



Collaborating Organizations

Ministry of National Health Services, Regulation and Coordination

Islamabad – Pakistan

Pakistan Health Research Council

Islamabad – Pakistan

Diabetic Association of Pakistan; WHO Collaborating Centre

Karachi – Pakistan

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University

Karachi – Pakistan

ISBN: 978-969-499-009-5

2nd National Diabetes Survey of Pakistan (NDSP) 2016-2017

Published: 2018

This report describes the findings of the 2nd NDSP 2016-2017 conducted under the authority of the Ministry of National Health Services, Regulation and Coordination and implemented by Pakistan Health Research Council, Diabetic Association of Pakistan and Baqai Institute of Diabetology and Endocrinology. The document may be freely reviewed, abstracted, reproduced and translated, in part or in whole, but is not for sale or use in conjunction with commercial purposes.

Suggested citation as follow:

2nd National Diabetes Survey of Pakistan 2016-2017, Page (.....).

Authors

Prof. Abdul Basit

Baqai Institute of Diabetology and Endocrinology, Karachi - Pakistan

Dr. Asher Fawwad

Baqai Institute of Diabetology and Endocrinology, Karachi - Pakistan

Dr. Huma Qureshi

Pakistan Health Research Council, Islamabad – Pakistan

Dr. Muhammad Arif Nadeem Saqib

Pakistan Health Research Council, Islamabad – Pakistan

Mr. Ibrar Rafique

Pakistan Health Research Council, Islamabad – Pakistan

Prof. AS Shera

Diabetic Association of Pakistan and WHO Collaborating Centre, Karachi - Pakistan

Contributors

NDSP Contributors (with surnames in alphabetical order);

Dr. Mujeeb Ur Rehman Abro

Shaheed Mohtarma Benazir Bhutto Medical University, Larkana – Sindh

Dr. Khawaja Ishfaq Ahmed

Pakistan Institute of Medical Sciences, Islamabad – Punjab

Dr. Khurshid Ahmed

Zahid Medical Centre, Hub – Baluchistan

Dr. Waqar uddin Ahmed

Deputy Director (HSR), Pakistan Health Research Council

Dr. Sobia Sabir Ali

Lady Reading Hospital, Peshawar - Khyber Pakhtunkhwa

Prof. Ahmed Bilal

Faisalabad Medical College, Faisalabad – Punjab

Dr. Anam Butt

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University, Karachi – Sindh

Prof. Bikha Ram Devrajani

Sindh Institute of Endocrinology and Diabetes, Liaquat University of Medical and Health Sciences, Jamshoro - Sindh

Mr. Ejaz Hayder

Pakistan Health Research Council, Karachi - Sindh

Dr. Yasir Humayun

DHO Office, Mansehra - Khyber Pakhtunkhwa

Mrs. Rabia Irshad

Pakistan Health Research Council, Karachi – Sindh

Dr. Riasat Ali Khan

Canada Medical Group Hospital, Defence, Karachi – Sindh

Dr. Asima Khan

Sindh Government Hospital, New Karachi, Karachi - Sindh

Dr. Aamir Akram Khowaja

Sindh Government Qatar Hospital, Karachi – Sindh

Dr. Raheela Khowaja

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University, Karachi – Sindh

Prof. Qazi Masroor

Quaid-e-Azam Medical College, Bahawalpur – Punjab

Dr. Magsood Mehmood

Fatmatu Zahra Hospital, Gujranwala - Punjab

Mr. Hassan Moin

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University, Karachi - Sindh

Dr. Wasif Noor

Akhuwat Health Services Diabetes Centre, Lahore – Punjab

Dr. Tahir Rasool

Akhuwat Health Services Diabetes Centre, Lahore – Punjab

Mrs. Rubina Sabir

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University, Karachi - Sindh

Dr. Pir Alam Said

DHQ, Sawabi - Khyber Pakhtunkhwa

Prof. Abrar Shaikh

Ghulam Muhammad Mahar Medical College, Sukkur – Sindh

Mr. Bilal Tahir

Baqai Institute of Diabetology and Endocrinology, Baqai Medical University, Karachi – Sindh

Prof. Salma Tanveer

Nishter Medical University, Multan – Punjab

Prof. Bilal Bin Younus

Sakeena Institute of Diabetes & Endocrine Research, Lahore – Punjab

Prof. Jamal Zafar

Pakistan Institute of Medical Sciences, Islamabad - Punjab

Foreword

Diabetes is a major public health issue resulting in morbidity and mortality worldwide. The complications including stroke, cardiovascular diseases, renal failure etc resulting from diabetes are major cause of premature deaths globally. It is estimated that a person died every 6th second due to diabetic complications. Major risk factors for diabetes are physical inactivity, obesity and unhealthy diet. According to International Diabetes Federation, about 425 million people are living with diabetes and more than 70% cases are in low and middle-income countries. Diabetes is common in South East Asia constituting 19.3% of total world's diabetic population. It is estimated that in Pakistan, about 7.5 million people have diabetes and this number is predicted to increase to 16.1 million by the year 2045. About 1.8% of the global GDP (1.31 trillion USD) is being spent on the management of diabetes.

First National Diabetes Survey of Pakistan was conducted in 1994-98. Keeping in view the changing demography of the country, a need was felt to have renewed data on diabetes in term of its prevalence in different age groups, gender and urban rural setting. It aimed to help policy makers in making informed policies on the prevention and control of this deadly disease.

To achieve above goal, Pakistan Health Research Council (PHRC) together with Diabetic Association of Pakistan (DAP) and Baqai Institute of Diabetology and Endocrinology (BIDE) planned 2nd National Diabetes Survey of Pakistan (NDSP) 2016-17 under the auspices of the Ministry of National Health Services, Regulation and Coordination (NHSRC). The Survey was conducted in all four Provinces of Pakistan i.e. Punjab, Sindh, Khyber Pakhtunkhwa and Baluchistan to provide national level data on the prevalence of diabetes and its risk factors in the country.

I congratulate PHRC, DAP and BIDE; for successfully completing the 2nd NDSP 2016-2017 and bringing out report which I believe would be highly beneficial to policy makers in planning for prevention and control of diabetes in Pakistan. I am confident that academicians, policy makers, healthcare experts and all relevant stakeholders would make optimal use of this report.

MUHAMMAD ALI SHAHZADA

Additional Secretary Ministry of National Health Service, Regulation and Coordination

Source: International Diabetes Federation (IDF) Atlas 8th Edition 2017



Prof. Nam H. Cho
President
International Diabetes
Federation

As the President of International Diabetes Federation (IDF), it gives me great pleasure to express my feelings at the accomplishment of "National Diabetes Survey of Pakistan (NDSP) 2016–2017"

The mission of the International Diabetes Federation (IDF) is to promote diabetes care, prevention, and cure worldwide with an estimated number of people with diabetes worldwide close to 425 million and projections indicating well over 629 million in 2045. The current diabetes epidemics is of great concern to those affected, their families and all the concerned governments and social security systems.

This survey will help to understand the current epidemic situation in one out of the seven IDF Regions, namely the Middle East and North Africa (MENA) Region and will contribute to identify solutions in the Region with special emphasis to Pakistani population were the number of people affected is particularly very high.

I on behalf of IDF, congratulate all the stakeholders in this regard and I am confident that this survey will help in early identification and effective management. Moreover, nationwide preventive strategies will be focused to combat the situation.



Prof. Akhtar HussainPresident
Diabetes in Asia Study Group

I feel honored and privileged to write a message at the successful completion of "National Diabetes Survey of Pakistan 2016–2017". The survey comprehensively looks at the prevalence of diabetes, hypertension, obesity and dyslipidemia along with their associated risk factors in all provinces of the country.

Globally, diabetes is a major cause of morbidity and mortality. A progressive rise of diabetes in the productive age group specifically in Asia foretells a serious public health issue in this part of the world. There is enough evidence on preventability of the disease by applying sets of effective preventive and curative measures.

Pakistan had an estimated 7.5 million people with diabetes and this number was predicted to be increased up to 16.1 million by the year 2045 according to an International Diabetes Federation. This number was extrapolated on the basis of previous National Diabetes Survey of Pakistan conducted in 1994-1999. I congratulate, Ministry of National Health Services, Regulation and Coordination, Pakistan Health Research Council, Diabetic Association of Pakistan, Baqai Institute of Diabetology and Endocrinology and the whole NDSP team on the successful completion of this survey.

These figures are alarming and emphasizing the urgent need for National strategies for early diagnosis, comprehensive management along with cost effective preventive measures. I wish them all the best in this regard.



Dr. S. Abbas RazaPresident

South Asian Federation of

Endocrine Societies

On behalf of the South Asian Federation of endocrine Societies (SAFES), it gives me an immense pleasure to write a message for the 2nd National Diabetes Survey of Pakistan (NDSP 2016 – 2017), conducted in all four provinces of Pakistan simultaneously.

The mission of the South Asian Federation of endocrine Societies (SAFES) is to focus the region-specific health related issues, particularly diabetes and other endocrine disorders and to facilitate their standardized treatment as per the available region-specific evidences and guidelines.

At present, nearly half a billion people live with diabetes worldwide. Low and middle-income countries share almost 80% of the diabetes burden. Rapid urbanization, unhealthy diets and increasingly sedentary lifestyles have resulted in previously unheard higher rates of diabetes. Diabetes can be successfully managed and complications can be prevented especially when detected early. Even better, by making lifestyle changes the risk of developing diabetes can be reduced markedly. Many countries still lack prevalence studies and many populations are not systematically surveyed. Still, more multi-dimensional and multi-sectoral research is needed to tackle the diabetes epidemic.

NDSP was well planned community-based survey providing current estimates of people suffering from diabetes in different strata of the Pakistani population. I would like to express my deepest gratitude to Ministry of National Health Services, Regulation and Coordination, Pakistan Health Research Council, Diabetic Association of Pakistan, Baqai Institute of Diabetology and Endocrinology and the whole NDSP team along with it's participants on conducting and successful completion of this National Survey and at the same time I on behalf of SAFES would like to extend our unconditional and full support to combat this epidemic support in Pakistan.



Prof. Ali JawaPresident
Pakistan Endocrine Society

It gives me great pleasure to write this message on behalf of Pakistan Endocrine Society (PES) for the 2nd National Diabetes Survey of Pakistan (NDSP) 2016–2017. Pakistan Endocrine Society promotes educational and research activities in Endocrinology, Diabetes and Metabolism and has always supported the initiatives taken to address the National level health related issues, specifically in the field of diabetology and endocrinology.

Diabetes is a global health crisis and its chronic nature can lead to its debilitating complications. Globally, Governments are struggling to meet the cost of diabetes care and the financial burden will continue to expand due to the growing number of people developing diabetes. International and National researchers are convinced that prevention at all levels is the only solution to halt this rising epidemic.

I take this opportunity to extend my heartiest congratulations to all the contributors and stakeholders of NDSP on this successful accomplishment.

Keeping the scale of the problem in mind, we certainly need ongoing research in the field of Diabetes and Endocrinology. It indeed is my sincerest hope that scientists and researchers redouble their efforts to refine our understanding of diabetes epidemic especially in Pakistan not only to benefit local population but ultimately the greater humanity as a whole.

Acknowledgements

The 2ndNational Diabetes Survey of Pakistan (NDSP) 2016 - 2017 was successfully completed due to the efforts and involvement of numerous organizations and individuals at different stages of the survey. We would like to thank everyone who participated and make this survey a success.

First of all, we are grateful to the Ministry of National Health Services, Regulation and Coordination (NHSRC) in Pakistan for its support. We would like to extend Special thanks to Capt® Zahid Saeed, the Secretary Ministry of National Health Services Regulation and Coordination. We would also like to thank Mr. Naveed Kamran Baloch and Mr. Muhammad Ayub Sheikh, former Secretaries, Ministry of NHSRC and Dr. Assad Hafeez, Director General (Health) for their valuable and constant support throughout the survey.

We wish to express our special thanks to Pakistan Health Research Council and Diabetic Association of Pakistan for their support and facilitation in this survey. We sincerely appreciate the efforts of Ms. Rabia Irshad and Mr. Ejaz Haider at PHRC Research Centre JPMC, Karachi for performing laboratory tests for the survey. We also pay special thanks to Dr. Faiza Bashir, Ms. Saima Naz and Mrs.Sumera Abid for their contribution and efforts.

In addition, the untiring efforts of survey Coordinators and field supervisors are highly acknowledged. We appreciate the efforts of the monitoring teams who despite the difficult field conditions visited the field areas and monitored the survey activities.

We acknowledge the support of Research and Laboratory department of Baqai Institute of Diabetology and Endocrinology (BIDE), Karachi for data management. We are extremely grateful to the Seventeen teaching hospitals and/or diabetes centers participated in the 2ndNDSP 2016 - 2017.

Last but not the least we would like to express our sincere thanks and gratitude to the Survey Coordinators, survey teams, data entry teams and above all the household members/community who participated in the survey.

Prof. Abdul Basit

Chair, IDF - MENA Region
Director,
Baqai Institute of Diabetology and Endocrinology,
Baqai Medical University, Karachi

Contents

| \mathbf{E} | XECUTIVE SUMMARY | 21 |
|--------------|---------------------------------------|----|
| F | ACT SHEET | 24 |
| | Fact sheet; Gender | 25 |
| | Fact Sheet; Provinces | 25 |
| Η | IGHLIGHTS | 27 |
| 1. | INTRODUCTION | 31 |
| 2. | MATERIALS AND METHODS | 35 |
| | 2.1 Ethical approval | 35 |
| | 2.2 Study area and population | 35 |
| | 2.3 Inclusion and Exclusion criteria: | 35 |
| | 2.4 Sample Size | 35 |
| | 2.4.1 Sample size calculation | 35 |
| | 2.4.2 Sampling technique | 36 |
| | 2.4.3 Sample size distribution. | 36 |
| | 2.5 Survey team | 36 |
| | 2.6 Data collection method. | 37 |
| | 2.7 Measurements | 37 |
| | 2.8 Blood Sampling and Testing | 38 |
| | 2.9 Definitions | 38 |
| | 2.9.1 Diabetes | 38 |
| | 2.9.2 Hypertension | 38 |
| | 2.9.3 Body Mass Index | 38 |
| | 2.9.4 Dyslipidemia | 39 |
| | 2.10 Data Analysis | 39 |
| 3. | RESULTS | 45 |
| | 3.1 Demographic characteristics | 45 |
| | 3.1.1 Age and gender | 47 |
| | 3.1.2 Education Level | |
| | 3.1.3 Marital Status | 48 |
| | 3.1.4 Tobacco Addiction | 48 |
| | 3.2 Measurements | 49 |
| | | |

| 3.2.1 Body Mass Index (BMI) | 49 |
|--|----|
| 3.2.2 Hypertension | 49 |
| 3.3 Diabetes | 51 |
| 3.3.1 Normal glucose tolerance | 51 |
| 3.3.2 Pre-diabetes | 51 |
| 3.3.3 Prevalence of Newly diagnosed diabetes mellitus | 51 |
| 3.3.4 Prevalence of known diabetes | 51 |
| 3.3.5 Prevalence of diabetes mellitus | 51 |
| 3.3.6 Prevalence of diabetes and pre-diabetes, by Province and area of Pakistan | 51 |
| 3.3.7 Age, gender and area specific prevalence of pre-diabetes and diabetes in four provinces of Pakistan: | 51 |
| 3.4 Dyslipidemia | 61 |
| 3.4.1 Raised cholesterol | 61 |
| 3.4.2 Hyper triglyceride | 61 |
| 3.4.3 High low-density lipoprotein | 61 |
| 3.4.4 Low high-density lipoprotein | 61 |
| 3. 5. Risk Factor Analysis | 65 |
| 3. 5.1 Risk factor of diabetes, dyslipidemia, hypertension and obesity | 65 |
| 4. CONCLUSION | 69 |
| 5. RECOMMENDATIONS | 69 |
| 6. REFERENCES | 71 |
| 7. APPENDICES | 74 |
| Appendix I | 74 |
| Cluster (district) and sub-clusters (tehsils/towns) distribution | 74 |
| Appendix II | 75 |
| Informed consent (English) | 75 |
| Informed consent (Urdu) | 77 |
| Appendix III | 79 |
| Data Collection Form (English) | 79 |
| Data Collection Form (Urdu) | 81 |
| Appendix IV | 83 |
| Specimen Collection and Handling | 83 |

| Appendix V | 86 |
|--------------------|----|
| Testing Procedures | 86 |
| Appendix VI | |
| Ethical Clearance | |
| Appendix VII | 88 |
| Gallery | |
| J | |

List of Tables

| Table 1: Baseline characteristics of the study participants | 31 |
|--|----|
| Table 2: Age and gender distribution of respondents | 32 |
| Table 3: Educational level of respondents | 32 |
| Table 4: Marital status of the respondents | 33 |
| Table 5: Tobacco addiction among respondents | 33 |
| Table 6: BMI classification of the respondents by Asian cutoff | 34 |
| Table 7: BMI classification of the respondents by WHO criteria | 35 |
| Table 8: Hypertension status of the respondents | 35 |
| Table 9: Prevalence of normal glucose tolerance | 37 |
| Table 10: Prevalence of pre-diabetes | 38 |
| Table 11: Prevalence of newly diagnosed Diabetes Mellitus | 39 |
| Table 12: Prevalence of known diabetes | 40 |
| Table 13: Prevalence of Diabetes Mellitus | 41 |
| Table 14: Prevalence of diabetes and pre-diabetes; by province and areas of Pakistan | 42 |
| Table 15: Hypercholesterolemia status of the respondents | 46 |
| Table 16: Hypertriglyceridemia status of the respondents | 47 |
| Table 17: Status of high low-density lipoprotein of the respondents | 47 |
| Table 18: Low high-density lipoprotein of the respondents | 47 |
| Table 19: Risk factor of Diabetes, Dyslipidemia, Hypertension and Obesity | 48 |

List of Figures

| Figure 1: Step by step approach for the 2 nd NDSP 2016 - 2017 | 27 |
|---|----|
| Figure 2: Structure of survey team | 28 |
| Figure 3: Flow of 2 nd NDSP 2016 - 2017 team | 29 |
| Figure 4: Age-stratified prevalence of diabetes among men and women with urban and rural | |
| distribution | 45 |
| Figure 5: Age-stratified prevalence of pre-diabetes among men and women with urban and rura | al |
| distribution | 45 |

Diabetes – A growing national crisis





EXECUTIVE SUMMARY

Diabetes mellitus is major public health issue of 21st century, especially for lower middle-income countries (LMIC). According to estimation, number of people suffering from diabetes is anticipated to be increase to 629 million people worldwide by 2045 as compared to 425 million people in 2015. The complications resulting from diabetes are major cause of premature deaths round the globe and it is estimated that a person died every 6th second due to diabetic complications. In 2015 only, approximately 5 million deaths were reported from developing countries.

Pakistan has an estimated 7.5 million people with diabetes and this number is predicted to increase to 16.1 million by the year 2045. This number is extrapolated on the basis of surveys done by the Diabetic Association of Pakistan (DAP) and World Health Organization (WHO) collaborating center in 1994 – 1998 (1st National Diabetes survey of Pakistan, 1st NDSP). Hence, a repeat survey is needed to estimate the recent numbers of people suffering from type 2 diabetes so as to take a step forward in developing strategies for cost effective management and primary prevention. Therefore, 2nd NDSP 2016 – 2017 was designed to obtain the recent number of type 2 diabetics in urban and rural areas of Pakistan.

The survey was conducted using three stage sampling technique. In the first stage, population was stratified into urban and rural areas. In the second stage, clusters were randomly selected using probability proportional to size (PPS) technique. In the third stage, sub-clusters based on tehsils/towns were selected randomly.

A total of 27 clusters were selected from all four Provinces of Pakistan and 46 sub-clusters (21 urban and 25 rural). The sample size was distributed into clusters and sub-clusters based on probability proportionate to size technique so that cluster with larger population size will have larger sample size.

Data was collected in two phases. In the 1st phase, a pre-survey visit was made for the selection of houses following systematic sampling technique. The first house in the lane was selected randomly and afterwards every tenth house was identified. The selected household members were requested to come after an overnight fast (at least 8 hours) to the camp on the specific day. Participants were enrolled after taking informed consent. The fasting sample and sample for

glucose tolerance test (two hours after giving 75 grams of anhydrous glucose load) were taken. Height, weight, waist circumference, waist-hip ratio and blood pressure were measured as per the standardized procedures.

Plasma glucose, lipid profile (total cholesterol, high density lipoprotein cholesterol, low density lipoprotein cholesterol) and HbA1c was performed using standard biochemical methods. Fasting and random (2-hour post 75gm glucose load) plasma glucose levels were performed at the site while HbA1c and lipid profile was done in laboratory.

A total of 10834 persons were enrolled (43.9% males, 56.1% females). More than half (53.5%) had at least primary level education. Almost 30.2% had positive family history of diabetes.

According to Asian BMI cut off values, 27.7% respondents had normal (BMI <23), 14.2 % were overweight (BMI 23-24.9) and 58.1% were obese (BMI \geq 25). Mean BMI of participants was $27.2 \pm 6.0 \text{ kg/m}^2$ which were almost similar in all Provinces.

Overall, mean systolic and diastolic blood pressure was 126 ± 19 mmHg and 84 ± 14 mm/Hg respectively. Highest mean blood pressure was seen in Baluchistan and lowest in Khyber-Pakhtunkhwa. Around half of the population (46.2%) had hypertension. The prevalence of hypertension increased with age and was highest (65.7%) in age group 60 years or above.

As per oral glucose tolerance test (OGTT) criteria, overall age adjusted weighted prevalence of diabetes was 26.3%. Among them, 19.2% were known diabetic cases while 7.1% were diagnosed during this survey. Prevalence of diabetes in urban and rural areas was 28.3% and 25.3%, respectively. Highest prevalence of diabetes was observed in Sindh (32.3%) followed by Punjab (30.2%), Baluchistan (29.5%) and KPK (13.2%). Prevalence of pre-diabetes was 14.4%, urban and rural distribution was 15.5% and 13.9%, respectively.

Overall glycemic dysregulation (diabetes, plus pre-diabetes) was 43.8% and 39.2% in urban and rural areas, respectively. Prevalence of pre-diabetes and newly diagnosed diabetes was higher in Baluchistan as compared to other provinces.

Urban women showed significantly higher prevalence of diabetes than rural women above the age of 40 years (p<0.05). On the other hand, urban men in the age group 30-39 years showed significantly (p<0.05) lower prevalence of diabetes than rural men. Rural men showed significantly higher prevalence of pre-diabetes then urban men for the age group of 40-49 years

while for women, significant difference was seen in urban as compared to rural population for the age group of 30-39 years. Age (greater than or equal to 43 years), family history of diabetes, hypertension, obesity and dyslipidemia were significant risk factors for diabetes ($p \le 0.0001$).

Overall, 60.7% population had normal cholesterol levels. Deranged cholesterol was mostly seen in people aged between 50-59 years followed by those aged 40-49 years. High triglycerides were seen in 48.9% respondents and this was mostly seen in individuals who were between 50-59 years of age. Overall 39.7% respondents had high levels of low density lipoprotein. Majority of the population had low values of high density lipoprotein (83.5% men, 90% women).

FACT SHEET

The 2ndNational Diabetes Survey of Pakistan (NDSP) 2016-2017 was a population-based survey of adults aged 20 years above which was carried out from February 2016 to August 2017. A three-stage sampling technique was used to produce representative data for that age range in Pakistan. A total of 10834 adults participated in the Survey and information were collected on demographics and physical measurements such as height, weight and blood pressure as well as testing of blood samples for fasting and random(2-hours oral glucose tolerance test) plasma blood glucose levels, HbA1c and lipid profile was also done. A repeat survey is planned after 05 years if funds permit.

| Fact sheet; Gender | | | | | | |
|---|-------------------------------|---------------------------------|----------------------|--|--|--|
| Descriptions | Both Sexes (%) (95% CI) | Males (%) (95% CI) | Females (%) (95% CI) | | | |
| Normal glucose tolerance | | | | | | |
| Oral glucose tolerance test (OGTT) | 60.0 | 61.6 | 58.9 | | | |
| | (59.0-60.9) | (60.2-62.9) | (57.6-60.1) | | | |
| HbA1c | 64.5 | 63.4 | 65.3 | | | |
| | (63.6-65.4) | (62.0-64.7) | (64.1-66.5) | | | |
| Pre-diabetes | | | | | | |
| Oral glucose tolerance test (OGTT) | 14.4 | 12.8 | 15.6 | | | |
| | (13.7-15.0) | (11.8-13.7) | (14.6-16.5) | | | |
| HbA1c | 5.5 | 5.1 | 5.9 | | | |
| | (5.0-5.9) | (4.4-5.7) | (5.3-6.4) | | | |
| Prevalence of newly diagnosed diabetes mellitus | | | | | | |
| Oral glucose tolerance test (OGTT) | 7.1 | 7.3 | 6.9 | | | |
| | (6.6-7.5) | (6.5-8.0) | (6.2-7.5) | | | |
| HbA1c | 7.4 | 8.5 | 6.7 | | | |
| | (6.9-7.8) | (7.7-9.2) | (6.0-7.3) | | | |
| Prevalence of known diabetes | | | | | | |

| Fact sheet; Gender | | | | | | |
|------------------------------------|-------------------------------|--------------------------|----------------------|--|--|--|
| Descriptions | Both Sexes (%) (95% CI) | Males (%) (95% CI) | Females (%) (95% CI) | | | |
| Oral glucose tolerance test (OGTT) | 19.2 | 18.9 | 19.4 | | | |
| | (18.4-19.9) | (17.7-20.0) | (18.4-20.3) | | | |
| HbA1c | 22.4 | 22.9 | 22.1 | | | |
| | (21.6-23.1) | (21.7-24.0) | (21.0-23.1) | | | |
| Prevalence of diabetes mellitus | | | | | | |
| Oral glucose tolerance test (OGTT) | 26.3 | 26.2 | 26.3 | | | |
| | (25.4-27.1) | (24.9-27.4) | (25.1-27.4) | | | |
| HbA1c | 29.9 | 31.5 | 28.8 | | | |
| | (29.0-30.7) | (30.1-32.8) | (27.6-29.9) | | | |
| Lipid profile | | | | | | |
| Hypercholesterolemia | 39.3 | 39.1 | 39.5 | | | |
| | (38.3-40.2) | (37.7-40.4) | (38.2-40.7) | | | |
| High low density lipoprotein | 87.4 | 83.5 | 90 | | | |
| | (86.7-88.0) | (82.4-84.5) | (89.2-90.7) | | | |
| Low high density lipoprotein | 38.7 | 38.9 | 40.4 | | | |
| | (37.7-39.6) | (37.5-40.2) | (39.1-41.6) | | | |
| Hypertriglyceridemia | 48.9 | 54.1 | 45.6 | | | |
| | (48.0-49.8) | (53.0-55.5) | (44.3-46.8) | | | |

| Fact sheet; Province | | | | | | |
|--|---------------------------|--------------------------|--|--------------------------------|--|--|
| Description | Punjab (%) (95% CI) | Sindh (%) (95% CI) | Khyber Pakhtunkhwa (%) (95% CI) | Baluchistan (%) (95% CI) | | |
| Diabetes | | | | | | |
| Known diabetes | 23.7 | 23.6 | 10.2 | 19.1 | | |
| | (22.6-24.7) | (21.9-25.2) | (8.7-11.7) | (15.7-22.4) | | |
| Fasting glucose | 3.3 | 3.6 | 1 | 3.1 | | |
| | (2.8-3.7) | (2.8-4.3) | (0.5-1.5) | (1.6-4.5) | | |
| 2 hours oral glucose | 1.1 | 1.4 | 0.4 | 4.7 | | |
| tolerance test* | (0.8-1.3) | (0.9-1.8) | (0.1-0.7) | (2.9-6.4) | | |
| Both fasting glucose and 2 hours glucose tolerance | 1.1 | 3.7 | 1.6 | 2.6 | | |
| | (0.8-1.3) | (2.9-4.4) | (0.9-2.2) | (1.2-3.9) | | |

| Fact Sheet; Provinces | | | | | | |
|--|---------------------------|--------------------------|--|--------------------------------|--|--|
| Description | Punjab (%) (95% CI) | Sindh (%) (95% CI) | Khyber Pakhtunkhwa (%) (95% CI) | Baluchistan (%) (95% CI) | | |
| Newly diagnosed | 6.5 | 8.7 | 3 | 10.4 | | |
| | (5.8-7.1) | (7.6-9.8) | (2.1-3.8) | (7.8-12.9) | | |
| Total diabetes | 30.2 | 32.3 | 13.2 | 29.5 | | |
| | (29.0-31.3) | (30.4-34.1) | (11.5-14.8) | (25.6-33.3) | | |
| Pre-Diabetes | | | | | | |
| Fasting glucose | 1.8 | 1.8 | 1.1 | 0.4 | | |
| | (1.4-2.1) | (1.2-2.3) | (0.5-1.6) | (0.1-0.9) | | |
| 2 hours glucose tolerance test | 11.2 | 8.4 | 2.9 | 42.4 | | |
| | (10.4-11.9) | (7.3-9.4) | (2.0-3.7) | (38.2-46.5) | | |
| Both fasting glucose and 2 hours glucose tolerance | 2.1 | 1.5 | 0.5 | 8.7 | | |
| | (1.7-2.4) | (1.0-1.9) | (0.1-0.8) | (6.3-11.0) | | |
| Total Pre-diabetes | 15.1 | 11.7 | 4.5 | 51.5 | | |
| | (14.2-15.9) | (10.4-12.9) | (3.4-5.5) | (47.2-55.7) | | |

HIGHLIGHTS

The 2nd National Diabetes Survey of Pakistan (NDSP) 2016-2017 was a population-based survey of adults aged 20 years above which was carried out from February 2016 to August 2017. The data was weighted to give a representative figure for the country. The burden was calculated using recent census findings 2017.

HIGHLIGHTS

PREVALENCE OF DIABETES

- Overall 26.3% were diabetic (27.4 million people aged 20 and above)
- 1 in 4 persons aged 20 and above had diabetes
- Glycemic dysregulation (diabetes, pre-diabetes) was more in urban (43.8%) as compared to rural (39.2%).

MEASUREMENTS:

- 14.2% were overweight (14.4 million people aged 20 and above)
- 43.9% were obese (44.6 million people aged 20 and above)
- 46.2% were hypertensive (47 million people aged 20 and above).

CHOLESTEROL

• Overall 39.3% had hypercholesterolemia (40 million people aged 20 and above)

Globally, about 42.9 million people have diabetes which is expected to rise 628.6 million by 2045.





1. INTRODUCTION

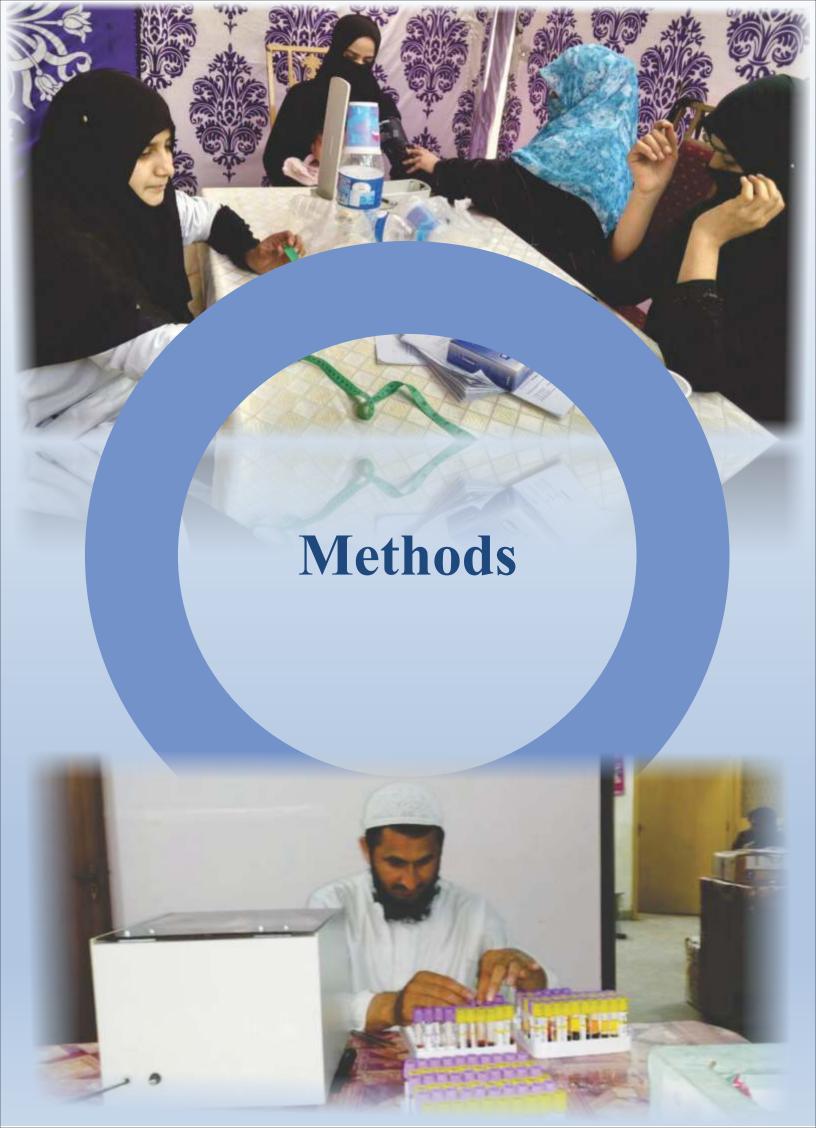
The epidemic of diabetes is an alarming public health issue of 21st century, especially for lower middle-income countries (LMIC).¹ It is predicted that between 2010-2030, the prevalence of diabetes in developing countries would become 67%.² Macrovascular and microvascular complications of diabetes are the major cause of premature deaths globally^{3,4} with one person dying every sixth second due to the complications of diabetes.^{4,5} Approximately, five million (2/3rd) diabetes related deaths were reported from developing countries in 2015.^{4,6}

Globally, according to World Health Organization (WHO), hypertension (13%), tobacco use (9%), diabetes (6%), physical inactivity (6%) and overweight & obesity (5%) are the major risk factors causing morbidity and mortality related to cardiovascular diseases (CVD).^{7,8} It is estimated that about 17 million deaths occur worldwide due to CVD each year, out of which hypertension alone accounts for 9.4 million mortalities.⁹ Although the prevalence of hypertension is rising worldwide¹⁰ but this increase is quite sharp in many LMIC.¹¹ Obesity is now recognized as a disease on its own¹² and according to WHO, obesity is one of the most serious and common, yet not well focused public health problem in both developed and developing countries.¹³ The World Health Statistics report 2012, reported that worldwide, one in six adult is obese and about 2.8 million subjects die each year due to the consequences of obesity.¹⁴

In Pakistan, the 1st National Diabetes Survey of Pakistan (NDSP) 1994-1998 revealed 8.7% prevalence of diabetes. Since then, many small-scale studies have done which showed that the prevalence of diabetes varies between 13.1% to 26.9%. In 2003, the prevalence of hypertension was reported as 20%. In 2016, Pakistan Medical Research Council (PMRC) conducted the NCDs survey in the Punjab and Sindh using WHO's STEPS methodology (no blood testing was done) in which prevalence of hypertension and obesity were 37% and 41.3% resepectively. Few other studies from Pakistan reported obesity between 27% to 49%. Dyslipidemia is a major risk factor for atherosclerosis leading to CVD. Few community-based studies from Pakistan have reported the prevalence of dyslipidemia to be around 30%. State of the prevalence of dyslipidemia to be around 30%.

Keeping in view above, there was a need to do a national level survey to determine the current prevalence of diabetes, hypertension, obesity and dyslipidemia in Pakistan. Therefore, this 2nd National Diabetes Survey of Pakistan (NDSP) 2016 - 2017 was planned which was conducted to

determine the prevalence of diabetes, hypertension, obesity and dyslipidemia in Pakistan including urban and rural populations of all four Provinces.





2. MATERIALS AND METHODS

2.1 Ethical approval

The ethical approval for the 2^{nd} NDSP (2016 – 2017) was obtained from National Bioethics Committee (NBC) of Pakistan.

2.1 Study area and population

This nationwide 2nd NDSP (2016 - 2017) was conducted over a period of eighteen months from February 2016 to August 2017 in all four Provinces of Pakistan (i.e. Punjab, Sindh, Khyber Pakhtunkhwa and Baluchistan). The survey was a joint collaboration of Ministry of National Health Services, Regulation and Coordination (MoNHSRC), Pakistan Health Research Council (PHRC), Diabetic Association of Pakistan (DAP) and Baqai Institute of Diabetology and Endocrinology (BIDE). The available census data of 1998 was obtained from Pakistan Bureau of Statistics (PBS) regarding information about actual population and area distribution of Pakistan³⁰.

2.3 Inclusion and Exclusion criteria:

The inclusion criteria were as follow;

 Healthy individual age 20 and above living in Pakistan (Punjab, Sindh, Khyber Pakhtunkhwa, and Baluchistan) with Pakistani nationality

The exclusion criteria were as follow;

- Pregnant cases
- Subjects who were non-residents of Pakistan or living army locations, Jail inmates and migratory population

2.4 Sample Size

The sample size details are given below.

2.4.1 Sample size calculation

In order to calculate the prevalence at national level, following formula was used to calculate the overall sample size.³¹

Sample size (n) =
$$\frac{Z^2 \times (\text{Expected prevalence}) \times (1 - \text{expected prevalence})}{(\text{Margin of error})^2} \times \text{design effect}$$

An estimated sample size of 10697 subjects was calculated using the above formula with reasonable margin of error and level of confidence ³².

2.4.2 Sampling technique

Multi-stage sampling technique was used. The first stage involved the stratification of population on the basis of urban and rural domains under Punjab, Sindh, Khyber Pakhtunkhwa, Balochistan and Islamabad as defined in census (1998). The second stage constituted of clusters made under each stratum. Clusters was randomly selected using probability proportional to size (PPS) technique and number of cluster to be selected from each province were based on "Rule of thumb" given as under ³².

Number of clusters (k) = (sample size of stratum/
$$2$$
) $^{\circ}$ 0.5

Third stage contained sub-clusters based on tehsils/towns which were selected randomly.

2.4.3 Sample size distribution

A total of 27 clusters were selected from all four Provinces of Pakistan. In the 3rd stage, sub-clusters based on tehsils/towns were selected randomly. There were 46 sub-clusters comprising of 21 tehsils/towns from urban and 25 tehsils/towns from rural areas that were finally selected. The overall sample size was divided into each stratum and then to each cluster according to their population sizes. Sample size was equally distributed to sub-clusters (tehsils/towns)(Appendix1). Therefore, stratum or cluster with larger population size will have larger sample size and greater probability of being selected.

2.5 Survey team

Seventeen teaching hospitals and/or diabetes centers participated in the 2nd NDSP 2016 - 2017. The training sessions of these teams were conducted from February 2016 to July 2016. The teams were trained to identify households, to fill the questionnaire, to take anthropometric measurements and to collect blood samples. Each team comprised of laboratory technicians, paramedical staff and survey officers, led by the physician as provincial coordinator of that cluster. A detailed structure of whole survey is given in Figure 1, 2 and 3.

2.6 Data collection method

Data collection method of this study was based on multiple indicator cluster survey manual³³. It was done in two phases. In the 1st phase, a pre-survey visit was made for the selection of houses following systematic sampling technique. The first house in the lane was selected randomly and afterwards every tenth house was identified. In case, residents of the identified household were not present or if they refused to participate, the next consecutive household was taken. Teams marked the houses and informed the adult residents about the survey and their participation. The selected household members were requested to come after an overnight fast (at least 8 hours) to the camp on the specific day. Only those who gave an informed written consent for interview, anthropometry, screening, blood sample and data collection were selected for the survey. Once fasting sample was taken then each participant was given 75 grams of anhydrous glucose load and were requested to stay in the screening facility for at least two hours. Meanwhile, their clinical and anthropometric data was collected by the trained paramedic staff under the supervision of survey officer.

2.7 Measurements

Height, weight, waist circumference, waist-hip ratio and blood pressure were measured as per the standardized procedure. ^{16, 22, 25} Individuals were requested to take 10 minutes rest in a sitting position before measurement of blood pressure and two readings was taken and mean of these reading was used.

2.8 Blood Sampling and Testing

Blood samples were collected by using sterilized disposable vacutainer tubes containing sodium fluoride (for glucose), EDTA K2 (for HbA1c) and gel (for lipids). Within 1 hour of blood collection, the samples were centrifuged, separated and sent to the laboratory. Plasma glucose was measured using the glucose oxidase peroxidase method, total cholesterol by CHOD-PAP method, triglycerides by GPO-PAP method, high density lipoprotein cholesterol (HDL-C) by homogeneous enzymatic calorimetric method, low density lipoprotein cholesterol (LDL-C) by CHOD-PAP method and HbA1c by high-performance liquid chromatography (HPLC) method. Fasting and random (2-hour post 75gm glucose load) plasma glucose levels were performed at the site while samples for HbA1c and lipid profile were transported as per protocol to BIDE

where they were kept in -80 freezer until transported to PHRC, Jinnah Postgraduate Medical Centre (JPMC), Karachi for analysis.

2.9 Definitions

Following definitions were used to interpret the results.

2.9.1 Diabetes

According to WHO diagnostic criteria, diabetes was diagnosed and the results of plasma glucose testing were categorized as follows: isolated impaired fasting glucose (fasting glucose level, ≥110 mg/dl and <126 mg/dl, 2-hour glucose level in the glucose tolerance test ≤140 mg/dl); isolated impaired glucose tolerance (fasting glucose level, <110 mg/dl and 2-hour glucose level, >140 and <200 mg/dl); combined impaired fasting glucose and impaired glucose tolerance (fasting glucose level, ≥110 and <126 mg/dl, and 2-hour glucose level >140 and <200 mg/dl); and undiagnosed diabetes (fasting glucose level, ≥126 mg/dl or 2-hour glucose level, ≥200 mg/dl or both).³⁵

For diagnosing diabetes, HbA1c was used as the diagnostic tool. According to the American Diabetes Association (ADA) standards of care HbA1c \geq to 6.5% (48 mmol/mol) was diagnosed as diabetic while HbA1c between 5.7 – 6.4 % (39 to 46 mmol/mol) was considered as prediabetic.³⁶

2.9.2 Hypertension

Participants were considered hypertensive if they were already diagnosed by a physician or if they were taking any anti-hypertensive medication or if the systolic blood pressure (SBP) was \geq 140 mmHg and/or diastolic blood pressure (DBP) \geq 90 mmHg.^{16,19}

2.9.3 Body Mass Index

Using WHO definition for obesity in Asian population, Body mass index (BMI) of 25 and 27 (kg/m²) was labeled as overweight and obesity respectively. Central obesity was defined as waist-to-hip ratio (WHR) \geq 0.9 and \geq 0.8, and/or waist circumference \geq 90 cm and \geq 80 cm in males and females respectively. 12, 13

2.9.4 Dyslipidemia

Using the Adult Treatment Panel III guidelines; dyslipidemia was classified as one or more of the following conditions in fasting state; serum cholesterol >200 mg/dl, serum low density lipoprotein cholesterol (LDL-C) >130 mg/dl, serum high density lipoprotein cholesterol (HDL-C) < 40 mg/dl and < 50 mg/dl for male and female respectively and serum triglycerides (TG) > 150 (mg/dl). Participants were also considered to have dyslipidemia if they were taking any anti-cholesterol medications.

2.10 Data Analysis

Data analysis was conducted on Statistical Package for Social Sciences (SPSS), version 20. Descriptive analysis included the estimation of mean values and standard deviations for continuous variables. Categorical variables and prevalence values were presented in the form of frequency and percentage. Risk factors of diabetes, hypertension, obesity and dyslipidemia, were examined using multivariate logistic regression. P-value of <0.01 was considered statistically significant. For all estimates, the study population was weighted to the latest available demographic information at Pakistan Bureau of Statistics.

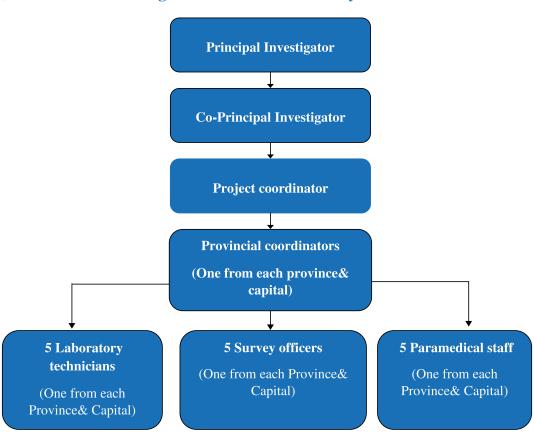
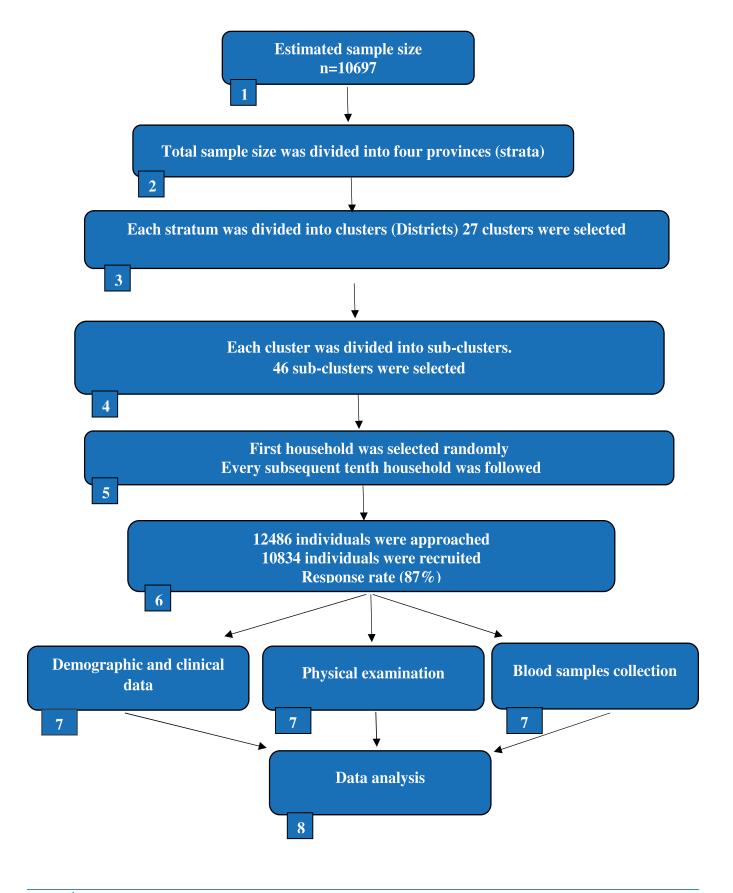
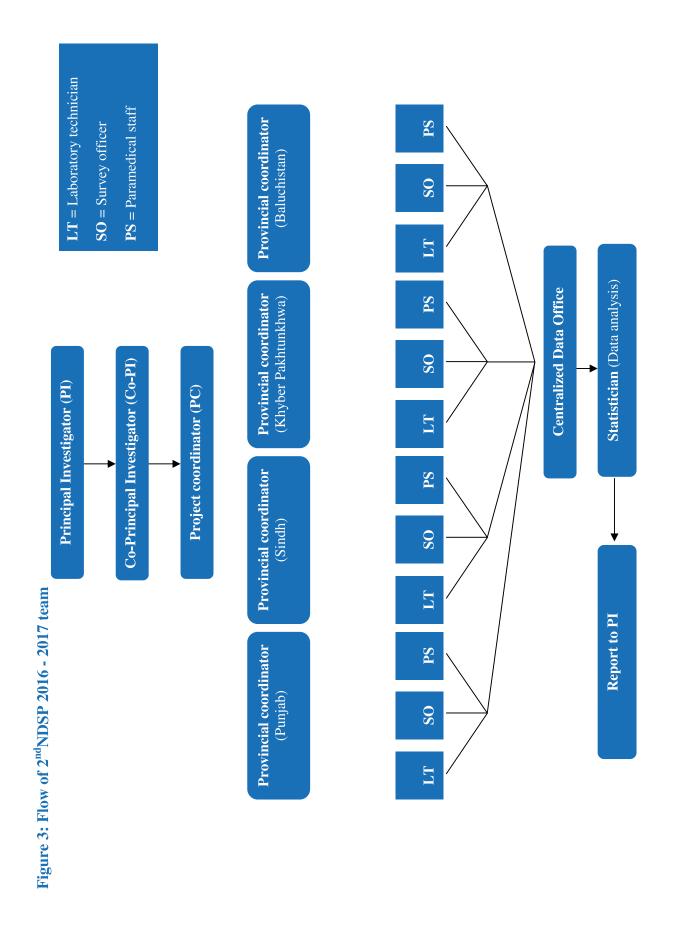
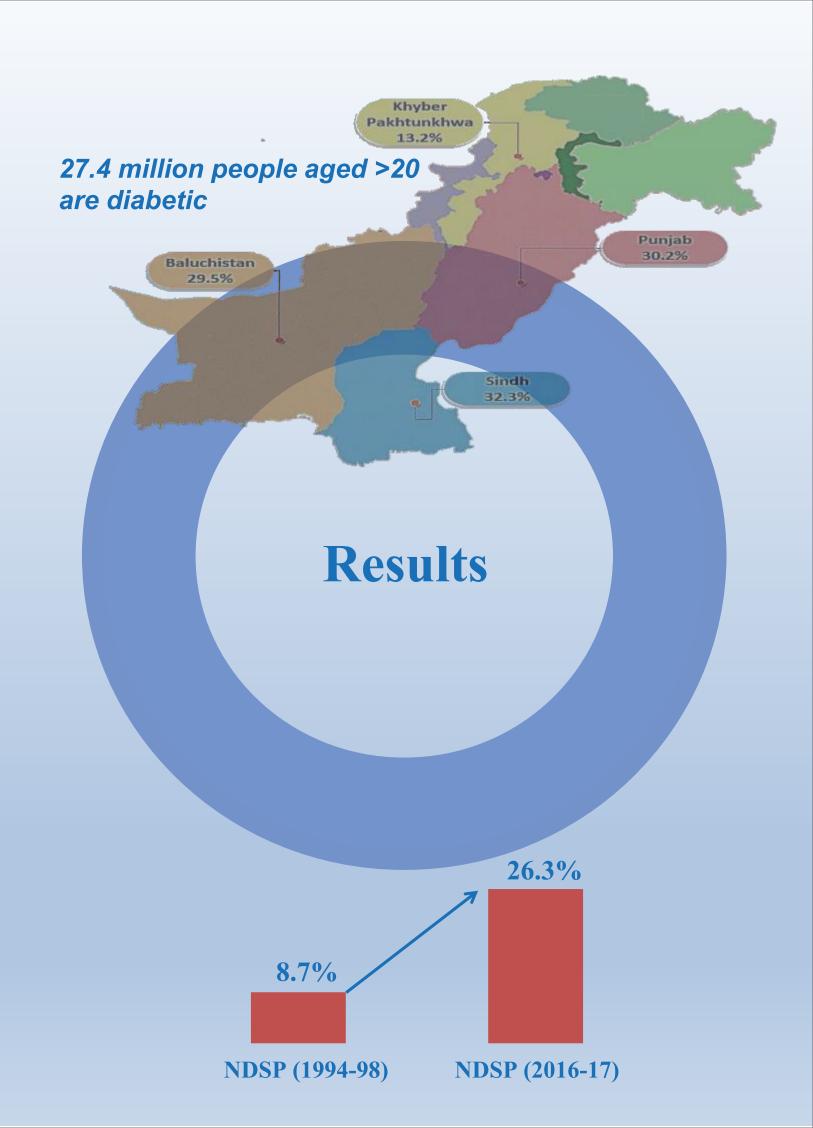


Figure 1: Structure of survey team

Figure 2: Step by step approach for the 2ndNDSP 2016 - 2017









3. RESULTS

3.1 Demographic characteristics

Basic characteristics of study population are presented in table 1. A total of 10834 persons were screened for diabetes; out of whom 43.9% were males and 56.1% were females. More than half of persons (53.5%) had at least primary level education. Almost, a third (30.2%) of the population had positive family history of diabetes and 14.5% were tobacco users. Mean BMI of participants was almost similar in all Provinces with an overall mean of 27.2 ± 6.0 kg/m². Highest mean blood pressure was seen in Baluchistan and lowest in Khyber-Pakhtunkhwa.

Table 1: Baseline characteristics of the study participants

| Description | Punjab | Sindh | Khyber Pakhtunkhwa | Baluchistan | Overall |
|-------------------------------------|------------------|------------------|--------------------|------------------|------------------|
| Number of participants | 6221 | 2531 | 1544 | 538 | 10834 |
| Age (years) | 43.5 ± 14.1 | 45.5 ± 14.2 | 40.3 ± 12.9 | 48.4 ± 12.81 | 43.8 ± 14.0 |
| Gender | | | | | |
| Male | 2457 (39.5) | 1192 (47.1) | 835 (54.1) | 257 (47.8) | 4756 (43.9) |
| Female | 3764 (60.5) | 1339 (52.9) | 709 (45.9) | 281 (52.2) | 6078 (56.1) |
| Primary education or more | 2675 (49.6) | 1353 (61.3) | 759 (55.6) | 278 (54.9) | 5065 (53.5) |
| Tobacco addiction | 614 (11.2) | 493 (22.2) | 117 (8.9) | 152 (29.1) | 1376 (14.5) |
| Positive family history of diahetes | 1509 (27.8) | 760 (37) | 240 (22.2) | 236 (43.9) | 2745 (30.2) |
| Body mass index (kg/m²) | 27.5 ± 6.1 | 26.5 ± 5.6 | 27.2 ± 6.1) | 26.7 ± 5.0 | 27.2 ± 6.0 |
| Male | 91.7 ± 14.5 | 92.3 ± 12.1 | 98.0 (12.4) | 96.7 ± 11.7 | 93.1 ± 13.8 |
| Female | 92.4 ± 15.0 | 94.3 ± 13.4 | 82.3 (11.7) | 101.1 ± 15.6 | 93.2 ± 14.5 |
| Blood pressure (mmHg) | | | | | |
| Systolic | 126.0 ± 19.8 | 127.3 ± 18.5 | 121.3 (14.1) | 135.7 ± 17.6 | 126.1 ± 18.9 |
| Diastolic | 84.5 ± 15.1 | 82.8 ± 11.8 | 79.8 (11.5) | 89.0 ± 11.2 | 83.7 ± 14.0 |

3.1.1 Age and gender

The age distribution is shown in table 2. Overall, 26.5% were between 40-49 years followed by 21.5% were between 30-39 years. In women, majority were between 30-39 years whereas, in men it was 60+ years.

Table 2: Age and gender distribution of respondents

| A go groung (voorg) | Men | Women | Both genders |
|---------------------|------|-------|---------------------|
| Age groups (years) | % | % | % |
| 20-29 | 44.2 | 55.8 | 16.7 |
| 30-39 | 39.8 | 60.2 | 21.5 |
| 40-49 | 40.9 | 59.1 | 26.5 |
| 50-59 | 44.1 | 55.9 | 19 |
| 60+ | 52.8 | 47.2 | 16.3 |
| Overall | 43.8 | 56.2 | 100 |

3.1.2 Education Level

Educational level showed that 38.6% were illiterate and only 9.3% had completed graduation (Table 3).

Table 3: Educational level of respondents

| | Level of education | | | | | | | | |
|-----------------------|--------------------|-------------------------|--------------|----------------|---------------|------------------------|--|--|--|
| Age groups (years) | Illiterate % | Can read/ write % | Primary % | Intermediate % | Graduate % | Post- graduate % | | | |
| 20-29 | 20.8 | 6.8 | 9.8 | 18.2 | 18.5 | 6.5 | | | |
| 30-39 | 32.7 | 8.4 | 14.3 | 10.2 | 9.4 | 5.6 | | | |
| 40-49 | 41.5 | 7.9 | 12.8 | 8.1 | 7.9 | 4 | | | |
| 50-59 | 43.6 | 8.4 | 16.6 | 6.1 | 6.7 | 3.4 | | | |
| 60+ | 53.8 | 7.4 | 13.6 | 3.9 | 5.3 | 2.3 | | | |
| Overall | 38.6 | 7.8 | 13.5 | 9.1 | 9.3 | 4.4 | | | |

3.1.4 Marital Status

Majority of participants were married (84.8%) followed by never married (12.9%) as given in table 4.

Table 4: Marital status of the respondents

| | Marital status | | | | |
|--------------------|----------------|---------|---------------------|-------|--|
| Age groups (years) | Single | Married | Divorced/ Separated | Widow | |
| | % | % | % | % | |
| 20-29 | 54.4 | 44.8 | 0.6 | 0.1 | |
| 30-39 | 9.2 | 90.2 | 0.3 | 0.2 | |
| 40-49 | 3.7 | 94.6 | 0.5 | 1.2 | |
| 50-59 | 2.8 | 94.3 | 0.6 | 2.3 | |
| 60+ | 2.5 | 90.8 | 0.5 | 95 | |
| Overall | 12.9 | 84.8 | 0.5 | 1.8 | |

3.1.5 Tobacco Addiction

The proportion of current tobacco users was 14.5%. The prevalence of tobacco addiction increased with age and was highest (20.7%) in those aged 60 years or above (Table 5).

Table 5: Tobacco addiction among respondents

| Age groups (years) | Current tobacco users | Ex- tobacco users | Non- tobacco users |
|--------------------|-----------------------|-------------------|--------------------|
| 20-29 | 9.8 | 0.9 | 89.3 |
| 30-39 | 11.8 | 1.8 | 86.3 |
| 40-49 | 13.5 | 3.1 | 83.4 |
| 50-59 | 17.9 | 3.9 | 78.2 |
| 60+ | 20.7 | 5 | 74.2 |
| Overall | 14.5 | 2.9 | 82.5 |

3.2 Measurements

3.2.1 Body Mass Index (BMI)

Table 6 presents the BMI score according to Asian cutoff. About 27.7% respondents had normal weight (BMI <23), 14.2 % were overweight (BMI 23-24.9) and 58.1% were obese (BMI ≥ 25).

Using WHO criteria, 14.2% were overweight (BMI 25-26) and 43.9% were obese (BMI> 27). Most obese population was seen in those aged between 40-49 years (Table 7).

3.2.2 Hypertension

Overall, 46.2 % population had hypertension. The prevalence of hypertension increased with age and was highest (65.7%) in age group 60 years or above (Table 8).

Table 6: BMI classification of the respondents by Asian cutoff

| | Body mass index | | | | |
|--------------------|--------------------------------------|---|-------------------------|--|--|
| Age groups (years) | Normal <23 kg/m ² % | Over-weight 23-24.9 kg/m ² % | Obese ≥25 kg/m² % | | |
| 20-29 | 42.6 | 14.1 | 43.3 | | |
| 30-39 | 21.8 | 15.4 | 62.8 | | |
| 40-49 | 16.1 | 12.2 | 71.7 | | |
| 50-59 | 17.7 | 14.3 | 68.0 | | |
| 60+ | 26.5 | 15.3 | 58.2 | | |
| Overall | 27.7 | 14.2 | 58.1 | | |

Table 7: BMI classification of the respondents by WHO criteria

| | | Body mass index (BMI) | | | | |
|-----------------------|--------------------------------------|---------------------------------------|-------------------------|--|--|--|
| Age groups (years) | Normal <23 kg/m ² % | Over-weight 23-24.9 kg/m ² | Obese ≥25 kg/m² % | | | |
| 20-29 | 56.7 | 13.6 | 29.8 | | | |
| 30-39 | 37.2 | 13.2 | 49.5 | | | |
| 40-49 | 28.3 | 14.7 | 56.9 | | | |
| 50-59 | 32.0 | 16.4 | 51.6 | | | |
| 60+ | 41.8 | 15.4 | 42.8 | | | |
| Overall | 41.9 | 14.2 | 43.9 | | | |

Table 8: Hypertension status of the respondents

| Age groups (years) | Hypertensive % | Non-hypertensive % |
|-----------------------|-------------------|-----------------------|
| 20-29 | 29.5 | 70.5 |
| 30-39 | 43.5 | 56.5 |
| 40-49 | 58.3 | 41.7 |
| 50-59 | 64.3 | 35.7 |
| 60+ | 65.7 | 34.3 |
| Overall | 46.2 | 53.8 |

3.3 Diabetes

3.3.1 Normal glucose tolerance

Overall prevalence of normal glucose tolerance was 60.09%. Normal glucose tolerance was higher in men than women. The prevalence of normal glucose tolerance with respect to age stratification showed a declining pattern (Table 9).

3.3.2 Pre-diabetes

Overall, 14.4% respondents were pre-diabetic. Pre-diabetes was more prevalent in women who were aged between 30-39 years (Table 10).

3.3.3 Prevalence of newly diagnosed diabetes mellitus

Overall, prevalence of newly diagnosed diabetes was 7.1% with slightly more preponderance (7.3%) in men than women (6.9%). Most newly diagnosed diabetics were aged 50 and above in both genders (Table 11).

3.3.4 Prevalence of known diabetes

Table 12 shows that almost 19.2 % of the respondents already knew that they were diabetics. Its prevalence was higher in women and older age.

3.3.5 Prevalence of diabetes mellitus

Overall prevalence of diabetes was 26.3% (Table 13). The prevalence increased with age and was maximally seen in 50-59 years' age (50.3% men and 49.3% women).

3.3.6 Prevalence of diabetes and pre-diabetes, by Province and area of Pakistan

Using the oral glucose tolerance test (OGTT) criteria, highest prevalence of diabetes was seen in Sindh followed by Punjab. Highest prevalence of pre-diabetes was seen in Baluchistan as compared to other provinces (Table 14).

3.3.7 Age, gender and area specific prevalence of pre-diabetes and diabetes in four provinces of Pakistan:

Urban women showed significantly higher prevalence of diabetes than rural women above the age of 40 years whilst in men these trends was seen in the age group of 60 years and above (p<0.05). On the other hand, urban men in the age group 30-39 years showed significantly (p<0.05) lower prevalence of diabetes than rural men (Figure 5).

Rural men showed significantly higher prevalence of pre-diabetes than urban men for the age group of 40-49 years whilst for women, significant difference was seen in urban as compared to rural population for the age group of 30-39 years (Figure 6).

Table 9: Prevalence of normal glucose tolerance

| | Gender | Age (years) | Weighted Prevalence (%) | Weighted Prevalence (%) | Overall Prevalence (%) |
|-------------|--------|----------------|-------------------------|-------------------------|------------------------|
| | | 20-29 | 82.4 | | |
| OGTT Female | | 30-39 | 64.6 | | |
| | Male | 40-49 | 50.1 | 61.6 | |
| | | 50-59 | 35.8 | | |
| | | 60+ | 34.9 | | 60.09 |
| | 20-29 | 79.6 | | 00.07 | |
| | | 30-39 | 60.6 | | |
| | Female | 40-49 | 46.3 | 58.9 | |
| | | 50-59 | 35.9 | | |
| | | 60+ | 35.3 | | |
| | | 20-29 | 85.6 | 63.4 | |
| | | 30-39 | 65.6 | | |
| | Male | 40-49 | 50.6 | | |
| | | 50-59 | 36.7 | | |
| ШЬА10 | | 60+ | 38.2 | | 64.525 |
| HbA1c _ | | 20-29 | 89.8 | | 04.323 |
| | | 30-39 | 67.7 | | |
| | Female | 40-49 | 47.7 | 65.3 | |
| | | 50-59 | 39.2 | | |
| | | 60+ | 40.4 | | |

Table 10: Prevalence of pre-diabetes

| | | A 90 | Weighted | Weighted | Overall |
|-------|--------|-------------|------------|------------|------------|
| | Gender | Age | Prevalence | Prevalence | Prevalence |
| | | (years) | (%) | (%) | (%) |
| | | 20-29 | 9.7 | | |
| | | 30-39 | 13.4 | | |
| | Male | 40-49 | 13.2 | 12.8 | |
| | | 50-59 | 14 | | |
| OGTT | | 60+ | 16.9 | | 14.4 |
| OGII | | 20-29 | 13.1 | | 14.4 |
| | | 30-39 | 17.3 | | |
| | Female | 40-49 | 14.7 | 15.6 | |
| | | 50-59 | 14.8 | | |
| | | 60+ | 16.3 | | |
| | | 20-29 | 2.1 | | |
| | | 30-39 | 5.9 | | |
| | Male | 40-49 | 5.5 | 5.1 | |
| | | 50-59 | 7.7 | | 5.5 |
| HbA1c | | 60+ | 9.5 | | |
| HbA1c | | 20-29 | 2.5 | | 5.5 |
| | | 30-39 | 7 | | |
| | Female | 40-49 | 7.2 | 5.9 | |
| | | 50-59 | 8.3 | | |
| | | 60+ | 9.6 | | |

Table 11: Prevalence of newly diagnosed diabetes mellitus

| | | Age | Weighted | Weighted | Overall |
|--------|--------|----------|------------|------------|------------|
| | Gender | (years) | Prevalence | Prevalence | Prevalence |
| | | (j cars) | (%) | (%) | (%) |
| | | 20-29 | 3.6 | | |
| | Male | 30-39 | 5.8 | | |
| | | 40-49 | 9.6 | 7.3 | |
| | | 50-59 | 13.4 | | |
| OCTT | | 60+ | 11.4 | | 7.1 |
| OGII | OGTT | 20-29 | 4.2 | | 7.1 |
| Female | | 30-39 | 6.3 | | |
| | Female | 40-49 | 8 | 6.9 | |
| | | 50-59 | 9.2 | | |
| | | 60+ | 12.2 | | |
| | | 20-29 | 6 | | |
| | | 30-39 | 7.8 | | |
| | Male | 40-49 | 10.6 | 8.5 | |
| | | 50-59 | 12.5 | | 7.4 |
| HbA1c | | 60+ | 10.4 | | |
| IIDAIC | | 20-29 | 3.9 | | 7,4 |
| | | 30-39 | 6.6 | | |
| | Female | 40-49 | 9.6 | 6.7 | |
| | | 50-59 | 7.9 | | |
| | | 60+ | 9.6 | | |

Table 12: Prevalence of known diabetes

| | | A 000 | Weighted | Weighted | Overall |
|-------|--------|--------------|------------|------------|------------------|
| | Gender | Age | Prevalence | Prevalence | Prevalence |
| | | (years) | (%) | (%) | (%) |
| | | 20-29 | 4.3 | | |
| | | 30-39 | 16.3 | | |
| | Male | 40-49 | 27 | 18.9 | |
| | | 50-59 | 36.9 | | |
| OGTT | | 60+ | 36.9 | | 19.2 |
| | | 20-29 | 3.1 | | 19.2 |
| Fen | | 30-39 | 15.8 | | |
| | Female | 40-49 | 31 | 19.4 | |
| | | 50-59 | 40.1 | | |
| | | 60+ | 36.1 | | |
| | | 20-29 | 6.2 | | |
| | | 30-39 | 20.7 | | |
| | Male | 40-49 | 33.3 | 22.9 | |
| | | 50-59 | 43.2 | | 22.4 |
| HbA1c | | 60+ | 42 | | |
| HbA1c | Female | 20-29 | 3.9 | | 22. T |
| | | 30-39 | 18.6 | | |
| | | 40-49 | 35.6 | 22.1 | |
| | | 50-59 | 44.7 | | |
| | | 60+ | 40.4 | | |

Table 13: Prevalence of diabetes mellitus

| | | Ago | Weighted | Weighted | Overall |
|-------|--------|---------|------------|------------|------------|
| | Gender | Age | Prevalence | Prevalence | Prevalence |
| | | (years) | (%) | (%) | (%) |
| | | 20-29 | 7.9 | | |
| | | 30-39 | 22 | | |
| | Male | 40-49 | 36.6 | 26.2 | |
| | | 50-59 | 50.3 | | |
| OGTT | | 60+ | 48.3 | | 26.3 |
| OGII | | 20-29 | 7.3 | | 20.3 |
| | | 30-39 | 22 | | |
| | Female | 40-49 | 39 | 26.3 | |
| | | 50-59 | 49.3 | | |
| | | 60+ | 48.3 | | |
| | | 20-29 | 12.3 | | |
| | | 30-39 | 28.5 | | |
| | Male | 40-49 | 43.9 | 31.5 | |
| | | 50-59 | 55.6 | | |
| HbA1c | | 60+ | 52.3 | | 29.9 |
| HDAIC | | 20-29 | 7.7 | | 29.9 |
| | | 30-39 | 25.2 | | |
| | Female | 40-49 | 45.1 | 28.8 | |
| | | 50-59 | 52.6 | | |
| | | 60+ | 50 | | |

Table 14: Prevalence of diabetes and pre-diabetes; by province and areas of Pakistan

| Description | Punjab | Sindh | Khyber Pakhtunkhwa | Baluchistan |
|----------------------------------|------------------|------------------|--------------------|---------------------|
| Overall (urban and rural) | % (95%CI) | % (95%CI) | % (95%CI) | % (95%CI) |
| Diabetes | | | | |
| Known diabetes | 23.7 (22.6-24.7) | 23.6 (21.9-25.2) | 10.2 (8.7-11.7) | 19.1 (15.7-22.4) |
| Fasting glucose (FG) | 3.3 (2.8-3.7) | 3.6 (2.8-4.3) | 1 (0.5-1.5) | 3.1 (1.6-4.5) |
| 2 hours glucose tolerance (2hGT) | 1.1 (0.8-1.3) | 1.4 (0.9-1.8) | 0.4 (0.1-0.7) | 4.7 (2.9-6.4) |
| Both FG and 2hGT | 2.1 (1.7-2.4) | 3.7 (2.9-4.4) | 1.6 (0.9-2.2) | 2.6 (1.2-3.9) |
| Newly diagnosed diabetes | 6.5 (5.8-7.1) | 8.7 (7.6-9.8) | 3 (2.1-3.8) | 10.4 (7.8-12.9) |
| Total diabetes | 30.2 (29.0-31.3) | 32.3 (30.4-34.1) | 13.2 (11.5-14.8) | 29.5 (25.6-33.3) |
| Pre-diabetes | | | | |
| Fasting glucose (FG) | 1.8 (1.4-2.1) | 1.8 (1.2-2.3) | 1.1 (0.5-1.6) | 0.4 (0.1-0.9) |
| 2 hours glucose tolerance (2hGT) | 11.2 (10.4-11.9) | 8.4 (7.3-9.4) | 2.9 (2.0-3.7) | 42.4 (38.2-46.5) |
| Both FG and 2hGT | 2.1 (1.7-2.4) | 1.5 (1.0-1.9) | 0.5 (0.1-0.8) | 8.7 (6.3-11.0) |
| Total Pre-diabetes | 15.1 (14.2-15.9) | 11.7 (10.4-12.9) | 4.5 (3.4-5.5) | 51.5 (47.2-55.7) |

| Lurban Diabetes 21.7 21.5 7.4 17.4 Known diabetes 21.7 21.5 7.4 17.4 Known diabetes (19.8-23.5) (19.2-23.7) (4.5-10.2) (11.8-22.9) Fasting glucose (FG) (4.4-6.5) (1.6-3.3) (0.0-2.4) (0.0-3.8) 2 hours glucose tolerance (2hGT) (1.8-3.4) (0.9-2.2) (0.0-2.2) (0.0-3.8) Both FG and 2hGT (1.8-3.3) (1.8-3.5) (0.2-1.4) (0.4-2.2) Newly diagnosed diabetes (9.9 6.8 2.9 8.5 Newly diagnosed diabetes (8.5-11.2) (5.4-8.1) (1.0-4.7) (4.4-12.5) Newly diagnosed Glabetes (8.5-11.2) (5.4-8.1) (1.0-4.7) (4.4-12.5) Pre-diabetes (2.9-4.33.7) (2.5-8-30.7) (6.9-13.6) (19.5-52.3) Pre-diabetes 2.1 (1.2-2.6) (0.0-6.0) (0.00.6) (0.00.6) (0.00.6) Pourrs glucose tolerance (2hGT) (11.9 (0.2-13.6) (0.2-13.6) (0.0-13.6) (0.0-13.6) (0.0-13.6) | Description | Punjab | Sindh | Khyber Pakhtunkhwa | Baluchistan |
|--|----------------------------------|---------------|---------------|--------------------|-------------|
| 21.7 21.5 7.4 (19.8-23.5) (19.8-23.7) (19.8-23.5) (19.2-23.7) (4.5-10.2) (5.5 1.2 1.2 1.2 1.3 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4 | Urban | | | | |
| 21.7 21.5 7.4 (19.8-23.5) (19.2-23.7) (4.5-10.2) 5.5 2.5 1.2 (4.4-6.5) (1.6-3.3) (0.0-2.4) 1.8 1.6 1.1 (1.2-2.4) (0.9-2.2) (0.0-2.2) 2.6 2.7 0.6 2.6 2.7 0.6 (1.8-3.3) (1.8-3.5) (0.2-1.4) 9.9 6.8 2.9 (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) (11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) (2.0-3.5) (0.4-1.5) (0.0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Diabetes | | | | |
| (19.8-23.5) (19.2-23.7) (4.5-10.2) (6.5-10.2) 5.5 2.5 1.2 (1.2.4) (1.6-3.3) (1.6-3.4) (1.2.4) 1.8 1.6 1.1 1.1 1.1 1.8 1.6 1.6 1.1 2.6 2.7 (0.0-2.2) (0.0-2.2) 2.6 2.7 (0.0-2.2) (0.0-2.2) 3.6 1.8-3.5) (1.8-3.5) (0.2-1.4) 9.9 6.8 2.9 6.8 8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 31.6 28.3 10.3 11.9 (25.8-30.7) (6.9-13.6) 11.9 7.7 (11-2.6) 11.9 7.7 (11-2.6) 11.9 7.7 (1.1-2.6) 11.9 7.7 (1.1-2.6) 11.9 7.7 (1.1-2.6) 11.9 7.7 (1.1-2.6) 11.9 7.7 (1.1-2.6) 11.9 0.2 0.2 11.9 0.2 0.2 11.9 | Known diabetes | 21.7 | 21.5 | 7.7 | 17.4 |
| 5.5 2.5 1.2 (4.4-6.5) (1.6-3.3) (0.0-2.4) 1.8 1.6 1.1 1.8 1.6 1.1 2.6 2.7 (0.0-2.2) 2.6 2.7 (0.0-2.2) 2.6 2.7 (0.0-2.2) 3.6 (1.8-3.3) (1.8-3.5) (0.2-1.4) 6.8 2.9 6.8 2.9 8.5-11.2) (5.4-8.1) (1.0-4.7) (1.0-4.7) 31.6 28.3 10.3 (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) (2.2-3.1) (2.0-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0.6-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (19.8-23.5) | (19.2-23.7) | (4.5-10.2) | (11.8-22.9) |
| 1.8 1.6 1.1 2.6 2.7 0.0-2.2) 2.6 (1.8-3.3) (1.8-3.5) 0.0-2.1.4) 9.9 6.8 2.9 (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 20.4-33.7) (25.8-30.7) (6.9-13.6) 21 1.9 0.2 21 1.9 0.0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.9 0.4-1.5) 0.0-0.6) 1.5.1-18.5) (8.9-12.2) 0.5-3.6) <th>Fasting glucose (FG)</th> <th>5.5 (4.4-6.5)</th> <th>2.5 (1.6-3.3)</th> <th>1.2 (0.0-2.4)</th> <th>1.9</th> | Fasting glucose (FG) | 5.5 (4.4-6.5) | 2.5 (1.6-3.3) | 1.2 (0.0-2.4) | 1.9 |
| 1.2-2.4) (0.9-2.2) (0.0-2.2) 2.6 2.7 0.6 2.6 (1.8-3.3) (1.8-3.5) (0.2-1.4) 9.9 6.8 2.9 6.8 2.9 2.9 6.8 2.9 2.9 31.6 28.3 10.3 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) 11.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 2.9 (0.4-1.5) (0.0-6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | 2 hours obucose tolerance (2hGT) | 1.8 | 1.6 | 1.1 | 5.7 |
| 2.6 2.7 0.6 (1.8-3.3) (1.8-3.5) (0.2-1.4) 9.9 6.8 2.9 (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 0.2 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (1.2-2.4) | (0.9-2.2) | (0.0-2.2) | (2.3-9.0) |
| (1.8-3.3) (1.8-3.5) (0.2-1.4) 9.9 6.8 2.9 (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) (20-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Both FG and 2hGT | 2.6 | 2.7 | 9.0 | 6.0 |
| 9.9 6.8 2.9 (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 20.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 2.8 1 0.2 16.8 10.6 2.1 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (1.8-3.3) | (1.8-3.5) | (0.2-1.4) | (0.4-2.2) |
| (8.5-11.2) (5.4-8.1) (1.0-4.7) 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Newly diagnosed diabetes | 6.6 | 8.9 | 2.9 | 8.5 |
| 31.6 28.3 10.3 (29.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (8.5-11.2) | (5.4-8.1) | (1.0-4.7) | (4.4-12.5) |
| 29.4-33.7) (25.8-30.7) (6.9-13.6) 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Total diabetes | 31.6 | 28.3 | 10.3 | 25.9 |
| 2.1 (1.4-2.7) (1.1-2.6) (0-0.6) (1.0-4-13.3) (6.2-9.1) (0.2-3.1) (0.2-3.1) (0.2-3.5) (0.4-1.5) (0.6-0.6) (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (29.4-33.7) | (25.8-30.7) | (6.9-13.6) | (19.5-32.3) |
| 2.1 1.9 0.2 (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Pre-diabetes | | | | |
| (1.4-2.7) (1.1-2.6) (0-0.6) 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Fasting olucose (FG) | 2.1 | 1.9 | 0.2 | 0 |
| 11.9 7.7 1.7 (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (1.4-2.7) | (1.1-2.6) | (0-0.6) | (0-0) |
| (10.4-13.3) (6.2-9.1) (0.2-3.1) 2.8 1 0.2 (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | 2 hours glucose tolerance (2hGT) | 11.9 | 7.7 | 1.7 | 45.9 |
| $\begin{array}{c ccccccccccccccccccccccccccccccccccc$ | | (10.4-13.3) | (6.2-9.1) | (0.2-3.1) | (38.6-53.1) |
| (2.0-3.5) (0.4-1.5) (0-0.6) 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | Both FG and 2hGT | 2.8 | 1 | 0.2 | 13.4 |
| 16.8 10.6 2.1 (15.1-18.5) (8.9-12.2) (0.5-3.6) | | (2.0-3.5) | (0.4-1.5) | (0-0.6) | (8.4-18.3) |
| (15.1-18.5) (8.9-12.2) (0.5-3.6) | Total Pre-diahetes | 16.8 | 10.6 | 2.1 | 203 |
| - | | (15.1-18.5) | (8.9-12.2) | (0.5-3.6) | (52.1-66.4) |

| Description | Punjab | Sindh | Khyber Pakhtunkhwa | Baluchistan |
|----------------------------------|---------------------|------------------|--------------------|------------------|
| Rural | | | | |
| Diabetes | | | | |
| Known diabetes | 21.4 (20.1-22.6) | 19.4 (17.0-21.7) | 10.3 (8.5-12.0) | 15.1 (11.3-18.8) |
| Fasting glucose (FG) | 2.3 (1.8-2.7) | 6.4 (4.9-7.8) | 1.7 (0.9-2.4) | 5.1 (2.8-7.3) |
| 2 hours glucose tolerance (2hGT) | 0.6 (0.3-0.8) | 1.5 (0.7-2.2) | 0.7 (0.2-1.1) | 5.7 (3.3-8.1) |
| Both FG and 2hGT | 1.7 (1.2-2.1) | 4.8 (3.5-6.0) | 2.7 (1.7-3.6) | 4.4 (2.2-6.5) |
| Newly diagnosed diabetes | 4.6 (3.9-5.2) | 12.7 (10.7-14.6) | 5.1 (3.8-6.3) | 15.2 (11.4-18.9) |
| Total diabetes | 26 (24.6-27.3) | 32.1 (29.3-34.8) | 15.4 (13.3-17.4) | 30.2 (25.4-34.9) |
| Pre-diabetes | , | | | |
| Fasting glucose (FG) | 1.4 (1.0-1.7) | 1.6 (0.8-2.3) | 1.7 (0.9-2.4) | 0.3 (0.0-0.8) |
| 2 hours glucose tolerance (2hGT) | 10.7 (9.7-11.6) | 9.1 (7.3-10.8) | 4.4 (3.2-5.5) | 37.1 (32.0-42.1) |
| Both FG and 2hGT | 1.4 (1.0-1.7) | 2.2 (1.3-3.0) | 0.9 | 6.4 (3.8-8.9) |
| Total Pre-diabetes | 13.5 (12.4-14.5) | 12.9 (10.9-14.8) | 7 (5.5-8.4) | 43.8 (38.6-48.9) |

OGTT criteria for diagnosis of pre-diabetes; Fasting $110-125~\mathrm{mg/dl}$ and/or RBS $140-199~\mathrm{mg/dl}$ OGTT criteria for diagnosis of diabetes; Fasting $\geq 126 \text{mg/dl}$ and/or 2-hours $\geq 200 \text{mg/dl}$

Figure 4: Age-stratified prevalence of diabetes among men and women with urban and rural distribution

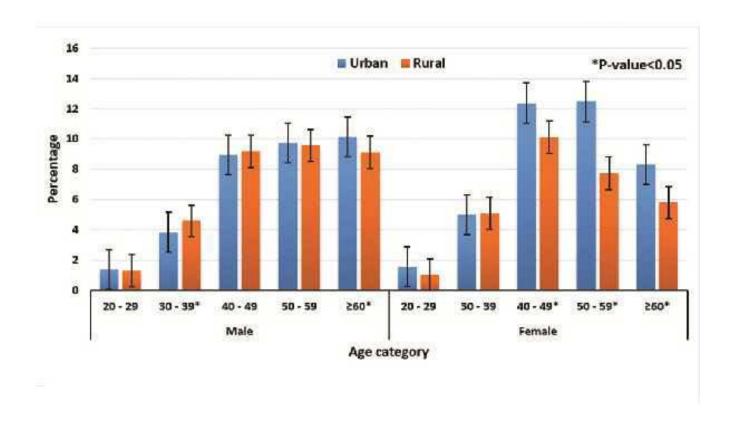
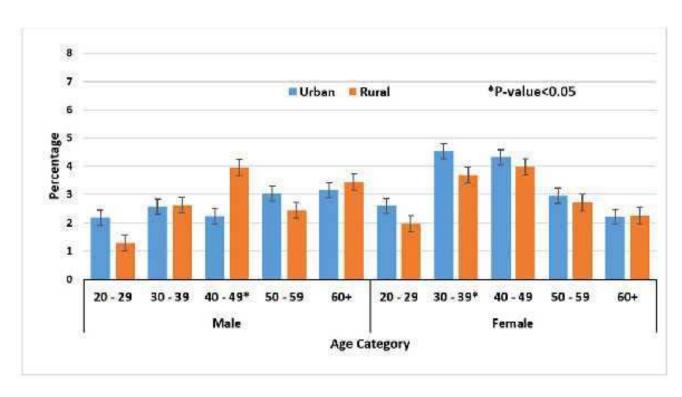


Figure 5: Age-stratified prevalence of pre-diabetes among men and women with urban and rural distribution



3.4 Dyslipidemia

3.4.1 Raised cholesterol

About, 39.3% population had raised cholesterol levels (>200mg/dl). Deranged cholesterol was mostly seen in people aged between 50-59 years followed by those aged 40-49 years (Table 15).

3.4.2 Hyper triglyceride

High triglycerides were seen in 51.5% respondents and this was mostly seen in individuals who were between 50-59 years of age followed by those who were between 40-49 years (Table 16).

3.4.3 High low-density lipoprotein

Overall, 39.7% respondents had high levels of low density lipoprotein. Majority of such respondents were aged between 40-49 years followed by 50-59 years (Table 17).

3.4.4 Low high-density lipoprotein

Majority of the population had low values of high density lipoprotein (83.5% men, 90% women). Among men, highest percentage of low high-density lipoprotein was seen in age group 40-49 years, while in women it was seen in age group 60+ years (Table 18).

Table 15: Hypercholesterolemia status of the respondents

| | Chole | sterol |
|--------------------|-----------------|-----------------|
| Age groups (years) | % ≤200 mg/dl | % >200 mg/dl |
| 20-29 | 70.7 | 29.3 |
| 30-39 | 58.3 | 41.7 |
| 40-49 | 53.4 | 46.6 |
| 50-59 | 51.3 | 48.7 |
| 60+ | 58.8 | 41.2 |
| Overall | 60.7 | 39.3 |

Table 16: Hypertriglyceridemia status of the respondents

| | Triglyc | eride |
|--------------------|-----------------|-----------------|
| Age groups (years) | % ≤150 mg/dl | % >150 mg/dl |
| 20-29 | 67.0 | 33.0 |
| 30-39 | 48.8 | 51.2 |
| 40-49 | 38.6 | 61.4 |
| 50-59 | 37.0 | 63.0 |
| 60+ | 48.0 | 52.0 |
| Overall | 51.1 | 48.9 |

Table 17: Status of high low-density lipoprotein of the respondents

| | LDL | |
|--------------------|-----------------|-----------------|
| Age groups (years) | % ≤130 mg/dl | % >130 mg/dl |
| 20-29 | 66.4 | 33.6 |
| 30-39 | 59.1 | 40.9 |
| 40-49 | 54.8 | 45.2 |
| 50-59 | 55.0 | 45.0 |
| 60+ | 59.9 | 40.1 |
| Overall | 60.3 | 39.7 |

Table 18: Low high-density lipoprotein of the respondents

| | | | HDL | |
|---------------------|-----------|-----------|-----------|-----------|
| Age groups (years) | M | en | W | omen |
| rigo groups (Jears) | % | % | % | % |
| | >40 mg/dl | ≤40 mg/dl | >50 mg/dl | ≤50 mg/dl |
| 20-29 | 15.7 | 84.3 | 13.7 | 86.3 |
| 30-39 | 17.1 | 82.9 | 8.5 | 91.5 |
| 40-49 | 15.2 | 84.8 | 8.3 | 91.7 |
| 50-59 | 17.0 | 83.0 | 8.0 | 92.0 |
| 60+ | 20.0 | 80.0 | 7.3 | 92.7 |
| Overall | 16.5 | 83.5 | 10 | 90 |



Risk Factors





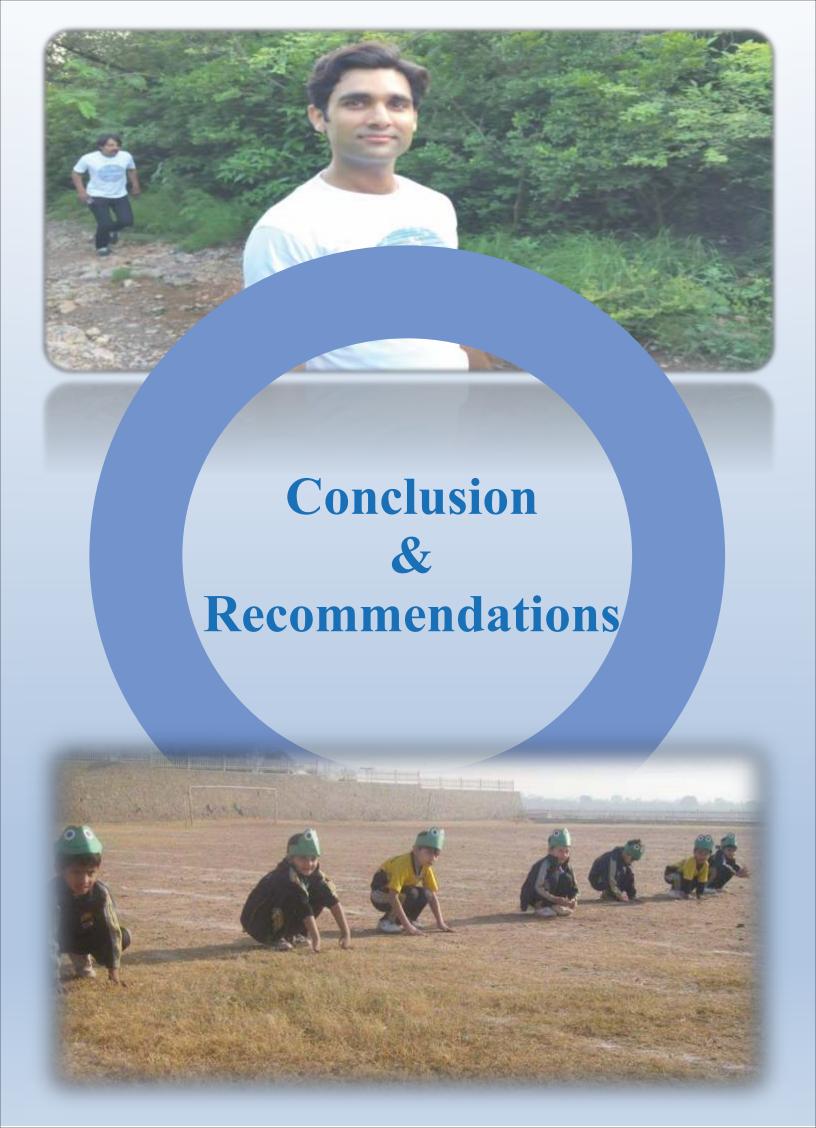
3. 5. Risk Factor Analysis

3. 5.1 Risk factor of diabetes, dyslipidemia, hypertension and obesity

Table 19 shows the multivariable logistic regression for identifying the associate risk factors for diabetes, pre-diabetes, hypertension, obesity and dyslipidemia. Age (greater than or equal to 43 years), family history of diabetes, hypertension, obesity and dyslipidemia were significant risk factors for diabetes (p \leq 0.0001). Similar were the risk factors for pre-diabetes including education with the exception of obesity (p \leq 0.05). Significant association was found between diabetes, hypertension, obesity and dyslipidemia ($p \le 0.05$).

Table 19: Risk factor of Diabetes, Dyslipidemia, Hypertension and Obesity

| | Diabetes | | Pre-diabetes | | Dyslipidemia | mia | Hypertension | nsion | Obesity | ty |
|------------------------------------|---------------------|---------|------------------------|-------------|-------------------|---------|------------------------|---------|------------------------|---------|
| Risk Factors | Adjusted Odds ratio | P-value | Adjusted Odds ratio v | P- value | P-value | P-value | Adjusted Odds ratio | P-value | Adjusted Odds ratio | P-value |
| Location (Rural) | | | | | 0.55 (0.4-0.7) | <0.0001 | | | 1.1 (1.0-1.3) | 0.032 |
| Age (≥43 years) | 3.3 (2.9-3.8) | <0.0001 | 1.8 (1.5-2.2) | | | | 2.3 (2.0-2.6) | <0.0001 | | |
| Gender (Female) | | | | | 0.53 (0.4-0.7) | <0.0001 | 1.6 (1.4-1.9) | <0.0001 | 1.4 (1.2-1.6) | <0.0001 |
| Education (primary or more) | | | 0.8 (0.7-0.9) 0.034 | | 0.70 (0.6-0.9) | 0.004 | | | | |
| Family history of diabetes | 1.9 (1.7-2.2) | <0.0001 | 1.5 (1.2-1.8) < 0.0001 | | | | 1.3 $(1.2-1.5)$ | <0.0001 | 1.2 (1.0-1.4) | 0.001 |
| Diabetes | | | | | 1.84 (1.3-2.4) | <0.0001 | 2.2 (2.0-2.6) | <0.0001 | 1.4 (1.2-1.6) | <0.0001 |
| Dyslipidemia | 1.8 (1.3-2.4) | <0.0001 | 1.5 (1.1-2.1) 0.011 | | 1 | 1 | 1.3 $(1.1-1.8)$ | 0.019 | 1.7 (1.3-2.1) | <0.0001 |
| Hypertension | 2.2 (1.9-2.5) | <0.0001 | 1.2 (1.0-1.4) 0.009 | | 1.3 (1.1-1.7) | 0.008 | _ | ı | 1.7 (1.5-1.9) | <0.0001 |
| Obesity | 1.4 (1.3-1.6) | <0.0001 | | | 1.7 (1.3-2.1) | <0.0001 | 1.7 (1.5-2.0) | <0.0001 | ı | ı |





4. CONCLUSION

The Survey revealed that overall prevalence of diabetes was 26.3%. This shows that there are 27.4 million people aged 20 and above with diabetes. The prevalence was slightly high in urban (28.3%) than rural area (25.3%). Similarly, overall glycemic dysregulation (diabetes, pre-diabetes) was also high (43.8%) in urban areas as compared to rural area (39.2%).

The Survey also revealed that 43.9% were obese (44.6 million), 46.2% (48.1 million) were hypertensive and 39.3% had high cholesterol level (41.3 million).

These findings of the Survey imply that diabetes and its associated conditions i.e. obesity, hypertension and dyslipidemia have reached to epidemic proportions.

5. RECOMMENDATIONS

Diabetes is one of the major leading causes of morbidity and mortality. Its prevalence in the country was 8.7% in 1st National Diabetes Survey which is now 26.7%; indicating a massive increase in the burden. Keeping in view this, it is important to take immediate measures for its prevention and control.

The following are suggestions which may be considered for future planning.

- There is a need to create awareness among masses about the causes and risk factors of the diseases i.e. unhealthy diet and lack of physical activity. To achieve this task, print and electronic media may be used. Similarly, community elders, religious leaders may also be encouraged to play their role for creating awareness among common people to adopt healthy life styles.
- Education departments (Federal and Provincial) need to be involved for promoting physical activity in schools and colleges so that next generation may adopt healthy lifestyle. Similarly, information about the risk factor and prevention of diabetes may be added in the curriculum of primary and secondary level. Besides this, the administrations of educational institutions must ensure availability of place for physical activity of the students as well as the availability of healthy food items in the cafeteria.
- Effective interventions are required at general population level to increase intake of fruits, vegetables and avoid intake of junk food as well as promotion of physical activities.

- Health care providers may be trained on the diagnostic and treatment of diabetes
 mellitus especially in primary health care units. Besides this, the facilities for early
 diagnosis and treatment of diabetes must be available at all levels including primary,
 secondary and tertiary health care.
- The non communicable diseases (NCDs) unit working at the Ministry of National Health Services, Regulation and Coordination may coordinate with the Provincial Health Departments for formulation of National Action Plan to prevent and control this deadly disease.

6. REFERENCES

- 1. Ogurtsova K, da Rocha Fernandes JD, Huang Y, et al., IDF Diabetes Atlas: Global estimates for the prevalence of diabetes for 2015 and 2040. Diabetes Res Clin Pract. 2017; 128: 40-50.
- 2. Shaw JE, Sicree RA, Zimmet PZ. Global estimates of the prevalence of diabetes for 2010 and 2030. Diabetes Res Clin Pract. 2010; 87(1): 4-14.
- 3. Global report on diabetes. World Health Organization (WHO) 2016.
- 4. International Diabetes Federation (IDF), Atlas 7th edition 2015.
- 5. Fact Sheet NCD (Diabetes). World Health Organization (WHO) South-East Asia Diabetes fact sheet, Department of Sustainable Development and Healthy Environments. 2014.
- 6. Narayan VKM, Zhang P, Williams D, et al. How should developing countries manage diabetes? Canadian Medical Association Journal. 2006; 175(7): 733-736.
- 7. Global health risks. Mortality and burden of disease attributable to selected major risks. Geneva, World Health Organization, 2009.
- 8. Sharma SK, Ghimire A, Radhakrishnan J, et al,. Prevalence of Hypertension, Obesity, Diabetes, and Metabolic Syndrome in Nepal, International Journal of Hypertension, 2011, Article ID 821971,
- 9. WHO. A Global Brief on Hypertension. Geneva, Switzerland: World Health Organization; 2013.
- 10. Olives C, Myerson R, Morkdad AH, et al. Prevalence, awareness, treatment, and control of hypertension in United States countries, 2001-2009. PLoS One. 2013;8:e60308.
- 11. Alwan A, Armstrong T, Bettcher D, et al. Global Status Report on Non communicable Diseases. Geneva, Switzerland: World Health Organization; 2011.
- 12. Pradeepa R, Anjana RM, Joshi SR, et al. Prevalence of generalized & abdominal obesity in urban & rural India- the ICMR INDIAB Study (Phase-I) [ICMR INDIAB-3]. The Indian Journal of Medical Research. 2015; 142(2):139-150.
- 13. World Health Organization (WHO). Obesity: preventing and managing the global epidemic. Report of a WHO consultation. (1-253). World Health Organ Tech Rep Ser. 2000; 894: i–xii.

- 14. World Health Statistics. World Health Organization (WHO), Geneva: 2012.
- 15. Shera AS, Jawad F, Maqsood A. Prevalence of diabetes in Pakistan. Diabetes Res Clin Pract 2007; 76: 219-22.
- 16. Shera AS, Rafique G, Khwaja IA, Ara J, Baqai S, King H. Pakistan national diabetes survey: prevalence of glucose intolerance and associated factors in Shikarpur, Sindh Province. Diabetic Med 1995; 12:1116-21.
- 17. Shera AS, Rafique G, Khwaja IA, Baqai S, Khan IA, King H. Pakistan National Diabetes Survey prevalence of glucose intolerance and associated factors in North West at Frontier Province (NWFP) of Pakistan. J Pak Med Assoc 1999; 49:206-11.
- 18. Shera AS, Rafique G, Khawaja IA, Baqai S, King H. Pakistan National Diabetes Survey: prevalence of glucose intolerance and associated factors in Baluchistan province. Diabetes Res Clin Pract 1999; 44:49-58.
- 19. Shera AS, Basit A, Fawwad A, et al,. Pakistan National Diabetes Survey: prevalence of glucose intolerance and associated factors in the Punjab Province of Pakistan. Prim Care Diabetes 2010; 4: 79-83.
- 20. Zafar J, Bhatti F, Akhtar N, et al. Prevalence and risk factors for diabetes mellitus in a selected urban population of a city in Punjab. J Pak Med Assoc. 2011; 61(1):40-7.
- 21. Zafar J, Nadeem D, Khan SA, Jawad Abbasi MM, Aziz F, Saeed S. Prevalence of diabetes and its correlates in urban population of Pakistan: A Cross-sectional survey. J Pak Med Assoc. 2016; 66(8):922-7.
- 22. Basit A, Alvi SFD, Fawwad A, Ahmed K, Ahmedani MY, Hakeem R. Temporal changes in the prevalence of diabetes, impaired fasting glucose and its associated risk factors in the rural area of Baluchistan. Diabetes Res ClinPract 2011;94 (Dec (3)):456–62.
- 23. Jafar TH, Levey AS, Jafary FH, et al,. Ethnic subgroup differences in hypertension in Pakistan. J Hypertens. 2003; 21(5):905-12.
- 24. Non-communicable diseases (NCDs) Risk factors survey, Pakistan. Pakistan Health Research Council 2016.
- 25. Fawwad A, Alvi SF, Basit A, Ahmed K, Ahmedani MY, Hakeem R. Changing pattern in the risk factors for diabetes in young adults from the rural area of Baluchistan. J Pak Med Assoc. 2013; 63(9):1089-93

- 26. Wang Y, Mi J, Shan XY, Wang QJ, Ge KY. Is China facing an obesity epidemic and the consequences? The trends in obesity and chronic disease in China. Int J Obes (London) 2007; 31: 177-88.
- 27. Tabatabaei-Malazy O, Qorbani M, Samavat T, Sharifi F, Larijani B, Fakhrzadeh H. Prevalence of Dyslipidemia in Iran: A Systematic Review and Meta-Analysis Study. International Journal of Preventive Medicine. 2014;5(4):373-393.
- 28. Zahid N, Claussen B, Hussain A. High prevalence of obesity, dyslipidemia and metabolic syndrome in a rural area in Pakistan. Diabetes & Metabolic Syndrome: Clinical Research & Reviews 2008; 2(1):13-19
- 29. Iffat W, Rahim N, Shakeel S, et al,. Prevalence of dyslipidemia through screening among karachities. International Journal of Current Research. 2017; 9(06): 52796-52799.
- 30. Pakistan Bureau of Statistics. Population Census 2017. Available at: http://www.pbscensus.gov.pk/sites/default/files/Population_Results.pdf (last assessed on October 10, 2017)
- 31. Daniel WW, editor. 7th ed. New York: John Wiley & Sons; 1999. Biostatistics: a foundation for analysis in the health sciences.
- 32. Pakistan Demographic and Health survey. National Institute of Population Studies Islamabad, Pakistan and MEASURE DHS ICF International Calverton, Maryland, USA. June 2013.
- 33. Multiple indicator cluster survey manual. Chapter 4: Designing and selecting the sample. Available at: mics.unicef.org/files?job...sha=3d97a05358bb0e37 (last assessed on October 12, 2017)
- 34. Fawwad A, Sabir R, Riaz M, Moin H, Basit A. Measured versus calculated LDL-cholesterol in subjects with type 2 diabetes. Pak J Med Sci. 2016; 32(4):955-60.
- 35. Definition and diagnosis of diabetes mellitus and intermediate hyperglycemia: report of a WHO/IDF consultation. World Health Organization 2006
- 36. American Diabetes Association (ADA). Standards of medical care in diabetes—2016. Diabetes Care. 2016;39 (suppl 1):S1-S106.
- 37. Executive Summary of The Third Report of The National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, And Treatment of High Blood Cholesterol In Adults (Adult Treatment Panel III). JAMA. 2001 May 16; 285(19):2486-97.

7. APPENDICES

Appendix I

Cluster (district) and sub-clusters (tehsils/towns) distribution

| Urban (3711) | | Rural (6986) | | |
|------------------|-----------------------|---------------------|----------------------|--|
| District | Tehsils/town | District | Tehsils/town | |
| 1 ((0) | Attock (30) | Islamabad (40) | Zone-IV (40) | |
| Attock (60) | Fateh Jung (30) | | | |
| Rawalpindi (390) | Rawalpindi (195) | | | |
| Kawaipinai (370) | Gujar Khan (195) | | | |
| Islamabad (80) | Zone-I G6 (80) | | | |
| Lahore (1110) | Nishtar town (555) | Gujranwala (1090) | Khilaishahpur (545) | |
| Lanore (1110) | Iqbal town (555) | Oujranwara (1090) | Aroop town (545) | |
| Multan (290) | Mumtazabad (145) | Hafizabad (390) | PindiBhattiyan (390) | |
| Withtair (290) | Shershah (145) | Sahiwal (990) | Okara (495) | |
| | | Saniwai (990) | Chichawatni (495) | |
| | | Muzaffargarh (1490) | Kotaddu (1000) | |
| | | Muzaffargarh (1480) | Alipur (480) | |
| Larkana (70) | Larkana (70) | Vhoimpun (510) | Nara (255) | |
| Nawabshah (40) | Nawabshah (40) | Khairpur (510) | Kotdiji (255) | |
| | | Thomaskas (290) | Mithi (190) | |
| | | Tharparkar (380) | Chachoro (190) | |
| | | Dodin (440) | Talhar (220) | |
| | Orangi town (397) | Badin (440) | Matli (220) | |
| Karachi (1191) | North Nazimabad (397) | | | |
| | Korangi town (397) | | | |
| Charsadda (40) | Charsadda (40) | Buner (290) | Daggar (145) | |
| D 1 (220) | Town-I (110) | | Gagra (145) | |
| Peshawar (220) | Town-II (110) | Mansehra (630) | Balakot (630) | |
| Kohat (40) | Kohat (40) | | Risalpur (124) | |
| | | Nowshera (370) | Nizampur (124) | |
| | | | Rashakai (124) | |
| Mastung (40) | Mastung (40) | Sibi (84) | Lehri (84) | |
| Lockello (140) | Hub City (70) | | Hub Goth (86) | |
| Lasbella (140) | Bela (70) | Lasbella (258) | Bella Goth (86) | |
| | | | Gadani Goth (86) | |
| | | Khuzdar (34) | Nall (34) | |

Appendix II

Informed consent (English)

Information and agreement paper between the participant and the researcher

Dear Sir/Madam,

Baqai Institute of Diabetology and Endocrinology (BIDE) has planned a nationwide survey to determine the prevalence of type 2 diabetes in Pakistan. The survey will be called Diabetic registry of patients with type 2 diabetes (DROP-2).

This letter of information is to invite you to participate in this survey. This project is being carried out with the permission from the ethical committee of our institute. You are selected randomly as a possible participant in this study. The result of the study will serve as the baseline data for further research regarding diabetes care in Pakistan. Moreover, the main aim of this survey is to find the prevalence of diabetes in Pakistan. To do this, we have to collect some information from you and also need to conduct some physical and clinical examinations which will help to fulfill the objectives of the project with your kind permission.

From every consented participant, 7ml blood will be collected in fasting condition for biochemical analysis (HbA1c, FBG and lipid profile). Additional 2ml blood (RBG) will be drawn after 2 hours of 75gram oral glucose intake. You are free to choose either to participate or not to participate in this project. You can trust that any information you will give us, including the results of your examinations, will all be treated very confidentially and will only be used for research purpose only. For that reason, it is important that the information you give be as correct and truthful as possible. You also have the right to withdraw from the project at any time during the study without providing reasons. This will have no negative consequences on you.

Maybe you cannot receive too much direct benefit from taking part in this research, but it will have a favorable impact on public health in future. Your participation will contribute greatly to our research. We thank you in advance for agreeing to help us out. If you have any question on concern about the study, please contact with us without hesitation.

| I have received the information about the study. I hereby | nave received the information about the study. I hereby consent to participate in it. | | | |
|---|---|--|--|--|
| Signature of the participant | Dated: | | | |

| Signature of the Principal investigator | Dated: |
|---|--------|
| Signature of the person obtaining Consent | Dated: |

• Any questions you may have will be answered by:

Name: Dr. Asher Fawwad

Phone Number: **021-36707179**

• In case of a research-related emergency, call:

Day Emergency Number: 021-36707179

Night Emergency Number: 021-6688897

Informed consent (Urdu)

فارم برائے تحریری رضامندی

محققین(researches) اور امیدوارے درمیان میں مطالعاتی پروگرام

محرم جناب / محرمه جناب عاليه!

پاکستان میں بقائی اسٹیوٹ آف ڈایا بیٹولوتی اینڈ انڈ وکر اکٹولوتی نے تو می سطح پر ذیا بیطس کی قتم دوم (Type II Diabetes) کا جائزہ لینے کا منصوبہ بنایا ہے۔ ذیا بیطس کے مریض کا بیجائزہ" قتم دوم ذیا بیطس کا عمل اندرا ہے (Diabetic Registry of Patient with Typ) کہایا جائےگا۔ DROP-2 یا 2 Diabetes) کہایا جائےگا۔

اس فارم کے ذریعے آپ سے اس جائزے میں شرکت کی اجازت کی جاتی ہے۔ اور ساتھ ہی آپ کو آگاہ کیا جاتا ہے کہ یہ project ادارے فی معارف survey کی ethical committee کی اجازت سے لیا جائےگا اس study میں مکندا میدوار کی حیثیت سے لیا جائےگا اس ethical committee کی اجازی اس متعلق معلومات جمع کرنا ہے تا کہ پاکستان میں ذیا بیطس کی احتیاط کے توالے سے مزید research کی متعلومات آپ جا سکے اور ذیا بیطس کے پھیلاؤ (Prevelance) کورو کئے کیلئے مکندا قد امات کئے جا سکے اسکے لئے آپ کے جرپورتعاون سے ہمیں پھیمعلومات آپ سے حاصل کرنا ہوگی اور آپ کے پچیج جسمانی (Physical) اور طبی (test (Clinical) کینے ہو نگے آپ کا بیتعاون ہمارے Project کے مقصد کو بردا کرنے میں مددگار ثابت ہوگئے آپ کا بیتعاون ہمارے Project کے مقصد کو بردا کرنے میں مددگار ثابت ہوگا۔

ہررضا مندامیدوارے نہارمنہ اس 7 خون کو حیاتی کیمیائی تجزیہ (RBG, HbA1c, Lipid Profile) کے لیے بھے کیا جائے گا اور پھر 75 کرام گلوکوز کی مقدار پالے نے کے گئے بعد منز پیا 10 نوب کو بھر (RBG) ایاجائے گا۔ آپ اس Project کے دوران آزادر جیں گے گہ آپ حصہ لیس گرام گلوکوز کی مقدار پالے نے کے گئے بعد منز پیا 10 علی جو معلومات بھی جمین ویں گے جس بھی آ کے Bio Chemical Test Result بھی شامل جیں۔ ان بیانہ لیس۔ آپ بھر وسد کر بھتے ہیں گیا آپ جو معلومات بھی جمین ویں استعمال کیا جائے گا۔ اس کی وجہ سے بین مروری ہے کہ جو معلومات بھی آپ جمین ویں وہ جہاں سے کوخفیہ طور پر صرف Project بین ہو تھا۔ آپ کے پاس یہ افتحار بھی ہوگا کہ آپ جب چاہیں بغیر کوئی وجہ بتائے اس Project سے نگل سکتے جس سال کیا جائے گا۔ اس کا آپ پر کوئی منفی اثر نہیں پڑیگا۔ بوسکتا ہے کہ بطاہر آپ کو اس Research سے کوئی فائد و حاصل نہ ہو مگر مشتقبل میں عام لوگوں پر اس کا مفید اثر پڑیگا۔ آپ کی شرکت ہمارے Project میں قابل احترام ہوگی۔ ہم آپ کی مدد کے بہت زیادہ شکر گڑا رہیں۔ اگر اس میں جو جو بیانی فرما کر بغیر کسی جھوک کے ہم سے دابط کریں۔

| ام امیدوار: | ومتخط اميدوار: |
|--------------------------------|---------------------|
| ارخ: | |
| | |
| نیل انوینفکیز کے دست <u>خط</u> | : * 72/5 |

تارخ:

رمنامندی قارم <u>لینے والے کے دیتھا:</u> موجود ہمعلو ماتی فررا کع

اس تحقیق ہے متعلق مزید سوالات کے جواب کے لئے رابط کریں نام: ڈاکٹراشعرفواد فون921-36707179

تحقیق (Research) معتقلق ایرجنسی میں رابط کے لئے

ون کے وقت ایر جنی تمبر 36707179-021 رات كے وقت ايرجنى نمبر36688897-021

| Data Collecti | ion Form (English) Appendix III | | | | | |
|--|--|--|--|--|--|--|
| Punjab Sindh Khyber Pakhtun Baluchistan | Survey no.: Date (dd-mm-yy): District: | | | | | |
| DEMOGRA | APHY | | | | | |
| Participant's 1 | Participant's name: Father's name: | | | | | |
| Address: | | | | | | |
| Date of birth (| (dd-mm-yy): Age (years): | | | | | |
| Gender: | Male 1; Female 2 | | | | | |
| Community: | Urdu speaking 1; Punjabi 2; Sindhi 3; Pathan 4; Balochi 5; Pashtuns 6; Seraiki 7; Others 8 | | | | | |
| Education: | Illiterate 1; Can read/write 2; Primary 3; Secondary 4; Intermediate 5; Graduate 6; | | | | | |
| | Post graduate 7 | | | | | |
| Occupation: | Administrative/professional 1; Business 2; Skilled labor 3; Manual labor 4; Home duties 5; | | | | | |
| | Unemployed 6; Pensioner 7 | | | | | |
| Addiction: | Tobacco: Yes 1; Ex-addict 2; No 3 | | | | | |
| | Alcohol: Yes 1; Ex-addict 2; No 3 | | | | | |
| Exercise: | Sedentary 1; Light 2; Moderate 3; Heavy 4 | | | | | |
| Marital status | : Single 1; Married 2; Divorced/separated 3; Widowed 4 | | | | | |
| FASTING S | STATUS | | | | | |
| How many hou | ars since last food or drink, except water? (in hours) | | | | | |
| MEDICAL | HISTORY | | | | | |
| Has a doctor ev | ver told you that you have diabetes? No 1; Yes 2; Don't know | | | | | |
| (If yes, year of | diagnosis:) | | | | | |
| Present, regular | r treatment of diabetes | | | | | |
| None (y/n) | Diet (y/n) Herbal(y/n) Oral drugs (y/n) Insulin (y/n) | | | | | |
| (if drugs or ins | ulin, specify type:) | | | | | |
| Have any of yo | our first degree relatives ever had diabetes? No 1; Yes 2; Don't know 3 | | | | | |
| (Specify family | y member(s):) | | | | | |

| Has a doctor ever told you that you have high blood pressure? | No 1; Yes 2; Don't know 3 |
|---|--|
| If yes, are you currently, regularly taking medication? | No 1; Yes 2; Don't know 3 |
| (Specify type:) | |
| Have any of your first degree relatives ever had high blood press | ure? No 1; Yes 2; Don't know 3 |
| (Specify family member(s):) | |
| Usual physical activity, occupational So | edentary 1; Light 2; Moderate 3; Heavy 4 |
| Usual physical activity, leisure | |
| Never 1; less than once a week 2; $1-2$ t | imes/week 3; More than 3 times/week 4 |
| OBSTETRIC HISTORY (women only) | |
| How many babies have you had born alive? | number |
| How many babies have you had born dead (stillbirths)? | number |
| How many miscarriages have you had? | number |
| How many surgical abortions have you had? | number |
| EXAMINATION | |
| Height (cm) We | eight (kg) |
| Waist circumference (cm) | o circumference (cm) |
| Blood pressure (mmHg) | |
| Systolic First | Second |
| Diastolic First | Second |
| BLOOD CHEMISTRY | |
| GTT – Overnight fasting (mg/dl) | GTT – After 2hrs glucose (mg/dl) |
| Cholesterol (mg/dl) | Triglycerides (mg/dl) |
| Low density lipoprotein (mg/dl) | High density lipoprotein (mg/dl) |
| HbA1c (%) | |

Data Collection Form (Urdu)

| 12-14 | دیا بیطس کا <i>ہ</i> | باناب ا |
|-------------------|--|--|
| ءرڻ: ٠ | | منده اير بالاتفاد |
| ضلع | | بلونچستان اسلام آباد |
| ويموكرافي | | |
| اميدواركانام: | | |
| شاخق كارونبر | | موبائل فبرز المساليات |
| :=; | | W 2 - 24 W. |
| تاريخ پيرائش: | | - D- 9 |
| جش: | مرد 1 ، الديت 2 | and the second s |
| پرادرگی: ژن | | ، بلوپی 5، پختون 6، مراکی 7، دوبرے 8 |
| تعليم: | جال 1، لكوارد علماب 2، يراتمرى 3، | 4 ، اعزمیڈیٹ 5، کرنجیٹ 6، |
| | پاسکریج پیت 7 | |
| - | پیشدور 1، کاروبار 2، بشرمندمزدور 3، وکر ونلینه یاب ملازم 7، طالبعلم8 | ا 4 ، فحرك كام كائ 5 ، يروزگار 6 ، |
| ك: | وقعيد ياب الأرام (، عا جسم ه تمباكو: بال 1، يسلي 2، ثو | |
| . = | عبور بال 1 يبلي 2 م أم شراب بال 1 يبلي 2 م أم | |
| از دوا آق حثیت : | غیرشادی شده 1، شادی شده 2، طالق یافته! | 4 |
| ررون دردش: | يرون در مياني د بلکي در درمياني د ، جماري 4 | 7.77. FW 17 |
| | SEEDING COMMITTER STATE STREET | 1 |
| نهادمند: ا | تحقی در پہلے آپ نے کھایایا بیاہے یائی کے علاو | |
| مجى معائف كيا | بحى آپ كوا اكنز نے بتايا كه آپ كوشوگر ؟ | فين 1، بان 2، عشين 3 |
| Ti) | رې توکې په ځا د د د د د د د د د د د د د د د د د ا | |
| | آپ مستقل علاج کرواتی بین؟ | |
| | اپ ن علاق خداق این: پ | ادويات انولين |
| | 내라면하 (~~)() | |
| | رکوئی دولیا انسولین ہے تو نام ہتا کیں۔۔۔۔۔۔ آپ کے خوٹی رشتے میں کے کی کوشوکرہے؟ | نين 1، بال 2، پيش |
| Principle Company | 15 183125 Just 11382 | 31 14 24 56 1 14 51 1 14 |

| | نیں 1، بال 2، پیٹیں 3 | كيا آپ كوكى داكش في بناياك آپ كوبلذي يشرب؟ |
|---|---|---|
| | فين 1، بال 2، يعين 3 | اگر ہے و آپ متقل ون کی دوالے دے ہیں؟ |
| | | (نام اختم بتائمیں۔۔۔۔۔) |
| | ديس 1، إلى 2، عِدْيِس 3 | كياآپ كۈنى رائىية مى كى كوبلى پريشر ب؟ |
| | | (اگر ہے قونام داننے کریں۔۔۔۔۔۔) |
| | بينص ربنا 1، باكا 2، اعتدال يهند 3، جمارى 4 | عام طور پر کام کے وقت جسمانی سرگری کیا ہے؟ |
| | | فارخ وقت میں جسمانی سرگری کیا ہے؟ |
| | 1-2مرتبد 3 ، التي من 3 عناد مرتبد 4 | كونين 1 ، تفتي ش اليك دفعد ا كم 2 ، تفتي من |
| (| (افتياري | حل كم متعلق موالات (مرز شادى شداقرا أن ك ك) |
| | تغداد | آپ کے کتے بچے زندہ بین؟ |
| | تعداد | آپ ك كنت يج بيدا بوت اي الحت بوك |
| | تعداد | آپ کے کتنے بچ خود ضائع ہو گئے؟ |
| | تعداد | آپ نے کتے بچے شائع کروائے؟ |
| | | يأش |
| | وزنو | cm &u |
| | cm | cm |
| | | فشارخوان |
| | mmHg // mmHg // | سئولک پېلا 🗆 |
| | mmHg mmHg | ۋايىشۇنگ پېيلا 🔲 |
| | | خون كشيث |
| | GTT گلوکوز کردو گفتے بعد | GTT نهارت |
| | ژاینگر انڈ(mg/dl) | کولسٹرول (mg/dl) |
| | باقی وشی کیپور و مین (mg/di) | لوژینسی لیچو پروغمن(mg/dl) |
| | | (%)HbA1C |
| | | |

Appendix IV

Specimen Collection and Handling

- 1. Wash your hands before and after each sample collection.
- 2. Identify the patient.
- 3. Verify that the patient in fasting.
- 4. Assemble necessary supplies like vacutainer needle with holder, vacutainer tubes (Nafl for glucose, gel for lipid profile and EDTA for HbA1c), sprite or alcohol swab, gauze pad, bandage and gloves.
- 5. Reassure the patient. Be calm and confident.
- 6. Position the patient in a chair and explain the procedure to patient.
- 7. Verify paperwork and select the tubes.
- 8. Ensure the patient's hand is closed.
- 9. Select a vein site.
- 10. Cleanse the vein puncture site with spirit in a circular motion and allow drying.
- 11. Apply or tied the tourniquet (No longer than one minute).
- 12. Inspect the needle. Use safety equipment.
- 13. Perform vein puncture using the correct order of the draw i-e first gel, then EDTA and in last Nafl tube.
- 14. Rotate Nafl and EDTA tubes gently to mix.
- 15. Do not rotate or mix gel tube.
- 16. Ensure patient's hand is open.
- 17. Release and remove the tourniquet.
- 18. Place the gauze pad over the puncture site.
- 19. Remove the needle. Never recap needle.
- 20. Apply pressure to vein puncture site until bleeding has stopped.
- 21. Bandage the patient's arm.
- 22. Dispose of the needle unit in a puncture resistant container.
- 23. Label the tubes and record the date and time of collection; collector's initials, name of patient, and lab number.
- 24. Clean up all supplies and waste to discard in appropriate containers.

- 25. In case on non-diabetic patient, give a load of 75gms glucose in 250-300 ml of water orally to patient and note time with first sip.
- 26. Instruct patient to sit for two hours and do not perform any physical work.
- 27. Send properly labeled blood collection tubes to the laboratory with appropriate slips.
- 28. After 2 hours of oral glucose draw RBS sample with same precaution and procedure.
- 29. Leave patient courteously.
- 30. Send properly labeled blood RBS sample tube to the laboratory with appropriate slip.

Separation of Serum / Plasma

- 1. Check the sample and enter in file/register.
- 2. Allow to clot the gel containing tube. 30 min at room temperature or 15 min in water bath at 37°C.
- 3. Centrifuge gel and Nafl tubes for 10-15 minutes at 3500-4000 rpm.
- 4. Then separate serum / plasma from cells in pre-labeledeppendorf tube.
- 5. Whole blood sample can be save in same tube or in eppendorf tube.

Specimen Storage

- 1. Serum / plasma will be stable on ambient temperature for 2-3 hours, in refrigerator for 1 week, at -20° C for 60 days and at -80° C for 2-5 year.
- 2. Refrigerate EDTA tube immediately. Whole blood sample can be stored at 2-10°C for 1 week, at -20°C for 60 days and at -80°C for 2-5 year.

| TEST | VACUTAINER | AMOUNT | INSTRUCTIONS | STORAGE |
|------------------|--|--------|--|--|
| Glucose | Grey top (Vacutainer containing Sodium fluoride) | 2 ml | Mix the tube contents by gently inverting the tube five times. Centrifuge for 10-15 minutes at 3500-4000 rpm. Then separate plasma from cells. | PLASMA: Ambient 2-3 hrs Refrigerated for 1 week -20°C for 60 days -80°C for 2-5 year |
| Lipid Profile | Yellow top (Gel containing Vacutainer) | 3-5 ml | Allow to clot. (do not mix)Centrifuge for 10-15 minutes at 3500-4000 rpm. Then separate serum from cells. | SERUM: Ambient 2-3 hrs Refrigerated for 1 week-20°C for 60 days -80°C for 2-5 year |
| HbA1c | Purple top (Vacutainer containing EDTA) | 2 ml | Mix the tube contents by gently inverting the tube five times. | Refrigerated immediately -20°C for 60 days -80°C for 2-5 year |

Transportation

- 1. **Primary receptacle:** Seal the lid of Eppendorf tube properly with the help of sticking tape.
- 2. **Secondary receptacle:** Place the sealed Eppendorf tube in durable, watertight, leak-proof rack. Several wrapped primary receptacles may be placed in one secondary receptacle. Seal the rack with lid.
- 3. **Tertiary receptacle or outer shipping package:** The secondary receptacle is placed in an outer shipping package with ice pack or dry ice which protects it and its contents from outside influences such as physical damage and water while in transit.

Appendix V

Testing Procedures

Procedure for Glucose, Cholesterol, Triglyceride, Hdl–Cholesterol (Direct), LDL–Cholesterol (Direct) and HbA1c Testing

Glucose (fasting / random) by GOD PAP, Cholesterol by CHOD-PAP, Triglyceride by GPO-PAP, HDL—cholesterol by Direct and LDL—cholesterol by direct method as follow

- 1. Perform everyday usage procedure of Selectra Pro S (Start up, calibration and controls).
- 2. Give sample ID to Selectra Pro S, select tests and allot a test number to specimen.
- 3. Place sample on sample trey at appropriate number.
- 4. Start analyzer and wait for results.
- 5. Note the result.

HbA1c by HPLC method:

- 1. Perform everyday usage procedure of D-10 (Start up, calibration and controls).
- 2. Set samples in sample rack.
- 3. Mix samples by gently inverting rack 3-5 times.
- 4. Insert samples in window of D-10.
- 5. Allow D-10 to scan samples.
- 6. Give sample ID according to position number in rack.
- 6. Start analyzer and wait for results.
- 7. Note the result.

Glucose (fasting / random) by GOD PAP on semi automated analyzers:

Take 4 test tubes and label as blank, standard, control and sample #

| | Blank | Std | Control | Test |
|-----------------|-------|-------|---------|-------|
| Reagent | 1 ml | 1 ml | 1 ml | 1 ml |
| Distilled water | 10 μ1 | | | |
| Standard | | 10 μ1 | | |
| Control sample | | | 10 μ1 | |
| Serum | | | | 10 μ1 |

Mix & incubate for 10 min at 37°C or for 15 min at room temperature.

Read at appropriate program.

Appendix VI

Ethical Clearance



National Bioethics Committee (NBC) Pakistan



Date: 36th January, 2017

Ref: No.4-87/17/NBC-226/NBC/ > L64

Passen

Minister of State, Ministry of Statement Health Services Regulation

Chaleperate

Secretary Afairmry of 30158/4/C. Operational of Pairiess

NASSAC, Compressed of Pakinter Pakister Health Rysupols Countil Pakis der Startte Romanch Committel Schendung E. 4.300-bit Persidente, College of Physiciania and Surgeon of Pakistan. Presidente, Pakistan Mudical and Ummit Connell, President Provident, Pakistan Association of

Executive Director, Polistan Hashin Broarch Countyl, Number Superlary WHO County Representative Provident, Separate Count Day

DG5 to G59 Surgran Graeral

Director General Bealth, Purple Director General Bealth, Sindle Director General Health, Khyber Director Houlth Services, FATA Director General Bestule.

Discussion Comment Beatch, A.J.C.

Singletrary Pakestan Norsing Cou

Manhers

Penf, Dr. Aselin Ahmad (Chromosomor)

Prof. Dr. Farius Montain

Prof. Dr About Akhter Salorest

Frot Dr Abstol Rorring Sable Dr. Amoir Mintafa Jufacey

Dir Assessitälleb

He Farult Oadle

Dr. Neboun About Tipo

für, Salma Pervais Jahal

Dr. Janished Akhtar

Dr. Forkhanda Ghafaor

Dr.Imgarullah Nemen

Br.Manter Arrest Klain Mr. Abdul Ghard Sauch

Prof. Dr. Abdul Basit

Bagai Institute of Diabetology and Endocrinology (BIDE) Bagai Medical University,

Plot No.1-2, II B. Nazimabad No. 2

Karachi,

Subject: Prevalence of type 2 diabetes in Pakistan.

Dear Dr. Prof. Abdul Basit,

This is with reference to your email dated 10th November 2016, requesting ethical review / approval of above cited project.

I am pleased to inform you that the above mentioned project has been cleared by the "Research Ethics Committee" of "National Bioethics Committee" for a period of one year.

For the continuation of project in the next years, you have to send a progress report and a formal request asking for continuation of projects (however, you do not need to submit REC application or pay any processing fee again).

Kindly keep the National Bioethics Committee, Secretariat updated about the progress of the project and submit the formal final report on completion.

Yours sincerely

(Prof. Dr. Assim Ahmad) Chairman

NBC- Research Ethics Committee

Pakistus Health Research Council, Stabula-e-Jamburat, Off Constitution Avenue, Sector G-572, Islamahad www.chepakaman.org.pk. e-mail:phepakastan.org//gmmil.com Fel: 92-51-9224325, 9216793, Fox 9216774. Gallery Appendix VII











