

**UNIVERSITY OF JOS**  
**MEDICAL RESEARCH PROJECT**

**ANNUAL REPORTS**

**1983**

**1984**

**1985**

**University of Jos**  
**Japan International Cooperation Agency(JICA)**

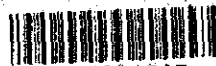
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FIRST ANNUAL REPORT: DECEMBER 1983

UNIVERSITY OF JOS/JAPANESE INTERNATIONAL CO-OPERATION AGENCY  
RESEARCH PROJECT

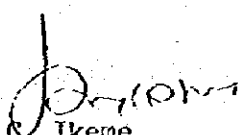
1. The Master Plan for this project was approved and the Record of Discussion signed in June 1982. Members of The Japanese Team arrived in October 1982.

Following the arrival of the members of The Team, the Co-ordinating Committee was formed under the Chairmanship of The Dean, Faculty of Medical Sciences. The function of this Body was to define the various research groups and co-ordinate their activities from time to time.

2. From the period January 1983 to October 1983, the various research groups planned their projects and in a number of cases conducted pilot activities. The Research groups broke themselves up into a number of sub-groups concerned with specific projects. With the arrival of equipment in September 1983 most of the groups were able to commence work in their various projects and the enclosed reports represent a summary of their activities in the last three months.
3. These activities reflect the Faculty objectives - that the Faculty will organise means of studying the health needs of the population and of providing practical solutions to them. In organising to achieve these objectives a wider participation than the Faculty has been involved. The following Departments have been actively involved in the research projects under this scheme and the attached reports emanate directly from them:

- a) Department of Community Health
- b) Department of Microbiology
- c) Department of Chemical Pathology
- d) Department of Paediatrics
- e) Department of Biochemistry
- f) Department of Zoology
- g) Department of Chemistry
- h) Department of Geology and Mining
- i) JICA Project Team

It is envisaged that at a later stage other Departments such as Medicine, Pathology, Social Sciences etc. may become involved in the Project.

  
A.C. Ikeme  
January 1984

cc: Vice Chancellor

ACI/eitn

*Shirley T. Jones*  
*Director*

## ANNUAL REPORT

### UNIVERSITY OF JOS/JICA RESEARCH PROJECT ON DISTRIBUTION OF ENVIRONMENTAL TRACE ELEMENTS IN PLATEAU STATE AND ITS IMPACT ON THE HEALTH OF LOCAL INHABITANTS

#### 1 INTRODUCTION

This study is made up of -

- 1.1 Trace element analysis in surface water in Plateau State by Potentiometric and voltametric methods
- 1.2 Trace element distribution in bed rock, latrite and stream sediments of the Jos Plateau, and its environmental significance
- 1.3 An analysis of the trace element composition of various categories of natural and artificial water bodies in the Jos Plateau and the bioaccumulation of the elements in some aquatic organisms in relation to water sources and mining activities
- 1.4 Plasma and tissue levels of important trace elements and the metallo carrier-proteins among the inhabitants of the mining district of Plateau State
- 1.5 The evaluation of the trace element levels of city water supplies for the Jos metropolis and their metabolic/health significance

In the 1982/83 academic session, some work had commenced on Sub-Project 2 & 3 whereas Sub-project 1 is just about taking off with preliminary sampling and analysis. Sub-project 4 is designed to take substantial bearing from preliminary results arising from the other parts of this study.

#### 2 ACTIVITIES OF THE GROUPS

- 2.1 Sub-Project 1 Trace element analysis in surface water in Plateau State by Potentiometric and Voltametric methods

##### 2.1.1 Investigators

Mrs E Akueshi	Department of Chemistry
Mr C F Oduoza	Department of Chemistry
Dr A Russer	Department of Chemistry

In respect of this project, work is concentrating on standardization of methods for analysis of water from the various studies mentioned on Projects 2 & 3, using Iron Selective Electrodes.

### 2.1.2 Preliminary results

For the calibration of bromine selective electrode, (OP-Br No 802 from 7MD radelkis BUDAPEST), bromide standard solutions ranging from  $10^{-1}M$  to  $10^{-5}M$  were prepared. Potassium bromide (Analytical grade from MERCK) dried between  $100^{\circ}C$  and  $115^{\circ}C$  in the oven was used in preparing the standard solutions. Salt bridge was prepared using ammonium nitrate and agar powder.

Using the expanded scale pH meter (Model 290 Mk 2 pH Meter from PYE UNICAM) or the digital pH meter (PW 9409 digital pH meter, Philips from PYE UNICAM) and the calomel reference electrode, (EIL No 1370710), the bromine selective electrode will be calibrated and in about ten days time the water samples collected will be analysed.

Parallel to these interference effect between  $Br^{-}$  anions and other anions present in solutions will be investigated in order to compute the true activity of  $Br^{-}$  anions in samples.

## 2.2 Sub Project 2 - Trace element distribution in bed rock, latrite and stream sediments of the Jos Plateau, and its environmental significance

### 2.2.1 Investigators

Dr A E O Ogezi	Department of Geology and Mining
Dr M Alam	Department of Geology and Mining
Mrs M E Adiku-Brown	Department of Geology and Mining

This project is aimed at examining the effects of mining and mineral processing on the distribution of certain environmental sensitive trace elements, such as lead, arsenic and mercury, examining the distribution of economically important elements for their mining potential and environmental impact, the effects of weathering and sedimentation on the different elements, and a source of geochemical baseline information for mining, industrialisation and determination of sites for dams, urban water supply and irrigation.

### 2.2.2 Preliminary results

More than 250 sediments and rock samples have been collected. Most of them are from around Jos city, especially from Gurum River, Delimi River, ponds and lakes from Rayfield area. The rest of the samples are scattered over the State (see Map and localities).

Almost 150 of these samples have been analysed from sedimentological point of view. Silt and clay fractions of each of these samples have been separated and are ready waiting chemical analyses.

Chemical analysis of the sediments and hard rock samples are being delayed because of lack of chemicals, platinum and teflon beaker ware, as well as lack of centrifuge and rock crushing equipment. Study of the fine-grained sediments, especially their hydrodynamic behaviour and their rates of absorbing and releasing chemicals into the water mass and the biota is also being delayed due to lack of necessary equipment.

Since the beginning of the year, a systematic water, sediments and rock sampling has been carried out with other Zoology Group handling Project 3. Water collected from these trips are being analyzed using the Department of Geology and Mining Atomic Absorption Unit. So far water, sediments and rock samples have been collected from Jos International Brewery Lake, Rayfield Sailing Club's Lake, Shendam, Jing River, Helpang, Shendam Dam, Pan-Dam, Dep River, Bukuru Water Supply Lake, Mine Lakes and rivers between Bukuru and Pankshin Panyam Fish Farm ponds and Pankshin Water reservoir.

A number of elements for which AAS Lamps are available are being analysed at this moment. More standards and lamps for AAS not available for other elements would facilitate the investigation.

In some cases, it may be necessary to concentrate the water because the concentration levels of certain elements may be too low to be detected by the AAS.

#### 2.2.3 Constraints

This group is encountering difficulties in carrying out chemical analysis of the sediments and bed rock samples. The following alternative materials for digestion of the samples are needed -

- A (1) Hydrofluoric Acid - 4 litres
- (2) Teflon Vessels - Hydrofluoric Acid resistance - 30 ml capacity

OR

- B (1) Sodium Carbonate - 2 kg
- (2) Nickel Crucible - 30 ml capacity

OR

- C (1) Sodium Peroxide - 2 kg
- (2) Nickel Crucible - 30 ml capacity

OR

- D (1) Lithium Tetraborate
- (2) Platinum Crucible (95% platinum coated with 5% gold)?

2.3 Sub-Project 3 - An analysis of the trace element composition of various categories of natural and artificial water bodies in the Jos Plateau and the bioaccumulation of the elements in some aquatic organisms in relation to water sources and mining activities.

2.3.1 Investigators

Prof Chiweyite Ejike    Department of Zoology  
Dr D I Anadu            Department of Zoology  
Mr I J Chidobem        Department of Zoology  
Mr E C Kemdirim (MPhil/PhD candidate) Department of Zoology

Investigations on the general biological aspects of this work commenced during 1983. Three categories of waters have been examined for their levels of lead, zinc, cadmium and copper and the associated distribution of planktons, macro-invertebrates and fish. These water types are -

- (1) The Hill Stream (the Delimi) which develops into a third other stream that courses through the city of Jos.
- (2) The Reservoir for city water supply arising from damming of the Shen River.
- (3) Ponds in artificial fish farm at Panyam.

2.3.2 Preliminary results

The investigations show fairly high levels of Pb, Zn, Cd and Cu in water sediments, planktons and macro-invertebrates. During the latter part of the year 1983 and since the beginning of 1984 the scope of this Sub-Project has been expanded and sampling locations/sites chosen. Parallel areas representing mining(M) & non-mining(N) areas have been chosen. Twenty such sites which come under three broad zones of (a) Jos area, (b) Highlands outside Jos, and (c) Lowlands, have been delimited (see Fig I and Table I, showing the Map of the general area of study and sampling locations and sites).

Systematic collection of samples has commenced and so far water sediments, plankton, fish food organisms, eg aquatic insect larvae and other macro-invertebrates including some fish where possible have been taken from the following sites which can be identified from the figure and table referred to above -

- (1) Old Mine Lake from which the local Brewery takes its water.
- (2) Old Mine Lake being used for recreational activities by a Sailing Club.
- (3) The Shen River reservoir.

- 5-
- (4) Old Mine Lake used for city water supply for Bukuru.
  - (5) Down stream rivers, draining from old mine lakes and used for irrigation of vegetable farms at Bisichi and Kassa respectively.
  - (6) Kassa Dam.
  - (7) Panyam Reservoir for city water supply on the River Rim.
  - (8) Pankshin Water Supply Dam.
  - (9) River Jing at Zong.
  - (10) The City Water Supply Reservoir under construction at the city of Shendam and on river Shamanka.
  - (11) Pan Dam Lake.
  - (12) Dep River.
  - (13) City Water supply, Akwanga.
  - (14) Ansoh River, southern edge of the Plateau.

Twenty-five batches of samples representing surface water, sediment, planktons and macro-invertebrates have been collected. Water samples are being analysed using the JICA Atomic Absorption Spectrophotometer for trace elements. In addition, the biological samples, namely plankton, macro-invertebrates and fish are being pre-treated for evaluation of distribution of trace elements.

As anticipated, a good range of variations is noted in relation to concentration of the trace elements as well as in other water quality characteristics, but analysis of data is not yet in a stage to allow for correlative generalizations.

### 2.3.3 Constraints and Projections

The work has generally no major constraint except for the availability of the urgently required reagents as contained in our request to the Team Leader, Japanese Mission, Dr H Takahashi. The original protocol has been slightly modified to ~~create~~ a separate dimension to be covered in Sub-project 5.

## 2.4 Sub-Project 4 - Plasma tissue levels of important trace elements and the metallo carrier-proteins among the inhabitants of the mining district of Plateau State

### 2.4.1 Investigators

Dr U P Isichei    Dept of Clinical Chemical Pathology  
Dr S C Das        Dept of Clinical Chemical Pathology

The research plans as outlined in the paper presented by the Department to the Convener for the Trace Element Project, Prof Chiweyite Ejike on 8 November 1983 remains unaltered.

The list of the reagents and other laboratory equipment needed for the investigation has been submitted to Dr H Takahashi and are being expected.

Sample collection is in progress and would be processed periodically in batches.

During the last vacation when Mr Noda was giving the senior staff members of the Trace Element Group some instruction on how to use the atomic absorption spectrophotometer our most senior technologists were away from the Department. Mr Noda returned to Japan in December but our technologists are now back and would learn what they missed from their colleagues.

### 2.4.2 Preliminary results

The collection of samples have now commenced and the baseline picture in apparently normal people in Plateau State is being determined. Thereafter, we hope to carry out an assessment of the trace element pattern among the mining population in Plateau State. In addition, blood and tissue samples would be collected from various population groups of interest after the assessment of the data collected from other teams in the Trace Element Group from the Faculty of Natural Sciences and the Department of Biochemistry. An assessment of the plasma and tissue trace element levels among some patients with certain clinical conditions, eg liver cirrhosis, hepatomegaly and cardiomegaly, etc would then be conducted in the final stages of the study.

### 2.4.3 Projection

As mentioned in the project plan which was submitted last year, we intend to compare the trace element levels derived from the above population groups with the pattern in the low-lying non-mining district of some suitable area in the southern part of the country.

2.5 Sub Project 5 - The evaluation of the trace element levels of city water supplies for the Jos metropolis and their metabolic/health significance.

2.5.1 Investigators

Dr K K Sen     Department of Biochemistry  
Dr G A Ubom    Department of Biochemistry

This report represents the preliminary work and data obtained by the Biochemistry Department, Faculty of Medical Sciences, as part of its participation in the JICA project for the year ending December 1983. All the data were obtained using the atomic absorption spectrometer which was successfully installed and tested around November 1983.

2.5.2 Preliminary results

Other than the test runs which were necessary to ascertain the reliability of the AAS, water samples from several sources - rivers, treatment reservoirs, pipe-borne and wells - have been analyzed for health-related cations content. These have included Sn, Mg, Ca, Pb, Mn, Cu, Na, K, Zn, Fe and Cd. There appears to be some disparity in the levels of these metals in treated tap water and that of well water. The levels in these water samples are to be compared with acceptable international levels to verify the suitability of these waters for drinking. Previous water analysis studies have implicated some of these cations as being responsible for some of the diseases arising from pollutants and deficiencies of the elements. The results are too preliminary to permit such association.

Crops and vegetables have been collected from gardens irrigated with industrial and mining wastes. The wastes from these mining and industrial areas are being analyzed. The crops and vegetables from these gardens have been digested and are to be analyzed in order to establish if the heavy metal pollutants found in the industrial and mining effluents are incorporated into the vegetables. Analysis on the crops and vegetables harvested from farms irrigated with the industrial and tin-mining effluents are at present not complete.

2.5.3 Projection

These preliminary studies have provided data that will guide us in designing and achieving our goals of (i) determining the concentrations of health-related inorganic (cations) constituents of drinking water, food, soils and plants in Jos (Plateau) area, (ii) generating a water quality map for the Jos metropolis under studies and (iii) establishing the effects of the detected cationic levels on animals. The monitoring of the levels of pollutants found in the industrial and mining effluents, as they are transferred to man is a necessary corollary of this investigation and will be pursued in Sub-Project 4. Titration techniques are being worked out for the determination of Hg, As and I. This has become necessary because the AAS which is the major equipment for the "trace" elements assay is not presently set-up for their determination.

This map illustrates the Plateau State region in Nigeria, highlighting its geographical features and administrative boundaries. The state is shaded with a stippled pattern. Major towns and cities are marked with dots, including Kaduna, Bauchi, Wukari, Makurdi, Lafia, Keffi, Shendam, and others. The map shows a network of rivers, including the Benue, Gongola, Kaduna, and others. State boundaries are indicated by dashed lines. A scale bar in the bottom left corner shows a distance of 60 km, and a north arrow is located in the top left corner.

Table I

UNIVERSITY OF JOS/JICA TRACE ELEMENT STUDIES CHOICE OF  
SAMPLING LOCATIONS AND SITES.

	LOCATION	AREA	ROCKTYPE	MINING ACTIVITY	ASSOC. RIVER/STR.
A JOS AREA	JOS 1	GOLD BASE GRUM RAPIN JAKI SHEN RAYFIELD MIANGO ROAD MIANGO AREA	BIOTITE GRANITE    BIOTITE GRANITE	M     N	KARAMI DELEMI SHEN R.  RAPIN LAWA NGELL RIVER
	JOS 2	HELIPANG AREA PANYAM	BASALT BASALT	M N	
	JOS 3	BARKIN - GANGERE DOGON GARBA	OLDER - GRANITE  OLDER - GRANITE	M  N	KASSA R.  QUREE R.
	PANKINSHIN 1		SYENITE SYENITE	COYKES -DYKES	
	PANKINSHIN 2		MIGMATITE MIGMATITE	OLD MINING LOCATION N	
	AKWANGA 1	AKWANGA- (WAMBA) ABU  GUDI	MIGMATITE GNEISSES  ✓	M  N	ABU  GUDI
B HIGHLANDS	AKWANGA 2	NASARAWA EGON	YOUNGER GRANITES	M(?) N(?)	MADA TRIB & MAGAMA TRIB
	NASARAWA	ODEGI (?)	BIOTITE	M N	OKONA MADA
	LAFIA	AZARA PANDAM	SHALES SANDSTONES GNEISSES	M N	DEEP RIVER
	SHANDAM	WASE  SHANDAM	SEDIMENT SANDSTONES & SHALES LEAD, PBS, ZINC	M  N	WASE RIVER
C LOWLANDS					

## UNIVERSITY OF JOS/JICA RESEARCH PROJECT

### REPORT FOR 1983 ON ENDEMIC GOITRE STUDIES

The overall objectives of JICA-UNIJOS Research Project on Endemic Goitre are as follows:-

1. To study the prevalence and geographical distribution of endemic goitre in Plateau State.
2. To study the possible etiological factors of endemic goitre in the area.
3. To study the prevalence of such conditions as deaf mutism, endemic cretinism, hypothyroidism etc. which are known to be associated with endemic goitre prevalence.
4. To study the chemico-pathological, histopathological, immunological and clinical profile of endemic goitre cases in the area.

The above objectives are aimed to be achieved by a multi-disciplinary study phased out over a period of five years. The following workers are currently involved in the study:-

#### Department of Community Health

1. Dr. I.C. Tiwari
2. Ms. G. Onyejuruwa
3. Mr. Kba Okoronkwo

#### Dept. of Chemical Pathology

1. Dr. U.P. Isichei
2. Dr. S.C. Das

#### Phase I of the Study:

A WHO seminar on Endemic Goitre\* has suggested "A prevalence of 5 percent and above of enlargement of thyroid in pre-adolescent age group" as the criterion for endemicity of endemic goitre.

Phase I of the present study therefore envisages goitre survey among secondary school children of Plateau State. A total of 6000 boys and girls of adolescent and pre-adolescent age group will be examined for the purpose using the following WHO criteria for goitre enlargement:-

<u>Grade of thyroid Size</u>	<u>Description of the gland</u>
0a . . . . .	Normal thyroid gland
0b . . . . .	Gland distinctly enlarged on palpation but not usually visible when head extended.
1 . . . . .	Gland enlarged on palpation, usually visible when head thrown back and neck extended.

\*Perez, c. et al, Bulletin of World Health Organisation, 18, 217, 1953

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2 . . . . . Goitre visible with head in normal position, palpation not necessary to establish enlargement.

3 . . . . . Goitre grossly visible at a distance.

The survey was started in the month of December 1983 in the Secondary Schools of Bassa Local Government Area.

The following is the analysis of data of goitre survey carried out in the school children of Zagun and Binchi Secondary Schools covering students of Form 1 to 5 -

Table 1: Age and Sex-wise Prevalence of Endemic Goitre Among Secondary School Children

Age (Years)	Male			Female		
	Examined	With Goitre	%	Examined	With Goitre	%
Below 16	43	6	13.9	19	4	21.1
16 and Above	180	28	15.5	53	33	62.3
Total	223	34	15.2	72	37	51.4

Table 2: Sex-wise Prevalence of Different Grades of Endemic Goitre Among Secondary School Children

	Male		Female		Both		Sex
	No.	%	No.	%	No.	%	%
0a (Normal)	189	84.8	35	48.7	224	75.9	
0b (Enlarged on palpation)	32	14.3	29	40.2	61	20.6	
1 (Enlarged Visible)	2	0.9	8	11.1	10	3.5	
Total	223	100	72	100	295	100	

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Table 3: Prevalence of Endemic Goitre According to Source of Water Supply

Source of Water	Children Examined	With Goitre	%
Well	199	43	21.1
Stream	96	28	29.1

Observations - From the data analysis the following observations can be made, although the data is not sufficient to make any conclusions at this stage -

1. Of 295 Secondary School Children examined 61 (20.6%) had grade 0b and 10 (3.5%) had grade 1 enlargement of thyroid giving a total prevalence rate of 24.1%
2. Amongst male children the prevalence of endemic goitre was 15.2 as compared to 51.3% in female children a difference of more than 3 times. However grade one enlargement was more than 10 times higher in female children than male children.
3. Prevalence of endemic goitre was 17.0% in children below 16 years age as compared to 25.7% in children 16 years and above. While among male children difference in prevalence between two groups was marginal (13.9% against 15.5% respectively) the difference in female children was three times i.e. 21.1% in female children below 16 years as against 62.3% in female children 16 years and above.
4. Children consuming stream water has higher prevalence of endemic goitre as compared to children consuming well water for drinking (29.1% against 21.1% respectively).

Plan of Action during 1984:

1. The goitre survey amongst school children is expected to be completed by May/June 1984. Based on the survey results it would be possible to identify geographical areas which are endemic for goitre.
2. Survey for endemic goitre and other clinical conditions viz. deaf mutism, endemic cretinism etc. associated with endemic goitre will be carried out in the general population where goitre endemicity is established by school survey.

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The goitre survey among general population will be started in July 1984 and will continue during 1984/1985 depending on the size of geographical and population distribution of endemic goitre as determined by school survey data.

3. Along with goitre survey of general population studies on iodine and other trace elements content of water supply, food and soil is planned to be carried out in collaboration with JICA research group on trace elements.
4. The departments of Chemical Pathology and other disciplines are expected to join the goitre study in their respective fields once the population survey is started.

List of equipments:

This has already been submitted by the Departments of Community Health and Chemical Pathology to JICA for their action.

DEPARTMENT OF MEDICAL MICROBIOLOGY  
FACULTY OF MEDICAL SCIENCES  
UNIVERSITY OF JOS

FIRST ANNUAL PROGRESS REPORT OF THE JICA/UNIVERSITY OF JOS  
RESEARCH COLLABORATIVE PROJECT ON DIARRHOEAL DISEASES IN  
CHILDREN UNDER 5 - YEARS IN JOS PLATEAU STATE, NIGERIA:  
UNIVERSITY OF JOS/JICA/MEDICAL MICROBIOLOGY/COMMUNITY  
MEDICINE/PAEDIATRICS GROUP PROJECT 1983

1. List of Participants/Investigators

ACADEMIC RESEARCH STAFF: DIARRHOEAL DISEASES RESEARCH GROUP

R.A.O. Shonekan	Ag. Head: Dept. of Medical Microbiology	Principal Investigator Convener
Dr. W. Kozak	Senior Lecturer (Bacteriology)	
Dr. V. Kumar	Senior Lecturer (Virology/Imm.)	
Dr. I.C. Tiwari	Head of Dept. (Community Health)	Principal Investigator
Dr. U.P. Nwene	Lecturer I (Community Health)	
Prof. I. Swiatkowska	Paediatrics	Principal Investigator
Dr. Z. Lazowski	Ag. Head Paediatrics	
Mr. I. Takahashi	J.I.C.A. TEAM	Collaborative Research:
Mr. H. Saida	J.I.C.A.	Med. Microbiology Laboratory

1.1 TECHNICAL STAFF

Miss Agatha Ani	Lab. Technologist (Bacteriology)	Seconded full-time to JICA/UNI JOS Medical Microbiology collaborative
Mr. Joseph Ona	Lab. Attendant Med. Microb. Unijos	Research Laboratory.

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2. PREPARATORY STUDIES FOR STANDARDISATION  
AND COLLECTION OF BASIC STATISTICAL DATA

- 2.1 While awaiting the arrival of the bulk of JICA Equipments and Reagents consigned from JAPAN, preparatory studies were carried out from March - June 1983 on samples collected from all diarrhoeic patients at the Jos University Teaching Hospital. Attention was concentrated on narrowing discrepancies observed in the diagnostic results of same sample by reviewing and standardising the methods of sampling, treatment of materials, testing the activity and potency of Sera as well as Reference Strains of Cultures. As a result a working protocol was drawn up which has since become valuable for the surveillance carried out on JUTH Clinical samples from August - December 1983, and thereafter on the on-going Research Project.
- 2.2 Following on the final presentation to the University by H.E. the Japanese Ambassador in Nigeria of the J.I.C.A. Equipment in October 1983, the modern Research Laboratory was taken over completely and fitted with newly arrived equipment for the JICA/UNIJOS Medical Microbiology Research Team to commence work promptly on Infantile Diarrhoeal Disease project. The protocol drawn up by the Committee after several consultative meetings formed the basis for the investigations.
- 2.3 The first phase of Year I of the Research Project commenced in October 1983 with planned investigation on ten samples collected daily from the children with Clinical Diarrhoea from the Paediatric Ward of J.U.T.H. After the first month the multi-disciplinary investigation - involving the Departments of Medical Microbiology, Paediatrics, Community Health and J.I.C.A. Team - suffered initial set back with drastic reduction in patients' population which led to small numbers and irregular flow of samples. Plans have now been concluded to augment samples from the Paediatrics Ward of the Plateau Hospital from February 1st, 1984, to be supplemented later from the S.I.N. Jankwano Hospital at Jos and the VOM S.U.M. Hospital.

3. OTHER ACTIVITIES: Results

- 3.1 Recruitment of two personnel are on hand to meet our crucial needs, namely:
- i) A "Collector" from the Dept. of Community Medicine who would also ensure adequate completion of data forms.
  - ii) A Lab. Technologist qualified in Parasitology to handle efficaciously, the investigations on Diarrhoea Stools and Blood for concomittant Malaria infections.

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- 3.2 The results of the single stool specimens collected from each of the children have been collated and are attached as Appendices as follows:-

- Appendix I      Bacteriological data from Infantile Diarrhoeal Cases from October - December 1983.
- Appendix II    Parasitological investigations from Diarrhoeal Cases of Children 5 - years and under treated at JUTH from August - December 1983.
- Appendix III   Epidemiological Data on 88 cases of Diarrhoea in children 5 - years and under examined between October - December 1983.

- 3.3 The data gathered from this pilot study - characterisation of causative agents & their prevalence etc - will serve as a basis by which the results of projected activities may be subsequently judged - especially of any changes; in meeting the useful goal of documenting the prevalence of faecally transmitted microbiological agents and parasites; in assessing activities designed to improve sanitation and intensify health education; to provide interim as well as long - term drug prophylaxis directed against the infective agents.

4. PROGRAMME FOR 1984

- 4.1 For the remaining period of the year I Research project, i.e. January to September 1984 - sampling would be increased to about 20 daily and extended to VOM S.U.M. Hospital. Closer relations will be maintained between the three departmental groups Microbiology/Paediatrics/Epidemiology so that results could be integrated. The investigation into the Malarial aspect of Infantile Diarrhoeal Diseases will be introduced and the typing of the actological agents would be more detailed.
- 4.2 Year II Programme will take on the Viral Aspects investigation into the Diarrhoeal Diseases of children. The draft protocol (Appendix IV) has been prepared and a list of the equipment and reagents required have been forwarded to the Team Leader of JICA, Dr. Takahashi. A Lab. Technologist (Virology) has been interviewed and processing of his appointment being pursued. Subject to arrival of the equipment and reagents, the inclusion of Viral investigations should commence by May 1984.
- 4.3 In September 1983 - the combined results of the year I project will be studied and a joint review of the progress made so that account will be taken of experience gained to introduce appropriate modifications to the plans for Year II programme.

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- 4.4 Details of the pilot project as planned for the under - five children on hospital - based communities in two contrasting areas - Highland; Urban/Rural, and Lowland: Rural/Urban - will be worked out after item No. 3 above has been implemented.

R.A.O. Shonekan,  
Principal Investigator,  
Medical Microbiology.

## APPENDIX I

### JICA PROJECT: DIARRHOEA IN CHILDREN PRELIMINARY REPORT - BACTERIOLOGICAL DATA-BACTERIOLOGY TEAM - DR. W. KOZAK

During the period October - December 1983, 88 single stool samples of children below 5 years of age suffering from diarrhoea, were collected and bacteriologically examined. From these samples, 75/85% were obtained from children's Health Clinic, and OPD, JUTH, and the remaining 13/15% of samples were provided by Paediatric Ward of the same hospital.

Collection of specimens and laboratory diagnosis were carried out as described in an earlier submitted protocol.

From specimens examined, just about 11% yielded *Escherichia*, *Salmonella* and *Shigella* strains, which are well documented as causative agents of diarrhoea. In 4 cases, various bacterial strains were isolated, which in some conditions can be responsible for that disease.

The quality of the number of specimens examined cannot permit a definite conclusion yet on the prevalence of bacterial diarrhoea nor the frequency of a particular causative agents of this disease among children. Curtailment of specimens arose at JUTH from recent introduction of fees paid by patients. However, the working team secured co-operation in this respect for additional samples from paediatric clinics. These should result in increase in number of available specimens for investigation.

## APPENDIX II

### JICA PROJECT: DIARRHOEA IN CHILDREN UNDER 5 YEARS PARASITOLOGICAL DATA/PRELIMINARY REPORT OCTOBER - DECEMBER 1983 - R.A.O. SHONEKAN

A total of 49 samples ( 55% ) only were available from October - December 1983 for Parasitological study as compared with 88 specimens which were employed for Bacteriological and Clinico - Epidemiological study reported above. The disparity in numbers resulted from failure to obtain stools in 39 children.

Of the 49 diarrhoeic samples examined the analysis of the samples age grouping showed that 57% were 1 - 6 months old, 30% were among 7 - 11 months old, while 13.0% were those of 1 - 3 years. Result of the examination showed only two positive cases of the protozoan Entamoeba coli representing 4% prevalence.

Comments: Although the sample number is too small for analysis, the meagre recovery of intestinal parasites from the 49 samples submitted raised questions of possible faulty presentation of stool, or inadequacy of the stipulated routine techniques viz. recovery of ova and cysts by concentration methods. This has become manifestly obvious when compared with the results hereunder of the base-line data whose results for the corresponding period showed among the under-five year a 14% overall prevalence of intestinal parasitism.

Harboured in descending order of prevalence were:-  
Ascaris lumbricoides, Hookworm, T. enia spp, H. nana,  
Trichuris trichuria and Strongyloides stercoralis  
among the Helminths; while among the Protozoans, they  
were in similar order T. hominis, Giardia lamblia and E. coli.

## APPENDIX IIA

### PARASITOLOGICAL FINDINGS - CHILDREN UNDER FIVE TREATED AT OPD & CHILD HEALTH CLINIC JUTH AUGUST - DECEMBER 1983:

#### BASE - LINE DATA STUDY - R.A.O. SHONEKAN

This study was conducted from August - December 1983 to obtain base - line data on parasitological aetiological agents. The total number of single stool specimens examined was 327 from children under five - with diarrhoea and/or abdominal disturbances - who had attended the O.P.D. and Children's Clinic of the Jos University Teaching Hospital. The overall prevalence rate was 26% (85) while the distribution according to sex of the positive samples was 56% for Boys and 44% for Girls.

The sequence of helminthic prevalence was Ascaris lumbricoides, Hookworms, Hymenolepis nana and S. mansoni, Trichuris trichuria, Taenia spp and Strongyloides stercoralis.

The four protozoan species detected showed the prevalence rate in sequence with Trichomonas hominis, Entamoeba histolytica, Giardia intestinalis and Entamoeba coli.

The picture of the earliest detection of infections with intestinal parasitism recorded were:-

- 1 - 2 month old: Trichomonas hominis and Giardia lamblia
- 3 - 5 month old: Ascaris lumbricoides.
- 6 - 11 month old: Hookworm.
- 1 - 3 year old: H. nana & S. mansoni + above

Evidence of multiple infections was noted on eight of the children.

The above are furnished merely as progress report but the base - line data will form eventually useful controls for the rest of year I pilot study of the project when completed.

### APPENDIX III

#### JICA-UNIJOS RESEARCH PROJECT ON DIARRHOEAS IN CHILDREN UNDER FIVE

##### CLINICO-EPIDEMIOLOGICAL DATA

The study on diarrhoeal diseases in Children under five was started in October, 1983. In all 104 cases of diarrhoeas attending the O.P.D., Child Health Clinic and Paediatric ward were studied. Of these, 2 cases were rejected as they did not fall in the study group.

The following tables show the clinical and socio-economic, environmental characteristics of the cases studied. It is not epidemiological data as neither the population nor the data from controls (for case-control study) are available.

Table 1: Age and Sex-wise Distribution of Children  
Suffering from Diarrhoea

Age	Male (n=62)		Female (n = 40)		Both Sex (n=102)	
	No.	%	No.	%	No.	%
Below 7 months	36	58.06	26	65.00	62	60.78
7 - 12 months	18	29.03	9	22.50	27	26.47
2 - 3 years	8	12.90	5	12.50	13	12.75

Table 2 Environmental Sanitation in the Families of  
Children Suffering from Diarrhoea

Water Supply (n = 102)	Tap		Well		Not Recorded	
	No.	%	No.	%	No.	%
	8	80.39	9	8.82	11	10.78
Toilet Facility (n = 102)	Available		Not available		Not recorded	
	No.	%	No.	%	No.	%
	76	75.51	15	14.71	11	10.78

Table 3      Literacy Status of Mother

Literacy	No. (n = 102)	%
Illiterate	50	49.0
Primary School	26	25.5
Secondary & above	16	15.7
Not recorded	10	9.80
Total	102	100

Table 4      Method of Feeding of Children under Examination

Feeding Method	No. (n = 102)	%
Breast alone	37	36.27
Breast & Bottle	51 *	51.00
Solids	3	2.94
Not recorded	12	11.76
	103	

\*One child was also on solids, so he has been double counted raising the total to 103

Table 5      History of other cases of Diarrhoea in the family

Other Cases of Diarrhoeas	No. (n= 102)	%
Yes	14	13.73
No	60	59.82
Not recorded	28	27.45
Total	102	

..../3

Table 6      Consistency of Stool in Diarrhoeal Cases

Consistency	No. (n = 102)	%
Soft	13	12.75
Watery	85	83.33
Not recorded	4	3.92

\*59 Cases had mucus while 8 cases had blood in the stool.

Table 7      Associated Symptoms in the Diarrhoeal Cases

Symptoms	No. (n = 102)	%
Vomiting	44	43.14
Abdominal Pain	28	27.45
Fever	40	39.22
No other symptom	25	24.51

Some children had more than one symptoms.

Table 8      Treatment Received by the Child before coming to The Hospital

Treatment	No. (n = 102)	%
No Treatment	9	8.82
Home/Native	16	15.69
Modern Medicine	37	36.27
Not recorded	40	39.22
102		

..../4

Observations:-

Based on the above data (Table 1 - 8) the following observations are made:

- 1) Sixty percent of the children examined for diarrhoea were below 7 months age. This was not due to higher incidence of diarrhoea in this group but because majority of children examined were from child Health Clinic where mostly young children are brought by the mothers.
- 2) Eighty percent of the families had tap water supply and 75.5% had toilet facilities. Most of the children examined were from Jos metropolis where water supply and toilet facilities are available to the households.
- 3) Almost half (49%) of the mothers were illiterate while another one quarter were educated up to primary school only. Low literacy among mothers is known to be associated with poor child rearing practices leading to higher morbidities among children.
- 4) More than fifty percent women were practicing bottle and breast feeding both. But when this data is contrasted with the fact that about 60% of the children examined were 6 months and below, it is clear that a large proportion of children are put on bottle feeding quite early. This may have implications in terms of diarrhoeal diseases.
5. In about 14% of cases history of another case of diarrhoea in the family was present. It would therefore worthwhile to screen other members of the family for carrier status, sub-clinical and mild cases. The source of infection may also be possible to detect on screening of human cases, animals and environment of the case thus compiling the epidemiological picture of diarrhoea among children.
6. In majority (83.3%) cases children had watery stools. A large number of cases (about 60%) had mucus in their stool.
7. While one quarter of the cases did not have any other symptom, vomiting, abdominal pain and fever were commonly seen to be associated with diarrhoea.
8. More than one third of the cases were already on modern anti-diarrhoeal drugs while another 16% had home/native treatment. Because of introduction of drug, the bacteriological isolation of cases may be adversely influenced.

Dr. Tiwari )  
Dr. Hwene ) Dept. of Community Health

UNIVERSITY OF JOS/JICA RESEARCH PROJECT

FIRST ANNUAL PROGRESS REPORT OF THE RESEARCH COLLABORATIVE  
PROJECT TEAM ON MEDICAL ENTOMOLOGY AND PARASITOLOGY

re: JAPANESE INTERNATIONAL COLLABORATIVE AGENCY (J.I.C.A.)  
PROJECTS 1983

The following is the position so far with respect to the JICA - UNIJOS collaborative projects on Medical Entomology and Parasitology, for the period 1982/83.

A total of seven (7) Academic/Research staff of the University and J.I.C.A. Project were involved in the Team, plus some two (2) Postgraduate Research students.

Academic/Research Staff

Prof. M.O.E. Iwuala	-	Department of Zoology
Dr. D.M. Roberts	-	" " "
Dr. R. Irving Bell	-	" " "
Dr. C.O.E. Onwuliri	-	" " "
Mr. R.O.A. Shonekan	-	Dept. of Medical Microbiology
Dr. I.C. Tiwari	-	" " Community Medicine
Dr. H. Takahashi	-	J.I.C.A. Research Project

Postgraduate Students:

Mr. M. Maduabum	-	Department of Zoology
Mr. D. Beakye	-	" " "

Six sub-projects were clearly mapped out as detailed in the original proposal. The position so far regarding each sub-project is as follows:

PROJECT 1: "STUDY OF DISTRIBUTION OF BLACK-FLIES (SIMULIUM SPECIES)  
ON THE JOS PLATEAU" - PROF. M.O.E. IWUALA, MR. M. MADUABUM  
AND R.H. TAKAHASHI

Following some preliminary work, this project was commenced actively by April 1983.

For some eight months (April - November) regular bi-monthly samples have been taken from various river systems in the Jos Plateau area, to establish presence or absence of immature stages of Silulium species.

Altogether 13 rivers have been consistently sampled, and of these black flies have been found to breed in three rivers viz: River Assob, River Farin Ruwa, and Cambo River located respectively in Riyom, Bokkos and Bassa Local Government Areas of Plateau State.

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Simulium breeding in the rivers was more noticeable in the mid-rainy season months of June to August, and so far six species of blackflies have been identified from the samples viz: S. aloocki; S. aureosimile; and S. cervicornotum C. damnosum s.l; S. hargreavesi; and S. mcrahani.

It is intended to continue the sampling exercise over some 18 to 24 months' period and subsequently to investigate in due course the factors conditioning the distribution and breeding of the various Simulium species.

PROJECT 2: "EFFECT OF WATER VELOCITY ON BLACK-FLY  
RELATIVE ABUNDANCE": DR. D.M. ROBERTS

This project is still awaiting the arrival of much needed equipment, especially the water velocity meter. A suitable site has been chosen in the River Assob for the study, but preliminary sampling cannot be made without the velocity meter.

PROJECT 3: "EFFECT OF TEMPERATURE AND HUMIDITY ON PUPAL SURVIVAL":  
DR. D.M. ROBERTS & MR. D. BOAKYE

This project was started in January 1983, but progress of work is being delayed by lack of cooled incubators in the Zoology Department (2 are still being awaited from JICA). Consequently, so far only the four low temperatures earlier listed in the Research Proposal have been tested at a saturated humidity. The following results were obtained:-

Laboratory studies were conducted to compare pupal survival, adult emergence and adult longevity after pupae had been exposed to four temperatures:- 0, 4, and 12°C; for different time periods:- 1 - 7 days. The greatest mean survival were obtained at 8 and 12°C and the least at 0 and 4°C for old pupae (pupae with dark coloured cuticle) and for young pupae (pupae with pale coloured cuticle) respectively. At 0 and 8°C, there was no significant difference between the mean pupal survival for both pupal ages. Young pupae survived significantly longer than old pupae when exposed to 12°C, while old pupae survived better at 4°C. Generally, 8°C had the best survival rate, and 0°C the poorest. Adult emergence occurred mainly during the day and extended over 4 days. Mean adult longevity varied among adults from pupae exposed to different temperatures and time periods, but were generally non-significantly different. In the field, pupation times were similar to that in the laboratory, which was mainly diurnal with a major peak in the morning (6 - 9.00 h) and a minor peak in the evening.

PROJECTS 4 & 5: "DISTRIBUTION OF IMMATURE MOSQUITOES ASSOCIATED WITH  
THE RIVER SYSTEMS OF THE PLATEAU" AND "SEASONAL AND  
VERTICAL DISTRIBUTION OF TREE-HOLE MOSQUITO BREEDING"  
- DR. R.J. IRVING-BELL

Shortened taxonomic keys have now been composed for the identification to species of the adults and larvae of the major genera present in the northern guinea savannah of Nigeria.

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Collections of mosquito larvae and preliminary observations have been made in the following areas with a view to choosing specific study sites:-

Ngell River near Miango: Has a limited range of habitats owing to lack of trees.

Assob River: This is a possible study site although tree holes are present together with the usual rock pools and mud pools.

Kurra Falls complex: The area is atypical owing to the reservoir system and Eucalyptus plantings.

Kagoro Forest: Provides a wide range of habitate including many tree-holes; however the river with its rain forest border is not typical of the savannah region.

Other rivers need to be visited before a selection of the two sites required in the proposal can be made.

In addition the following requirements must be satisfied before the projects can be run fully effectively:

- a) Funds for locally-bought materials and transport.
- b) The arrival of specialist items requested from JICA, in particular, larval preservative fluid (lactophenol).
- c) Student researchers willing to undertake the projects as part of their higher degree programmes.

PROJECT 6: THE PREVALENCE AND DISTRIBUTION OF ONCHOCERCIASIS -  
C.O.E. ONWULIRI, I.C. TIWARI, AND R.O.A. SHONEKAN

Strategies for the above investigation have been mapped out although actual field surveys of the endemic areas have not been carried out. However, it is hoped that the greater part of the work in Phase 1 (as specified in the original proposal) would be finished before the end of December 1984. The delay in the take-off of the actual field survey was due mainly to the absence of a Corneosceral punch needed for taking skin-snips.

PROF. M.O.E. IWUALA  
UNIJOS/JICA MEDICAL  
ENTOMOLOGY RESEARCH GROUP





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SECOND ANNUAL REPORT - DECEMBER 1984

University of Jos/Japanese International Cooperation Agency  
Research Project.

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- 1) Following the signed Agreement of the University of Jos/JICA Research Projects, the members of the Japanese Team arrived Jos in October 1982 and research began on the various sub-groups of the research projects.

The first Annual Report of these Research Projects was compiled by Professor A.C. Ikeme in December 1983. This volume is the Second Annual Report incorporating the preliminary reports of December 1983 and the mid-term results from January 1984 to December 1984.

Sub-Group (1): Project: Trace Element Analysis

- (a) Dr. G.A. Ubom and Mr. C. Noda - Preliminary report on the trace elements detection in natural sources in Plateau State, Nigeria.
- (b) Prof. C. Ejike, Dr. D. Anadu and Dr. J. Chidobem - Distribution of trace Elements in some surface waters.
- (c) Mrs. M. Adiuku-Brown and Dr. A. Ogezi - Comparative sample de-composition techniques for the analysis of simple sulphide minerals.

- (d) Dr. A. E. Ogezi, Mrs. M. E. Adiuku-Brown and Dr. M. Alam-  
Environmental and economic studies of trace elements  
associated with mining and mineral processing in  
parts of Plateau State, Nigeria. a preliminary  
report.

Sub-Group (2): Project - Goitre

- (a) Prof. I. Tiwari and Mr. M. Okoronkwo - JICA/UniJos  
project on endemic goitre.

Sub-Group (3): Project - Diarrhoea

- (a) Dr. W. Kozak and Mr. R. Shonekan - Bacteriological  
investigation of diarrhoea in children.  
(b) Miss. A. Ani and Mr. M. Takahashi - Aetiological  
studies of infantile diarrhoea disease seen in  
Jos University Teaching Hospital. A preliminary  
report.  
(c) Dr. U. P. Nwene - Data on clinical presentation and  
socio-demographic profile of diarrhoeal cases.

Sub-Group (4): Project - Medical Entomology

- (a) Prof. M. Iwuala, Mr. Maduabun and Dr. H. Takahashi.-  
Study of distribution of black flies (*Simulium*  
species) on the Jos Plateau  
(b) Dr. D. Roberts and an MSC student - Effect of  
water velocity on the relative abundance of  
immature black flies  
(c) Dr. D. Roberts, Mr. D. Boakye and Mr. J. Ogidi -  
Effect of temperature and humidity on black fly  
pupal survival  
(d) Dr. R. Irving-Bell and Mr. I. Sessay -  
Distribution of immature mosquitoes associated  
with two riverine sites.

- (e) Dr. R.J. Irbing-Bell and Mr. I. Sesay -  
Mosquitoe breeding habitats: Distribution and relative abundance of species at a Kagoro forest stream in Northern Nigeria.
- (f) Dr. C. onwuliri, Dr. I. Lawal and Mr. B. Nwoke -  
Prevalence of distribution of onchocerciasis on the Jos Plateau.
- (g) Prof. M. Iwuala, M. M. Maduabum and Dr. H. Takahashi -  
Study of the seasonal abundance and population characters of immature stages of black flies in the Jos Plateau area
- (h) Dr. D. Roberts and Mr. J. Davis-Cole - Effect of predators and type of substrate on the relative abundance of immature black flies.
- (i) Dr. D. Roberts and Dr. Irving-Bell - Dispersal of Adult black flies
- (j) Dr. R. Irving-Bell - Niche distribution in relation to physiological state of adult populations of black flies and mosquitoes.

The Faculty objective of identifying the health needs of the population and providing solutions to these is continually kept in mind and in these research. Expansion of the research to include other Faculties, we feel will provide a broader base for data acquisition for the solution of these problems.

24, December 1984.

Revd Prof. E. O. Oji  
Dean, Faculty of Medical Sciences

PRELIMINARY REPORT ON THE TRACE ELEMENTS  
DETECTION IN NATURAL SOURCES IN PLATEAU STATE,  
NIGERIA.

Gregory Abraham UBOM and Chiyoichi NODA  
Department of Biochemistry, Faculty of Medical Sciences,  
University of Jos

This report is a summary of work and data collected by the Biochemistry Department, Faculty of Medical Sciences, as part of the participation in the JICA project up to the period ending December 1984. The cationic concentration data were obtained using the Atomic Absorption Spectrometer(AAS) which was successfully installed and tested in the Department by November, 1983. The Anionic levels-studies were performed with the Ion Chromatographic Analyzer(ICA) which was installed in the Department around June, 1984. The remaining data - total organic carbon (TOC), soluble silicates ( $\text{SiO}_2$ ), ammonia ( $\text{NH}_3$ ) etc, were determined using the facilities at the Tokyo Metropolitan Research Laboratories of Public Health by Dr. Ubom during his JICA sponsored leave in Japan between October and December, 1984.

Table 1 and 2 are comparative cationic and anionic concentration data on water samples from goitrous and non-goitrous locations.

Table 3 shows the trace and heavy metals composition in water samples collected from chemically treated(tap) and untreated sources (streams, wells, dumps and industrial effluents) and Table 4 is the organic and inorganic composition of water samples collected from different locations of Jos Metropolis.

The observation in all cases of analyzed waters is that water whether treated or untreated is soft (ie. low in  $\text{Mg}^{++}$  and  $\text{Ca}^{++}$ ). The quality of water analyzed is generally good and falls within the internationally specified limits with the exception of high iron concentration in some industrial runoffs. The organic components, oxygen (Biochemical and chemical) demands, disinfectant quality of these waters should be monitored in order to be able to make broader assessments of the water.

Table 1. Anions, trace and heavy metals concentrations in waters at the Goitrous villages.

Location	unit; $\mu\text{g/l}$ , rest $\text{mg/l}$																			Total hardness
	I <sup>-</sup>	F <sup>-</sup>	Cl <sup>-</sup>	NO <sub>3</sub> <sup>-</sup>	NO <sub>2</sub> <sup>-</sup>	Br <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	K	Ca	Mg	Na	Zn	Cu	Fe	Mn	Cd				
ASSAK (ST)	0.5	0.1	0.5	0.0	0.3	0.0	0.6	0.7	0.2	0.1	1.3	0.011	0.002	0.45	0.016	0.000	0.9			
BIMBOP 1 (W)	0.0	0.1	0.6	0.0	4.4	0.0	1.2	4.1	2.5	0.2	4.2	0.005	0.001	0.37	0.050	0.000	7.1			
BIMBOP 2 (W)	0.0	0.7	14.2	0.0	52.6	0.2	0.0	9.1	3.6	17.9	1.6	8.1	0.000	0.000	0.70	0.023	0.000	51.5		
BINCHI 1 (W)	0.0	0.3	0.9	0.0	1.9	0.2	0.0	1.2	2.1	4.3	2.7	8.3	0.012	0.006	0.54	0.062	0.002	22.2		
BINCHI 2 (W)	2.4	0.2	3.6	0.0	0.1	0.0	0.0	6.9	7.1	24.5	2.2	8.8	0.018	0.000	2.75	0.141	0.000	70.6		
DUTSENKURA (ST)	0.0	0.1	0.2	0.0	1.7	0.0	0.0	0.4	0.6	0.2	0.1	1.4	0.000	-	0.15	0.009	0.000	0.9		
IBANDI (ST)	0.0	0.2	0.7	0.0	0.3	0.0	0.4	1.3	1.0	0.9	2.9	-	0.000	1.60	-	0.000	6.4			
KISANCHI 1 (SP)	0.0	0.2	1.7	0.0	0.9	0.0	1.7	1.1	3.1	1.4	7.9	0.003	0.000	0.45	0.014	0.000	13.7			
KISANCHI 2 (W)	0.0	0.2	3.9	0.0	0.5	0.2	0.0	1.0	2.5	2.9	1.1	6.0	0.005	0.000	0.50	0.027	0.000	11.9		
KISHI-II 1 (W)	0.0	0.3	3.0	0.0	9.7	0.5	0.0	2.2	2.1	3.2	0.9	8.3	0.014	0.000	1.03	0.028	0.002	11.8		
KISHI-II 2 (W)	0.0	0.2	6.7	0.0	3.9	0.5	0.0	0.6	2.3	3.1	0.5	8.2	0.010	0.000	0.57	0.018	0.000	9.9		
Mean	0.3	0.2	3.3	0.0	6.9	0.2	0.0	2.3	2.5	5.7	1.1	6.0	0.008	0.001	0.83	0.039	0.000	18.3		

Note: Figures are average of 2-5 tests.

ST-stream

W-well

SP-spring

Table 2. Anions, trace and heavy metals concentrations in waters in Jos area

Location	unit: $\mu\text{g/l}$ , rest mg/l																	
	I <sup>-</sup>	F <sup>-</sup>	Cl <sup>-</sup>	NO <sub>2</sub> <sup>-</sup>	NO <sub>3</sub> <sup>-</sup>	PO <sub>4</sub> <sup>3-</sup>	Br <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	K	Ca	Mg	Na	Zn	Cu	Fe	Mn	Cd	Total hardness
DOGONDUTSE (W)	0.0	0.2	1.5	0.0	4.5	0.0	0.0	1.1	1.5	0.1	0.1	4.0	0.008	0.000	0.05	0.013	0.003	0.7
JARAWA (W)	2.7	0.1	4.2	0.0	14.1	0.0	0.0	1.0	2.9	1.8	0.3	6.9	0.020	0.000	0.19	0.036	0.000	5.8
JOS RIVER (R)	0.0	0.1	1.1	0.0	1.5	0.0	0.0	0.7	1.5	0.2	0.1	2.3	0.005	0.000	0.19	0.004	0.003	0.9
LIBERTY DAM (D)	1.5	0.2	0.8	0.0	0.1	0.0	0.0	0.6	1.3	0.8	0.3	1.1	0.000	0.000	0.40	0.011	0.000	3.3
NARAGUTA (ST)	3.9	0.1	1.7	0.0	0.4	0.0	0.0	0.6	1.4	1.0	0.5	3.8	0.007	0.005	0.61	0.065	0.002	4.6
TIN MINING 1 (P)	7.0	0.2	0.6	0.0	0.1	0.0	0.0	0.5	0.6	0.8	0.2	0.9	0.005	0.003	0.16	0.016	0.000	2.9
TIN MINING 2 (P)	1.3	0.2	0.4	0.0	0.0	0.0	0.0	0.3	0.7	0.9	0.3	0.9	0.010	0.014	0.26	0.039	0.000	3.5
Mean	2.3	0.2	1.5	0.0	3.0	0.0	0.0	0.7	1.4	0.8	0.3	2.8	0.008	0.003	0.27	0.026	0.001	3.1

Note: Figures are average of 2-5 tests.

W-well

R-river

D-Dam

ST-stream

P-pond

Table 3.1, Comparative trace and heavy metal contents in treated and untreated waters in Jos and its suburbs.

LOCATION	Sn (PPB)	Mg (PPM)	Ca (PPM)	Pb (PPB)	Mn (PPB)	Cu (PPB)	Na <sup>M</sup> (PPB)	K (PPM)	Zn (PPM)	Fe (PPB)	Cd (PPB)
1. Gangare Well (W)	206.	1.0	3.0	5.4	8.6	15.9	11.0	5.5	0.0	162.7	0.2
2. Gangare Jos Well (W)	138.6	2.0	14.0	7.0	3.2	7.0	16.0	7.0	0.0	255.8	0.4
3. Aze Gangare Well (W)	160.9	2.0	8.0	5.3	14.8	21.4	16.8	14.4	0.0	262.1	0.3
4. Zaria Road Tap (T)	195.9	0.5	7.0	4.7	3.0	3.1	2.2	1.8	0.1	104.7	0.5
5. Warden Barrack(Blk) Well (W)	147.6	0.0	1.3	3.4	5708.4	26.0	1.2	1.3	0.0	17,769.6	0.4
6. Warden Barrack(Blk) Tap (T)	141.3	28.0	228.0	5.5	6.4	19.6	42.0	9.8	0.7	332.5	2.6
7. St. Louis College Jos- Tap.	129.2	0.3	1.0	7.8	6.7	6.8	1.3	1.3	5.6	7,152.0	0.5
8. Gada Biu Tap (T)	132.1	0.5	2.0	5.3	10.6	8.9	1.1	1.2	0.3	335.3	0.5
9. St. Louis College Jos- Tap	140.6	0.2	0.8	7.8	780.0	41.5	10.5	1.5	0.3	42.0	0.5
10. Gada Biu Well (W)	120.1	2.0	12.0	8.6	16.6	31.5	30.0	6.3	0.0	0.0	0.3
11. Mr. Ali Stream (S)	130.5	1.5	2.5	7.1	41.0	43.3	9.0	1.0	0.0	3892.0	0.3
12. Mr. Ali Well (W)	139.7	0.3	3.0	7.1	15.4	35.1	2.8	0.8	0.0	316.0	0.4
W - Well S - STREAM											

Table 3-2

LOCATION	Sn (PPB)	Mg (PPM)	Ca (PPM)	Pb (PPB)	Mn (PPB)	Cu (PPB)	Na (PPM)	K (PPM)	Zn (PPM)	Fe (PPB)	Cd (PPB)
13. Katakoto Well Water (W)	215.8	1.5	1.6	5.9	16.8	238.2	24.5	3.0	0.0	638.0	0.8
14. Katakoto Street Tap (T)	173.8	1.5	14.0	11.1	7.8	18.7	1.2	1.3	0.0	201.0	0.5
15. JZ 54 H/L Tap (T)	125.9	0.5	2.0	5.9	5.5	23.3	1.0	1.1	0.7	94.0	0.6
16. JZ 28 Apeta Well (W)	136.8	1.0	3.5	7.6	148.0	42.6	23.0	7.2	0.7	169.9	0.5
17. Lake Chad	160.2	5.5	7.5	10.1	780.0	19.6	6.5	3.0	0.0	7,192.0	0.5
18. Pan Well (W)	179.4	5.0	6.5	6.2	54.0	70.4	12.0	2.5	0.0	702.0	0.7
19. Pan Dam (D)	140.9	5.0	6.0	5.2	1248.0	0.0	9.5	2.7	0.5	3684.4	0.6
20. Pan Un (D)	158.1	1.5	7.5	7.2	36.4	10.4	10.0	3.2	0.0	345.4	0.6
21. Treated Brewery (I)	198.0	1.5	6.0	6.6	21.3	36.0	2.1	10.8	0.0	1028.0	0.7
22. Brewery Exhaust (I)	3613.0	0.2	0.8	8.2	218.0	6.8	17.5	1.8	0.0	5420.0	0.7
23. Human Activity (I)	202.1	0.2	3.5	34.5	2.2	0.6	14.4	1.8	0.2	96.0	0.7
24. Bukuru Sn Mining (I)	277.6	1.0	3.5	9.2	2.4	1.2	1.8	2.0	0.0	61.0	0.5

Table 3.5

LOCATION											
	Sn (PPB)	Wg (PPM)	Cd (PPM)	Pb (PPB)	Mn (PPB)	Cu (PPB)	Na (PPM)	K (PPM)	Zn (PPM)	Fe (PPB)	Cd (PPB)
25. Inside Water (Waste)	220.8	1.5	2.0	8.4	8.8	8.9	30.0	2.0	0.0	239.0	0.7
26. Treated Water	201.1	1.0	2.0	10.8	32.4	15.0	2.4	1.3	0.0	2040.0	0.0
27. Ind. Exhaust (I)	220.1	1.0	2.0	7.5	10.4	63.2	2.5	1.8	0.6	203.0	0.8
28. PAN SAME	371.3	1.0	1.5	9.6	34.1	104.4	2.6	1.5	0.3	374.0	0.8
29. Dytex Exhaust (I)	144.0	1.5	10.8	349	212.0	179.9	3.9	10.5	0.2	607.0	0.7
30. Coca Cola Exhaust (I)	186.5	3.0	102.0	475	84.0	53.3	3.8	42.9	0.3	16280.0	0.8
31. Combined Brewery - Treated + Exhaust (I)	306.9	0.2	32.0	35.8	31.0	83.9	3.5	2.5	0.2	3200.0	1.3
32. Bukure #2 Mining (I)	220.1	6.5	32.0	576	1488.0	173.0	18.5	7.0	0.8	1160.0	0.9
33. Coca Cola Mxt. Exa. (I)	292.4	3.9	1.0	218	760.0	54.0	6.0	22.0	0.2	18040.0	0.7
34. Coca Cola Exhaust (I)	253.7	1.0	1.9	177	12.2	30.0	3.5	1.5	0.1	12744.0	0.7

Table 4. Organic and inorganic composition of water samples collected from Jos Metropolis.

LOCATION	PH	Na	K	AS	CD	Pb	Mn	Mg	Cu	Zn	Sn	Fe	SiO <sub>2</sub>	TDS	F	CL	NO <sub>3</sub>	SO <sub>4</sub>	CaCO <sub>3</sub>		
	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM	PPM		
1. NARAGUTA STREAM.	6.8	5.6	1.7	ND	ND	1.5	0.7	0.7	2.9	0.01	0.025	ND	0.19	0.24	0.14	1.9	0.4	1.4	0.6		
2. TUDUN WADA WELL (STATE SECRETARIAT)	6.0	5.9	2.9	ND	0.2	ND	0.9	0.36	6.2	0.02	0.04	ND	0.04	0.09	0.23	1.2	1.2	ND	15.4		
3. TIN MINING NGI ROAD	6.5	2.4	0.28	ND	ND	ND	1.4	0.2	1.1	0.007	0.01	ND	0.03	0.102	0.23	0.6	0.2	0.2	ND	3.0	
4. " " NGI 2 "	6.6	2.2	0.28	ND	ND	ND	0.4	0.3	1.3	0.004	0.005	ND	0.03	0.153	0.27	0.4	1.3	0.3	ND	5.0	
5. JOS TAP WATER.	8.6	3.0	1.50	ND	0.2	ND	0.3	0.3	10.0	0.161	0.05	ND	0.06	8.3	0.49	0.2	0.9	1.5	22.3	28.4	
6. LIBERTY DAM.	6.6	2.4	1.32	ND	0.3	ND	0.9	0.4	1.6	0.007	0.005	ND	0.16	8.3	0.27	0.16	0.4	0.16	0.2	ND	7.6
7. LAMINGO DAM.	6.4	3.0	1.32	ND	ND	3	ND	0.3	1.7	ND	0.005	2	0.18	11.1	0.27	0.13	0.4	0.16	6.1	ND	5.0
8. RAIN WATER (FROM ROCK) JOS.	1.8	0.28	ND	0.2	ND	4.3	0.2	2.0	3.2	0.05	3.0	0.36	1.2	0.29							

**PROJECTIONS:** Eventhough the AAS and ICA are installed in the Biochemistry Department, there was an initial understanding that other Departments including Chemical Pathology, Zoology and Geology would send representatives to study and master the use of these equipments. So far, the goal of training representatives has not been attained because the various Departments have not availed themselves of this opportunity. It is envisaged that this training will be continued till the end of May 1985 when the Japanese expert (Mr. Noda), attached to the project, will be leaving for home. Training in the maintenance and repair procedures on these equipments are to be intensified, especially to the Biochemistry Department Nigerian counterpart who may take full responsibility for these equipments after the Japanese expert leaves.

Goitre studies will be extended to include analysis on foods and soils along with the water analysis which we have been surveying in the "goitre villages". This extended studies may allow us to make the correct associations between cations, anions and the incidence of goitre - which cationic or anionic deficiencies potentiate goitre.

The impact of human activity on drinking water will be studied using variables that include ammonia, nitrate, nitrite and phosphate detection. The effects of disinfection of drinking water and the correlation between the lack of disinfection with diarrhoea and dysentery will be investigated. Comparative studies will be continued between treated purified and untreated waters.

Studies on the heavy metal content of vegetables, crops harvested in Jos tin mining and industrial effluents will be continued. The concentration patterns for these metals starting from the soil, water, crops, vegetables and animals will be established. Our results so far show that for nearby all the elements studies, the elemental concentrations are as much as 100 fold higher in the crops than in the waste effluents. This work will be pursued further to establish the extent of elemental concentration in mammals.

We anticipate to start monitoring the organic constituents of health significance in water and foods. Our interest would involve the determinations of the concentration of benzene, chlorinated alkanes and alkenes, chlorophenols,

polynuclear aromatic hydrocarbons, triholomethanes and pesticides in our waters and foods. Most of these organic species are presursors to carcinogens or are carcinogens themselves and should be detected and controlled even when they occur in low levels in our environment. For this phase of the research we request that JICA supply us with 1) a gas chromatograph (GC) - the type that is sensitive enough to meet our needs. 2) A mini electrophoretic equipment to be used in monitoring any changes in protein pattern and concentration in mammalian sera as a result of the presence of those organic toxins in the human system. 3) Reagents for the determination of dissolved oxygen in water and industrial wastes. 4) Reagents for the determination of turbidity, colour, soluble silicates ( $\text{SiO}_2$ ) and ammonia.

PRELIMINARY REPORT ON THE DISTRIBUTION OF TRACE ELEMENTS  
IN SOME SURFACE WATERS STREAM SEDIMENTS AND TISSUE DIGESTS  
OF AQUATIC BIOTA IN WATERS OF JOS PLATEAU.

By

EJIKE, C., D. I. ANADU & J. I. CHIDOBEM

INTRODUCTION

abatement

Effective/of aquatic trace elements pollution depends not only on treatment and control of waste waters but largely on the efficient monitoring of the general aquatic environment. Monitoring inland freshwater ecosystems involve two basic approaches.

1. Direct measurement of the spatial and temporal concentration of the trace-element or of important biologically active substances such as oxygen which are depleted by pollution
11. The use of biological indicators which range from micro-organisms, macroinvertebrates and fish and physiological indices such as Biological Oxygen Demand (BOD) measurements to total community indicators such as changes in community structure.

The latter is widely employed in monitoring aquatic pollution. It includes such indices as the general species indicator index, community structure index, oxygen production (P) and utilisation (R) index and the consequent P/R ratios.

Defining the maximum permissible level of trace element in aquatic systems is of fundamental importance in monitoring trace-element distribution. High concentrations of heavy metals are probably responsible at times for reducing faunastic diversity in aquatic system. Often a mixture of trace-elements are present in varying levels in a freshwater body simultaneously and it is then extremely difficult to estimate the contribution of individual or component trace-elements.

Extensive methodologies have been developed for evaluating the effects of discrete trace elements toxicants on various aquatic organisms. Many of these have been amplified by concern over the environmental implication of bioconcentration of these trace elements. Hazards to aquatic biota as well as to human health caused by local contamination of the aquatic environment have been reported on by various workers.

METHODS

Analysis of trace-elements and pretreatment of sediments and Macroinvertebrates were carried out after the methods of Thomas and Chamberlin (1974). Digestion of water was done as recommended in APHA (1974). Trace elements levels in water, substrate and tissue digests were then monitored by the Atomic absorption spectrophotometry. The study design allowed for sampling of streams, mine lakes and reservoirs. For the first two, selection was made from mining and non-mining zones and as far as possible from upland and lowland waters (See first JICA report).

Special attention was paid to the following elements Si, P, Mn, Fe, Cu, Zn, As, Se, Cd and Pb. While animal tissue digests of aquatic insects, decapod crustaceans, gastropod molluscs and fishes were investigated.

Because rainy season values are complicated by changing dilution factors greater reliance is placed on dry season values which apart from being more stable are likely to impact more on the life of the aquatic forms because of their higher levels of concentration.

Below is a tabulated form of the animal groups collected at the different stations in the dry season of 1984. Most of the sampled rivers and lakes drain into the Southern rivers which finally are drained into the Atlantic. The Delimi and a few other rivers North of Jos town drain into Shari river which is finally drained into the Lake Chad.

**Table 1: Dominant animal groups during the dry season in some Jos Plateau Waters.**

Sampling date	Site description and Name	Rocktype	Drainage type	Common Aquatic Animals	Bottom type
29/1/84	Pandam Lake		Into R/Benue through Dep river	Freshwater cray fish (large number) (Requires full identification) Ephemeroptera larvae Corixidae, the dragon fly larvae Anisoptera, Coleoptera, Hemiptera Diff. Fish species including <u>Alestes</u> , <u>Labeo</u> , <u>Citharinus</u> etc.	
29/1/84	Shendam Impoundment	Sandstones and shales		Cray Fish (requires full identification) Hemichromis, Water snails of the gastropod type	
29/1/84	Shendam bypass Stream with weirs & bridge at Jing		Into River Benue	Odonata (diff. types) Water beetles Coleoptera Blompharia	Sand stones
12/2/84	Kuru BaBa River		Possibly R/Benue	<u>Tillapia zilli</u> <u>Garra waterloti</u> <u>Water Scorpion</u> Dragon Fly, Damselfly larvae, Coleoptera, Corixidae	Gravels
12/2/84 8	Asob at Gangere Kibo		River Benue	<u>Garra waterloti</u> <u>Aphyosemlon</u> sp Corixidae Dragon Fly larvae, Hemiptera Ephemeroptera nymphs.	Boulder & gravels

Table 1 Cont'd.

Sampling date	Site description and Name	Rock type	Drainage type	Common Aquatic Animals	Bottom type
12/2/84	Riom river	Weathered basalt	R. Benue	<u>Garra waterloti</u> <u>Aphyosemion sp</u> Hemiptera (stick-insects) Dragon Fly larvae Coleoptera, Crab.	Sandy
12/2/84	Asob on Jos-Akwanga Rd		R. Benue	Cray Fish (Large numbers) Dragon Fly larvae <u>Garra waterloti</u> Ephemeroptera Coleoptera Damselfly larvae, Trichoptera	Boulder
12/2/84	Barmo Cement Kaduna/Plateau		R. Benue	<u>Garra sp</u> Enough fish for population studies & electro-fishing collection Hemiptera, Ephemeroptera Dragon fly larvae	Sandy
8/3/84	Miango Bukuru Goro river	Granite & sand	R. Benue	Odonata, Damselfly larvae, Hemiptera, Water stridders & skaters <u>Garra sp</u>	Boulder & brown colour
8/3/84	Miango reservoir			Damselflies, Ephemeroptera, Fish-Lates <u>niloticus</u> .	
8/3/84	River Delimi at Tilindin Fulani	Migmatite	L. Chad	Chara <u>Garra waterloti</u> <u>Berillius sp.</u> Ephemeroptera Dragon Fly larvae Damselfly larvae Chironomid larvae Biomphalaria (snails).	Sand & Boulder
12/3/84	Stream supplying Lafia Fish pond on Pankshin Rd		Benue River	Crayfish <u>Astacus?</u> Lots of Catfish probably of Ephemeroptera Dragonfly larvae.	

### Results resume

#### 2. Trace elements levels

The results so far indicate that some of the surface waters especially those located in mining zones and therefore likely to receive some form of effluents from mine wastes, contain more than the recommended levels of trace elements. The following elements have been identified as presenting possible hazards.

**Table 2. Observed range in Jos Plateau Waters (1) PPb;  
(2) ppm**

1	Cd	(1) 0.2 - 2.5
2	Zn	(1) 7.0 - 2,000.
3	Cu	(1) 3.0 - 700
4	Pb	(1) 1 - 45
5	Mn	(1) 7 - 20,000
6	Se	(1) T - 40
7	Fe	(2) 0.05 - 3.0

The data for sediments and tissue digests will be presented in a future report.

#### REFERENCES

- APHA (1974) Standard Methods for examination of Water and Waste Water American Public Health Association Washington, D.C.
- Thomas and Chamberlin (1974) Colorimetric Coemical Analytical methods. Tintometer L.T.D. England.

COMPARATIVE SAMPLE DECOMPOSITION TECHNIQUES  
FOR THE ANALYSIS OF SIMPLE SULPHIDE MINERALS

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Abstract

Although a number of analytical techniques exist for the analysis of the metal content of sulphide minerals, one of the most favoured methods is by atomic absorption spectrophotometry (AAS). This is due to its precision, accuracy and convenience. The various sample decomposition techniques described in the literature for the analysis of sulphide minerals using atomic absorption spectrophotometry include the use of single acids of various strengths, such as nitric acid; a mixture of two strong acids, such as aqua regia; and a combination of more than two acids, such as hydrofluoric/perchloric/nitric and hydrochloric acids.

In this investigation, seven standard and modified standard procedures were used in the analysis of ten sulphide minerals from different parts of Nigeria with a view to finding out the relative advantages of the methods in terms of time, convenience, cost, accuracy and precision as a basis for a larger geochemical and environmental programme involving sulphides and other minerals associated with mining and mineral processing. The effects of the digestion techniques on the liberation and subsequent determination of eleven elements - Au, Ag, Cd, Ni, Cr, Co, Mn, Cu, Fe, Zn and Pb - were investigated. The results show that most of the elements could be efficiently, cheaply and conveniently leached using simple acids, such as 50 percent nitric acid, which is one of the simplest of the leaching techniques.

Introduction

In view of the importance of geochemical analysis in geology, several geochemical techniques have been developed for the elemental analysis of geochemical samples (Stenton, 1976; Westwood & Cooper, 1973).

These include gravimetric and titrimetric (classical) methods of analysis, molecular fluorimetry, radiometric and radioactivation methods, mass spectrometry and spark-source spectrography, x-ray emission spectrometry, flame emission and atomic absorption spectrophotometry.

Since geochemical prospecting techniques are critically dependent on precise and reasonably accurate analytical methods that are rapid enough to permit thousands of analyses to be carried out in a short time, most analysts have resorted to the use of the atomic absorption spectrophotometry. This is due largely to its precision, accuracy, convenience and the relatively less operator skill required in the use of the instrument (Reeves and Brooks, 1978).

Most substances requiring geochemical analysis are complex and consist of several elements or compounds. Quite frequently, these materials fall short of the ideal in terms of solubility, volatility, stability or homogeneity, and all determinations in which the atomic absorption spectrophotometer (AAS) is used require that the element to be determined be in solution, several sample decomposition techniques prior to analysis have appeared in the literature (e.g. Stanton, 1976). The methods described usually differ depending on the nature of the sample and elements to be analysed.

Reeves and Brooks (1978) recommended the use of hot or cold hydrochloric acid ( $\text{HCl}$ ), for the decomposition of most carbonates; solutions of nitric acids ( $\text{HNO}_3$ ), for attacking ores of metals, like cobalt, copper, manganese, nickel and lead; and boiling concentrated  $\text{HNO}_3$  for leaching sulphides of elements, such as lead and zinc from a silicate matrix. They also suggested the use of a mixture of  $\text{HCl}$  and  $\text{HNO}_3$  (aqua regia) for facilitating the decomposition of carbonates, and the dissolution of sulphides using aqua regia or making the nitric acid medium more oxidizing through the addition of bromine.

In the decomposition of sulphide minerals, Rubeska (1968) specifically suggested the addition of mercuric nitrate solution to a mixture of nitric, sulphuric and tartaric acids in order to analyse for silver, zinc, cadmium-copper and lead in sphalerite, chalcopyrite, pyrite and galena. Donaldson (1902), in her "Methods for the Analysis of Ores and Related Materials," suggested the use of  $\text{HNO}_3$ ,  $\text{HCl}$ ,  $\text{HClO}_4$  and  $\text{HF}$  in the analysis of sulphide minerals for copper, nickel, zinc, cobalt, cadmium and iron. Other analysts (e.g. Jeffery and Hutchison, 1981) recommended the use of fusion methods in the analysis of alkali metals, carbonates, hydroxides, peroxides, nitrates and borates. In this study, some of the above methods were employed in the analysis of some common sulphide minerals collected from several locations and mines, largely in sedimentary rocks in the Benue Trough of Nigeria, with a view to finding out precisely if the very simple methods yield data comparable with the most complicated ones in the analysis of lead, copper, iron, nickel, cobalt, manganese, zinc, gold, silver, chromium and cadmium using the same sample solutions. Such a study would contribute to reducing the risks involved in geochemical analysis using mixtures of several concentrated acids and encourage more analysts to study real geochemical samples. The use of the simple acids might also be faster, cheaper and easier for use by geoscientists with little formal training in analytical chemistry. This report, which is part of a major geochemical and environmental programme, discusses scientific, technical, economic and geochemical significance of the data.

### Sampling

All the samples used were obtained from staff of the Department of Geology and Mining, University of Jos, and from the members of the public requesting for free advice on mineral resources of Nigeria. Therefore, no rigorous sampling procedure was undertaken in the classification and collection of the samples. Several samples are composed of mixed sulphides with gangue minerals, such as quartz, carbonates and sulphates.

All samples were crushed to very fine sizes, usually 100 mesh, using porcelain mortars. Details of the samples and their localities are given in Table 1.

#### Analytical Methods

The major equipment used in the determination was a manually operated Perkin-Elmer Atomic Absorption Spectrophotometer, Model 2380, with a digital readout. In addition, some equipment of the Nigerian Mining Corporation was used. Samples were weighed into 500cm<sup>3</sup> conical flasks or teflon beakers and the required acids added. The mixtures were covered and heated on electric heating blocks or sand bath in a fume cupboard.

All reagents used were analytical grade reagents, unless otherwise stated. Solutions for standard calibration were obtained from the American Bureau of Standards and standard mineral samples were also analysed and used as standards.

Table 1: Details of Sulphide Samples Used

Sample Number	Sample Description	Sample Locality
A1	Altered sphalerite with minor galena	New Zurak Mine, Plateau State.
B2	Less altered sphalerite	New Zurak Mine, Plateau State.
C3	Galena	Ishagu, near Abakaliki, Anambra State.
D4	Sphalerite	New Zurak Mine, Plateau State.
E5	Sphalerite	Ishagu, near Abakaliki, Anambra State.
F6	High Pure Powder (White)	New Zurak, Plateau State.
G7	Galena	Azara, Near Lafia (probably)
H8	Sphalerite	Abakaliki, Anambra State.
I9	Chalcopyrite	"
J10	Arenopyrite	Probably from Riehl, Bauchi State.

All glassware and plastic containers were washed using acid washing liquid containing one percent nitric acid solution.

About 0.2g of each mineral sample was accurately weighed into a 500cm<sup>3</sup> clean, dry, conical flask and decomposed according to the methods briefly described below. Blank solutions were also prepared during each procedure and digestions were carried out in fume cupboards.

Method 1: 50 percent HCl/Conc HNO<sub>3</sub> (Reeves and Brooks, 1978) Modified

To 0.2g of each sample weighed into a conical flask, was added 10cm<sup>3</sup> of 50 percent hydrochloric acid. The mixture was heated on a hot plate for about 30 minutes. This boiling was to ensure that sulphur escaped as hydrogen sulphide to avoid precipitation of lead as sulphate. This was followed by the addition of 20cm<sup>3</sup> of concentrated nitric acid. After cooling, the mixture was boiled until the quantity of acid left in the conical flask was reduced to about 2cm<sup>3</sup>.

The conical flask was then brought down, the sides washed with 25cm<sup>3</sup> distilled water and the contents of the flask returned to the hot plate and heated to boiling. The digest/solution was left to cool, transferred to a 100cm<sup>3</sup> volumetric flask and diluted to volume. Appropriate dilutions of these stock solutions were made where necessary prior to the determination of the elements concerned.

Method 2: 50 percent HCl/50 percent HNO<sub>3</sub> (Reeves and Brooks, 1978)

In this modified method of Reeves and Brooks (1978), the procedure followed was exactly the same as in the above method except that both acids were of 50 percent dilution.

Method 3: Aqua Regia (3:1 HCl:HNO<sub>3</sub>) (Stanton, 1966)

The aqua regia used was prepared just before use. 30cm<sup>3</sup> of the mixture was used for the digestion of each 0.2g of sample. The rest of the procedure was as described in Method 1 above.

Method 4: Concentrated Nitric Acid (Reeves and Brooks, 1978)

In this method, 30cm<sup>3</sup> of concentrated nitric acid was used on each 0.2g of sample digested.

Method 5: 50 percent Nitric Acid (Stanton, 1966)

In this modified method after Stanton (1966), 30cm<sup>3</sup> of 50 percent HCl was used on each 0.2g of sample digested.

Method 6: Tartaric Acid/Nitric Acid/Mercuric Nitrate (Rubeska, 1968)

In this method, 10cm<sup>3</sup> of 10 percent tartaric acid was added to 0.2g of each sample in a conical flask followed by the addition of 6cm<sup>3</sup> of concentrated HNO<sub>3</sub>. The mixture was left to stand at room temperature overnight. The following day, the mixture was heated on a steam bath for one hour, cooled and 3cm<sup>3</sup> of mercuric nitrate containing 10mg Hg/ml was added. The mixture was returned to the steam bath and heated until the sulphur turned white. The content of the flask was filtered and the undissolved residue and sulphur thoroughly washed with distilled water, collected into a 100cm<sup>3</sup> volumetric flask, and made up to volume.

Method 7: Concentrated HCl/HNO<sub>3</sub>/HClO<sub>4</sub>/HF Mixture (Donaldson, 1982)

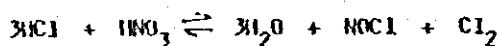
0.2g of powdered sample was transferred into a teflon beaker and 10cm<sup>3</sup> of HCl added. The beaker was covered and boiled for several minutes. The beaker was removed from the hot plate and 10cm<sup>3</sup> of concentrated HNO<sub>3</sub> was added and the mixture boiled until most of the sulphide minerals were decomposed. Then 10cm<sup>3</sup> of perchloric acid and 5cm<sup>3</sup> of hydrofluoric acid were added. The solution was boiled for several minutes after which the cover was removed and the solution evaporated to white fumes of perchloric acid. The solution was cooled, the sides of the beaker washed down with distilled water and the solution evaporated to a moist paste. To the moist paste, 3cm<sup>3</sup> of concentrated nitric acid and 25cm<sup>3</sup> of water were added and the mixture heated to dissolve the soluble salts. Some high quality dry paper pulp was added to the solution, where necessary, using Whatman No. 40 filter paper, the solution was filtered into a 100cm<sup>3</sup> volumetric flask. The paper and residue were washed properly and the solution made up to volume.

Results and Discussions

The results of this study are as presented in Tables 2 to 12. In general, the results show that iron, copper, nickel, chromium,

cobalt, manganese, cadmium and zinc were easily leached out using any of the acid digestion techniques. The most complicated methods, such as those utilising  $\text{HCl}$ ,  $\text{HNO}_3$ ,  $\text{HClO}_4$  and  $\text{HF}$  in one digestion mixture did not always produce the highest and most consistent results.

In the analysis for lead, it was observed that lead sulphate easily precipitated out of solution, giving the erratic results shown in Table 10. However, a repeat of the analysis with a smaller sample weight 0.4g rather than 0.2g proved very suitable as seen in Table 12. Table 12 also shows that all the seven digestion methods gave comparative results which are in agreement with earlier suggestions by several analysts. The results on Table 3 show that Methods 1, 2, 4, 5 and 7 gave silver contents very close to those of Method 6. Method 6, suggested by Rubenka (1967), describes a particular modification for the analysis of silver. In general, low results were obtained in the analysis for silver using aqua regia (Method No. 3). This is because this acid mixture proceeds as follows:



Nascent chlorine is expected to attack precious metals and some sulphides taking most metals into solution either as their simple chlorides or as complex chloro-anions. However, as silver chloride is not very soluble, this may have contributed to the low values obtained for silver using the aqua regia method of sample attack.

It was generally observed that there is the need for a little quantity of the acid to remain in the digestion flask at the end of the digestion. In cases where the acids were allowed to completely dry up, lower results were obtained, particularly for lead. The precipitates formed were insoluble in water, probably because of oxidation. It was also observed that the sphalerite samples were digested very easily using any of the methods, and as would be expected, they did not create the problem of producing insoluble precipitates even if the solutions dried up. Therefore, the published general analytical descriptions or methods for sulphides are not of general applicability since some sulphides create less digestion problems than others. As noted from above, lead sulphides probably give most problems.

For example, the starting quantity in the analysis for lead in galena should not be the same as the starting quantity for the analysis of zinc in zinc sulphide-sphalerite. For sphalerite, the major, minor and trace elements can easily be determined in the same sample solution, but for lead, which requires a smaller amount of sample to be digested, an analyst may run the risk of not being able to detect the trace elements if analysis for lead must be done on the same sample solution as precipitates of lead salts may co-precipitate other trace elements.

Another important factor which needs mentioning here is that the digestion flask should be covered during sample digestion to avoid the mixture drying up even before the sample is completely digested. Since drying up of samples would lead to the addition of more acids to the digested samples, this would raise the possibility of contamination and the amounts of elements present in the blanks.

In all the samples analysed, gold could not be detected probably because of the low contents of gold in the samples. To assay for gold, higher sample weights would have been digested for this specific purpose, in view of the generally lower gold contents in these types of sulphides. Although detailed geochemical and genetic discussions will be carried out elsewhere, the data presented in the tables show that the Zurak and Lafia(?) area sulphides contain the highest amounts of silver, and that, in general, the trace element contents reflect the environment from which the samples were obtained.

#### Conclusion from the Results

It can be concluded that 50 percent nitric acid is a convenient, simple and cheaper method for the digestion of simple sulphide minerals prior to the determination of lead, iron, copper, zinc, manganese, chromium, silver, nickel, cobalt and cadmium, using atomic absorption spectrophotometry. The use of concentrated acids or a mixture of concentrated acids in addition to the possible hazards

and high cost involved, does not always produce exceptionally different results from those obtained using 50 percent nitric acid.

This study suggests that simple, easy to obtain and work with acids should be used for the digestion of sulphide samples rather than wait for the purchase of platinum crucibles or teflon beakers at exorbitant prices before sulphide samples can be decomposed for analysis with a mixture of hydrofluoric, hydrochloric, perchloric and nitric acids as suggested by Donaldson (1982). In this era of recession and high cost of imported materials, such laboratories can conveniently carry out digestions using beakers or conical flasks with concentrated nitric acid, or better, cheaper and safer still, with 50 percent nitric acid.

#### Acknowledgement

We thank colleagues in the Department, the Faculty of Natural Sciences, University of Jos, the Nigerian Mining Corporation, as well as the Japanese International Cooperation Agency (JICA) and Dr. H. Takahashi for their support and encouragement. Some analyses were carried out in the laboratories of the Technical University of Berlin with the assistance of Dr. G. Hothel and the Volkswagen Foundation of the Federal Republic of Germany. I.U. Onyemachi typed the manuscripts.

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**Table 2: Iron Content (Weight Percent) in some Nigerian Sulphides**

**DIGESTION METHODS**

Samples	1	2	3	4	5	6	7	Sample Locality and Type
	50% HCl/ Conc HNO <sub>3</sub>	50% HCl/ 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/ HNO <sub>3</sub> / HClH <sub>4</sub> HF	
A	2.00	2.00	2.00	2.00	2.00	2.00	1.80	Zurak sphalerite (altered)
B	2.00	2.00	2.00	1.80	2.60	2.00	2.20	"
C	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Ishagu galena
D	8.60	8.60	9.40	9.40	8.60	9.60	9.00	Zurak sphalerite
E	1.001	1.00	1.00	1.00	1.00	1.00	1.00	Ishagu sphalerite
F	1.40	1.40	1.40	1.40	1.40	1.20	1.40	Zurak white mine dump powder
G	3.80	3.80	3.80	3.60	3.60	3.20	2.60	Iefia-Azara galena
H	4.80	4.60	4.40	4.80	4.80	4.80	4.40	Abakaliki sphalerite
I	35.40	35.80	36.40	36.70	36.20	33.00	31.60	"
J	27.00	26.80	26.80	26.80	20.60	14.00	27.40	Rishi-Bauchi arsenopyrite

n.d. = not detected.

**Table J: Silver Content (in parts per million) in some Nigerian Sulphides**

**DIGESTION METHODS**

Samples	1	2	3	4	5	6	7	Sample Locality and Type
	50% HCl/ Conc HNO <sub>3</sub>	50% HCl + 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/HNO <sub>3</sub> / HClO <sub>4</sub> / HF	
A	836	758	26	1117	930	1172	883	Zurak sphalerite
B	125	94	31	172	156	164	172	"
C	172	172	156	227	219	211	294	Ishlagu galena
D	94	109	47	148	102	109	94	Zurak sphalerite
E	258	273	16	289	281	273	297	Ishlagu sphalerite
F	438	404	94	461	469	453	547	Zurak white mine dump powder
G	1430	1461	55	1461	1445	1781	1075	Lafia-Azara galena
H	31	36	8	31	36	39	31	Abakaliki sphalerite
I	195	211	94	188	195	211	211	Abakaliki chalcopyrit
J	102	102	86	94	94	109	102	Rishi Bauchi arseno- pyrite(?)

Table 4: Copper content (ppm) in some Nigerian Sulphide Samples

DIGESTION METHODS

Samples	1	2	3	4	5	6	7	Sample Locality and Type
	50% HCl/ Conc. HNO <sub>3</sub>	50% HCl + 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tart- aric/ Nitric/ Herc. Nitrate HF	HCl/ HNO <sub>3</sub> / HClO <sub>4</sub> / HF	
A	536	536	571	571	554	554	607	Zurak sphalerite
B	1429	1429	1429	1464	1429	729	1500	"
C	36	36	36	36	37	36	37	Ishiegu (Abakaliki) galena
D	714	730	730	732	714	732	732	Zurak sphalerite
E	643	536	661	607	625	661	571	Ishiegu (Abakaliki) sphalerite
F	536	571	571	571	579	554	554	Zurak white mine dum powder
G	79057	78929	78929	78929	79057	60357	78571(7)	Lafia-Azara galena
H	571	589	607	607	607	625	643	Abakaliki sphalerite
I	245536	251706	251786	248214	250000	2393	2527	Abakaliki chalcopyrite
J	212	214	214	179	196	214	196	Rishi- Bauchi arseno- pyrite(7)

Table 5: Nickel contents (ppm) in some Nigerian Sulphides

Samples	DIGESTION METHODS							Sample Locality and Type
	1	2	3	4	5	6	7	
	50% HCl/ Conc HNO <sub>3</sub>	50% HCl/ 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/HNO <sub>3</sub> / HCl/HF	
A	45	45	30	30	20	20	30	Zurak sphalerite (altered)
B	45	40	35	35	40	15	15	"
C	20	20	n.d.	20	20	20	20	Ishiegu (Abakaliki) galena
D	20	20	20	20	20	20	20	Zurak sphalerite
E	15	15	15	15	15	15	10	Ishiegu (Abakaliki) sphalerite
F	50	50	50	55	65	50	50	Zurak sphalerite
G	n.d.	n.d.	n.d.	20	20	n.d.	n.d.	Lafia-Azura galena
H	25	20	n.d.	25	25	n.d.	n.d.	Abakaliki sphalerite
I	210	275	27	260	255	210	220	Abakaliki chalcopyrite
J	55	55	n.d.	40	50	n.d.	20	Rishi-Nauchi arsenopyrite(?)

n.d. = not detected.

Table 61 Chromium Content (ppm)

DIGESTION METHODS

Samples	1	2	3	4	5	6	7	Sample Locality and Type
	50% HCl/ Conc HNO <sub>3</sub>	50% HCl/ 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/HNO <sub>3</sub> / HClO <sub>4</sub> /H <sub>2</sub> O	
A	30	30	15	35	35	35	40	Zurak sphalerite (altered)
B	30	30	10	25	30	30	30	"
C	35	35	5	30	30	30	35	Ishisu (Abakoliki) galena
D	20	20	20	20	20	20	20	Zurak sphalerite
E	25	25	5	15	20	20	20	Ishisu (Abakoliki) sphalerite
F	30	30	5	20	20	25	20	Zurak white mine dump powder
G	10	10	n.d.	10	10	10	5	Lafin-Azara galena
H	5	5	n.d.	5	5	5	5	Abakoliki sphalerite
I	10	10	n.d.	n.d.	5	10	10	Abakoliki chalcopryite
J	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Richi-Bauchi arsenopyrite

n.d. = not detected.

**Table 7: Cobalt Content (ppm) in some Nigerian Sulphide Samples**

Samples	DIGESTION METHODS							Sample Locality and Type
	1	2	3	4	5	6	7	
	50% HCl/ Conc HNO <sub>3</sub>	50% HCl/ 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/HNO <sub>3</sub> / HClO <sub>4</sub> /HF	
A	335	335	335	335	335	335	335	Zurak sphalerite (altered)
B	775	775	775	775	775	775	775	"
C	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Ishiagu (Abakaliki) galena
D	835	835	835	835	835	775	775	Zurak sphalerite
E	335	335	335	335	335	335	335	Ishiagu (Abakaliki) sphalerite
F	335	335	335	335	335	335	335	Zurak white mine dump powder
G	110	110	110	110	110	110	110	Lafia-Azara galena
H	500	500	445	500	500	390	400	Abakaliki sphalerite
I	335	335	335	335	335	335	335	Abakaliki chalcocysite
J	665	665	720	665	665	555	665	Rishi-Bauchi arsenopyrite

n.d. = not detected.

Table 8: Mercury Content (ppm) in some Nigerian Sulphide Samples

Samples	DIGESTION METHOD							Sample Locality and Type
	1	2	3	4	5	6	7	
	50% HCl Conc $\text{HNO}_3$	50% HCl 50% $\text{HNO}_3$	Aqua Regia	Conc. $\text{HNO}_3$	50% $\text{HNO}_3$	Tartaric/Nitric/Hydrochloric Nitrate	HCl/ $\text{HNO}_3$ / $\text{HClO}_4$ /HF	
A	555	575	555	600	605	600	520	Zurak sphalerite (altered)
B	130	130	130	130	125	135	120	"
C	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Ishaga (Abakiliki) galena
D	2760	2775	2760	2840	2755	2900	2770	Zurak sphalerite
E	5	5	5	5	5	5	5	Ishaga (Abakiliki) sphalerite
F	745	740	745	750	760	760	610	Zurak white mine clay powder
G	40	40	35	35	35	35	35	Lafia-Arara galena
H	840	840	845	805	800	820	835	Abakiliki sphalerite
I	13750	14000	13500	13750	13500	13750	14250	Abakiliki chalcopyrite
J	25	25	20	30	25	20	25	Rishi Bauchi arsenopyrite

n.d. = not detected.

Table 9: Cadmium Content (ppm) in some Nigerian Sulphide Samples

Samples	DIGESTION METHODS							Sample Locality and Type
	1 50% HCl Con HNO <sub>3</sub>	2 50% HCl 50% HNO <sub>3</sub>	3 Aqua Regia	4 Conc. HNO <sub>3</sub>	5 50% HNO <sub>3</sub>	6 Tartaric/ Nitric/ Mercuric Nitrate	7 HCl/HNO <sub>3</sub> / HClO <sub>4</sub> /HF	
A	607	634	598	589	643	616	536	Zurak sphalerite (altered)
B	2197	2197	2232	2260	2270	2116	2045	"
C	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Ishiqw (Abakaliki) galena
D	1366	1348	1331	1322	1340	1241	1215	Zurak sphalerite
E	1697	1652	1623	1643	1652	1661	1590	Ishiqw (Abakaliki)
F	696	674	679	679	696	706	652	Zurak white mine clump powder
G	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	Lafia-Azara galena
H	2241	2241	2206	2232	2293	2259	2277	Abakaliki sphalerite
I	90	90	80	80	80	n.d.	10	Abakaliki chalcopyrite
J	27	27	27	27	27	27	27	Rishi-Bauchl greenopyrite

n.d. = not detected.

Table 10: Lead Content (weight percent) in some Nigerian Sulphides

DIGESTION METHODS

Samples	1	2	3	4	5	6	7	Sample Locality and Type
	50% HCl (conc. HNO <sub>3</sub> )	50% HCl 50% HNO <sub>3</sub>	Aqua Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	100% HNO <sub>3</sub> H <sub>2</sub> O <sub>2</sub> /NaOH	100% HNO <sub>3</sub> H <sub>2</sub> O <sub>2</sub> /NaOH	
A	21.00	44.72	18.00	33.80	17.42	30.54	38.74	Zurik sphalerite (altered)
B	0.03	0.03	0.03	0.03	0.03	0.03	0.03	"
C	44.72	56.60	25.22	30.40	47.32	44.04	72.30	Ishaga (Abakaliki) galena
D	0.03	0.02	0.01	0.03	0.03	0.03	0.03	Zurik sphalerite
E	0.03	0.03	0.02	0.03	0.03	0.03	0.03	Ishaga (Abakaliki) sphalerite
F	7.67	7.81	7.00	7.50	7.15	7.12	7.00	Zurik white siliceous chert nodules
G	62.40	67.34	15.11	63.20	45.34	44.50	69.00	Lafia-Azara galena
H	1.47	1.47	1.45	1.47	1.54	1.55	1.51	Abakaliki sphalerite
I	1.00	1.01	1.21	1.00	1.03	1.02	1.00	Abakaliki chalcopyrite
J	0.04	0.06	0.04	0.05	0.04	0.04	0.04	Richt-Asuchi chalcopyrite(?)

Table 11: Zinc Content (weight percent) in nine Nigerian Sulphide Samples

Samples	DIGESTION METHODS							Sample Locality and Type
	1	2	3	4	5	6	7	
	50% HCl Conc. HNO <sub>3</sub>	50% HCl 50% HNO <sub>3</sub>	Aqua. Regia	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Mercuric Nitrate	HCl/HNO <sub>3</sub> / HClO <sub>4</sub> /HF	
A	26.00	26.00	26.05	27.00	27.00	27.70	26.05	Zurak sphalerite (altered)
B	52.20	52.10	52.11	52.24	52.20	52.23	52.23	"
C	0.01	0.01	0.01	0.01	0.01	0.01	0.01	Ishaga (Abakaliki) galena
D	49.73	49.73	49.70	49.73	49.73	49.73	40.05	Zurak sphalerite
E	62.72	62.72	62.71	62.70	62.74	62.73	62.60	Ishaga (Abakaliki) sphalerite
F	30.43	30.43	30.32	30.60	30.50	30.50	30.04	Zurak white mine dump powder
G	0.11	0.11	0.11	0.12	0.11	0.11	0.12	Lafia-Azara galena
H	61.73	61.70	61.70	62.90	62.01	61.98	61.70	Abakaliki sphalerite
I	0.09	0.05	0.05	0.05	0.05	0.05	0.05	Abakaliki chalcopyrite
J	0.004	0.004	0.004	0.004	0.004	0.004	0.004	Pishi-Uuchi arsenopyrite(?)

**Table 12: Lead Content (weight percent) in some Nigerian Sulphide Samples using Soxhlet (0.1 gm) Sample results for digestion**

**DIGESTION METHODS**

Sample	Digestion Methods							Sample Locality and Type
	1	2	3	4	5	6	7	
	50% HCl Conc HNO <sub>3</sub>	50% HCl 50% HNO <sub>3</sub>	Aqua Regia HNO <sub>3</sub>	Conc. HNO <sub>3</sub>	50% HNO <sub>3</sub>	Tartaric/ Nitric/ Hydrofluoric Nitrate	HCl/HNO <sub>3</sub> / HCl/HF	
A	38.90	38.90	38.90	36.10	37.50	35.50	38.70	Zurak sphalerite (altered)
C	83.30	83.30	83.30	77.90	77.90	76.40	76.30	Ishaku (Abakaliki) galena
F	8.30	8.30	7.90	7.90	7.90	7.00	8.80	Zurak white mine ultrapowder
G	68.30	61.00	61.00	69.00	69.70	68.30	69.20	Lafin-Azana galena

ENVIRONMENTAL AND ECONOMIC STUDIES OF TRACE ELEMENTS  
ASSOCIATED WITH MINING AND MINERAL PROCESSING IN  
PARTS OF PLATEAU STATE, NIGERIA - A PRELIMINARY REPORT\*

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ABSTRACT.

The Plateau State of Nigeria has a surface area of about 58000 km<sup>2</sup> and elevation above mean sea level ranges from about 200 to 1800 metres. A number of major tributary river systems which flow to the Rivers Niger and Benue and to Lake Chad rise from the Plateau. The State is covered by igneous, metamorphic and sedimentary rocks of Precambrian to Recent age within which, under favourable geological conditions, are several mineral occurrences and economic deposits. Economic deposits of cassiterite, columbite-tantalite, lead-zinc, thorite, zircon, barytes, salt, clays, mineral water, limestone, coal, iron ore, marble and gemstones are known in the State. A number of these have been mined, processed and smelted on large- or small-scale basis.

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\*Research supported by the University of Jos, the Japanese International Cooperation Agency (JICA) and the Volkswagen Foundation of the Federal Republic of Germany.

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A large number of samples of rocks, minerals, stream sediments, soils and laterites, mine dumps, slags, water and plant materials have been carefully collected from areas of known mineral occurrences and mineral processing plants and also from apparently 'barren' areas and analysed for major, minor and trace elements. This forms the basis for students' training and systematic baseline trace element data used for a multidisciplinary study in mineral exploration, fundamental studies of crustal evolution and ore-forming processes, and environmental studies, including those in agriculture, medical geography, pollution studies and ecological fields of interests since major metal pollution originates from geological weathering, industrial processing of ores and metals, the use of metals, leaching of metals from domestic and solid waste dumps, and animal and human excretions.

Preliminary results from this on-going study show that although mining and mineral processing have contributed a lot to the national economy and the growth of a number of towns in Plateau State, it has also contributed immensely to environmental pollution. However, with careful monitoring and mineral exploitation using modern techniques of large- and small-scale mining and mineral processing, the mining industry in Plateau State could continue to provide jobs, save and earn foreign exchange and supply major alloy metals.

(such as tin, tantalum, niobium, molybdenum, tungsten and titanium) and probably by-product metals (such as silver, antimony, cobalt, germanium, cadmium, copper, uranium, thorium, hafnium and zirconium). However, under the current practice without strict environmental protection laws and enforcement, major environmental and health hazards for the State and other parts of Nigeria which are drained by rivers flowing from the high plateau might result.

## 1 INTRODUCTION

Metals make up the largest group of the chemical elements, but their characteristics differ greatly within the biosphere. It is now widely recognised that the important factor determining the effects of a metal on organisms is not generally its total concentration but rather its speciation - the specific forms and nature of the element. Metals may occur as hydrated ions, as complex inorganic or organic ligands, as adsorbed and absorbed ions, as colloids, as precipitates, and in live and dead biota. Some metals may also be essential to human life, whereby as trace elements they are always present in certain proportions in all healthy living tissues of the same species and deficiency symptoms may be noted with their depletion or removal which can be attributed to distinct

biochemical defects. Examples include potassium, sodium, magnesium and calcium. When the supply of some essential trace elements, such as zinc, becomes excessive, they become toxic. Toxic levels are generally 40- to 200-fold of the minimum levels required for correct nutritional activity in the organism. Hence, in the studies of trace elements from the possible sources to man, the general trend followed is that an undersupply leads to element deficiency, sufficient supply results in optimum conditions, and oversupply results in toxic effects and, possibly, death in the end.

The essential metals, whose ions are used by biological systems must of necessity be both relatively abundant in nature and readily available as soluble species. However, when considered from environmental pollution point of view, metals may be classified (i) as non-critical, such as Na, K, Mg, and Ca, which are essential metals and hard acceptors of electrons (Lewis acids) and which form stable bonds with hard donors (Lewis bases), such as  $H_2O$ ,  $OH^-$ ,  $Cl^-$ , etc; (ii) toxic, but very soluble or very rare; and (iii) very toxic and relatively accessible metals. The two last groups are generally soft acceptors which form stable bonds with soft donors, such as -SH groups and other active sites of proteins. Some common elements which, at toxic levels,

become major health hazards are As, Be, Cd, Cr, Cu, Pb, Hg, Mo, Se, Ag, Th, and Zn. In some cases, unusual concentrations have produced intoxicating or catastrophic effects, as in the cases of mercury poisoning in Minamata Bay, Japan; Cadmium poisoning causing the "itai itai" disease in the Japanese villages on the banks of the Jintsu River, Toyama, Japan; lead poisoning, as in Leipzig, Germany, during the summer of 1930, etc. (Forstner and Wittmann, 1983). A large number of such studies have been documented from Japan, Europe and U.S.A., but similar studies are yet to be published on Nigeria.

In a number of developed countries, variable but enforceable national standards that determine the quality of water for different uses have been set. Although no such enforceable standards or environmental protection laws currently exist in Nigeria, the quality of Nigeria's surface and underground waters can be compared with these international standards so as to establish concentration levels below which harmful effects may be produced. Toxic metals may occur in aqueous ionic forms derived from minerals and from man-made sources, such as herbicides, fertilisers, detergents, metal processing and industrial processing and products. Permissible limits for waters must take cognisance of the accumulation by organisms through the food chain and

the environmental impact of waters discharged to the environment, especially for domestic, recreational, agricultural and industrial purposes. Drinking water quality criteria have been proposed by the World Health Organisation (WHO) and several developed countries. These limits show wide fluctuations which often vary by up to a factor of 200.

In general, there are five different major sources from which metal pollution of the environment originates. These are geological weathering of rocks, industrial processing of ores and metals, the widespread use of metals and metal components, leaching of metals from domestic refuse and solid waste dumps, and animal and human excretions, which generally contain heavy metals. Plateau State has several mineral occurrences, most representatives of Nigerian igneous, metamorphic and sedimentary rock types, and has a long history of mining, mineral processing and smelting. Hence, the resultant extensive environmental and human pollution through geological and human activities provides a key area for the environmental and economic geological study of trace element mobility from baseline or background sources in rocks and minerals through geological, human and biological activities to man via the food chain.

In areas characterised by metal-bearing rocks, these trace elements will necessarily occur at much higher levels

than the baseline or natural background levels in the water, soil, plants and bottom sediments of the area. Areas with economic deposits are exploited to recover and process the ore. This in turn leads to the disposal of tailings, discharge of effluents to specially constructed lakes or natural drainage systems, discharge of effluents and possibly processing and smelting operations. These processes result in atmospheric and hydrological pollution. The general problem which, therefore, arises is how to distinguish between natural geological weathering and metal enrichment attributable to human and industrial activities.

This study is, therefore, aimed at determining the trace element contents in rocks, stream sediments, minerals, soils, plant and water samples from the tin and lead-zinc mining and non-mining areas of Plateau State with a view to establishing their negative and positive environmental effects and economic significance (Ogezi, 1984). Another aim is that since critical pollution, geochemical and mineral exploration studies are often hampered by cumbersome, costly and often dangerous analytical techniques, simple, cheap and reasonably accurate and precise methods of analyses will be devised and developed to facilitate the large number of analyses involved (Adiuku-Brown and Ogezi, 1984 and this volume). Another important aspect of the study is

scientific students' training for the acquisition of new knowledge and skills and updating of ideas, especially in the fields of environmental geochemistry and mineral exploration (Adiuku-Brown, 1983; Agada, 1983; Bob-Manuel, 1983; Ekpo, 1984; Hamza, 1984; Ochonogor, 1984; Okeke, 1984; Onyeji, 1984). It is also hoped that in collaboration with the metal mining, processing and fabrication organisations, through the investigation of mineral processing of tin and its associated by-products and lead and zinc, cheap methods of extraction will be found for such metals as niobium and tantalum from tin and associated ores, and silver, cobalt, lead, zinc, etc. from lead-zinc ores. This would provide suitable indigenous alloying metals for virile Nigerian metal-based industries as well as save and earn foreign exchange, provide jobs and aid in the acquisition of technical skills.

## 2. SAMPLING AND SAMPLE PREPARATION

In view of their significance, it was decided to use samples of rocks, minerals, waters, stream sediments, mine dumps, ores, slags, soils, laterites and plant material for this study. Based on a detailed knowledge of the geology of the State, the possible sampling sites covering the high and low altitude areas of Plateau State were selected to include areas with different rock types, different types of water bodies, mineral occurrences, economic mineral deposits, mines, mineral processing and apparently barren areas. These areas cover the whole State. Based on this, about 10 areas and about 22 sampling localities were selected for coverage. In spite of this pre-sampling decision based on regional geology, a detailed geological reconnaissance was carried out at each sampling locality to deduce the local geology, evidence of mineral exploitation, topography, vegetation, drainage and other features of the area. Samples were labelled in the field and details recorded in field notebooks and on geological and topographic maps. A summary of the type and approximate number of samples collected is presented in Table 1.

At each point where a water sample was taken, air and water temperatures, conductivity, pH, Eh, turbidity and the colour of the water was recorded prior to a minimum

**Table 1: Summary of Types and Localities of Samples Collected for Preliminary Regional Study and Numbers treated so far.**

S/No.	Sample Types	Approximate Total No. Collected	Approximate No. Partially Collected
1	Rocks	200	40 - for mineralogy, major and trace elements
2	Stream sediments	100	25 - for mineralogy, trace elements and sizes
3	Soils and laterites	50	10 - for size fractions
4	Minerals	20	15 - for mineralogy, major and trace elements
5	Mine dumps	10	5 - for mineralogy
6	Smelting slags	5	-
7	Stream and lake waters	120	50 - for Ca, Mg, Cu, Fe, Zn, Mn and Pb
8	Plants	20	-
9	Well waters	20	15 - for pH, T and trace elements

of 250-500 ml of water being taken into clean, carefully rinsed-out plastic water bottles which were previously washed with dilute nitric acid.

Water samples were preserved in a refrigerator and analysed shortly after collection. To minimise the effects of organic substances, bring all the ions to the same form or oxidation state in solution and minimise metal absorption onto the walls of containers, all water samples were acidified with analytical reagent grade nitric acid to a pH of about 2. The possibility of sample contamination by substances leached from the container walls and absorption of constituents onto container walls were considered and tested for, but negligible. Standardised strict water sample collection, preparation and analytical procedures as well as the use of blanks and standard determinations are being devised for the detailed stage of this investigation. Similar procedures and basic equipment for pore waters in sediments and for plant materials are being collated.

For rock samples, fresh large samples weighing a minimum of 2 kg are collected, and representative fresh portions sawn for thin section preparation or crushed into powder for analysis. Sulphide minerals and stream sediments were leached using hot dilute nitric acid or acid mixtures as described in Adiuku-Brown and Ogezi (1984, and this volume).

For the major and trace element analysis of rocks and laterites carried out at the Technical University of Berlin,

the lithium tetraborate fusion technique was used. To do this,  $0.3000 \pm 0.0002$  g finely ground ( $\sim 120$  mesh) material was weighed and completely homogenised for at least 10 minutes in an agate mortar with  $1.2000 \pm 0.0002$  g lithium tetraborate. After complete homogenisation, 1.2500 g of the mixture was weighed into a  $6 \text{ cm}^3$  graphite crucible and put in the furnace set at  $1000^\circ\text{C}$  for 30 minutes for fusion to be complete. The molten mixture was quickly and completely transferred into a beaker, on a hot plate, containing 30 ml of 4 per cent nitric acid (made from 10 ml  $\text{HNO}_3$  and 165 ml distilled water) and the mixture stirred until completely dissolved. The solution was quickly filtered, the filter paper thoroughly washed and the filtrate made up to 50 ml, thereby giving a 200-fold dilution for the atomic absorption spectrophotometric determination of trace elements. For the determination of major elements, a 2000-fold dilution was used. Although the lithium tetraborate digestion technique is simple, effective and cheap when compared with other digestion techniques, the major problems are those of complete recovery of the molten mixture from the crucible and consistent analytical results.

## PERFORMANCE AND RESULTS

So far, a review of and the acquisition of relevant literature and analytical techniques have been completed. However, in view of the limited sensitivity of the sample preparation and analytical facilities and materials available, progress has been necessarily slow. Preliminary analysis of over 50 water samples using the Departmental Manual Perkin-Elmer 2380 Atomic Absorption Spectrophotometer (AAS) has been completed for Ca, Mg, Cu, Fe, Zn, Mn, Pb and Cd (Tables 1 to 5). This has largely confirmed the necessity for preconcentration of samples prior to AAS analysis of samples or analysis using more sensitive equipment that can analyse concentrations at parts per billion (ppb) levels instead of parts per million (ppm) possible with the AAS.

Since the inception of the project, seven B.Sc. (Honours) theses projects have been completed (Agada, 1983; Ekpo, 1984; Hamza, 1984; Manuel, 1983; Ochonogor, 1984; Okeke, 1984; Onyeji, 1984) and one research (M.Phil/Ph.D.) project has been initiated (Adiuku-Brown, 1983). In addition, two conferences papers have been presented in which it was shown that sulphides could be leached using simple dilute acids and that mining activity poses major environmental hazards in the Zurak area (Adiuku-Brown and Ogezi, 1984 and this volume; Ogezi, 1984).

In a study of an area of about 28 km<sup>2</sup> of the area around Delimi River Basin, off the Jos-Bukuru Road from Narabi to Tilden Fulani in Bauchi State, Agada (1983) carried out a sedimentological analysis of the upper reaches of the Delimi, which flows several hundred kilometres to Lake Chad in the northeast. He showed that the sediments are poorly- to moderately-sorted down the river basin and composed of coarse- to medium-grained sands which were deposited in one phase of deposition. The moderate rounding was as a result of the corroding ability of the channel bedrock. The winning of economic deposits of cassiterite, sand and gravel and shallow groundwater from alluvial sediments recharged by the Delimi River, which drains Jos, could have major effects on water quality. Detrital minerals and minor radioactive element mineralisation in the sediments of parts of Bauchi and Bornu States, several hundred kilometres north-east of Jos, have been ascribed to the rocks of the Younger Granite Province (C. Okujeni, oral communication, 1983). Pollution could similarly extend downstream towards Lake Chad.

In a similar geological, mineralogical and geochemical study of the northeast part of Naraguta Sheet 168, covering the Permanent Site of the University of Jos, Ekpo (1984) showed that the major rocks of the area are the migmatites, gneisses and the Jurassic Younger Granites, together with

minor intrusions of basic dykes and aplo-pegmatitic gneisses. By studying the geochemistry and mineralogy of the rocks, stream sediments and water, it was shown that the finest-clay-size fractions had the highest concentration of iron, copper, manganese, cobalt, lead, calcium and zinc. Concentrations could be correlated with the mineralogy, the geology and environmental factors. Elements, such as lead, cobalt and copper, detected could be correlated with pollution from metropolitan Jos.

Hamza (1984), in a similar study of the area south of the Jos-Rukuba Road, in Plateau State, studied the geology, trace element content and mineralogy of stream sediments within the area to provide a basis for predicting the environmental hazards and mineralisations of the area. Although the raw water samples had relatively low trace element contents, Hamza (1984) showed that in the stream sediments, the coarsest-size fractions formed the bulk of the samples and that the relative proportions decreased with progressively finer fractions. This may be due to sampling, or more likely, due to nearness of stream sediments to source rocks. After hot extraction with 4M  $\text{HNO}_3$ , it was found that, with very few exceptions, the finest or clay-size fractions of stream sediments had the highest contents of Fe, Cu, Mn, Pb, Ca, Mg and Zn (Table 2).

This was attributed largely to adsorption to clay fractions, but individual variations along the same streams of trace element concentrations was correlated with rock type and environmental factors, such as mining, mineral processing and human activities (Table 2).

Table 2: Trace Element Contents (ppm) of Different Stream sediment near Jos after Hot Extraction with 4M HNO<sub>3</sub> (after Hamza, 1984, Table 3 p. 47, and Table 7.1<sup>3</sup>, p. 58). The highest concentrations are in the finest size fractions. Note lower concentration in water samples.

Sample No.	Mesh Size of Stream Sediments	Trace Elements (ppm)							
		Fe	Cu	Mn	Co	Pb	Ca	Mg	Zn
STS-1	+0.5mm	200	0.5	3.0	0.3	0.5	0.0	1.8	1.1
	-0.5+0.25mm	275.7	0.3	3.8	0.3	0.6	0.1	1.8	1.6
	-0.25+0.125mm	298.6	1.4	5.8	0.4	1.0	0.3	66.3	2.2
	-0.125mm	586.5	16.9	14.0	0.7	2.0	0.8	12.4	12.5
	Water	4.0	n.d	1.0	n.d	n.d	3.5	2.7	0.3
STS-2	+0.5mm	119.0	0.6	1.0	0.4	0.5	1.1	0.4	70.0
	-0.5+0.25mm	118.9	0.0	1.0	0.4	0.8	1.0	0.3	45.0
	-0.25+0.125mm	167.6	0.6	1.0	0.4	0.8	1.9	0.5	90.0
	-0.125mm	275.7	3.9	3.0	0.6	3.6	2.1	0.6	5.0
	Water	4.0	n.d	n.d	n.d	n.d	3.5	2.7	n.d
STS-3	+0.5mm	127	1.3	1.6	0.1	0.6	0.1	2.8	1.1
	-0.5+0.25mm	137.8	0.3	1.8	0.1	0.9	0.1	1.8	0.9
	-0.25+0.125mm	210.8	0.7	2.8	0.3	2.0	0.2	2.8	1.4
	-0.125mm	437.8	7.1	8.0	0.5	2.5	0.4	9.8	4.0
	Water	3.3	n.d	n.d	n.d	n.d	3.3	2.7	n.d

n.d. = not detected.

Manuel (1983), in a stratigraphic and sedimentological study of the tin mineralisation in parts of the Gold and Base Company Mines in the Rayfield-Barakin Delimi areas, Jos, showed that the sediments are poorly sorted, angular to sub-angular, medium- to coarse-grained river sands deposited in an oxidizing environment in stream channels averaging 6 metres each in depth and width. Manuel (1983) also showed, from stratigraphic correlations and heavy mineral analysis, that the highest concentration of cassiterite and associated ores occurred at sediment depths ranging from 4 to 8 metres, due largely to their higher density. Detailed sedimentological analysis showed that the sediments were mainly saltated by the river currents and the poorly-sorted angular nature was due to the action of eddy currents on these materials deposited close to source areas.

Ochonogor (1984), in a geological and environmental geochemical study of the Rayfield-Bukuru area, where there was extensive past alluvial mining and mineral processing, showed that the concentration of most trace elements in the waters were within permissible international standards. Although a much larger sample population was required for firmer conclusions, Ochonogor (1984) tentatively ascribed the variations in elemental concentrations to environmental, human and geological factors, and recommended detailed multi-

-disciplinary study, especially since endemic goitre and other diseases, which occur in the general area, may be due to geochemical factors. Ochonogor also obtained trends of trace element concentrations in stream sediments similar to those of Ekpo (1984) and Hamza (1984), whereby trace elements generally increased with decrease in grain sizes of sediments (Table 3). Well water samples contained significantly lower Ca, Mg, Fe, Zn and Mn than adjacent stream waters, hence well waters would provide purer drinking waters (Table 3). However, in the absence of thick weathered superficial cover wells dry up during the dry season.

Table 3: Trace Element Contents of Stream Sediments (S), Stream Waters (SW), and Well Waters (WW) from the Rayfield-Bukuru area, near Jos (after Ochonogor, 1984; Table 3, p. 45, and Table 4, p. 47). Compare data with Table 2.

Sample	Stream Sediment Mesh Size (Mm)	Trace Elements (ppm)							
		Fe	Cu	Mn	Co	Pb	Ca	Mg	Zn
S1	+0.5mm	62.2	0.0	1.1	0.1	0.0	0.3	0.4	0.3
	-0.5+0.25mm	180.1	0.1	2.2	0.1	0.0	0.6	1.2	0.7
	-0.25 - 0.125mm	173.0	0.1	3.2	0.2	0.0	0.6	1.8	1.1
	-0.125mm	200.0	1.4	8.0	0.2	0.3	32.0	7.0	5.0
S2	+0.5mm	91.9	0.1	0.6	0.1	0.0	0.1	0.3	0.4
	-0.5 + 0.25mm	127.0	0.5	1.0	0.1	0.1	0.1	0.6	0.9
	-0.25+0.125mm	105.4	0.5	1.8	0.2	0.2	0.4	1.2	1.4
	-0.125mm	78.4	2.8	3.1	0.4	0.5	0.8	2.0	3.0
S3	+0.5mm	59.5	0.0	0.6	0.2	0.0	0.1	0.3	0.5
	-0.5 + 0.25mm	75.7	0.1	0.9	0.2	0.1	0.1	0.7	0.6
	-0.25+0.125mm	87.8	0.1	1.2	0.4	0.0	0.3	0.9	1.2
	-0.125mm	229.7	4.5	4.3	0.8	0.0	1.6	3.1	7.7
SW1	Stream Water	9.7	n.m	0.2	n.m	n.m	2.5	0.1	0.0
SW2	" "	8.7	n.m	0.3	n.m	n.m	16.0	2.7	0.0
SW3	" "	9.0	n.m	0.4	n.m	n.m	11.8	1.8	0.1
SW4	" "	9.7	n.m	0.2	n.m	n.m	2.4	0.1	0.0
WW1	Well Water	0.0	n.m	0.0	n.m	n.m	0.0	0.0	0.0
WW2	" "	0.0	n.m	0.0	n.m	n.m	0.0	0.0	0.0

n.m = not measured.

Okeke (1984), in a geological and hydrogeological study of the northern part of Naraguta Sheet 168 NE, between the Jos-Bukuru and the Rukuba Younger Granite Complexes of the Jos area, identified three hydrogeological units consisting of the Quaternary sedimentary deposits, the weathered zones in the crystalline igneous and metamorphic rocks, and the tectonically fractured zones in the crystalline rocks. The thickness of each unit varies and water levels fluctuates widely, resulting in several dry wells at the peak of the dry season, about April/May. Okeke correlated the chemical quality of the groundwater in the study area with the geology. When compared with World Health Organisation (WHO) standards for drinking water, both the surface and groundwater of the area have total hardness, pH, cations and anions, etc., within permissible limits. However, the colour and turbidity of the Delimi River and some wells, as well as their ammonia, taste and amount of total dissolved solids are largely, from preliminary examinations, excessive. These waters need treatment prior to domestic and industrial uses as they have been polluted by agricultural practice, the extensive use, especially along stream valleys of animal and manufactured fertilisers, disposal of solid and animal waste into and along the banks of the Delimi River and its tributaries

(such as near Gada Bauchi and Gada Biyu), leachates from the extensive past mining activities in the Haraguta area, and poor sanitation from cemeteries, septic tanks and liquid waste from open gutters draining Jos, especially as storm run-off. In areas of relatively thin superficial cover, or excessive concentration of water wells, where over-abstraction of water occurs, the water wells, which supply purer water than the stream water, most wells become dry during the dry season. A complete assessment of the quality of the surface and subsurface waters must include bacteriological studies.

Mineralogical examination of the Bisichi-Jantar mine dumps shows that the dump contains zircon, monazite, thorite, and opaque ore minerals, such as magnetite with ilmenite in oriented intergrowth, magnetite martitised, or replaced by limonite or replaced by rutile, pseudorutile or leucoxene. The oxidation and leaching is caused by the presence of water. Columbite-tantalite, and rare chromite and tapiolite are also present (A. Mücke, personal comm., 1984). Preliminary examination of these mine dumps and stream sediments indicate high radioactivity and mobile trace element contents.

In a geological study, with particular reference to the sedimentation pattern of the Delimi River Valley north of Jos, especially in order to delineate the parameters which determine the sedimentation of tin-bearing minerals, Onyeji (1984) identified tectonic structures, such as joints, veins, dykes, foliations and lineations in crystalline rocks, and graded bedding, cross bedding, massive bedding and mudcracks in stream-laid sedimentary rocks. The Delimi River is erosional and highly channelised when it passes through ridges. Grain size analysis shows largely unimodal distribution but with symmetrical and asymmetrical distribution. Samples with asymmetrical distribution of grains contain large amounts of heavy minerals, including cassiterite, and these samples largely occur upstream of channel bars. Mineralogical and textural features of grains suggest that the sources of the sediments are very close by, largely from the surrounding Younger Granite and Basement Complex rocks.

Like the proportional distribution found by Hamza (1984), Onyeji's more detailed sieving showed that although the greatest portion (over 80 percent) were the 2mm to 0.5mm fractions, there was also systematic decrease in the relative proportions from the 0.5 to 0.125mm mesh fractions. The major components of grains were 60-80 percent quartz, 10-30 per cent feldspars, 5-15 percent opaque oxides and up to 10 percent rock fragments for the light fractions, while the heavy minerals included monazite, zircon, magnetite, ilmenite and cassiterite.

Adiuku-Brown and Ogezi (1984 ; and this volume) analysed ten different sulphide minerals, consisting largely of galena, sphalerite and arsenopyrite, from different parts of Nigeria using acids, acid mixtures and different acid strengths and found that dilute nitric acid, which is one of the cheapest, safest and simplest of the leaching solutions used, gave results comparable to those using stronger and more dangerous acids or acid mixtures, or using the specific methods for sulphides. These sulphides could be correlated with their geological environments and could provide significant by - product elements, such as Cd, Sb, Cu, Ag from the sphalerite and galena, Sb from galena, and Ge and Cd from the sphalerite (Table 4). Elements which could also cause impurities and metallurgical problems are also discernible (Table 4). In view of the large number of samples that may be treated for a comprehensive study, as well as considerations of safety, procurement and technical problems, the use of simple dilute leachates is of vital importance, especially

Table 4: Major and Trace Element Contents of Samples from the New Zurak Lead-Zinc Mines, SW Plateau State, Nigeria.

Sample No.	Sample Type	Majors (wt.percent)			Trace Elements (ppm)									
		Fe	Zn	Pb	Cu	Sb	Ag	Cd	Ni	Cr	Co	Mn	Ge	
1	A1 Altered Sphalerite	2.0	26.0	30.8	564	-	950	603	40	34	335	575	-	
2	B2 Less altered Sphalerite	2.1	52.2	0.0	1447	-	158	2190	40	30	775	130	-	
3	D4 Less altered Sphalerite	9.0	49.7	0.0	726	-	109	1340	20	20	820	2795	-	
4	F6 White Mine Dump powder	1.4	30.5	8.0	562	-	475	683	52	25	335	750	-	
5	7663 'S' Galena	-	0.1	65.0	200	900	1900	-	-	-	-	-	-	
6	7665 'M' Galena	-	0.5	70.0	400	2200	500	-	-	-	-	-	-	
7	7660 'S' Sphalerite	2.5	58.62	2.0	-	-	200	2380	-	-	-	-	-	
8	7664 'M' Sphalerite	2.6	59.0	1.0	-	-	180	2000	-	-	-	-	-	
9	7659 'P' Occurrence Galena	-	0.0	73.0	200	500	600	-	-	-	-	-	-	
10	7651 'O' Occurrence Sphalerite	3.5	58.3	0.8	-	-	-	2600	-	-	-	-	-	
11	7652 'O' Occurrence Galena	-	0.5	70.0	500	300	450	-	-	-	-	-	-	

Notes: Serial Nos. 1-4 = Mean values of analyses from Adiyku Brown and Ogezi (this vol.)  
 Serial Nos. 5-11 = From NECCO Consultancy Report for Nigerian Mining Corporation, Jos (1976) on Geological Investigations of Lead-Zinc mineralisation in the areas of Abakaliki and Zurak, Nigeria (p137 - p138)(p137-p138)

- = Not determined or measured  
 n.d. = Not detected.

in developing countries short of foreign exchange, technical personnel and equipment.

In an economic and environmental study of the lead-zinc mining area of Zurak, south-eastern Plateau State, (Ogezi, 1984; and in Prep) showed that apart from the major economic significance of the Zurak mine, which has been closed for about 45 years since 1937, because of the metallurgical and infrastructural problems and depressed metal markets, the dumps are a major source of pollution to surface and underground water: as well as plant and animal life in the area (Table 4).

The mineralogy of the sphalerite - and galena-rich sulphides from the Zurak Pb-Zn mine consists of galena ( $PbS$ ), sphalerite ( $ZnS$ ), marcasite and pyrite ( $FeS_2$ ), and minor bravovite ( $Ni, Fe$ ) $S_2$ , chalcocopyrite ( $CuFeS_2$ ), bournonite ( $CuPbSbS_3$ ) and bravoite boulangerite ( $Pb_5 Sb_4 S_{11}$ ). White, highly corrosive mine dump powder is scattered over the mining area and has resulted from the partial decomposition of the sulphides to altered sphalerite, galena, chalcocopyrite, pyrrhotite, graphite and some sulphates. (A. Mücke, pers. comm., 1984). Some of these minerals are reported for the first time in Nigerian sulphide ores.

The Zurak mine pit water used for domestic purposes by the inhabitants of New Zurak is green in colour, has typical sulphide smell, and largely devoid of animal life and generally contains much higher concentrations of Ca, Mg, Zn, Pb and Cd than Jos tap water, streams and reservoir waters in the Jos area and the Zurak

area pond water not polluted by human activity (Table 5).

A multidisciplinary study of the area is planned to assess the total effects of mining and mineral processing on the small rural community of New Zurak Village and on plant and animal life in the area.

Apart from the investigations reported above, a preliminary investigation of the effects of human activity on the chemical quality of surface water has been carried out on samples of water from the Lamingo (Liberty) Dam and along a tributary stream to the Delimi from Lamingo Dam to the bridge on the Ring Road, near the Gospel Faith Church, Jarawa, Jos (Table 5). Apart from Mn, the dam water contains lower or similar amounts of Ca, Mg, Na, Zn and Fe. Stagnant waters with significant biological activity and areas along the streams defecated or used for laundry contained significantly higher Ca, Fe, and Na (Table 5). This type of study is being conducted throughout Jos area in streams and wells, but the necessity for integrated bacteriological studies cannot be overemphasised.

#### 4. CONCLUSIONS AND RECOMMENDATIONS

The results of the preliminary stages of this research project reported above has, apart from providing invaluable students' training, brought out clearly a number of problems and suggested a number of lines that could be followed in the next stage of the research:

- 1) Standardised instructions covering sample collection, preparation, treatment, storage and analysis procedures

Table 5: Concentration in parts per million (ppm) of Trace Elements in Liberty Dam Reservoir and Upper Delimi Tributary Waters (LD) near Jos, Jos City Tap Water, Zurak Mine Pit Water and Sabon Gari Pond Water, from unmineralised area near Zurak.

Field Sample No.	Sample Type and Locality	Trace Elements (ppm)								
		Ca	Mg	Mn	Na	Zn	Fe	Pb	Cd	Cu
1 LD1	Eastern bank. Liberty Dam Reservoir	1.00	0.30	38.0	2.12	0.01	0.2	-	-	-
2 LD2	Small northern overflow west of Sherre Hills	0.91	0.28	41.6	1.79	0.01	0.1	-	-	-
3 LD3	Small southern overflow west of Sherre Hills	8.40	0.34	11.5	2.10	0.02	0.1	-	-	-
4 LD4	Stagnant confluence of LD2 and LD3	3.18	0.37	15.0	2.10	0.02	0.1	-	-	-
5 LD5	Stream "laundry" ca 50m	4.95	0.54	26.1	2.60	0.01	1.2	-	-	-
6 LD6		0.61	0.09	26.4	3.20	0.01	1.3	-	-	-
7 LD7	Small spring to stream from NW	1.59	0.35	2.7	5.10	0.03	2.0	-	-	-
8 LD8	Water west (downstream) of Ring Road bridge near Gospel Faith Church, Jarawa, Jos	1.46	0.32	1.2	5.03	0.02	2.1	-	-	-
9 J2A	Jos Tap Water, Unijos	n.d.	2	n.d.	-	n.d.	n.d.	n.d.	n.d.	n.d.
10 ZW1	Zurak Mine Pit Water, south pit	2	111	3	-	14	n.d.	3	1	n.d.
11 ZW2	" , north pit	30	135	n.d.	-	98	n.d.	2	4	n.d.
12 ZW3	Sabon Gari, Zurak Pond Water	2	27	n.d.	-	n.d.	1	n.d.	n.d.	n.d.

- = not determined or measured

n.d. = not detected.

should be prepared, together with details of field determinations and observations, preferably in a card form.

- ii) The forms of ions to be determined and their significance should be looked into rather than the total ions being determined as at present since the ions which move with water are those which are either soluble in water or those that occur as components of stable suspensions as cations and anions, uncharged atoms, organic complexes, as colloids and suspended matter, and as ions adsorbed on suspended matter. Their toxicity and effects depend on their speciation or the forms and nature of the atoms.
- iii) In view of the relatively low concentrations of most trace elements in the water samples analysed and of (ii) above, preconcentration of atoms by ion exchange, solvent extraction, etc., or the use of more sensitive methods and equipment should be considered. Seasonal variations should also be checked in all waters.
- iv) With the acquisition of more basic equipment and systematic reagents, more sample collection, preparation, digestion and analysis would be embarked upon, especially for stream sediment, water and rock samples.

The following tentative scientific and environmental recommendations and conclusions could be reached from the results of these preliminary investigations:

- v) Most of the alluvial stream sediment deposits are derived

from nearby sources and, often, contain minerals with high radioactivity, such as monazite, thorite and zircon. These minerals may have harmful effects when used for houses.

- vi) Although some elements may be leached from mining, mineral processing and mine dumps, etc., their concentrations in sediments, waters, soils, plants and animals depend on their solubility and speciation as well as the season. Some lake water not suitable for domestic and industrial uses could be investigated for recreational and agricultural purposes.
- vii) Dilute nitric acid is adequate for the extraction of most major and trace elements from sulphides and stream sediments.
- viii) The clay-size or finest fractions of stream sediments generally had the highest concentration of the environmentally significant trace elements, and most variations could be explained by geological and human factors.
- ix) The well waters generally contained less trace elements and are more suitable for domestic and industrial purposes than stream waters, but seasonal variations are to be investigated in both waters. In general, their trace element contents reflect season, and geological and human factors. Bacteriological tests are necessary on the waters to deduce their optimum uses.

- x) Some sulphide and other ore minerals contain significant amounts of by-product trace elements which could make their mining, processing and extraction more economic than as of now.

#### ACKNOWLEDGEMENTS

We gratefully acknowledge the material, moral and logistic support of our colleagues in the Department of Geology and Mining and the faculty of Natural Sciences, University of Jos, especially those of C. Ejike, J. Pastor, I. Pospisil, D. Anadu, and J. Chidobem. Invaluable support and encouragement was also provided by the Japanese International Cooperation Agency (JICA) and Dr. H. Takahashi, the University of Jos, Faculty of Natural Sciences Research Grants Committee, and the Nigerian Mining Corporation, Jos. The VW Foundation of the Federal Republic of Germany, and the Technical University of Berlin provided logistic and laboratory facilities through Dr. G. Matheis, and the University of Göttingen provided assistance in mineralogy and ore microscopy through Professor A. Mücke. The input of students, especially S. Agada, B. Ekpo, K. Hamza, M. Manuel, R. Ochonogor, B. Onyeji and F. Okeke is gratefully acknowledged, Messrs and T.U. Onyemachi, M. Abah and Mrs K. J. Yakubu typed various versions of the manuscript.

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The last report of JICA-UNIJOS Research Project on Endemic Goitre which covered the period for second half of 1983 had outlined the overall objectives of the study, the methodology employed for survey, classification of goitres phasing of the project, plan of action for 1984 etc.

The survey of endemic goitre among pre-adolescent children in schools of Bassa L.G.A. of Plateau State was carried out. The following work has been completed up to the 1st half of 1984, ending June 1984.

1. A total of 1104 school children in the Government Secondary Schools of Binchi, Zangun, Jengre and Miango have been surveyed. (See enclosed map). The results of survey have been outlined in Tables 1 - 5 in Appendix A.
2. A total of 81 blood samples from the school children, both with and without thyroid enlargement, have been collected by the department of Chemical Pathology for Thyroid Function Tests. These samples have been frozen and stored for analysis at a later date.
3. The village leaders in Bassa L.G.A. (the Chief of Bassa and Heads of the villages) have been contacted to elicit their support and co-operation for the purpose of extending the goitre survey work to the general population in the area. The response have been most encouraging and support for the survey work for endemic goitre in villages has been promised.
4. A total of 19 drinking water samples from as many villages in the area has been collected for physico-chemical examination of water samples as well as determination of iodine (iodide ions by ion chromatography analyser) in the drinking water. Some samples have already been analysed and the results on the remaining are expected to be available soon.

Constraints in Carrying out the work:

According to the time schedule outlined in the initial report of the project a total of 6000 students were expected to be covered up by the survey team by the end of June 1984 as part of phase I of the study. Due to non-availability of the vehicle for survey work from April '84 onwards no survey could be carried out during the last quarter and the target of coverage of 6000 school children has not been achieved. Meanwhile the

schools have closed down for long vacation till September 1984. The equipment and chemicals needed by Chemical Pathology Department for thyroid function tests have not yet arrived and work has not been possible to be started.

Plan of action during Second Half of 1984:-

On the basis of experience gained till date the study team feels that within the multiple pressures of work on the academic staff involved in the study it would not be possible to cover the entire Plateau State under the survey work. This would be particularly so in view of the distances of different L.G.A. from Jos. The resource position for carrying out the study is also not expected to improve dramatically in near future. In view of the above, it is therefore necessary to modify the targets for achievement mentioned in earlier reports. The study team proposes the following changes -

- (i) The original target of coverage of 6000 school children for survey will be reduced to about 3000.
- (ii) A total of about 2000 school children yet to be covered up will be surveyed as soon as the schools open after long vacation.
- (iii) However, the remaining children will be covered up in two L.G.A. other than Bassa, the selection being made on the basis of their topographical location and physical features. The department of Land Survey of Plateau State has been approached and they promised to make available necessary data to help select the L.G.A. for the study.
- (iv) The community response for the survey of general population for endemic goitre has been very encouraging in Bassa L.G.A. This will be fully exploited to carry out the study of the prevalence of endemic goitre and the related clinical entities viz. cretinisms, hypothyroidism etc. Blood samples for study of thyroid function test will also be collected.
- (v) Water and soil samples from the area will be collected and subjected to iodine estimation. Thus a composite data on prevalence, clinical picture, thyroid function and iodine availability for at least one L.G.A. (Bassa) will be collected by the end of 1984, thus devetailing phase I and phase II of the study.