Children also represent a nonsmoking and non-professional group who have spent the greatest part of their lives in the same environment (Saric, 1978).

Recommendation

- 1-Relocating the charcoal kilns from their current locations to new locations far from populated areas and traffic lines.**
- 2- Promote plantation in areas surrounding the charcoal kilns, which will decrease the gas pollution that comes out from the charcoal kilns.**
- 3- The workers must use safety clothes, shoes and masks must be needed as a mandatory safety means.**
- 4- Modern techniques with higher efficiency and less hazards must be adopted.**
- 5-Use big tanks for the production of charcoal instead of traditional ways, in this way we can use filters to decrease the pollution coming from the charcoal kilns.**
- 6- Use oven for the production of charcoal.**
- 7- Enforce legislative regulations and codes for plantation of forests to be used as wood source for charcoal production to prevent deforestation.**
- 8- Water tanks in surrounded areas must be closed tightly to minimize pollution of soluble gasses and suspended particulate.**
- 9- More research studies about the occupational health of the workers.**

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APPENDICES

APPENDIX A

Table A.1 The concentration of CO, H₂S, NO₂, SO₂ and CO₂ . RH, temperature
During the period of May 2-11,1999

Time 8h)	CO ppm	RH	Tem °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	CO ₂ %
8	0.01	32.94	25.36	0.72	1.29	1.57	1.84
16	0.13	53.76	19.72	0.16	0.26	0.21	1.33
24	0.11	48.65	21.42	0.22	0.23	0.21	1.16
32	0.14	36.45	28.72	0.69	0.59	0.65	3.12
40	0.42	53.21	22.54	0.31	0.40	0.38	1.84
48	0.34	51.62	19.15	0.12	0.16	0.15	2.21
56	0.28	41.78	21.23	0.53	0.39	0.43	2.26
64	0.12	32.97	28.86	0.71	0.67	0.69	2.86
72	0.11	45.45	23.58	0.40	0.33	0.37	0.75
80	0.32	41.96	21.02	0.49	0.35	0.41	0.16
88	0.21	43.38	22.68	0.35	0.35	0.34	1.47
96	0.50	18.84	33.66	0.75	0.83	0.85	2.77
104	0.07	25.18	28.33	0.70	0.69	0.69	1.42
112	0.04	48.76	22.48	0.35	0.42	0.38	1.99
120	0.27	80.05	20.61	0.15	0.31	0.24	2.99
128	0.06	48.44	28.00	0.37	0.22	0.28	3.23
136	0.23	63.51	21.21	0.26	0.41	0.33	1.09
144	0.14	55.51	18.93	0.13	0.19	0.17	2.83
152	0.31	72.84	19.59	0.16	0.32	0.26	2.84
160	0.19	42.79	25.27	0.53	0.47	0.48	4.14
168	0.16	80.18	18.56	0.17	0.39	0.31	2.60
176	0.12	90.98	16.36	0.04	0.30	0.20	1.13
184	0.26	76.69	18.22	0.08	0.28	0.21	4.61
192	0.18	61.38	21.74	0.24	0.23	0.20	4.34
200	0.12	75.33	17.11	0.11	0.34	0.25	2.03
208	0.28	82.27	15.14	0.00	0.23	0.18	0.42
216	0.25	71.95	16.50	0.06	0.26	0.20	2.23
224	0.13	54.61	19.80	0.15	0.21	0.18	3.94

Table A.2 The 8-hour concentration of CO, H₂S, NO₂ and SO₂ gas and the 8-hour reading of RH, air temperature and wind speed during the period of May 31-June 9, 1998

time (8h)	CO ppm	RH	Air Temp	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Win direction degree	Win speed M/s
8	8.14	63.62	21.79	2.70	6.12	5.98	265.01	12.58
12	5.12	75.34	19.49	7.75	11.35	9.55	263.33	9.85
16	4.57	81.73	18.25	1.58	7.78	5.80	223.91	4.93
20	5.23	77.97	19.11	7.28	17.20	12.49	227.09	6.04
24	2.12	65.32	21.83	3.72	4.81	4.63	258.94	11.26
28	10.7	77.79	18.53	7.22	9.23	6.97	259.23	8.58
32	12.4	84.54	17.16	3.35	9.33	8.91	229.05	4.44
36	9.54	75.52	19.19	11.04	16.39	12.43	227.88	5.10
40	13.5	61.03	22.69	8.48	1.10	1.36	264.59	9.96
44	12.54	77.92	19.06	14.15	10.19	8.65	203.94	8.26
48	12.21	86.68	17.16	3.15	10.01	8.93	167.60	6.31
52	10.25	73.74	19.01	3.68	8.67	8.49	165.02	5.38
56	8.97	59.27	24.30	2.68	4.08	0.69	218.46	9.77
60	13.84	78.54	20.12	0.88	2.90	2.59	223.28	5.17
64	7.51	91.59	17.45	2.10	8.21	5.55	235.99	1.09
68	11.69	77.65	19.76	2.88	8.98	4.13	191.78	4.04
72	6.52	54.59	24.70	2.16	0.68	1.00	226.72	10.77
76	14.2	80.09	19.88	7.35	11.01	8.05	216.07	6.21
80	12.5	91.59	18.01	1.03	13.30	6.67	226.94	3.59
84	15.47	79.55	20.35	2.47	16.23	19.82	227.33	6.57
88	13.85	60.42	24.81	14.70	5.73	4.43	236.14	11.14
92	14.25	78.58	21.04	10.66	17.61	6.54	227.98	7.53
96	12.54	93.37	18.43	1.60	25.03	13.79	233.43	5.24
100	10.25	80.46	20.09	4.62	14.64	6.14	185.96	6.25
104	6.25	54.84	25.08	4.10	0.97	1.62	220.78	11.89
108	8.36	65.81	21.69	7.60	5.15	5.64	225.51	6.75
112	8.98	59.53	19.71	1.84	11.19	12.03	148.58	1.40
116	13.28	54.64	21.48	13.19	10.25	8.73	231.58	3.66
120	7.96	40.50	26.82	3.87	1.16	1.10	222.70	10.05
124	9.85	61.07	21.98	6.83	5.71	4.49	228.03	5.04
128	15.27	90.10	18.24	3.42	23.84	22.40	180.44	0.85
132	13.68	80.09	20.06	0.97	2.43	2.80	231.94	3.95
136	12.47	60.39	24.88	0.96	3.79	0.87	242.27	12.73
140	9.97	78.92	21.04	3.63	3.53	4.68	246.25	7.61
144	8.65	93.62	18.11	0.18	0.52	0.37	214.27	4.62
148	9.67	85.29	19.46	0.29	0.34	0.37	170.78	5.25
152	6.19	63.06	24.24	1.47	2.34	1.02	228.33	11.51
156	7.91	64.04	24.03	4.71	6.36	1.53	224.89	11.99

Table A.3 The 8-hour concentration of CO, H₂S, and NO₂, SO₂ and CO₂ in addition to RH, temperature and wind speed. During the period of September 27-29, 1998

Time (8h)	CO ppm	RH	Tem °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Win Sp. MH	CO ₂ ppm
8	10.2	74.41	23.06	12.14	12.76	23.45	2.37	0.28
16	13.8	78.75	21.84	11.06	5.70	7.73	1.16	0.01
24	9.8	78.82	20.1	10.44	8.21	8.37	1.78	0.11
32	12.4	79.12	19.22	12.22	9.32	9.71	2.11	0.20

Table A.4 The concentration of CO and H₂S gases and Temperature during the period of June 16-21, 1999

Time 8 (h)	CO ppm	Temp °C	H ₂ S ppm
8	31.65	16.02	22.85
16	37.65	17.03	21.79
24	35.79	15.31	20.67
32	32.44	16.94	19.17
40	32.21	15.68	18.06
48	39.35	17.44	18.44
56	33.24	17.27	20.37
64	50.48	18.72	4.25
72	38.09	19.45	0.54
80	39.57	19.46	0.56
88	39.09	19.71	0.55
96	40.94	19.26	16.65
104	37.15	17.78	23.54
112	42.53	19.32	22.75
120	39.74	17.57	22.59
128	37.53	19.02	22.23
136	39.50	19.11	22.20
144	35.97	18.49	25.03
152	37.03	18.61	26.97

Table A.5 The concentration of CO, NO₂ and SO₂ gases and RH and wind speed
During the period of July 2-5,1999

time (8h)	RH	CO ppm	NO ₂ ppm	SO ₂ ppm	Wind. S. M\S
8	85.40	20.86	10.32	1.82	1.88
16	87.57	35.60	9.15	1.03	1.01
24	92.30	50.55	11.12	1.86	2.32
32	84.60	60.20	8.47	2.13	2.16
40	86.03	169.17	14.98	1.07	1.31
48	76.49	94.35	7.10	1.98	2.42
56	85.40	161.66	13.2	1.87	1.18
64	80.40	144.99	15.7	1.85	2.52
72	83.07	167.75	15.2	2.2	2.37

Table A.6 The concentration of H₂S, NO₂, SO₂ and CO₂ gases in addition to RH, temperature during the period of March 19-April 22, 1999

Time (8h)	RH	Tem. °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	CO ₂ %
8.00	68.70	14.29	5.43	1.51	1.89	1.10
16.00	80.81	11.91		0.12	0.10	1.89
24.00	82.45	13.02		0.18	0.13	6.55
32.00	83.17	13.44	0.10	0.27	0.20	3.88
40.00	86.54	12.23		0.14	0.12	1.51
48.00	86.15	10.66		0.10	0.10	1.02
56.00	77.55	13.23	0.10	0.19	0.16	4.48
64.00	67.55	16.10	0.10	0.29	0.22	2.84
72.00	85.32	11.94	0.10	0.18	0.15	0.71
80.00	83.59	10.61		0.10		0.10
88.00	63.65	13.96		0.11	0.10	2.68
96.00	49.29	18.08	0.27	0.28	0.28	2.54
104.00	75.48	12.96	0.10	0.20	0.16	1.70
112.00	77.48	11.15		0.10	0.10	0.20
120.00	63.06	14.56	0.44	0.21	0.23	1.43
128.00	39.42	18.66	0.52	0.46	0.50	1.90
136.00	60.08	13.49	0.20	0.12	0.11	0.85
144.00	39.95	13.81	0.41	0.32	0.37	1.28
152.00	37.21	18.25	0.48	0.38	0.42	2.31
160.00	21.41	25.56	0.64	0.63	0.65	2.43
168.00	29.55	19.47	0.58	0.51	0.50	0.99
176.00	35.84	18.10	0.53	0.43	0.45	2.86
184.00	68.69	17.13	0.10	0.29	0.22	7.92
192.00	70.29	16.90	0.10	0.33	0.25	4.72
200.00	84.63	12.31		0.17	0.13	2.91
208.00	88.49	11.05		0.10	0.10	1.12
216.00	80.05	13.37	0.10	0.24	0.16	3.21
224.00	81.08	11.54	0.10	0.24	0.16	2.80
232.00	83.93	10.02		0.10	0.10	0.83
240.00	86.12	7.39		0.10		1.06
248.00	79.51	14.36	0.10	0.23	0.17	3.32
256.00	75.94	15.54	0.10	0.31	0.23	2.91
264.00	91.74	12.59		0.16	0.11	1.53
272.00	94.31	11.89		0.10	0.10	1.67
280.00	74.33	15.67	0.10	0.22	0.15	2.97
288.00	70.06	16.62	0.13	0.29	0.21	2.08
296.00	89.02	12.72		0.14	0.11	0.82
304.00	92.49	11.47		0.10	0.10	0.77
312.00	64.36	17.53	0.37	0.25	0.24	0.79
320.00	61.52	19.32	0.37	0.38	0.34	1.27
328.00	87.59	13.13	0.10	0.17	0.14	0.20
336.00	74.81	14.01		0.10	0.10	0.47
344.00	51.26	21.20	0.46	0.34	0.34	1.11
352.00	50.36	20.38	0.25	0.24	0.20	1.99
360.00	52.51	15.92	0.46	0.29	0.26	1.40

368.00	82.53	13.82		0.17	0.11	2.07
376.00	72.31	17.32	0.16	0.26	0.20	3.63
384.00	72.42	16.09	0.14	0.31	0.23	3.39
392.00	85.68	12.59		0.16	0.11	1.44
400.00	88.96	12.18		0.12	0.10	1.04
408.00	69.66	15.86	0.10	0.26	0.19	3.74
416.00	71.08	15.31	0.10	0.27	0.21	3.37
424.00	85.38	12.59		0.14	0.10	1.13
432.00	83.70	12.71		0.14	0.10	1.42
440.00	59.81	17.43	0.20	0.21	0.19	3.84
448.00	56.52	19.35	0.19	0.25	0.18	1.95
456.00	32.42	18.63	0.56	0.45	0.46	1.65
464.00	39.57	17.87	0.52	0.37	0.36	1.44
472.00	39.96	22.84	0.52	0.41	0.46	4.79
480.00	52.36	18.51	0.46	0.44	0.42	1.63
488.00	89.27	13.20		0.17	0.11	0.16
496.00	91.46	12.38		0.12	0.11	0.73
504.00	58.71	20.34	0.27	0.19	0.15	1.65
512.00	73.48	17.94	0.24	0.32	0.26	2.57
520.00	92.28	13.52		0.18	0.11	1.01
528.00	90.16	13.94	0.10	0.19	0.12	0.75
536.00	38.70	23.78	0.58	0.48	0.49	2.27
544.00	31.10	21.96	0.65	0.55	0.55	2.09
552.00	44.85	17.93	0.46	0.29	0.31	1.46
560.00	33.16	20.70	0.58	0.48	0.50	1.51
568.00	26.18	30.06	0.74	0.70	0.76	1.50
576.00	27.65	25.82	0.71	0.63	0.65	2.14
584.00	21.08	24.26	0.61	0.50	0.50	0.52
592.00	23.48	25.45	0.62	0.52	0.53	0.92
600.00	32.69	27.05	0.64	0.57	0.61	2.40
608.00	31.72	22.62	0.61	0.52	0.53	1.25
616.00	71.43	17.03	0.22	0.33	0.27	1.76
624.00	82.72	15.68	0.10	0.26	0.20	2.13
632.00	57.33	19.48	0.19	0.23	0.17	3.01
640.00	66.83	15.75	0.15	0.26	0.21	0.81
648.00	84.21	12.91		0.15	0.10	0.49
656.00	80.99	14.08	0.10	0.21	0.18	1.17
664.00	58.36	18.44	0.13	0.24	0.16	2.89
672.00	73.34	14.98	0.13	0.28	0.20	1.27
680.00	78.18	12.84		0.15	0.10	0.60
688.00	74.54	13.96	0.10	0.19	0.15	2.31
696.00	56.19	19.19	0.14	0.22	0.15	3.93
704.00	75.79	15.41	0.12	0.30	0.20	1.21
712.00	87.33	12.99		0.16	0.10	0.88
720.00	79.89	14.42		0.16	0.12	0.70
728.00	44.46	23.06	0.47	0.36	0.41	2.07
736.00	53.02	17.54	0.30	0.21	0.22	1.75
744.00	61.96	15.01		0.11	0.10	1.30
752.00	51.99	18.11	0.51	0.26	0.25	1.06
760.00	33.27	25.26	0.65	0.57	0.60	1.84
768.00	72.75	17.54	0.24	0.38	0.30	2.39
776.00	94.37	14.88		0.26	0.18	2.09

784.00	82.49	16.60	0.14	0.26	0.17	1.45
792.00	55.95	23.40	0.27	0.20	0.17	2.37
800.00	76.76	18.02	0.18	0.35	0.26	1.78
808.00	91.69	15.17		0.25	0.16	0.52
816.00	75.98	18.49	0.16	0.25	0.18	2.29
824.00	58.22	24.03	0.29	0.24	0.18	2.83
832.00	79.15	17.13	0.15	0.37	0.28	2.12
840.00	87.32	15.72	0.10	0.24	0.17	2.22
848.00	72.97	19.60	0.17	0.24	0.18	2.05
856.00	36.69	26.19	0.63	0.47	0.53	3.56
864.00	65.97	18.26	0.17	0.34	0.26	1.23
872.00	86.23	15.57	0.10	0.25	0.16	1.82
880.00	73.57	18.52	0.13	0.26	0.20	3.48
888.00	48.09	22.68	0.43	0.33	0.35	3.38
896.00	75.41	15.69	0.21	0.28	0.21	1.45
904.00	91.02	13.21		0.17	0.10	1.51
912.00	69.26	18.15	0.15	0.23	0.16	5.73
920.00	55.80	20.96	0.23	0.25	0.21	3.15
928.00	83.27	15.90	0.13	0.32	0.23	0.80
936.00	88.25	14.61		0.20	0.12	1.22
944.00	69.29	19.21	0.20	0.24	0.20	1.13
952.00	56.35	24.12	0.30	0.26	0.19	2.47
960.00	84.69	17.13	0.16	0.37	0.28	1.62
968.00	94.99	14.78	0.10	0.26	0.15	0.45
976.00	49.50	23.22	0.51	0.40	0.38	2.37
984.00	20.86	30.59	0.85	0.83	0.86	2.07
992.00	25.72	24.59	0.63	0.54	0.53	2.15
1000.00	34.82	21.02	0.50	0.45	0.44	1.06
1008.00	39.55	25.28	0.53	0.47	0.49	2.24
1016.00	25.93	30.29	0.77	0.75	0.79	2.97

Table A.7 The concentration of H₂S, NO₂ and SO₂ and the 8-hour average reading of RH, temperature wind speed and wind direction, during the period of May 28-30, 1998 .

Time (h)	RH	Tem. °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Win sp m/s
8	61.89	20.77	1.54	6.32	6.12	12.21
16	70.27	17.29	9.51	11.04	10.29	7.93
24	75.44	18.23	1.36	6.66	4.37	4.93
32	75.38	12.92	8.87	15.29	9.55	6.37
40	59.35	20.88	2.50	3.63	4.33	9.93
48	71.00	16.07	7.14	9.60	8.62	7.77
56	84.50	17.20	3.28	9.05	7.28	4.22
64	68.29	14.15	10.70	14.54	9.58	5.36
72	53.80	22.80	7.89	1.00	1.16	8.47

Table A.8 The concentration of H₂S, NO₂, SO₂ and CO₂ gases and RH, wind speed, temperature and wind direction during the period of Aug. 17-24,1998

time (8 h)	RH	Temp °c	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Wind W. Deg.	Wind.S.m/S	CO ₂ %
8	72.06	25.49	34.50	26.09	34.48	212.14	4.83	0.63
16	93.88	23.11	30.41	27.17	19.49	174.73	2.47	1.66
24	78.13	25.42	34.68	28.44	27.84	163.37	5.89	0.90
32	56.20	21.40	1.99	4.21	4.37	217.13	9.24	0.31
40	73.73	25.91	22.40	22.55	19.47	233.39	5.91	0.30
48	82.11	23.31	15.88	8.73	7.08	173.16	2.09	0.29
56	78.45	25.36	45.94	20.52	23.64	150.35	5.91	0.42
64	47.94	9.97	3.08	6.80	6.34	187.16	8.76	0.53
72	73.04	25.36	22.10	19.66	17.44	223.63	4.40	0.28
80	87.95	23.06	14.04	13.52	11.38	171.83	2.44	0.26
88	75.94	24.89	19.75	20.27	18.24	141.84	6.13	0.27
96	55.98	29.21	1.87	9.75	6.59	181.56	10.25	0.48
104	74.13	24.45	9.70	16.96	18.46	214.66	5.59	0.29
112	86.13	22.62	4.57	10.18	10.67	127.77	3.22	0.41
120	69.30	25.16	11.77	22.62	15.58	140.98	6.43	0.35
128	60.89	28.96	1.85	4.33	6.10	190.58	9.36	0.33
136	75.78	24.24	8.35	21.03	14.02	185.92	5.27	0.23
144	87.81	22.54	1.23	2.60	5.20	162.36	3.99	0.18
152	69.31	25.28	9.50	19.03	24.68	143.48	7.68	0.33
160	56.56	28.79	2.24	4.91	6.01	196.58	10.40	0.38
168	68.77	24.22	7.45	18.81	26.01	182.72	6.90	0.43
176	83.89	22.86	2.96	12.03	10.80	142.90	5.01	0.24
184	68.23	25.44	7.84	18.74	24.96	154.39	5.46	0.45
192	52.20	22.17	1.97	5.03	5.08	234.10	10.94	0.52
200	63.78	24.37	10.14	15.79	12.82	210.09	5.36	0.43
208	85.32	22.64	5.13	7.48	5.25	190.13	1.93	0.32
216	61.98	25.59	7.91	20.84	19.80	140.62	3.97	0.47
224	53.14	28.54	2.38	7.94	10.62	212.17	8.61	0.52

Table A.9 The 8-hour concentration of H₂S, NO₂, SO₂ and CO₂ gases and RH, temperature and wind speed during the period of Sept. 1-8,1998

Time(8h)	RH	Tem °c	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	CO ₂ %	WinSp.m\S
8	82.65	25.49	36.30	29.09	37.98	0.28	4.96
16	93.95	23.10	38.31	28.05	27.46	1.69	5.10
24	81.58	25.64	39.15	28.44	27.22	0.83	6.09
32	57.92	30.91	2.02	3.56	3.94	0.03	11.50
40	80.39	25.74	24.56	24.61	18.87	0.13	6.45
48	91.76	23.25	14.52	11.62	7.20	0.16	2.07
56	78.77	25.71	46.83	22.10	23.21	0.23	6.04
64	53.46	30.55	3.18	5.58	7.28	0.05	11.54
72	80.44	25.06	24.03	21.73	21.61	0.20	8.60
80	88.11	23.01	15.49	14.11	12.00	0.08	2.50
88	75.95	25.24	19.44	19.25	20.98	0.13	6.46
96	56.34	29.14	1.55	9.33	6.59	0.05	11.53
104	79.48	24.17	12.86	19.14	18.46	0.15	5.59
112	86.17	22.58	5.17	10.40	9.21	0.15	3.19
120	71.75	24.77	11.50	20.74	16.18	0.17	7.35
128	61.54	28.86	1.91	3.64	4.72	0.03	11.47
136	83.02	23.99	7.69	21.03	15.70	0.09	5.54
144	87.86	22.49	1.30	3.42	6.12	0.11	3.88
152	71.19	25.69	8.94	21.06	23.57	0.15	8.09
160	60.25	28.66	2.13	4.59	5.49	0.04	11.99
168	79.96	24.03	8.58	25.93	25.99	0.10	6.63
176	84.00	22.79	2.06	11.68	8.97	0.09	4.92
184	69.06	25.91	9.44	18.15	23.28	0.15	5.77
192	55.00	29.08	2.28	4.59	3.90	0.06	11.80
200	79.36	24.07	10.57	17.84	13.56	0.08	4.58
208	85.75	22.52	6.37	8.80	7.26	0.11	1.85
216	61.88	27.20	6.51	20.17	18.95	0.06	6.11
224	56.22	29.24	3.19	10.06	9.59	0.05	10.71

Table A.10 The concentration of H₂S, NO₂, SO₂ and CO₂ gases .RH, and temperature during the period of April 22-29, 1999

time (8h)	RH	Tem. °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	CO ₂ (%)
8	57.23	21.69	0.27	0.36	0.29	4.10
16	63.63	20.17	0.11	0.30	0.26	3.2
24	64.75	19.28	0.10	0.29	0.26	2.4
32	63.68	19.76	0.10	0.30	0.20	3.1
40	56.21	22.75	0.16	0.26	0.19	1.5
48	67.08	20.41	0.13	0.31	0.24	1.2
56	68.33	19.18	0.10	0.30	0.25	1
64	60.76	20.28	0.11	0.27	0.20	0.15
72	54.08	23.26	0.16	0.23	0.19	0.35
80	62.68	21.01	0.12	0.30	0.22	0.91
88	60.45	19.45	0.10	0.29	0.21	1.2
96	60.54	19.93	0.11	0.28	0.20	0.85
104	56.99	23.26	0.16	0.35	0.18	0.86
112	67.06	20.96	0.13	1.16	0.22	
120	67.02	20.23	0.19	1.06	0.26	
128	66.16	21.08	0.12	0.96	0.20	
136	62.30	23.05	0.22	0.94	0.23	
144	68.86	21.06	0.20	1.10	0.26	
152	68.63	20.08	0.29	8.90	2.18	
160	63.59	21.13	6.36	42.77	26.27	
168	60.01	23.40	8.02	48.59	23.14	
176	67.38	21.68	3.97	38.02	25.91	
184	68.29	20.59	1.81	50.04	23.65	
192	65.48	22.19	7.73	41.91	27.08	
200	62.62	24.64	9.19	48.89	29.98	
208	64.35	22.93	9.11	34.34	25.45	
216	66.92	21.52	5.48	43.25	25.86	
224	68.52	21.77	3.08	42.74	30.92	

Table A.11 The concentration of H₂S, NO₂, and SO₂ gases and RH, temperature During the period of May 12-19,1999

Time (8h)	RH	Tem. °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Win.Dir. Degree
8	57.23	21.69	0.26	0.34	0.29	189.48
16	63.65	20.17	0.11	0.30	0.26	72.19
24	64.75	19.28	0.09	0.29	0.26	8.01
32	63.68	19.76	0.09	0.30	0.20	96.89
40	56.21	22.75	0.16	0.26	0.19	157.53
48	67.08	20.41	0.13	0.31	0.24	160.72
56	68.33	19.18	0.10	0.30	0.25	160.72
64	60.76	20.28	0.11	0.27	0.20	266.42
72	54.08	23.26	0.17	0.23	0.19	295.49
80	62.68	21.01	0.12	0.30	0.22	293.19
88	60.45	19.46	0.10	0.29	0.21	293.17
96	60.54	19.93	0.11	0.28	0.20	293.17
104	56.99	23.26	0.16	0.27	0.18	269.94
112	67.06	20.97	0.12	0.71	0.21	200.54
120	67.02	20.23	0.11	0.50	0.26	190.39
128	66.16	21.08	0.12	0.62	0.20	216.82
136	62.30	23.05	0.21	0.30	0.19	275.59
144	68.86	21.07	0.14	0.10	0.25	275.58
152	68.63	20.11	0.23	1.57	2.05	275.55
160	63.61	21.13	2.99	25.16	10.04	247.28
168	60.02	23.40	5.42	22.86	13.62	245.56
176	67.39	21.69	1.64	12.30	16.00	245.52
184	68.29	20.61	0.80	19.13	13.24	245.52
192	65.48	22.19	3.41	19.72	12.74	204.19
200	62.62	24.71	3.51	21.61	16.76	175.12
208	64.35	22.93	5.09	14.14	14.97	175.14
216	66.94	21.52	1.94	23.23	13.69	175.15
224	68.48	21.81	1.09	21.38	15.46	175.14

Table A.12 The concentration of H₂S, NO₂, and SO₂ gases and wind direction, temperature and wind speed during Sep. 28- Oct. 5, 1999

Time B (h)	Air Temp °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	win s. m/s
16	23	5	0.1	20.1	4.7
24	22.9	16.8	6.9	16	3.2
32	22.9	12.4	19.8	5.1	1.6
40	22.9	65.7	3	15.8	1.8
48	22.8	13.4	4.6	16.6	2
56	22.6	43	25	20.1	3.8
64	22.5	10.1	4.8	18.9	5.2
72	22.4	12.5	12.1	12.3	1.5
80	22.3	14.6	22.3	9.3	2.4
88	22.2	25.3	12.4	9.9	1.3
96	22.2	25.7	19.5	17.8	2.4
104	22	20.4	18.8	19.2	2.4
112	22	11.3	17.9	12.7	2.3
120	22	30.7	7.1	16.9	0.4
128	21.9	31.5	10.9	7.2	0.5
136	21.8	18.7		9.9	0.3
144	21.8	33.5		12.1	0.9
152	21.7	47.1		16	1.9
160	21.6	47.5		12.1	0.4
168	21.6	52.5		13.8	1.1
176	21.6	41.4		8.7	1
184	21.5	3.4		10.5	0.3

Table A.13 The concentration of CO, H₂S, NO₂ and SO₂ and the 8-hour concentration of RH, temperature Wind direction and Wind speed during the period of May 19-20,1999

Time (8h)	CO ppm	RH	Tem. °C	H ₂ S ppm	NO ₂ ppm	SO ₂ ppm	Win.Dir. degree	Win.Sp. M\S
8	6.24	53.85	27.66	0.33	0.25	0.29	203.98	2.64
16	3.41	66.01	23.52	12.71	6.53	0.31	129.79	3.65

APPENDIX B

B.1

بسم الله الرحمن الرحيم

استبانته

يقوم الطالب ياسر كايد طالب الماجستير في برنامج العلوم البيئية في جامعة النجاح الوطنية بإجراء بحث حول تلوث الهواء الناجم عن المقاحم و أثره على سكان المنطقة .
لدى يرجى تعبئة هذه إلا استبانة علما بان المعلومات التي تعطى سوف تستخدم لأغراض البحث العلمي فقط.

التاريخ : المدرسة : البلدة :
الاسم : تاريخ الميلاد :

الجنس : ١ - ذكر ب - أنثى.....مكان السكن:

ضع / ضعي دائرة حول رمز الإجابة المناسبة:

١. عدد مرات زيارة الطبيب في الأربعة أسابيع الأخيرة ١- لم أقم بذلك ب- مرة واحدة ج مرتان د- ثلاث مرات أو أكثر
٢. هل تشعر بلهته عند المشي ١- لا ب- خفيفة ج - متوسطة د- قوية
٣. هل تخرج بلغم عند السعال ١- لا ب- خفيف ج - متوسط د- قوي
٤. هل شعرة بخشخشة في الصدر ١- لا ب- خفيفة ج - متوسطة د - قوية
٥. هل شعرت بضيق تنفس على شكل ضغط على الصدر ١- لا ب- أحيانا ج- عند المشي القصير د - عند المشي لمسافات
٦. هل شعر بإعياء أو تعب عند المشي ١- لا ب- تعب خفيف ج - تعب متوسط د - تعب شديد في صباح اليوم:
٧. هل سعلت ١- لا ب- مرة واحدة ج - مرتان د - ثلاث مرات أو أكثر
٨. هل عطست ١- لا ب- مرة واحدة ج مرتان د- ثلاث مرات أو أكثر
٩. هل عانيت من سيلان المخاط ١- لا ب - خيف ج- متوسط د- قوي
١٠. هل شعرت بحرقة في العين : ١- لا ب - بشكل خفيف ج - متوسطة د- قوية
١١. هل شعرت بحكة في الأنف ١- لا ب - حكة خفيفة ج- حكة متوسطة د- حكة قوية
١٢. هل شعرت بألم في الحلق ١- لا ب - بشكل خفيف ج - متوسط د - قوي
١٣. هل شعرت بالحصى : ١- لا ب -حصى خفيفة ج- حصى متوسطة د -حصى مرتفعة

APPENDIX C

**Table C.1.A Effect of Place ,F-Test when Alpha =5%
Variable I (Q 1-7)**

Source of variation	D.F	S.S	M.S.S	F-values
Due to places	6	21.0014	3.5	11.4816
Due to Errors	909	277.1156	0.3049	
Total	915	298.117		

$F_c > F_{prop}$.

There is a different in the respiratory symptoms due to place

Table C.1.B F-test To know the difference between which place

Mean		1	2	3	4	5	6	7
2.0485	Nazlet Zead (1)							
1.8235	Zabadch (2)							
1.6167	Yabed (3)	*						
1.3839	Arabeh (4)	*	*	*				
1.5238	Ancn (5)	*						
1.7532	Kferet (6)							
1.3839	Am archan (7)	*						

*= There is a difference to the greater mean

Table C.2.A Effect of Place ,F-Test when Alpha =5%
Variable II (Q 8-14)

Source of variation	D.F	S.S	M.S.S	F-values
Due to places	6	47.4525	7.9088	28.4199
Due to Errors	909	252.9586	.2783	
Total	915	300.4111		

$F_c > F_{prop}$.

There is a difference in the respiratory symptoms due to place

Table C.2.B F-test To know the difference between which place

Mean		1	2	3	4	5	6	7
2.3208	Nazlet Zead (1)							
1.4020	Zabadch (2)	*						
1.5027	Yabed (3)	*						
1.2606	Arabch (4)	*		*		*		
1.7222	Anen (5)	*						
1.4949	Kferet (6)	*						
1.4479	Am archan (7)	*						

* = there is a difference to the grater mean

Table C.3 T-test between polluted and non polluted villages
Group I (Q 1-7)

Area	Number of cases	mean	SD	T-calculated	D.F	T-table
Polluted	685	1.6669	0.588	5.61	914	1.96
Non polluted	231	1.4273	0.475			

T-calculated is greater than T-table there is a difference toward the polluted villages

Table C.4 T-test between polluted and non polluted villages
Group II (Q 8-14)

Area	Number of cases	mean	SD	T-calculated	D.F	T-table
Polluted	685	1.5606	0.586	3.58	914	1.96
Non polluted	231	1.4055	0.516			

T-calculated is greater than T-table there is a difference toward the polluted villages

Table C.5 Effect of sex, F-Test at alpha = .05

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Variable	Hypoth. ss	Error ss	Hypoth. MS	Error MS	F Sig	
I	2.22694	283.63610	2.22694	0.31100	7.16048	
II	1.52475	292.57704	1.52475	0.32081	4.75284	

There was difference between the sex factor

قياس التغيرات اليومية في تراكيز ملوثات الهواء وعلاقة ذلك
بانتشار أمراض الجهاز التنفسي لدى أطفال المدارس في منطقة
المفاحم في يعبد

ياسر عمر حافظ كايد

المشرف

بروفيسور محمد سليم علي اشتية

و

دكتور معتصم بعباع

في هذه الدراسة تمت دراسة تأثير إنتاج الفحم النباتي في منطقة يعبد وذلك بقياس تركيز الغازات المنبعثة من عملية الكربنة (التفحم) والغازات هي $CO, CO_2, NO_2, SO_2, H_2S$ وذلك على مدى ١٤ شهرا (أيار ١٩٩٨ - آب ١٩٩٩)، باستخدام محطة رصد بيئي ELE 8000/EMS 1417 Environmental Data logger ، حيث تم اختيار ثلاثة مواقع لقياس تراكيز الغازات فيها وكانت المواقع كما يلي الموقع الأول يبعد حوالي ١٨٠م عن تجمع المفاحم القريب من بلدة يعبد إلى الشمال الشرقي منه ، والموقع الثاني يبعد حوالي ٧٠م، مقابل للمفاحم وللأدخنة الصادرة منها، أما الموقع الثالث كان في تجمع المفاحم ،هناك عوامل فيزيائية أخرى تم قياسها مثل سرعة الرياح، اتجاه الرياح، درجة الحرارة والرطوبة، أما دراسة تأثير الغازات المنبعثة عن المفاحم على

الجهاز التنفسي للسكان في منطقة يعبد فقد تمت من خلال استنباهه مكونة من ١٤ سؤال وزعت على طلاب الصف الخامس في أربعة تجمعات سكنية يعتقد أنها ملوثة وثلاثة مجتمعات سكنية يعتقد أنها اقل تلوثا، ومن نتائج الدراسة وجود ارتفاع في تراكيز الغازات في المواقع الثلاث، و أن تركيز الغازات يرتبط بشكل معكوس مع المسافة عن المفاحم ، وكان أعلى تركيز في الموقع الثالث حيث كان تركيز الغازات (CO100.57ppm,NO₂11.7 ppm and SO₂1.75ppm) أما في الموقع الأول فقد كانت تراكيز الغازات منخفضة جدا حيث كان معدلها (CO 0.19ppm, NO₂ 0.29 ppm, SO₂ 0.26ppm,CO₂ 1.6% and H₂S 0.36ppm) بينما كانت التراكيز في الموقع الثاني اقل من الموقع الثالث أعلى من الموقع الأول: CO 37.9ppm, NO₂ 14.9ppm, SO₂ 14.7ppm, CO₂ 1.98% and H₂S 12.2ppm). تجاوزت التراكيز الحدود القصوى المسموح بها دوليا بينما لم تتجاوزها في الموقع الأول. كما أوضحت التحليلات الإحصائية باستخدام ANOVA and F-test أن هناك فرق واضح في تراكيز الغازات في المواقع الثلاثة، كما أظهرت التحليلات الإحصائية في هذه الدراسة تأثير واضح لعملية إنتاج الفحم على طلاب المدارس القريبة من المفاحم، كما أظهرت أن هناك فرق واضح بين تآثر الفتيات والأولاد في المدارس القريبة من المفاحم حيث تبين أن الأولاد أكثر تأثرا من الفتيات.