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## Cultural Change and Stress in Western Samoa: An application of lifestyle incongruity model by Thomas W McDade (Northwestern University)

#### <Points>

- Impact of cultural change on stress or immune function

- Field-friendly blood spot methods for assessing cell-mediated immune function (EBV antibodies) and non-specific current infection (C-reactive protein)

#### <Paper 1>

McDade TW et al. (2000) Culture Change and Stress in Western Samoan Youth: Methodological Issues in the Cross-Cultural Study of Stress and Immune Function. *American Journal of Human Biology*, 12: 792-802.

#### 1. Introduction

<u>Background</u>: (1) Westernization or urbanization resulted in adverse health outcomes in the populations under transition (most studies for adults, e.g., HT). (2) Psychosocial experience may suppress the human immune function.

<u>Hypothesis</u>: Westernization or urbanization suppressed immune function in Western Samoa.

<u>Another objective</u>: Apply "field-friendly" methods for assessing immune function and current infection.

## 2. Research Context: Western Samoa

Three distinct geographic regions:

- (1) Savaii: subsistence cultivation, most traditional forms of Samoan culture
- (2) Rural Upolu: easy access to Apia, economic and cultural transition zone
- (3) Apia: urban area, movie theater, disco, restaurants

## 3. EBV antibodies: marker of cell-mediated immune function

<u>Epstein-Barr virus</u>: ubiquitous; 100% infection rate by the age of 5 years; once infected, individuals harbor the virus for life in infected cells; under adequate cell-mediated immune function, the virus is maintained in a latent state; immunosuppression allows EBV to reactivate and release viral antigens, to which a antibody response emerge; indirect measure of cell-mediated immune function (Fig. 1).

<u>Previous studies</u>: Increase of EBV antibodies was associated with (1) medical school examination, (2) poor-quality marriage, (3) chronic stress with careing a amily member with Alzheimer's disease, (4) loneliness, defensiveness, and anxiety.

#### 4. C-reactive protein: marker of current infection

<u>C-reactive protein</u>: central component of the acute phase response, non-specific response to infection or injury; body's first line of defence against pathogens; concentration increase  $\times 100$ -1000 during the 24-72 hr following an injury or infectious challenges; level remain elevated during the course of infection; >5mg/l plasma = cut-off that indicates the "current infection"

#### 4. Methods

<u>Participants</u>: N=760 individuals, ages of 4 and 20 years; comparable age and sex distribution in three regions (Table 1).

## <u>Data</u>:

- (1) Dried spots of whole blood for measurement of EBV antibodies (marker for immune function) and C-reactive protein (marker for current infection).
- (2) Anthropometric measures
- (3) Observable symptoms of respiratory diseases
- (4) Demographic and psychosocial information
- (5) Housing style, ownershiop of western goods, travel experience, occupation of their parents

Analytical Framework:

Y: Level of EBV antibodies (marker of cell-mediated immune function) X: Region (level of urbanization/westernization)

Confounders: (1) nutritional status (NS), (2) Current infection (respiratory disease, C-reactive protein level >=5mg/l), age

## 5. Results

<u>Regional difference in lifestyle</u> (Table 2):

Apita (urban)-Upolu (transitional)- Savaii (rural)

<u>Potential confounders:</u>
(1) Nutritional status did not

- Nutritional status did not differ by region; undernutrition is not a problem; no association between nutritional status and EBV antibodies (Tab2, Fig2/3)
   Commut infaction (Preprint term infaction on CPD) = 7 mm) (Fig. 4);
- (2) Current infection (Respiratory infection or CRP>=5 mg) (Fig 4):

Apia > Upolu > Savaii; individuals with current infection removed (144 of 760) <u>Regional differences in immune function</u>:

PROC GLM (SAS): Y=EBV antibody, X1=age, X2=sex, X3=region; interaction terms were NS.

- Significant predictors of EBV antibodies were sex and region
- Post-hoc Sheffe pairwise comparisons: <u>EBV (Apia/Upolu) > EBV (Savaii)</u>

## 6. Discussion

Lower level of cell-mediated immune function in urban and transitional areas in Western Samoa: Catecholamine was also higher in urban than in rural areas in W Samoa; BP and distance from urban areas correlated after controlling confounders in A Samoa; Psychological stress caused by westernization or urbanization played the role to suppress the immune function of the people in W Samoa.

<u>Type of psychosocial stress</u>: challenge associated with managing a more western life style (Apia); Confronting contradictions between old and new ways of living (rural Upolu); lifestyle incongruity theory??

<Paper 2>

McDade TW (2001) Lifestyle incongruity, social integration, and immune function in Samoan adolescents. Social Science & Medicine, 53: 1351-1362. <Analysis at individual level>

#### 1. Model that explain the relationship between social change and disease risk

<u>Status inconsistency models (a term of Medical Sociology)/ lifestyle incongruity</u> <u>model:</u> Mechanisms: Individuals are motivated to accumulate elements of lifestyles in order for them to display their self-defined place in the system of social stratification. So, the individual struggling to maintain a higher style of life in the context of low SES feels stress. The individuals who are status-incongruent are continually scanning the social field searching to determine if they are being responded to with the sense of respect that they desire. <u>"Previous eight field studies:</u> (West Indies, Mexico, Brazil, USA, Samoa, England) all showed that the higher lifestyle incongruity was associated with higher BP after controlling for age, sex and anthropometric indicators (e.g., BMI). <u>Model structure:</u>

$$Y = a + b_1(x_1 + x_2) + b_2(x_1 - x_2) + e$$

Y: Biological outcomes hypothesized to be influenced by the acquisition of westernized lifestyles (e.g., stress, immunosuppression), x1: lifestyle (e.g., material goods, information), x2: household economic resources (e.g., occupation, education).

#### 2. Methods

<u>Subjects</u>: N=230, ages of 10-20 years living in Apia, Upolu, or Savaii Model:

(1) Material score (x1): Table 2, range 2-17;

(2) SES score (x2): the occupational rank of father/mother and frequency of receiving remittance, range 2-9

Each score was standardized to a mean of 50 and SD of 10.

X1-X2=lifestylel incongruity socre X1+X2=overall measure of lifestyle Confounders: age, sex, region

#### 3. Results

(1) Lifestyle incongruity explained the variation in EBV antibody level after controlling the effect of age, sex and region. (Table 3; Fig 1)

(2) The extent of social integration modify the relation between lifestyle incongruity and EBV antibody level?

Social integration=rely on family/friends when in need of help + satisfied with the relationship with family/friends: "traditional way of life in W Samoa"

Among the individuals who did not rely on family/friends and who were not satisfied with the relationship with family/friends, the effect of lifestyle incongruity on EBV antibody diminished. "Traditional Samoans" who have lifestyle incongruity may experience the EBV antibody increase or immuno-suppression most markedly. (Table 4; Fig 2)

# Field-friendly methods of EBV antibody and C-reactive protein measurements (papers 3,4,5)

- 1-1. Blood spot sample collection on filter paper (see Slides)
- 1-2. Quantification of Epstein-Barr (EB) virus antibodies (see Slides)
- 1-3. Quantification of C-reactive protein (see Slides)

## <References>

- 1. Mei JV et al. (2001) **Use of Filter Paper for the Collection and Analysis of Human Whole Blood Specimens.** *Journal of Nutrition*, 131: 1631S-1636S. <Validation of using filter paper for the collection of whole blood specimens>
- 2. McDade TW et al. (2000) Epstein-Barr Virus Antibodies in Whole Blood Spots: A Minimally Invasive Method for Assessing an Aspect of Cell-Mediated Immunity. *Psychosomatic Medicine*, 62: 560-567. <Validation of quantification of EBV antibodies in dried blood spots >
- 3. McDade TW et al. (2004) High-Sensitivity Enzyme Immunoassay for C-Reactive Protein in Dried Blood Spots. *Clinical Chemistry*, 50: 652-653. <Validation of quantification of C-reactive protein in dried blood spots >