before achieving LTC, and 49% achieved LTC. The relative hazard of death before achieving LTC compared to surviving without achieving LTC decreased for those with a history of opiate substitution therapy (OST) (HR 0.19, CI 0.10 to 0.34) and increased for HIV positive participants (HR 6.2, CI 3.6 to 10.6), those who started injecting after 1985 (HR 2.5, CI 1.3 to 4.8), those aged over 18 years at first injection (HR 2.2, CI 1.4 to 3.6), and those with a history of overdose (HR 2.0, CI 1.3 to 3.2). The relative hazard of achieving LTC compared to surviving without achieving LTC decreased for those with a history of OST (HR 0.39, CI 0.27 to 0.56), those who started injecting after 1985 (HR 0.56, CI 0.39 to 0.79) and those with a prison history (HR 0.69, CI 0.54 to 0.89); and increased for those aged over 18 years at first injection (HR 1.6, CI 1.2 to 2.1).

Conclusions: Few cohorts have sufficient follow-up to measure long-term cessation. The Edinburgh Addiction Cohort (EAC) suggests that exposure to OST is protective, reducing the risk of death before long term cessation, but OST also seems to increase duration of injecting drug use, reducing the likelihood of long term cessation.

CVD and metabolic syndrome

085 STATINS FOR THE PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE: CAUTION REQUIRED

¹JP Casas, ¹FC Taylor, ¹KJ Ward, ²M Burke, ³T Moore, ²GD Smith, ¹S Ebrahim. ¹Non communicable Disease Epidemiology Unit, London School of Hygiene and Tropical Medicine, London, UK; ²Department of Social Medicine, University of Bristol, Bristol, UK; ³Department of Community Medicine, University of Bristol, Bristol, UK

doi:10.1136/jech.2009.096735g

Background: Reviews of the effects of statins highlight the benefits of their use, leading expert committees to promote statin treatment on a global scale. However, most reviews have not distinguished between findings in primary and secondary prevention. Of the reviews which have attempted to look at the evidence for primary prevention, the role of statins is contradictory, leading to some scepticism among the cardiovascular community.

Objectives: To assess the effects, both benefits and harms, of statins in people without a history of CVD.

Methods: Systematic review of randomised trials comparing statins with usual care or placebo where duration of treatment was one year and follow up was six months. We searched Cochrane Central Register of Controlled Trials (CENTRAL), MEDLINE and EMBASE, until 2007. Data were extracted by two reviewers independently. Relative risk (RR) was calculated for dichotomous data and we used random effects models.

Results: Thirteen trials (14 arms) dating from 1995–2005 were located. A total of 43 150 participants were observed for up to 5.3 years; the mean age was 57 years (range 28–50 years), 71.7% were male, 92.8% were Caucasian. Three of the larger trials were stopped prematurely because significant reductions in primary outcomes between the intervention and placebo had been observed. Total mortality was reduced with the use of statins RR 0.84 (95% CI 0.75 to 0.94) as were all of the combined outcomes: fatal and nonfatal CVD events; RR 0.80 (95% CI 0.71 to 0.90); fatal and non-fatal CHD events RR 0.72 (95% CI 0.66 to 0.79) fatal and nonfatal stroke events RR 0.78 (95% CI 0.67 to 0.91). However, there was no strong evidence of benefit when single outcomes were evaluated. Of the seven trials reporting on adverse events, statins posed little harm. The majority of trials received industry sponsorship.

Conclusion: Composite endpoints were reported in preference to single end points and adverse events outcomes were not fully reported. Trials that were stopped prematurely may have contributed to an over-estimation of treatment effects. It is possible but unlikely that the results may not be generalisable to women, non-white people and those in old age. Caution in interpreting the results is required.

086 METABOLIC HEALTH CHANGES IN MIGRANTS MOVING FROM A RURAL TO AN URBAN ENVIRONMENT IN TANZANIA

N Unwin, P James, R McNally, on behalf of the UMOST investigators *Institute of Health and Society, Newcastle University, Newcastle, UK*

doi:10.1136/jech.2009.096735h

Background: High levels of rural to urban migration are a feature of most developing countries, and are thought to be associated with an increased risk of chronic non communicable diseases.

Aim: To investigate in adult rural to urban migrants in Tanzania changes in health related behaviours, BMI, lipids and blood pressure in the first 12 months following migration.

Methods: Through village key informants, men and women, aged 15 to 59, from Morogoro rural region intending to migrate to Dar es Salaam for at least 6 months were identified. Prior to migration and regularly (1 to 3 monthly) after migration, measurements were made, blood taken for lipids, and data on socio economic circumstances and aspects of life style collected by interview. For each migrant an age, sex and village matched non migrant was also assessed at baseline and 12 months later.

Results: Two hundred and nine migrants, 103 men and 106 women, had measurements prior to migration, mean age 28.0 (SD 11) and 29.5 (11) years respectively. At 12 months contact was maintained with 132 (63.2%) of the migrants. Following migration there were significant changes in diet, with migrants consuming more meat, fresh vegetables, coconut oil and margarine. Self reported regular physical activity declined, from 79% of men to 27% (95% CIs 39% to 58%) and 38% of women to 14% (7% to 36%). At 12 months migrants, compared to the non-migrants, had a higher BMI (by 0.64 kgm⁻², 95% CIs 0.28 to 1.0) and serum cholesterol (0.57 mmol l⁻¹, 0.27 to 0.88), but lower systolic (5.2 mm Hg, 1.7 to 8.5) and diastolic blood pressure (7.4 mm Hg, 5.1 to 9.7). Triglycerides were lower in migrants at 6 months (0.31 mmol l⁻¹, 0.06 to 0.58) but not at 12 months. Multiple linear regression was used to identify predictors of change in biological variables following migration. Associations (p<0.05) were found with aspects of diet for BMI, blood pressure, cholesterol and triglycerides, and increasing BMI predicting increasing triglycerides.

Conclusion: This relatively small study of rural to urban migrants in Tanzania found changes with mixed consequences for health following migration. Despite falls in physical activity and an overall tendency to increasing weight and cholesterol, there were apparently significant falls in blood pressure and (over the first 6 months) in triglycerides. Our tentative hypothesis is that changes in diet, from one dominated by carbohydrate to one of greater diversity, lead to favourable triglyceride and blood pressure changes, but that as weight increases these changes will be reversed.

087 HIGH BURDEN OF CARDIOVASCULAR DISEASE AND RISK PROFILE IN AN ELDERLY EASTERN GERMAN GENERAL POPULATION—POTENTIAL EXPLANATION FOR AN EAST–WEST GRADIENT OF CARDIOVASCULAR MORTALITY: THE CARLA STUDY 2002–2006

¹KH Greiser, ¹A Kluttig, ¹B Schumann, ¹O Kuss, ²JA Kors, ³K Werdan, ⁴CA Swenne, ¹J Haerting. ¹Institute of Medical Epidemiology, Biostatistics, and Informatics, Martin-Luther-University Halle-Wittenberg, Halle (Saale), Germany; ²Department of Medical Informatics, Erasmus Medical Center, Rotterdam, The Netherlands; ³Department of Internal Medicine III, Martin-Luther-University Halle-Wittenberg, Halle (Saale), Germany; ⁴Department of Cardiology, Leiden University Medical Center, Leiden, The Netherlands

doi:10.1136/jech.2009.096735i

Background: For cardiovascular diseases (CVD), an east-west mortality gradient across Europe has been described, which could not fully be explained by established risk factors. Likewise, the cause of the higher CVD mortality in eastern as compared to

western Germany has not yet been elucidated due to the scarcity of population-based data on CVD morbidity and risk factors.

Objective: To provide data on the distribution of CVD and its risk factors in an elderly general population in eastern Germany as compared to a western German population.

Methods: Cross-sectional data of 1779 East German participants of the population-based CARLA Study aged 45–83 at baseline were used to calculate sex- and age-specific means of risk factors, disease prevalence, and expected 10-year risk of fatal CVD to be compared with the data of 4261 participants of the south-west German KORA study. Risk of fatal CVD was calculated using the Systematic Coronary Risk Evaluation (SCORE) Germany algorithm which is based on sex, age, systolic blood pressure (SBP), current smoking, and cholesterol levels.

Results: There were no clear differences in age-specific mean body mass index (BMI) and smoking prevalence between CARLA and KORA subjects (except for higher smoking prevalence in 45–54 year old CARLA subjects), and only a slightly higher predicted 10-year CVD mortality in CARLA as compared to KORA subjects. Mean 10-year risk of fatal CVD in CARLA increased from 0.57% (95% CI 0.39 to 0.75%) in 45–49 year-old to 3.5% (3.0 to 3.9%) in 60–64 year-old women as compared to 0.37% (0.33 to 0.41%) and 2.91% (2.71 to 3.12%) in KORA respectively. However, the prevalence of diabetes and hypertension, and mean SBP were considerably higher in CARLA subjects across all sex-age-groups as compared to KORA subjects. For example, the prevalence of hypertension in CARLA increased from 55.4% (48.6 to 63.3%) in 45–54 year-old to 84.2% (79.3 to 89.2%) in 65–74 year-old women, while it was 36.7% (32.3 to 41.1%) to 61.5% (56.5 to 66.5%) in KORA, respectively. The prevalence of diabetes across 10-year age-groups in CARLA women increased from 4.9% (1.9 to 7.9%) in the 45–54 year-old to 19.1% (13.8 to 24.5%) in the 65–74 year-old as compared to 2.4% (1.0 to 3.8%) to 8.1% (5.3 to 10.9%) in KORA, respectively. Moreover, within narrow categories of BMI, CARLA subjects consistently showed a considerably higher waist-to-hip-ratio than KORA subjects.

Conclusions: Our results support the hypothesis that an especially high prevalence of diabetes and hypertension and other components of the metabolic syndrome may explain the excess CVD mortality in eastern Germany which is not accurately reflected in SCORE.

Methods in ethnicity research

088 INVESTIGATING THE ASSOCIATION BETWEEN ETHNICITY AND SURVIVAL FROM BREAST CANCER USING ROUTINELY COLLECTED HEALTH DATA: CHALLENGES AND POTENTIAL SOLUTIONS

¹A Downing, ^{1,2}D Forman, ¹JD Thomas, ¹RM West, ³G Lawrence, ¹MS Gilthorpe. ¹Centre for Epidemiology & Biostatistics, University of Leeds, Leeds, UK; ²Northern & Yorkshire Cancer Registry & Information Service, St James's Institute of Oncology, Leeds, UK; ³West Midlands Cancer Intelligence Unit, Public Health Building, University of Birmingham, Birmingham, UK

doi:10.1136/jech.2009.096735j

Background: Previous studies have reported differences in survival from breast cancer by ethnic group. Some of these studies have taken information on ethnicity from routinely collected data, such as Hospital Episode Statistics (HES). There are several problems associated with using ethnicity from HES data, such as multiple ethnicities being recorded for a single patient with multiple hospital visits, and missing data. This study will investigate methods to overcome these problems in order to assess the relationship between ethnicity and survival from breast cancer.

Data and Methods: 48 234 breast cancer patients diagnosed between 1997 and 2003 were identified from a linked cancer registry-HES dataset for two regions of the UK. Where multiple ethnicities were recorded for a patient a single ethnicity was allocated according to the last recorded and most popular code. The data were also expanded to include all

available hospital episodes (and all ethnicity information) for each patient (452 061 “episode-level” records). Ethnicity was missing in 16% of the patient-level records and 26% of the episode-level records. Multiple imputation (10 iterations) of missing ethnicity using age, stage, socioeconomic background and census area ethnic make-up as predictors was undertaken for the “last recorded”, “most popular” and “episode-level” data. Survival analysis (up to end 2006) was carried out using the imputed datasets.

Results: Across the two regions, 97.2% of the patients with a known ethnicity were White, 1.6% were South Asian and 0.8% were Black. White women were slightly older at diagnosis than the other groups, whilst Asian women had a higher proportion of early stage tumours, but these differences were not significant. Using “last recorded” ethnicity, unadjusted survival was higher in the Asian group compared to the White group (HR 0.77, 95% CI 0.66 to 0.92). After adjustment for age and stage this survival difference was no longer significant (HR 0.98, 95% CI 0.82 to 1.16). The results were similar using “most popular” ethnicity. Using the “episode-level” data to assign probabilities for each patient, unadjusted survival was again higher in the Asian group (HR 0.72, 95% CI 0.62 to 0.89) compared to the White group, but after adjustment survival was similar in the two groups. There was also some evidence of worse survival in the Black group compared to the White group (HR 0.98, 95% CI 0.98 to 1.39 after adjustment).

Conclusions: Assessment of the association between breast cancer survival and ethnicity presents many challenges. Previous research in this area may have reported biased results, because of missing data and the failure to use all available information.

089 RECRUITING SOUTH ASIANS TO A RANDOMISED TRIAL (PREVENTION OF DIABETES AND OBESITY IN SOUTH ASIANS) FOR THE PREVENTION OF DIABETES: THE CHALLENGES AND ACHIEVEMENTS

A Douglas, on behalf of the PODOSA Investigators Group *Public Health Sciences Section, University of Edinburgh, Edinburgh, UK*

doi:10.1136/jech.2009.096735k

Background: Despite recommendations to do so, few clinical trials focus on ethnic minority groups. There are concerns that data from general trials may not apply to them. PODOSA (Prevention of Diabetes and Obesity in South Asians) is a rare example of a prevention trial in European South Asians. The prevalence of type 2 diabetes among UK South Asian adults is extremely high.

Objectives of the Trial: To assess the feasibility and cost-effectiveness of the PODOSA lifestyle intervention (see below).

Methods: PODOSA is a cluster randomised controlled trial evaluating a family based, ethnically tailored, lifestyle intervention aiming to reduce the incidence of diabetes in people of Indian and Pakistani origin by reducing weight and increasing physical activity. Recruitment is via several channels. Eligible participants are those found to have impaired glucose levels (and therefore at high risk of developing diabetes) on an oral glucose tolerance test. The intervention group receives 15 contacts with a dietician over three years. The control group has 4 dietetic contacts. The dieticians' toolkit contains culturally tailored resources on diet and exercise.

Results: Recruitment commenced in July 2007 and plans to finish around August 2009. The trial has enjoyed support from individuals, community and religious organisations, media, leaders and health professionals. 122 families, with 135 people at high risk of diabetes and 101 family volunteers, have been recruited at the time of writing. Recruitment into the screening component of the trial has been slow, taxing and expensive. Referrals from NHS professionals have been few, and responses to radio, newspaper and website based publicity trivial. The response to written invitations has also been low. Face-to-face recruitment both with individuals and groups, has proved successful. 95% of those eligible have entered the trial.