

A Comparative Risk Assessment of Active Transportation in the United States

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Abstract

Comparative risk assessments to quantify the health potential of active transportation, e.g., walking and cycling, have been performed in the United States on a limited basis. Current methodologies, which rely on extensive data collection and user expertise, have estimated health benefits for select metropolitan areas under a few alternative scenarios. These analyses have largely been conducted using the Integrated Transport Health Impacts Model (ITHIM) which was developed in the UK and has been adapted for use in the US, primarily California. Here we present an implementation of the active transportation component of ITHIM to model the health impacts of modified active transportation levels. Using the newly developed software package *ITHIM* written in R, along with publicly available national datasets, we quantify the change in disability adjusted life years over a wide range of scenarios. The R package implementation of ITHIM’s physical activity component provides a robust environment for exploring the relationship between active transportation and positive health outcomes. A web-based user-interface is available for users who want a first estimate for the impact active transportation has on their state/region.

Introduction

Regular exercise has been shown to provide health benefits in terms of chronic disease prevention [1], cognitive function [2], and overall well being and satisfaction with life [3]. Still only half of Americans obtain the recommended levels of aerobic activity of 2.5 hours per week. There is considerable variability in the

attainment of this goal by location within the country. Supportive local built
and policy environments are important determinants of population-level activ-
ity patterns. Particularly, as they facilitate and foster routine and moderate
physical activity, such as walking, local decisions on design and policy can play
a huge role in improving the nation’s health.

Given the constraints and complexity of local governance and infrastructure
investments, land use, and human behavior there is a need for high quality
and reliable data and tools for decision-making related to urban transportaiton
systems. The lack of accurate and availbale estimates of the health impacts of
such decisions has limited the effectiveness of efforts to encourage non-motorized
transport in the US and in other countries. In the last decade new tools, ap-
plying comparative risk assessment methodologies, have made progress toward
quantifying the multiple health impacts of changes to the transportation system
and behaviors.

ITHIM is one such statistical model that integrates data on active transport,
physical activity, fine particulate matter and greenhouse gas emissions to pro-
vide estimates for the proportional change in mortality and morbidity for given
baseline and alternate travel scenarios. The model has been used to calculate
the health impacts of walking and bicycling short distances usually traveled by
car or driving low-emission automobiles [4, 5].

ITHIM uses a comparative risk assessment framework for the active trans-
port component of the model. We improve on the existing implementation
of the active transport component by including the distribution of non-travel-
related activity, improving numerical precision when computing the population
attributable fraction and creating a simple user-interface for the custom-built
R package *ITHIM*. We present this implementation here by using it to esti-
mate the relationship between overall, nationwide disease burden and active

transportation time in the United States.

Materials and Methods

As with the original ITHIM implementation we, employed a comparative risk assessment (CRA) across scenarios defined by mean active transportation time, i.e., combined walking and cycling. To do so we investigate the proportional change in national disease burden between the baseline model and a given alternative scenario, i.e., the population attributable fraction, PAF. The population attributable fraction is defined below and computed for each disease (ICD-10) included in the model, namely Breast Cancer (C50), Colon Cancer (C18-C21), Dementia (G30), Diabetes (E10-E14) and Cardiovascular disease (I10, I12, I15, I20-I25, I30-I31, I33, I40). We also include the disability impacts of depression, though it is not listed as a cause of death in CDC's data. For readability we omit an index indicating disease.

$$\text{PAF}_j = \frac{\int R_j(x)P(x) \, dx - \int R_j(x)Q(x) \, dx}{\int R_j(x)P(x) \, dx} \approx 1 - \frac{\sum_{i=1}^n R_j(x'_i)}{\sum_{i=1}^n R_j(x_i)} = 1 - \delta_j \quad (1)$$

Here $R(x)$ represents the risk when the individual has exposure x , physical activity (MET-hrs./week). $P(x)$ and $Q(x)$ denote the population density of individuals with exposure x , in the baseline and alternate scenarios, respectively. The exposure variable, x , is modeled as the sum of two independent random variables, travel-related and non-travel-related physical activity. \mathbf{x} and \mathbf{x}' represent the quantiles for the exposure distribution in baseline and alternate scenarios, respectively. δ_j is the proportional burden for disease j in the alternate scenario.

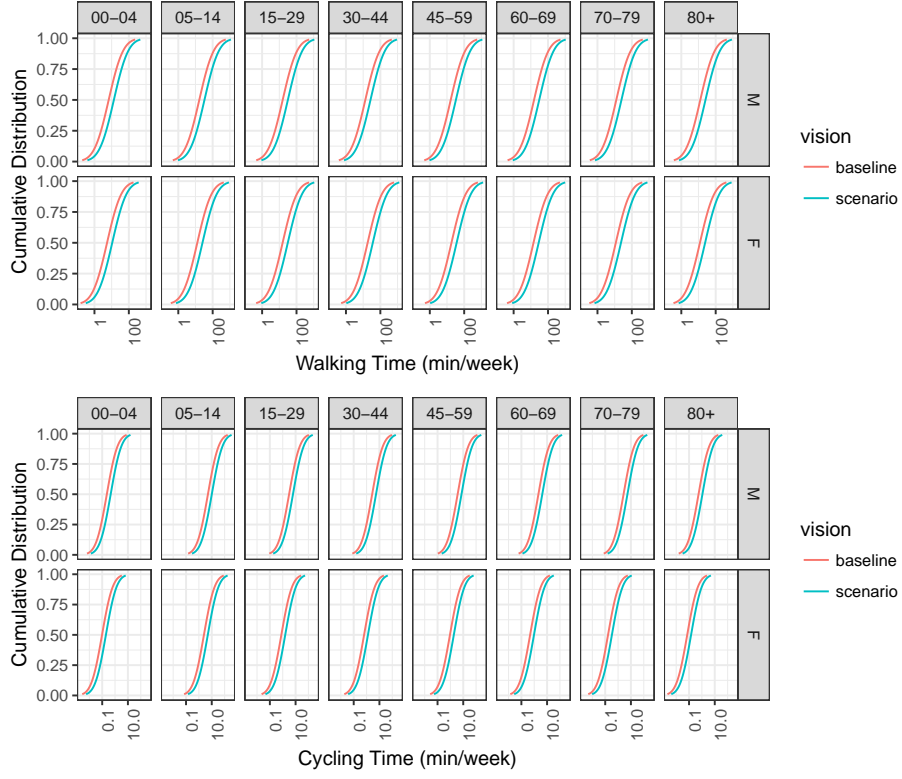


Figure 1: Distributions of walking and cycling transport time for baseline and scenario models, stratified by age and sex. In this example the scenario was created by doubling the overall walking and cycling means.

Travel-Related Physical Activity The distribution for travel-related exposure is estimated from the time spent walking or cycling, i.e., active transport time, which as in the original ITHIM model is assigned a log-normal distribution with constant coefficient of variation across age-sex classes. Travel-related exposure is estimated from active transport time using assumptions about how much physical activity is required for cycling and walking.

Non-Travel-Related Physical Activity With a simulated distribution for total physical activity in hand we compute the empirical percentiles in the baseline and alternate scenarios, \mathbf{x} and \mathbf{x}' . The proportional change in disease bur-

den, δ , may be approximated as follows, where R is the risk function for total
activity.

$$X_{ij}^{\text{non-travel}} \sim \log \mathcal{N}(\mu_{ij}, \gamma \mu_{ij}) \quad (2)$$

Within each age-sex class the non-travel-related exposure is modeled with a
log-normal distribution. The age-sex specific mean ratios, $r_{ij} = \frac{\mu_{ij}}{\mu_0}$ and γ are
estimated from the American Time Use Survey data [6].

Estimation of Disease Burden The disease burden, in this case DALYs, is
found for each of the quantiles using equation 3 using B_0 , the overall nation-
wide burden for the baseline. The overall burden is then computed as the sum
across quintiles, $B = \sum_{j=1}^n B_j$.

$$B_j = \delta B_0 \frac{\text{RR}_j}{\sum_{k=1}^n \text{RR}_j} \quad (3)$$

Risk Functions We assign risk based on dose-response curves for each of the
diseases included in the model.

$$\text{RR}_j(x) = e^{-\alpha_j \tilde{x}} \quad (4)$$

where $\tilde{x} = x^k$, $k = \frac{1}{2}$ and r varies by age-sex class and disease. For a table
of r values see supplementary material.

Nationwide Health Benefits to Active Transport For the purposes of
this implementation we limited our analyses to metropolitan regions of the
United States. To identify urban populations we used the National Center for
Health Statistics 2013 Urban-Rural Classification Scheme for Counties [7]. We
eliminated counties in the US with the classification of ‘nonmetropolitan’ (non-
core or micropolitan). We used this and similar classification schemes as well as

county and state identifiers in other datasets (NHTS, ATUS, CDC WONDER) 82
to ensure consistency in our estimates for input parameters. 83

To test the new implementation of ITHIM’s active transportation compo- 84
nent, we used the US national metro population estimates for active travel time 85
(walking and cycling) as the baseline. We obtained active travel time input 86
parameters for alternate scenarios using states with large sample sizes. From 87
these examples we were able to approximate low and high active travel scenar- 88
ios. Finally, we applied these active travel times to the US metro population to 89
perform the comparative risk assessment against the baseline US numbers. We 90
then examined the sensitivity of disease burden estimates to input parameters 91
including the active travel time and non-travel activity. 92

Value of Information 93

Insert analysis here. 94

Results 95

We report total change (averted or increased) in disability adjusted life years 96
(DALYs) as an estimate of disease burden change with alternative scenarios. 97

To estimate DALYs we use a methodology adopted from previous US imple- 98
mentations of ITHIM which scale Global Budern of Disease Estimates for the 99
US to smaller populations using mortality rate ratios [5]. Age, sex, and casuse 100
specific mortalities were obtained form the CDC WONDER database for US 101
metro counties for the years 2010-2014. Cells with mortality counts less than 102
10 deaths were suppressed for privacy and imputed (0 to 9) using R’s random 103
integer generator. 104

Our subset of the US population to only metropolitan counties consisted of 105
1,090 of 3,141 (34.7%) counties, or approximately 272M of 312M (87%) of the 106

total US population. The age and gender distribution of our subset population was slightly younger (making this up) than the national population. Baseline active transport times (walking and cycling) for the US across age and sex groupings was found to be approximately 43 min/week. Baseline non-travel activity time was found to be X (min/week) from the ATUS for US metropolitan populations.

Adapting ITHIM's methods to the R platform enables the rapid and thorough assessment of parameter influence within the model. Another important improvement enabled by the shift to a more computationally robust platform was to examine methodological assumptions of former implementations. One critical assumption involves the approximation of α used to calculate the attributable fraction (see Figure 1). Previous implementations used quintiles to approximate this value, essentially the area under the α curve. We improved upon those methods to use percentiles or 100 points to estimate the same values. We find our estimates to be much more stable and reliable at all values of active travel and non-travel activity parameters.

As exhibited in Figure 2, the variability in disease burden benefits varies to a much greater degree when using fewer points in the attributable fraction approximation. Importantly, we see greater stability when using the package and percentiles to approximate attributable fraction.

Comparing baseline to scenarios with increased active travel produced estimates of up to 360,000 fewer DALYs (with 300% increase in mean active travel time) for our US population subset. We then varied the mean non-travel activity time and active travel time to observe how this changed results. We observe the most dramatic responses in disease burden estimates to changes in active travel time when non-travel activity times are lowest. This result is consistent with the non-linear dose-response curve for physical activity and several

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