



Prevalence of metabolic syndrome among Roma: a comparative health examination survey in Hungary

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**Prevalence of metabolic syndrome among Roma: a comparative health examination survey in
Hungary**

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ABSTRACT

Objectives. The objective of our study was to compare the health status of the Roma people with that of the general population in Hungary.

Methods. A health examination survey to define the prevalence of metabolic syndrome and its components was performed in a representative random sample (n=646) of the Roma population aged 20-64 years living in segregated colonies and data were compared to that obtained in a representative random sample (n=1819) of the Hungarian population.

Results. The risks for central obesity, hypertension and raised triglyceride level among Roma adults were not differed from the Hungarian references, while raised fasting plasma glucose or known type 2 diabetes mellitus (OR=2.65, 95%CI 1.90-3.69), reduced HDL cholesterol level or treated lipid disorder (OR=2.15, 95%CI 1.65-2.79) were significantly more frequent in all age groups in the Roma sample. The prevalence of metabolic syndrome (OR=1.37, 95%CI 1.03-1.83) was also significantly higher among Roma than in the general Hungarian population.

Conclusions. Besides tackling the socioeconomic determinants of the poor health of Roma people, specific public health interventions considering increased genetic susceptibility to metabolic disturbances are needed to improve their health status.

KEYWORDS Roma, health examination survey, metabolic syndrome, genetic background

INTRODUCTION

Roma are the largest ethnic minority in Europe with an estimated number between 12 and 15 million¹. Their representation in the population is greatest in Bulgaria, Romania, Slovakia, Hungary, the Czech Republic and Slovenia, but the EU enlargements of 2004 and 2007 has enabled increasing numbers to migrate into and to settle in other countries of the EU. The Roma are concentrated in economically deprived regions, often living in segregated parts (colonies) characterized by severely unfavorable environmental conditions of human habitats.² Independently of the country where they live in, the common problems that this population group experiences are poverty, restricted access to education, high level of unemployment and social exclusion.³

On the base of the predominantly low socio-economic status of the Roma population and socio-economical status as determinant of health, it is a reasonable assumption that their health status is much worse and their average life expectancy is much shorter than that of the majority population, as is frequently mentioned both in research and public communications.^{4,5,6,7} Considering the fact that recording Roma ethnicity is not permitted in any kind of official documentation including medical records, birth and death certificates⁸ as well as some major obstacles that hinder or prevent the collection of reliable data on Roma and other minorities,⁹ these cannot be considered as evidence proved.

In a longitudinal study covering the entire population of Bulgaria between 1992-98 in 2011 Kohler and Preston¹⁰ presented “the first reliable life table measures and cause-specific mortality indicators according to ethnicity and religion” by linking data in the 1992 census to subsequent death records. Although identification of Roma found to be the least reliable among the groups considered and they were most likely to be misclassified resulting in undercounting, their mortality was found to be very high compared to all other ethnic/religious groups according to nearly all major causes of death.

Although many studies have documented high prevalence of communicable diseases,^{11,12} fewer have documented non-communicable diseases among the Roma people (see reviewed¹³). Even fewer

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3 studies have compared Roma health status with that of the majority population, and even if
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5 comparisons were made they were restricted to one or only a few (hypertension, diabetes) health
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7 indicators.^{14,15} In addition, the comparative studies were mainly questionnaire-based health interview
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9 surveys when self-assessed health status and functional limitations were considered as outcome
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11 indicators, and no medical examinations were carried out. The ethnic identification of individuals can
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13 also be contested.¹⁶

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15 The objective of our study was to compare the health status of the Roma people with that of the
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17 general population in Hungary. We conducted a study measuring the prevalence of metabolic
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19 syndrome and its components which overcomes the limitations with the existing evidence-base on
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21 Roma health noted above.

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23 - First, it is a definitive health examination survey from the epidemiological point of view.
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25 - Second, it targets health status of Roma in complexity by investigating the prevalence of
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27 metabolic syndrome as well as the prevalence of its components as it was defined by the
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29 International Diabetes Federation Consensus Group.¹⁷ Although the definition and clinical
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31 interpretation of metabolic syndrome is a subject of intense scientific discussion¹⁸ there is a
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33 general agreement that it is the most robust predictor of the increased susceptibility to
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35 different non-communicable diseases (cardiovascular diseases, type 2 diabetes, polycystic
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37 ovary syndrome, fatty liver, cholesterol gallstones, asthma, sleep disturbances, and some
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39 forms of cancer)¹⁹ most of them with high morbidity and mortality burden.
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41 - Third, data are compared with reference data of the majority population in Hungary.
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43 - Fourth, by involving representatives of the Roma population at all stages of the study high
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45 level validity has been reached through avoiding the misclassification of the study subjects
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47 and low response rate of selected Roma adults.
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METHODS

Sampling

A nationwide project surveyed the segregated colonies (SCs) in Hungary. Roma field workers nominated by the Roma non-governmental organizations identified the SCs. 94% of SCs' inhabitants declared themselves to be Roma.²⁰

The present investigation utilized this colony-registry in stratified multistep sampling. The study embraced 2 Hungarian counties (Hajdu-Bihar and Szabolcs-Szatmar-Bereg), where Roma colonies are accumulated. To focus the investigation to the highly segregated, closed Roma population, SCs with more than 100 inhabitants were considered as the study base. Of the 64 eligible SCs 40 and 25 households from each SC were randomly selected using the General Practitioners' validated (and corrected, if it was needed) household-lists. The 20-64 years old inhabitants of the resulting 1000 households comprised the final sampling frame, and 1 person from each household has been chosen by a member of the primary health care team using random table.

Roma Health Examination Survey

The participants were invited to the GPs' office where a questionnaire on socio-demographic factors, life-style and self-assessed health status was completed by GPs or practice nurses on the basis of interviewees' answers, and physical examination was carried out. The health status description utilized the former medical records of participants as well. Blood samples for laboratory investigations were taken. Informed consent from the participants was obtained. The data collection started in September 2011, and took 4 months, and it applied the methods of a former study on ~~the~~ metabolic syndrome (Hungarian Metabolic Syndrome Survey, HMSS).²⁰

Socio-demographic characteristics (age, gender, level of education), results of physical examination (body weight, height, waist circumference and blood pressure), serum concentrations of triglyceride,

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3 HDL-cholesterol and glucose assessed in fasting blood samples, and the medical history of lipid
4 disorders, hypertension and type 2 diabetes mellitus have been processed in the present investigation.

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8 The consensus definition of the International Diabetes Federation was used to determine the presence
9 of metabolic syndrome.¹⁷ Applied thresholds: central obesity ≥ 94 cm for men and ≥ 80 cm for women;
10 serum triglyceride concentration ≥ 1.7 mmol/l or specific treatment for it; serum HDL cholesterol
11 concentration < 1.03 mmol/l for men and < 1.29 mmol/l for women or specific treatment for it, systolic
12 blood pressure ≥ 130 mmHg or diastolic blood pressure ≥ 85 mmHg or specific treatment for it, fasting
13 plasma glucose level ≥ 5.6 mmol/l or previously diagnosed type 2 diabetes mellitus.

20 21 **Reference data set**

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24 The SCs' data have been compared to reference values determined by the above mentioned HMSS on
25 a representative sample of the Hungarian population (n= 1819).²⁰ The present investigation utilized the
26 1542 complete records of 20-64 years old adults from HMSS as reference dataset.

27 28 29 30 31 **Data analysis**

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34 The SC specific and the HMSS derived datasets ~~were have been~~ joined. The final SC-HMSS database
35 contained anonymized records. The prevalence of metabolic syndrome and that of its components
36 were calculated for different strata (age groups in both genders of the samples investigated). ~~The~~
37 prevalences of the metabolic syndrome and that of its components were estimated for different strata
38 (age groups in both genders of the samples investigated) and their 95% confidence intervals were
39 computed using the normal distribution. The chi square test was used for the comparison of
40 prevalences between both ethnicities and males and females, respectively. The statistical evaluation of
41 observed proportions has been carried out by their 95% confidence interval computed using the
42 normal distribution and by chi square test. The association of risk factors of age, gender, education
43 and ethnicity on the different health outcomes was analyzed using multiple logistic regression models.
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55 Stata 10.1 was used for the applied in statistical analysis.

RESULTS

Because of some non-collaborative GPs, the data collection was impossible in 3 SCs. Therefore, the sample contained 925 persons. The informed consent was signed, the questionnaire was completed, and the physical examination was undertaken by 725 adults. The records with any missing metabolic syndrome related data was excluded from the analysis. Finally, 646 SC-records were analyzed.

There were remarkable demographic differences between SC and Hungarian reference samples (table 1). The male proportion was much lower in studied SCs (39.16%) than in representative Hungarian sample (47.47%). The age distribution of SC sample was shifted towards the younger age groups, and strongly deviated from the Hungarian reference distribution. The level of education was considerably lower among SC inhabitants than the national reference. Both in the general Hungarian and the Roma populations the prevalence of the raised fasting plasma glucose concentration or formerly diagnosed diabetes mellitus, as well as that of the raised triglyceride level or treated lipid disorder were significantly more frequent among males, while the frequency of central obesity was higher among females. In the general population the prevalence of hypertension was higher among males, but the same difference can not be detected in the Roma group (table 2 A and B).

The central obesity ($p<0.001$) and the hypertension ($p<0.001$) were less frequent among SC inhabitants in both genders. Contrary, the reduced HDL cholesterol level ($p=0.029$) and the higher fasting blood glucose concentration ($p<0.001$) were significantly more frequent in the SC sample. The raised triglyceride level was similar in the studied samples ($p=0.084$). Altogether, the observed prevalence data of metabolic syndrome in SC (36.38%) and control Hungarian (34.96%) samples were not deviated from each other ($p=0.525$). (table 2 C, D and E)

The age specific prevalence estimates show that the central obesity and the hypertension were less frequent only among older CS inhabitants. On the other hand, the reduced HDL cholesterol levels and higher fasting blood glucose concentrations were manifested in almost every age group (figure 1).

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3 In multivariate model, the age proved to be significant risk factor for metabolic syndrome (OR=1.06,
4 95%CI 1.05-1.07) and for every of its components (table 3). Apart from the decreased HDL
5 cholesterol level, all the studied outcomes were significantly influenced by gender: the central obesity
6 was more frequent among women; other components and the metabolic syndrome (OR=0.75, 95%CI
7 0.62-0.91) were associated with the male gender. The studied outcomes were independent of the
8 education. Raised fasting plasma glucose or known type 2 diabetes mellitus (OR=2.65, 95%CI 1.90-
9 3.69), reduced HDL cholesterol level or treated lipid disorder (OR=2.15, 95%CI 1.65-2.79), and
10 consequently, the metabolic syndrome (OR=1.37, 95%CI 1.03-1.83) were more frequent among SCs'
11 inhabitants.
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DISCUSSION

Roma, the largest minority population of Europe shows an accumulation in the Central, Eastern and Southern (CES) European countries, so the problems related to this low educated, typically unemployed, marginalized population living in deep poverty were considered as regional challenges of CES countries. The opening of borders in the process of previous and ongoing EU expansion has enabled increasing numbers of Roma to settle in other parts of the EU and has focused attention on the need to address Roma exclusion not only at national, but also international level and has highlighted Roma problems as common European challenges.^{21,22} Recent changes to Canada's immigration legislation (Bill C-31) clearly show that the Roma problems do not respect even the continental borders.

Over the past decades, a series of national and international policy initiatives have been designed to improve the situation of the Roma, and EU-wide policy networks focusing on education, employment, housing and health have also been established to support Roma inclusion.^{21,23} Unfortunately, there has been limited assessment of actual outputs and results of the projects benefiting Roma inclusion and improving Roma health.²⁴ Additionally, new concern has been raised that the current economic crisis may disproportionately affect vulnerable communities, including the Roma.²⁵

Epidemiological studies on health of the Roma people were focused almost exclusively on communicable diseases and reproductive health before the turn of millennium,²⁶ and research has only recently extended to the field of non-communicable diseases and their risk factors. These studies are limited in number, and have severe uncertainties on the identification of Roma ethnicity and are restricted to one or a few indicators. Most of these studies can not be really conclusive, because no comparison was made with the overall population. Some others which make comparison report contradictory findings: some studies show no difference between Roma and non-Roma populations in the cardiovascular disease (CVD) occurrence,²⁷ while others report increased prevalence of various CVD risks among Roma.^{11,12,13,14}

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3 A comparative study of the CVD risk profile for a sample of 430 adult Roma living in rural Croatia
4 compared with those for the general Croatian population was recently published.²⁸ The findings
5 indicated that the Roma population bear a high CVD risk load related to smoking and high glucose
6 level; and a higher prevalence of CVD risks in women and the higher body mass index in younger age
7 group (18-34 years) characteristic for the Roma population were in contrast to the findings in the
8 general population of Croatia. However, although the components were targeted in the study, the
9 prevalence of metabolic syndrome was not defined.
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18 Until now only a single report²⁹ tried to estimate the prevalence of metabolic syndrome among Roma
19 and concluded that it is high. The small sample size (N=77), as well as the method used for sampling
20 (Roma people who visited the GPs with different complaints were included) excludes the possibility to
21 get scientifically acceptable estimates. The reported 50.6% prevalence can be interpreted as an
22 overestimate.
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30 In our present study the sample frame and size was almost identical with those we previously used in a
31 comparative health interview survey³⁰ and data obtained in the Roma sample have been compared to
32 reference values determined in the HMMS.²⁰ The prevalence of every components of metabolic
33 syndrome showed continuous elevation by age in the Hungarian reference population. Similar pattern
34 was observed for Roma only among 20 to 54 years old adults, but further elevation was not observable
35 in the 55-59 and 60-64 yrs age groups, values were unchanged (HDL cholesterol) or slightly decreased
36 (blood pressure, fasting serum glucose, triglyceride), while the prevalence of central obesity became
37 even significantly lower for the 55-59 yrs group of Roma people. The analysis of data with age-group
38 stratification clearly shows that the significantly lower prevalence of elevated blood pressure or treated
39 hypertension, as well as that of central obesity is the effect of their decreased prevalence in the older
40 age groups and not a characteristic of the whole Roma population; it is reasonable to suppose that the
41 decrease is a consequence of the deaths of people with risk factors combined before reaching age of
42 55. Considering the sex-composition of the Roma sample it can be suggested that mainly the men are
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3 The fact that the prevalence of elevated fasting glucose level and that of decreased HDL cholesterol
4 concentration is higher in all age groups of Roma strongly suggest that genetic background exists
5 behind these phenomena. A current review summarizing recently published reports on the genetic
6 architecture of lipid metabolism reports 52 genes responsible for HDL cholesterol level.³¹ In a large
7 scale epidemiological study (Tehran Lipid and Glucose Study) the estimates of age and gender
8 adjusted heritability for abdominal obesity, low HDL cholesterol, high triglyceride, high fasting blood
9 glucose and high blood pressure were 22, 40, 34, 38 and 23%, respectively ($p < 0.05$), i.e. the
10 contribution of genetic factors was highest to the development of low HDL cholesterol (40%) and high
11 fasting blood glucose (38%) levels.³² Concerning genetic determinants of carbohydrate metabolism
12 recent genome-wide association studies have identified and replicated 75 susceptibility loci associated
13 with type 2 diabetes and related metabolic traits (see reviewed in³³). Studies convincingly support the
14 hypothesis on the strong influence of the gene-environment interactions in the development of type 2
15 diabetes³⁴ and also demonstrate interactions of gene variants with measures of dietary intake and
16 exercise.^{35,36} Although it is generally accepted that in case of metabolic syndrome life-style changes
17 are the most adequate therapeutic interventions^{37,38} and presently there is insufficient support for
18 clinical application of gene-based prediction models in metabolic syndrome, there is direction and
19 encouraging progress in a rapidly moving field that is beginning to show clinical relevance.³⁹ It has to
20 be accepted that careful clinical trial programs are needed to determine which HDL raising therapeutic
21 interventions may indeed exert protective effect.⁴⁰

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42 The most obvious limitation of the current health examination survey of Roma people was that it was
43 not representative of the overall Hungarian Roma population. Those Roma who have assimilated with
44 the majority population were excluded, and consequently the survey captured the characteristics of the
45 most disadvantaged part of the Roma population. However, identification of the needs of this group is
46 the most important from a policy perspective. It is also important to note that the random sample
47 representative for the general Hungarian population of the Hungarian Metabolic Syndrome Survey
48 have included some people who are Roma, so it is possible that their inclusion was slightly dilute the
49 true difference between the populations.
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3 Our results show that metabolic syndrome strongly contribute to the development of the poor health
4 status of the Roma population. Among the components of metabolic syndrome decreased HDL-C and
5 elevated fasting blood glucose levels with genetic predisposition are the most prominent findings,
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7 which gives specific importance to the screening for these blood components on a regular basis.
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10 11 12 **Human Participant Protection**

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15 The study has been approved by the Ethical Committee of the Hungarian National Scientific Council
16 on Health (8907-O/2011-EKU, 285/PI/11).
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22
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24 Scientific Council on Health (ETT: 192/2009) and from the TÁMOP-4.2.2.A-11/1/KONV-2012-0031
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26 the Swiss Contribution Program (SH8/1). The contribution of fieldworkers, the Roma advocates and
27 other study personnel is gratefully acknowledged. Literature review was carried out as Activity 3 of
28 the WHO Vulnerability and Health Collaborating Centre's Program.
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37 **Conflicts of interest:** None declared.
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40 41 42 **Contributors**

43 R. Ádány had the original idea for the comparative Roma Health Examination Survey, participated in
44 the questionnaire and sampling design, interpreted the results and wrote the article. Z. Kósa and Á.
45 Moravcsik-Kornyicki participated in the questionnaire and sampling design, performed the sampling
46 and interpreted the results. Z. Szabó specified the diagnostic criteria and methods used to detect the
47 metabolic syndrome's components. J. Sándor planned the sampling, performed the statistical analysis
48 and interpreted the results. J. Diószegi and B. Roberts contributed to the writing of the manuscript.
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Keypoints

- This study shows that the prevalence of metabolic syndrome (MS) is significantly higher among Roma than in the general population in Hungary due to the much higher frequency of raised fasting plasma glucose (or known type 2 diabetes mellitus) and that of reduced HDL cholesterol level (or treated lipid disorder).
- Reduced HDL cholesterol and the higher fasting blood glucose concentrations were significantly more frequent in all age groups of the Roma sample in both genders, which may indicate a genetic background.
- These findings suggest that besides tackling the socioeconomic determinants of the poor health of Roma people, specific public health interventions considering increased susceptibility to disturbances both in carbohydrate and lipid metabolisms are needed to improve their health status.

REFERENCES

-
- 1 European Parliament. Resolution of 31 January 2008 on A European Strategy on the Roma. 2009/C
2
3
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55
56
57
58
59
60
68E06. P6 TA (2008) 0035. Brussels, Belgium: European Parliament; 2008.
 - 2 Kósa K, Daragó L, Ádány R. Environmental survey of segregated habitats of Roma in Hungary: a
way to be empowering and reliable in minority research. *Eur J Public Health* 2011;21:463-8.
 - 3 European Parliament. Resolution of 11 March 2009 on the social situation of the Roma and their
improved access to the labor market in the EU (2008/2137(INI)). Strasbourg, France: European
Parliament; 2009.
 - 4 Ginter E, Simko V, Wsolova L. Fall of the iron curtain: male life expectancy in Slovakia, in the
Czech Republic and in Europe. *Cent Eur J Public Health* 2009;17:171-4.
 - 5 Lenzi L. Roma People in Europe: A Long History of Discrimination. European Social Watch Report
2010. Italy: Amnesty International; 2010.
 - 6 Health and the Roma community analysis of the situation in Europe. Bulgaria, Czech Republic,
Greece, Portugal, Romania, Slovakia, Spain. Madrid, Spain: Fundacion Secretariado Gitano; 2009.
 - 7 Council conclusions on an EU Framework for National Roma Integration Strategies up to 2020.
Council of the European Union, 3089th Employment, Social Policy, Health and Consumer Affairs
Council meeting Brussels; 19 May 2011.
 - 8 Rimarova K. The health of the Roma people in Central and Eastern Europe. Kosice, Slovakia: Pavol
Jozef Safranik University; 2010.
 - 9 Kósa K, Ádány R. Studying vulnerable populations: lessons from the Roma minority. *Epidemiology*
2007;18:290-9.

1
2
3
4 10 Kohler IV, Preston SH. Ethnic and religious differentials in Bulgarian mortality, 1993-98. *Popul*
5
6 *Stud (Camp)* 2011;65:91-113.

7
8
9 11 Kabakchieva E, Amirkhanian YA, Kelly JA, McAuliffe TL, Vassileva S. High levels of sexual
10
11 HIV/STD risk behaviour among Roma (Gipsy) man in Bulgaria: patterns and predictors of risk in a
12
13 representative community sample. *Int J STD HIV* 2002;13:184-91.

14
15
16 12 Casals M, Pila P, Langohr K, Millet JP, Cayla JA. Incidence of infectious diseases and survival
17
18 among the Roma population. A longitudinal cohort study. *Eur J Public Health* 2012;22:262-6.

19
20
21 13 Parekh N, Rose T. Health inequalities of the Roma in Europe: a literature review. *Centr Eur J*
22
23 *Public Health* 2011;19:139-42.

24
25
26 14 Carrasco-Garrido P, López de Andres A, Barrera Hernandez V, Himenez-Trujillo I, Jimenez-
27
28 Garzia R. Health status of Roma woman in Spain. *Eur J Public Health* 2011;21:793-8.

29
30
31 15 Zeljko H, Skarić-Jurić T, Narancic NS, et al. Traditional CVD risk factors and socio-econimoc
32
33 deprivation in Roma minority population of Croatia. *Coll Antropol* 2008;32:667-76.

34
35
36 16 Masseria C, Mladovsky P, Hernandez-Quevedo C. The socio-economic determinants of the health
37
38 status of Roma in comparison with non-Roma in Bulgaria, Hungary and Romania. *Eur J Public Health*
39
40 2010;20:549-54.

41
42
43 17 Alberti KG, Zimmet P, Shaw J for the IDF Epidemiology Task Force Consensus Group. The
44
45 metabolic syndrome – a new worldwide definition. *Lancet* 2005;366:1059-62.

46
47
48 18 Kassi E, Pervanidou P, Kaltsas G, Chrousos G. Metabolic syndrome: definitions and controversies.
49
50 *BMC Med* doi:10.1186/1741-7015-9-48.

51
52
53 19 Grundy SM, Brewer HB Jr, Cleeman JI, Smith SC Jr, Lenfant C for the Conference Participants
54
55 Definition of Metabolic Syndrome. Report of the National Heart, Lung, and Blood Institute/American
56
57 Heart Association Conference on Scientific Issues Related to Definition. *Circulation* 2004;109:433–8.

1
2
3
4 20 Szigethy E, Széles G, Horváth A, et al. Epidemiology of the metabolic syndrome in Hungary.
5
6 Public Health 2012;126:143-9.
7

8
9 21 Measures to promote the situation of Roma EU citizens in the European Union. Brussels, Belgium:
10
11 Directorate-General for Internal Policies, Policy Department Citizens' Rights and Constitutional
12
13 Affairs; 2011.
14

15
16 22 EUR/RC62/8 Health 2020: policy framework and strategy. World Health Organization, Regional
17
18 Office for Europe; 2012.
19

20
21 23 Roma in an Expanding Europe. Challenges for the Future, Declaration of the Decade of Roma
22
23 Inclusion 2005-2015. Bulgaria, Sofia; 2 February 2005.
24

25
26 24 Fésüs G, Östlin P, McKee M, Ádány R. Policies to improve the health and well-being of Roma
27
28 people: the European experience. Health Policy 2012;105:25-32.
29

30
31 25 Lindert K, Schwarz A. Social Protection Responses to the Global Economic Crisis in ECA. World
32
33 Bank Knowledge brief. Washington, DC: World Bank; 2009.
34

35
36 26 Hajioff S, McKee M. The health of the Roma people: a review of the published literature. J
37
38 Epidemiol Community Health 2000;54:864-9.
39

40
41 27 Bogdanović D, Nikić D, Petrović B, et al. Mortality of Roma population in Serbia. Croat Med J
42
43 2007;48:720-6.
44

45
46 28 Zeljko HM, Skarić-Jurić T, Narančić NS, et al. Age trends in prevalence of cardiovascular risk
47
48 factors in Roma minority population of Croatia. Econ Hum Biol 2013 Jul;11:326-36. doi:
49
50 10.1016/j.ehb.2012.02.007. Epub 2012 May 15.
51

52
53 29 Hidvégi T, Hetyesi K, Biró L, Nádas J, Jermendy G. Screening for metabolic syndrome within a
54
55 minority ethnic group (adult Gypsy people) in Hungary. Bratisl Lek Listy 2012;113:721-4.
56
57
58
59
60

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- 1
2
3
4 30 Kósa Z, Széles G, Kardos L, et al. A comparative health survey of the inhabitants of Roma
5 settlements in Hungary. *Am J Public Health* 2007;97:853-9.
6
7
8
9 31 Asselbergs FW, Lovering RC, Drenos F. Progress in genetic association studies of plasma lipids.
10 *Curr Opin Lipidol* 2013;24:123-8.
11
12
13
14 32 Zarkesh M, Daneshpour MS, Faam B, et al. Heritability of the metabolic syndrome and its
15 components in the Tehran Lipid and Glucose Study (TLGS). *Genet Res (Camb)* 2012;94:331-7.
16
17
18
19 33 Sanghera DK, Blackett PR. Type 2 Diabetes Genetics: Beyond GWAS. *J Diabetes Metab*
20 2012;3(198):6948. doi:10.4172/2155-6156.1000198
21
22
23
24 34 Cornelis MC, Hu FB. Gene-Environment Interactions in the Development of Type 2 Diabetes:
25 Recent Progress and Continuing Challenges. *Annu Rev Nutr* 2012;32:245-59.
26
27
28
29 35 Brito EC, Lyssenko V, Renström F, et al. Previously associated type 2 diabetes variants may
30 interact with physical activity to modify the risk of impaired glucose regulation and type 2 diabetes: a
31 study of 16,003 Swedish adults. *Diabetes* 2009;58:1411-8.
32
33
34
35
36 36 Ruchat SM, Rankinen T, Weisnagel SJ, et al. Improvements in glucose homeostasis in response to
37 regular exercise are influenced by the PPARG Pro12Ala variant: results from the HERITAGE Family
38 Study. *Diabetologia* 2010;53:679-89.
39
40
41
42
43 37 Hu FB. Globalization of diabetes: the role of diet, lifestyle, and genes. *Diabetes Care*.
44 2011;34:1249-57.
45
46
47
48 38 Bo S, Ciccone G, Baldi C, et al. Effectiveness of a lifestyle intervention on metabolic syndrome. A
49 randomized controlled trial. *J Gen Intern Med* 2007;22:1695-703.
50
51
52
53 39 Blackett PR, Sanghera DK. Genetic determinants of cardiometabolic risk: A proposed model for
54 phenotype association and interaction. *J Clin Lipidol* 2013;7:65-81.
55
56
57
58
59
60

1
2
3
4 40 Landmesser U. High density lipoprotein - should we raise it? Curr Vasc Pharmacol 2012;10:718-9.
5
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For Review Only

Table 1
Socio-demographical characteristics of study samples representative for segregated Roma colonies and for Hungarian adult population.

Socio-demographical factors	Segregated Roma sample (N=646)	Representative Hungarian sample (N=1542)	
Age groups (years)	20-24	68 (10.53%)	93 (6.03%)
	25-29	59 (9.13%)	157 (10.18%)
	30-34	79 (12.23%)	161 (10.44%)
	35-39	101 (15.63%)	155 (10.05%)
	40-44	117 (18.11%)	171 (11.09%)
	45-49	72 (11.15%)	185 (12%)
	50-54	66 (10.22%)	237 (15.37%)
	55-59	47 (7.28%)	210 (13.62%)
	60-64	37 (5.73%)	173 (11.22%)
Sex	Female	393 (60.84%)	810 (52.53%)
	Male	253 (39.16%)	732 (47.47%)
Education	Less than primary	250 (38.7%)	31 (2.01%)
	Primary	304 (47.06%)	266 (17.25%)
	Vocational	75 (11.61%)	504 (32.68%)
	High school	15 (2.32%)	521 (33.79%)
	University	1 (0.15%)	204 (13.23%)
	Missing	1 (0.15%)	16 (1.04%)

Table 2

Ethnicity- and gender-specific prevalence of metabolic syndrome and its components among 20-64 years old Hungarians (A) and inhabitants of segregated Roma colonies (B), compared with each other (C, D, E).

Metabolic syndrome components	A. Representative Hungarian sample by gender			B. Segregated Roma sample by gender			C. Studied sample by ethnicity			D. Segregated Roma vs. Representative Hungarian males	E. Segregated Roma vs. Representative Hungarian females
	Females (N=810)	Males (N=732)	p	Females (N=393)	Males (N=253)	p	Segregated Roma sample (N=646)	Representative Hungarian sample (N=1542)	p	p	p
Central obesity	76.17% [73.25-79.09]	62.43% [58.94-65.92]	0.013	66.16% [61.5-70.81]	52.17% [46.05-58.30]	0.076	60.68% [56.91-64.45]	69.65% [67.35-71.94]	<0.001	0.004	<0.001
Raised blood pressure or treated hypertension	44.32% [40.92-47.72]	53.01% [49.41-56.60]	0.045	38.68% [33.89-43.47]	43.08% [37.01-49.15]	0.469	40.40% [36.62-44.19]	48.44% [45.95-50.94]	<0.001	0.007	0.063
Raised fasting plasma glucose concentration or formerly diagnosed diabetes mellitus	10.49% [8.39-12.59]	21.17% [18.23-24.12]	<0.001	21.88% [17.82-25.95]	35.18% [29.32-41.03]	0.005	27.09% [23.66-30.52]	15.56% [13.75-17.37]	<0.001	<0.001	<0.001
Raised triglyceride level or treated lipid disorder	31.36% [28.18-34.54]	44.95% [41.36-48.53]	<0.001	28.24% [23.82-32.67]	42.69% [36.62-48.75]	0.008	33.90% [30.25-37.55]	37.81% [35.39-40.23]	0.084	0.533	0.271
Reduced HDL cholesterol level or treated lipid disorder	34.2% [30.95-37.45]	38.11% [34.61-41.62]	0.273	55.73% [50.84-60.61]	51.38% [45.26-57.51]	0.554	54.02% [50.18-57.87]	36.06% [33.66-38.45]	0.029	<0.001	<0.001
Metabolic syndrome	32.72% [29.5-35.93]	37.43% [33.94-40.92]	0.179	34.86% [30.17-39.55]	38.74% [32.76-44.71]	0.496	36.38% [32.67-40.09]	34.95% [32.57-37.33]	0.525	0.712	0.460

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Table 3 Socio-demographical risk factors of metabolic syndrome and its components by multivariate logistic regression model for 20-64 years old adults in Hungary (odds ratios and 95% confidence intervals).

	Age (year)	Sex (Ffemale/ Mmale)	Education*				Ethnicity (Roma/ Hungarian)
			Pprimary	Vvocational	Hhigh school	Uuniversity	
Central obesity	1.06 [1.05-1.07]	1.93 [1.59-2.36]	1.36 [0.98-1.90]	1.44 [0.98-2.14]	1.45 [0.96-2.20]	1.06 [0.66-1.71]	0.84 [0.63-1.12]
Raised blood pressure or treated hypertension	1.09 [1.08-1.10]	0.62 [0.51-0.76]	1.09 [0.78-1.53]	1.02 [0.69-1.53]	0.85 [0.56-1.30]	1.01 [0.62-1.64]	0.94 [0.70-1.26]
Raised fasting plasma glucose concentration or known type 2 diabetes mellitus	1.07 [1.06-1.08]	0.40 [0.31-0.51]	1.31 [0.91-1.90]	1.03 [0.66-1.62]	0.72 [0.44-1.19]	0.59 [0.32-1.12]	2.65 [1.90-3.69]
Raised triglyceride level or treated lipid disorder	1.05 [1.04-1.06]	0.52 [0.43-0.62]	1.21 [0.87-1.68]	1.36 [0.93-2.00]	1.27 [0.85-1.90]	1.35 [0.85-2.16]	1.22 [0.93-1.62]
Reduced HDL cholesterol level or treated lipid disorder	1.02 [1.02-1.03]	0.93 [0.78-1.11]	0.96 [0.70-1.30]	0.96 [0.67-1.38]	0.80 [0.55-1.17]	0.82 [0.52-1.27]	2.15 [1.65-2.79]
Metabolic syndrome	1.07 [1.06-1.08]	0.75 [0.62-0.91]	1.04 [0.75-1.44]	0.99 [0.67-1.46]	0.89 [0.60-1.34]	0.87 [0.54-1.41]	1.37 [1.03-1.83]

* less than primary education serves as reference category

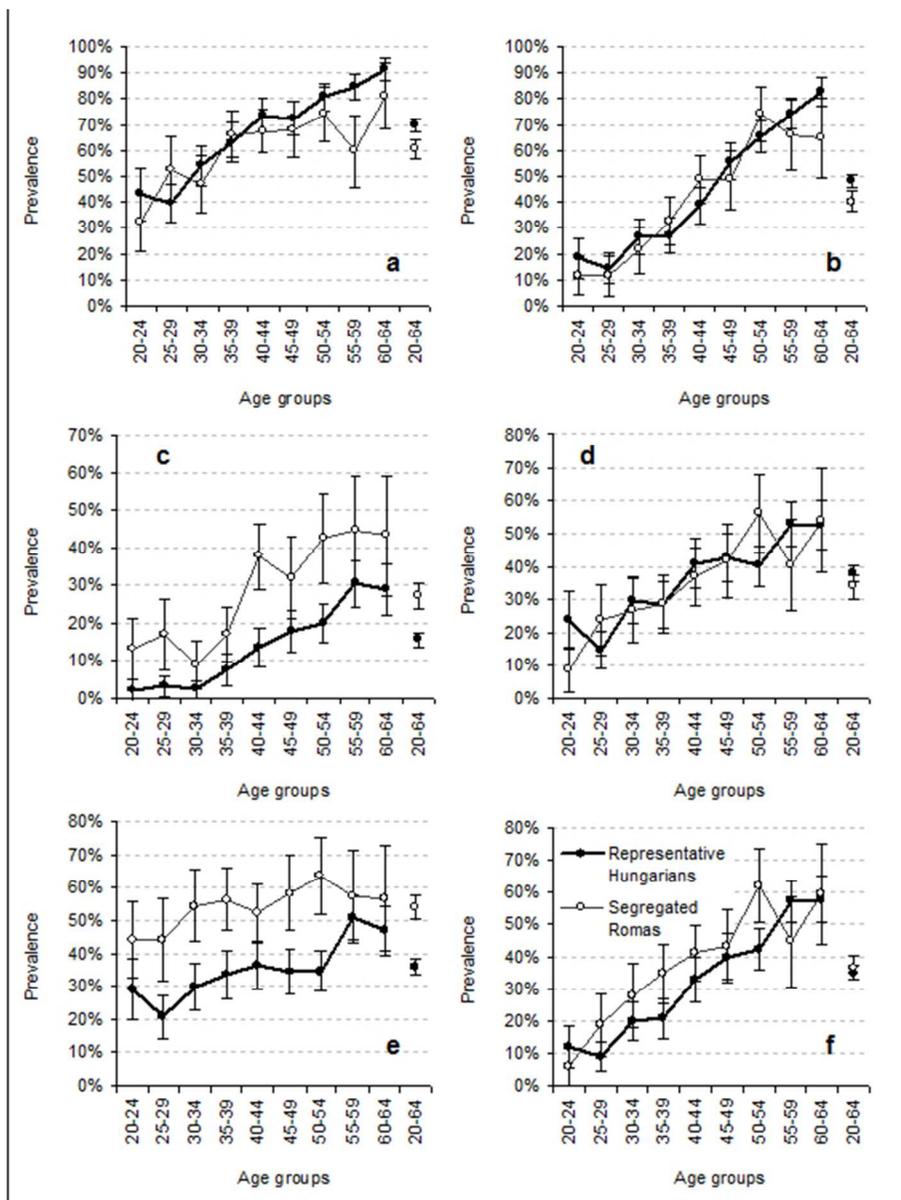
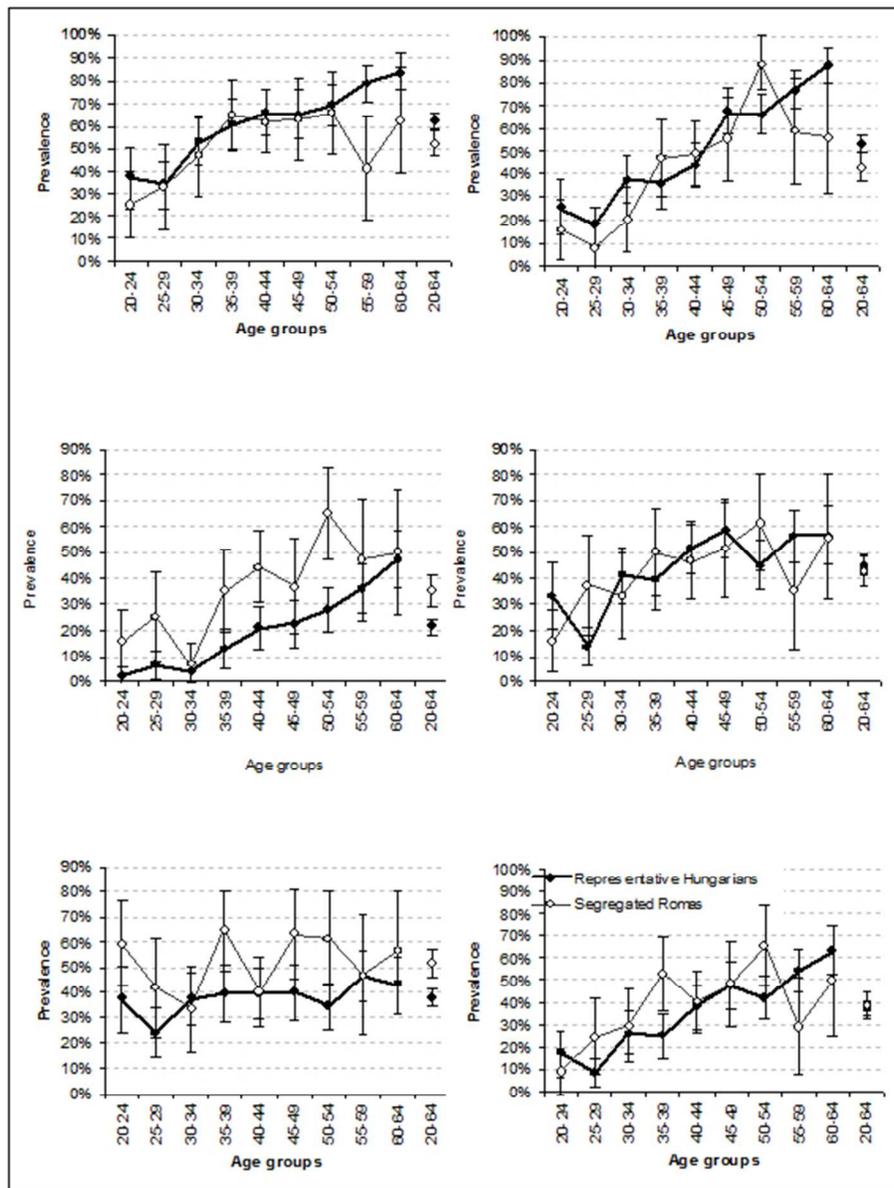
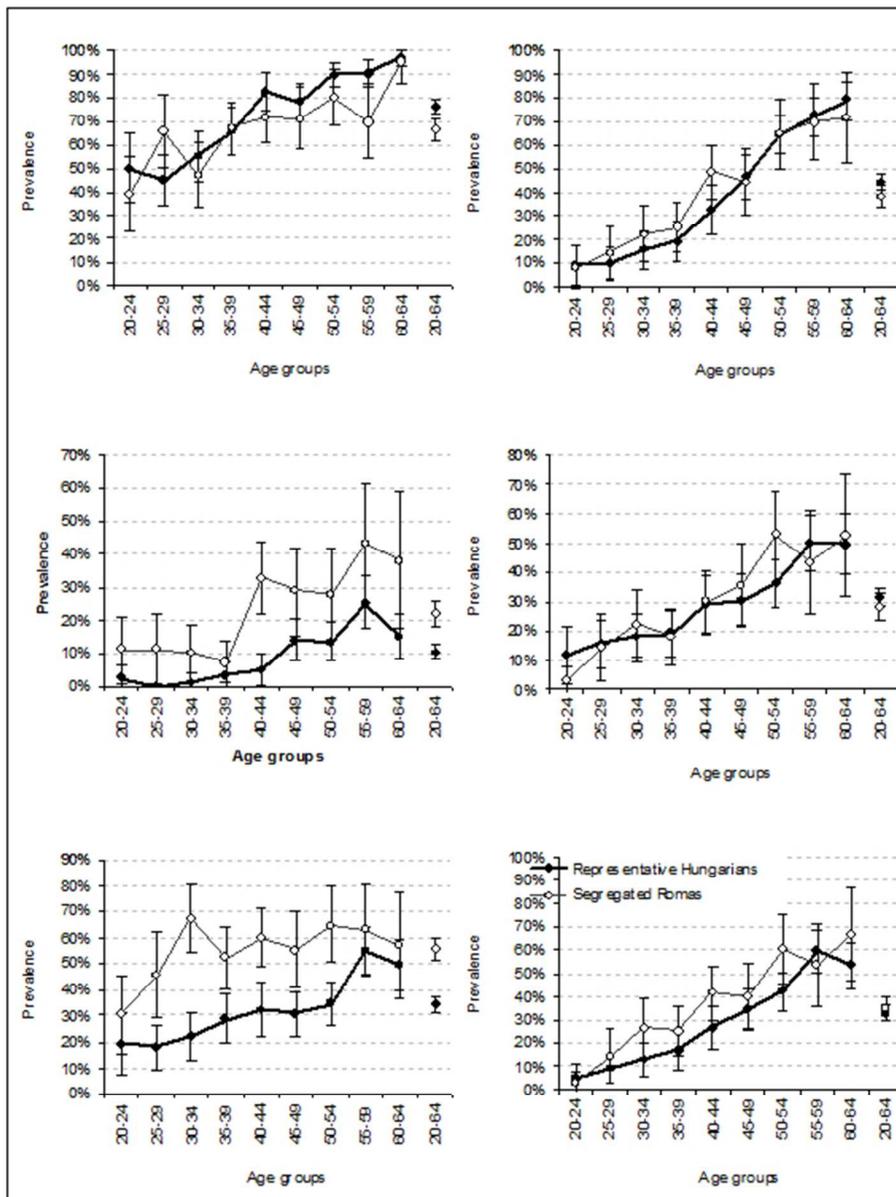


Figure 1

Age specific prevalence (with 95% confidence interval) of central obesity (a), raised blood pressure or treated hypertension (b), raised fasting serum glucose concentrations or formerly diagnosed type 2 diabetes mellitus (c), raised serum triglyceride levels or treated lipid disorders (d), reduced serum HDL cholesterol levels or treated lipid disorders (e), and metabolic syndrome (f) in 20-64 years old adults' samples representative for the population of Hungary and for the inhabitants of segregated Roma colonies.



Ad Figure 1 Age specific prevalence (with 95% confidence interval) of central obesity (a), raised blood pressure or treated hypertension (b), raised fasting serum glucose concentrations or formerly diagnosed type 2 diabetes mellitus (c), raised serum triglyceride levels or treated lipid disorders (d), reduced serum HDL cholesterol concentration or treated lipid disorders (e), and metabolic syndrome (f) in 20-64 years old males' samples representative for Hungary and for segregated Roma colonies in Hajdu-Bihar and Szabolcs-Szatmar-Bereg counties.



Ad Figure 1 Age specific prevalence (with 95% confidence interval) of central obesity (a), raised blood pressure or treated hypertension (b), raised fasting serum glucose concentrations or formerly diagnosed type 2 diabetes mellitus (c), raised serum triglyceride levels or treated lipid disorders (d), reduced serum HDL cholesterol concentration or treated lipid disorders (e), and metabolic syndrome (f) in 20-64 years old females' samples representative for Hungary and for segregated Roma colonies in Hajdu-Bihar and Szabolcs-Szatmar-Bereg counties.