

Using statistical methods and reproducible tools to gain new insights from biomedical and public health data

Ariel Mundo Ortiz

Centre de Recherches Mathématiques, Université de Montréal

MfPH Next Generation Seminar Series

3/15/23



Introduction

- Data is the core of research. However, data is not information, as it needs to be processed before we can get information from it.

Introduction

- Data is the core of research. However, data is not information, as it needs to be processed before we can get information from it.
- This is specially true in the case of health research: public health, or biomedical data can be complex, and decisions along the analysis can result in different interpretations.

Introduction

- Data is the core of research. However, data is not information, as it needs to be processed before we can get information from it.
- This is specially true in the case of health research: public health, or biomedical data can be complex, and decisions along the analysis can result in different interpretations.
- In this talk I will focus on two examples that showcase how we can get more insight from looking at data from a different perspective.

The Case of Public Health Data: COVID-19 Vaccination

COVID-19: Why?

- The pandemic is still ongoing

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: Why?

- The pandemic is still ongoing
- COVID-19 vaccination has been an important component of public health strategies aimed at managing the pandemic.

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: Why?

- The pandemic is still ongoing
- COVID-19 vaccination has been an important component of public health strategies aimed at managing the pandemic.
- However, COVID-19 vaccination has not been equal across different population segments.

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: Why?

- The pandemic is still ongoing
- COVID-19 vaccination has been an important component of public health strategies aimed at managing the pandemic.
- However, COVID-19 vaccination has not been equal across different population segments.

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: Why?

- The pandemic is still ongoing
- COVID-19 vaccination has been an important component of public health strategies aimed at managing the pandemic.
- However, COVID-19 vaccination has not been equal across different population segments.
- Individuals with lower income, and those belonging to a racial/ethnic minority have had lower vaccination uptake^{1,2}.

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: Why?

- The pandemic is still ongoing
- COVID-19 vaccination has been an important component of public health strategies aimed at managing the pandemic.
- However, COVID-19 vaccination has not been equal across different population segments.
- Individuals with lower income, and those belonging to a racial/ethnic minority have had lower vaccination uptake^{1,2}.
- This is important because these differences in vaccination uptake have implications on virus transmission.

¹Nafilyan et al. 2021.

²Gerretsen et al. 2021.

COVID-19: The Case of Ontario

- The Fields Institute collected some very nice data regarding COVID-19 vaccination in Ontario: the *Survey of COVID-19 related Behaviours and Attitudes*.

COVID-19: The Case of Ontario

- The Fields Institute collected some very nice data regarding COVID-19 vaccination in Ontario: the *Survey of COVID-19 related Behaviours and Attitudes*.
 - The survey ran between late 2021 and early 2022 and collected socio-demographic information along with self-reported vaccination status (“Have you received the first dose of the Covid vaccine?”)

COVID-19: The Case of Ontario

Table 1: Selected socio-economic factors from the survey

| Variable | Levels |
|----------------------|---|
| Age group | 16-34, 35-54, 55 and over |
| Income bracket (CAD) | under 25,000, 25,000-59,999, 60,000 and above |
| Race/ethnicity | Arab/Middle Eastern, Black, East Asian/Pacific Islander, Indigenous, Latin American, Mixed, South Asian, White Caucasian, Other |

COVID-19: The Case of Ontario

- Other studies have analyzed the dependency on vaccination status using socio-economic data.

COVID-19: The Case of Ontario

- Other studies have analyzed the dependency on vaccination status using socio-economic data.
- We could do the same, but what else can we get from this data?

COVID-19: The Case of Ontario

- Other studies have analyzed the dependency on vaccination status using socio-economic data.
- We could do the same, but what else can we get from this data?
 - There have been some interesting changes in Ontario with regard to healthcare.

COVID-19: The Case of Ontario

- Between 2006 and 2019, Ontario was geographically divided in “Local Health Integration Networks” (LHINs).

COVID-19: The Case of Ontario

- Between 2006 and 2019, Ontario was geographically divided in “Local Health Integration Networks” (LHINs).
- LHINs were essentially geographic intra-provincial divisions that determined where you could get health care.

COVID-19: The Case of Ontario

- Between 2006 and 2019, Ontario was geographically divided in “Local Health Integration Networks” (LHINs).
- LHINs were essentially geographic intra-provincial divisions that determined where you could get health care.
- There were 14 LHINs, with additional subdivisions.

COVID-19: The Case of Ontario

- Problems with the LHINs:

³Tsasis, Evans, and Owen 2012.

COVID-19: The Case of Ontario

- Problems with the LHINs:
- In multiple cases, the boundary of a LHIN did not match a municipal boundary.

³Tsasis, Evans, and Owen 2012.

COVID-19: The Case of Ontario

- Problems with the LHINs:
- In multiple cases, the boundary of a LHIN did not match a municipal boundary.
 - One part of a city would be in a LHIN whereas another part of it would be in another LHIN.

³Tsasis, Evans, and Owen 2012.

COVID-19: The Case of Ontario

- Problems with the LHINs:
- In multiple cases, the boundary of a LHIN did not match a municipal boundary.
 - One part of a city would be in a LHIN whereas another part of it would be in another LHIN.
 - Weakness in this approach due to complexity, lack of funding and bureaucracy were identified³.

³Tsasis, Evans, and Owen 2012.

COVID-19: The Case of Ontario

- In late 2019, Ontario adopted the Health Regions approach for healthcare and phased out the Local Health Integration Network (LHIN) approach.

COVID-19: The Case of Ontario

- In late 2019, Ontario adopted the Health Regions approach for healthcare and phased out the Local Health Integration Network (LHIN) approach.
- The change is relatively new. Multiple challenges:

COVID-19: The Case of Ontario

- In late 2019, Ontario adopted the Health Regions approach for healthcare and phased out the Local Health Integration Network (LHIN) approach.
- The change is relatively new. Multiple challenges:
 - Data for the Health Regions is not available from the Census.

COVID-19: The Case of Ontario

- In late 2019, Ontario adopted the Health Regions approach for healthcare and phased out the Local Health Integration Network (LHIN) approach.
- The change is relatively new. Multiple challenges:
 - Data for the Health Regions is not available from the Census.
 - **Have the Health Regions helped in reducing disparities in healthcare in the province?**

COVID-19: The Case of Ontario

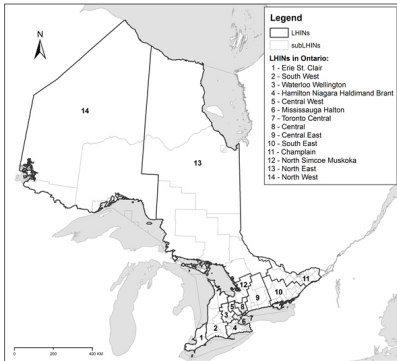


Figure 1: Ontario LHINs (Crighton et al. 2015)

COVID-19: The Case of Ontario

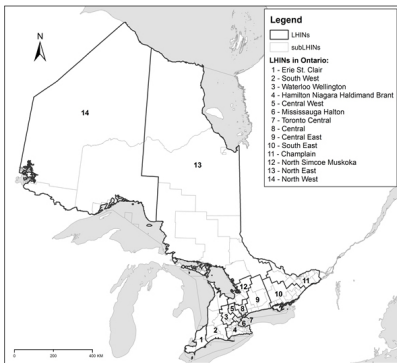


Figure 1: Ontario LHINs (Crighton et al. 2015)

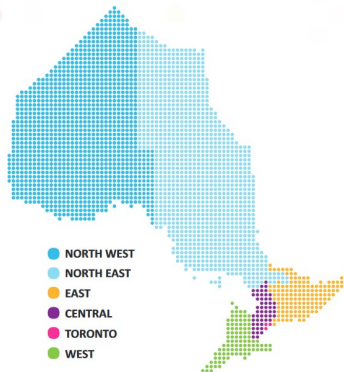


Figure 2: Ontario Health Regions (Ontario Business Health Plan 2022-2023)

COVID-19: The Case of Ontario

- Where in Ontario did responses come from?

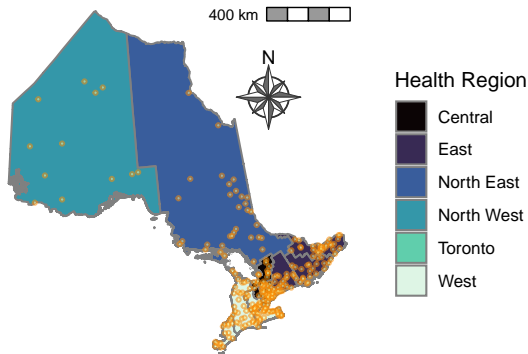


Figure 3: Geographic representation of the survey data collected by the Fields Institute

COVID-19: The Case of Ontario

- Therefore, we decided to integrate the different Health Regions in our analysis to determine the odds of vaccination.

$$\log \left(\frac{p(\text{vac})}{1 - p(\text{vac})} \right) = \beta_0 + \beta_1 (\text{Age group}) + \beta_2 \text{ Race} + \beta_3 \text{ Health Region} + \beta_4 \text{ Income} + \quad (1)$$

$$\beta_5 (\text{Health Region} \times \text{Race}) + \beta_6 (\text{Income} \times \text{Race})$$

Results

Table 2: **Selected** Multivariable Regression Results

| Characteristic | OR | 95% CI | p-value |
|---|------|------------|---------|
| Income (CAD) | | | |
| 60000 and above | — | — | |
| 25000-59999 | 0.59 | 0.39, 0.89 | 0.011 |
| under 25000 | 0.37 | 0.25, 0.56 | <0.001 |
| Race | | | |
| White/Caucasian | — | — | |
| Arab/Middle Eastern | 0.31 | 0.14, 0.69 | 0.004 |
| Black | 0.32 | 0.17, 0.60 | <0.001 |
| East Asian/Pacific Islander | 1.15 | 0.50, 2.66 | 0.7 |
| Indigenous | 0.44 | 0.19, 1.02 | 0.056 |
| Latin Aamerican | 0.28 | 0.11, 0.67 | 0.004 |
| Mixed | 0.64 | 0.25, 1.65 | 0.4 |
| Other | 0.22 | 0.12, 0.41 | <0.001 |
| South Asian | 0.91 | 0.49, 1.69 | 0.8 |
| Health Region | | | |
| Toronto | — | — | |
| Central | 1.47 | 0.92, 2.35 | 0.11 |
| East | 1.42 | 0.90, 2.23 | 0.13 |
| West | 1.55 | 1.05, 2.30 | 0.029 |
| Income and Race | | | |
| 25000-59999 * Arab/Middle Eastern | 1.79 | 0.67, 4.83 | 0.2 |
| under 25000 * Arab/Middle Eastern | 3.05 | 1.26, 7.39 | 0.013 |
| 25000-59999 * Black | 1.34 | 0.59, 3.05 | 0.5 |
| under 25000 * Black | 3.19 | 1.45, 6.99 | 0.004 |
| 25000-59999 * East Asian/Pacific Islander | 0.42 | 0.17, 1.05 | 0.062 |
| under 25000 * East Asian/Pacific Islander | 1.16 | 0.47, 2.86 | 0.8 |
| 25000-59999 * Indigenous | 1.36 | 0.48, 3.89 | 0.6 |
| under 25000 * Indigenous | 1.45 | 0.55, 3.80 | 0.5 |
| 25000-59999 * Latin American | 1.24 | 0.45, 3.43 | 0.7 |

Results

| Characteristic | OR | 95% CI | p-value |
|---------------------------------------|------|------------|---------|
| under 25000 * Latin American | 2.80 | 1.04, 7.51 | 0.041 |
| 25000-59999 * Mixed | 0.85 | 0.32, 2.26 | 0.7 |
| under 25000 * Mixed | 1.10 | 0.37, 3.27 | 0.9 |
| 25000-59999 * Other | 6.93 | 2.65, 18.1 | <0.001 |
| under 25000 * Other | 4.59 | 2.33, 9.05 | <0.001 |
| 25000-59999 * South Asian | 1.20 | 0.51, 2.85 | 0.7 |
| under 25000 * South Asian | 2.00 | 0.93, 4.30 | 0.077 |
| Race and Health Region | | | |
| Arab/Middle Eastern * Central | 0.66 | 0.26, 1.70 | 0.4 |
| Black * Central | 0.44 | 0.19, 0.98 | 0.046 |
| East Asian/Pacific Islander * Central | 0.98 | 0.38, 2.53 | >0.9 |
| Mixed * East | 0.91 | 0.28, 3.03 | 0.9 |
| other * East | 1.05 | 0.39, 2.83 | >0.9 |
| South Asian * East | 0.52 | 0.19, 1.45 | 0.2 |
| Arab/Middle Eastern * West | 1.00 | 0.37, 2.73 | >0.9 |
| Black * West | 0.76 | 0.32, 1.80 | 0.5 |
| East Asian/Pacific Islander * West | 0.52 | 0.20, 1.34 | 0.2 |
| Indigenous * West | 0.39 | 0.14, 1.09 | 0.073 |
| Latin American * West | 0.94 | 0.32, 2.72 | >0.9 |
| Mixed * West | 0.37 | 0.12, 1.16 | 0.089 |
| Other * West | 0.41 | 0.18, 0.93 | 0.032 |
| South Asian * West | 0.41 | 0.18, 0.95 | 0.037 |

¹ OR = Odds Ratio, CI = Confidence Interval

How do we interpret this?

- Our results show that there were disparities in vaccination uptake in Ontario.

How do we interpret this?

- Our results show that there were disparities in vaccination uptake in Ontario.
- People in certain racial minority groups had lower odds of vaccination than White/Caucasian individuals.

How do we interpret this?

- Our results show that there were disparities in vaccination uptake in Ontario.
- People in certain racial minority groups had lower odds of vaccination than White/Caucasian individuals.
- However, individuals that identified with a racial/ethnic minority and that were in a low household income bracket (<60k CAD) had higher odds of vaccination than individuals with a high household income.

How do we interpret this?

- Our results show that there were disparities in vaccination uptake in Ontario.
- People in certain racial minority groups had lower odds of vaccination than White/Caucasian individuals.
- However, individuals that identified with a racial/ethnic minority and that were in a low household income bracket (<60k CAD) had higher odds of vaccination than individuals with a high household income.
- This is likely caused by the type of occupation: people in racial minorities, and those with a low household income work in essential occupations⁴, and thus potentially got the vaccine to be able to work.

⁴Hawkins 2020.

How do we interpret this?

- But there are also intra-provincial differences in vaccine uptake within the Health Regions:

How do we interpret this?

- But there are also intra-provincial differences in vaccine uptake within the Health Regions:
 - For example, South Asian individuals in the West Health Region had lower odds of vaccination than in other Health Regions.

How do we interpret this?

- But there are also intra-provincial differences in vaccine uptake within the Health Regions:
 - For example, South Asian individuals in the West Health Region had lower odds of vaccination than in other Health Regions.
 - These results provide a more comprehensive assessment of COVID-19 vaccination rates within Ontario, as they showed that certain minority groups within specific income brackets and certain Health Regions had differences in vaccination.

Conclusions

- Data cleaning is **important**}

Conclusions

- Data cleaning is **important**
 - Unifying geographical data can be challenging

Conclusions

- Data cleaning is **important**
 - Unifying geographical data can be challenging
 - Specially because most data relies on legacy information from the LHINs

Conclusions

- Data cleaning is **important**
 - Unifying geographical data can be challenging
 - Specially because most data relies on legacy information from the LHINs
- A more granular view of data (in this case, examining differences within Health Region, Income and Race) can provide insight for public policy development.

Conclusions

- Data cleaning is **important**
 - Unifying geographical data can be challenging
 - Specially because most data relies on legacy information from the LHINs
- A more granular view of data (in this case, examining differences within Health Region, Income and Race) can provide insight for public policy development.
- There is a need for future studies that examine more in detail these differences and can provide a rationale.

The Case of Biomedical Data

Longitudinal Data

- Biomedical studies often collect longitudinal data to see the effect of an intervention over time:

Longitudinal Data

- Biomedical studies often collect longitudinal data to see the effect of an intervention over time:
 - How a chemotherapy treatment changes the metabolism of a tumor

Longitudinal Data

- Biomedical studies often collect longitudinal data to see the effect of an intervention over time:
 - How a chemotherapy treatment changes the metabolism of a tumor
 - How the concentration of a drug changes over time in the blood

Longitudinal Data

- Biomedical studies often collect longitudinal data to see the effect of an intervention over time:
 - How a chemotherapy treatment changes the metabolism of a tumor
 - How the concentration of a drug changes over time in the blood
- How is this data typically analyzed?

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + \beta_2 \times time_t + \beta_3 \times time_t \times treatment_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + \beta_2 \times time_t + \beta_3 \times time_t \times treatment_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

β_0 : the mean group value

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times \text{treatment}_j + \beta_2 \times \text{time}_t + \beta_3 \times \text{time}_t \times \text{treatment}_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

β_0 : the mean group value

$\text{time}_t, \text{treatment}_j$: fixed effects

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times \text{treatment}_j + \beta_2 \times \text{time}_t + \beta_3 \times \text{time}_t \times \text{treatment}_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

β_0 : the mean group value

$\text{time}_t, \text{treatment}_j$: fixed effects

β_1, β_2 and β_3 : linear slopes of the fixed effects.

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times \text{treatment}_j + \beta_2 \times \text{time}_t + \beta_3 \times \text{time}_t \times \text{treatment}_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

β_0 : the mean group value

$\text{time}_t, \text{treatment}_j$: fixed effects

β_1, β_2 and β_3 : linear slopes of the fixed effects.

ε_{ijt} : error, assumed to be $\sim N(0, \sigma^2)$

Linear Models

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + \beta_2 \times time_t + \beta_3 \times time_t \times treatment_j + \varepsilon_{ijt} \quad (2)$$

where,

y_{ijt} : is the response for subject i in treatment group j at time t

β_0 : the mean group value

$time_t, treatment_j$: fixed effects

β_1, β_2 and β_3 : linear slopes of the fixed effects.

ε_{ijt} : error, assumed to be $\sim N(0, \sigma^2)$

A LMEM follows the same exact structure, only incorporates a random effect α_{ij} , which allows for different intercepts.

Trends Over Time

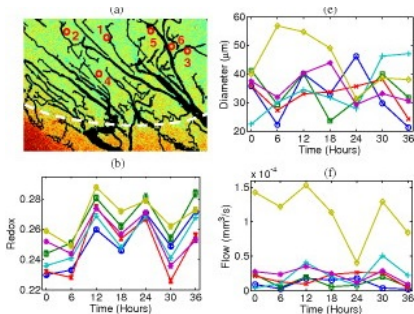


Figure 4: Tumor imaging data
(Skala et al. 2010)

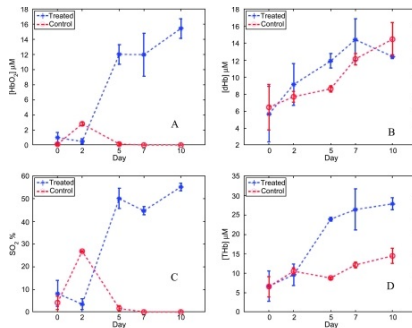


Figure 5: Tumor oxygenation data
(Vishwanath et al. 2009)

Trends Over Time

- The issue in those data is that the trends are not linear, and therefore, a linear model will miss changes in the signal where some metabolic or physiological relevant change is taking place.

⁵Beck and Jackman 1998.

Trends Over Time

- The issue in those data is that the trends are not linear, and therefore, a linear model will miss changes in the signal where some metabolic or physiological relevant change is taking place.
- Polynomial effects can be used, but they create biases at the boundaries of the covariates⁵.

⁵Beck and Jackman 1998.

Generalized Additive Models (GAMs)

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + f(time_t | \beta_j) + \varepsilon_{ijt} \quad (3)$$

- The change of y_{ijt} over time is represented by the *smooth function* $f(time_t | \beta_j)$ with inputs as the covariates $time_t$ and parameters β_j .

Generalized Additive Models (GAMs)

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + f(time_t | \beta_j) + \varepsilon_{ijt} \quad (3)$$

- The change of y_{ijt} over time is represented by the *smooth function* $f(time_t | \beta_j)$ with inputs as the covariates $time_t$ and parameters β_j .

Generalized Additive Models (GAMs)

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + f(time_t | \beta_j) + \varepsilon_{ijt} \quad (3)$$

- The change of y_{ijt} over time is represented by the *smooth function* $f(time_t | \beta_j)$ with inputs as the covariates $time_t$ and parameters β_j .
- We can use a *basis function* to estimate the smooth function.

Generalized Additive Models (GAMs)

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + f(time_t | \beta_j) + \varepsilon_{ijt} \quad (3)$$

- The change of y_{ijt} over time is represented by the *smooth function* $f(time_t | \beta_j)$ with inputs as the covariates $time_t$ and parameters β_j .
- We can use a *basis function* to estimate the smooth function.

Generalized Additive Models (GAMs)

$$y_{ijt} = \beta_0 + \beta_1 \times treatment_j + f(time_t | \beta_j) + \varepsilon_{ijt} \quad (3)$$

- The change of y_{ijt} over time is represented by the *smooth function* $f(time_t | \beta_j)$ with inputs as the covariates $time_t$ and parameters β_j .
- We can use a *basis function* to estimate the smooth function.
- Splines are helpful as basis functions: Thin plate regression splines (TPRS) are computationally efficient, and the underlying principle is that of polynomial pieces “joined” together

How GAMs work

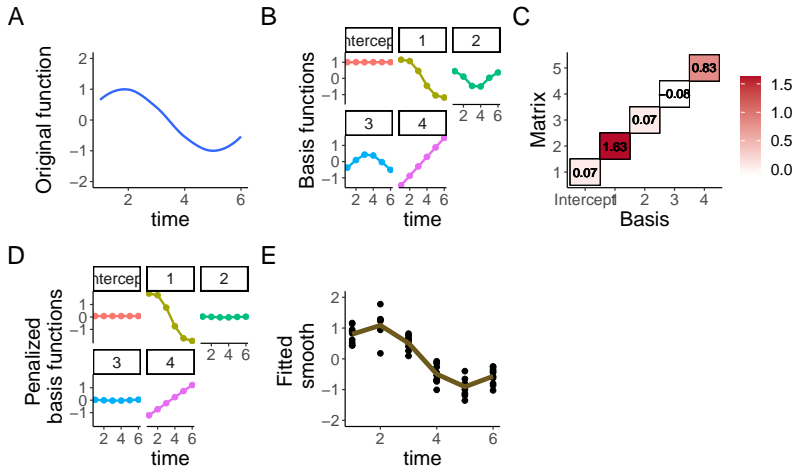


Figure 6: Fitting process of a GAM.

An Example

- Simulated data from a study on radiotherapy in a mouse model of melanoma⁶.

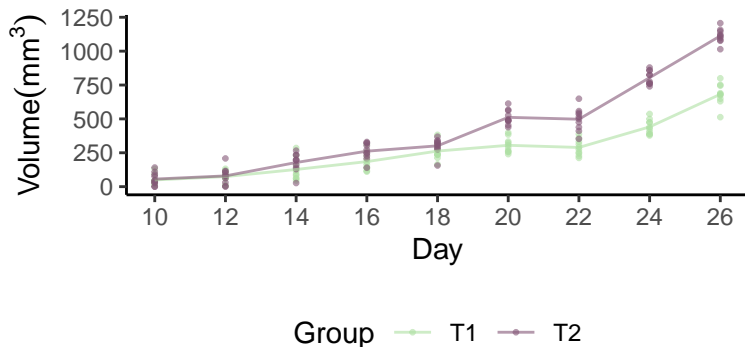


Figure 7: Tumor volume in two groups of tumors under radiotherapy

⁶Sen et al. 2011.

Fitting a GAM

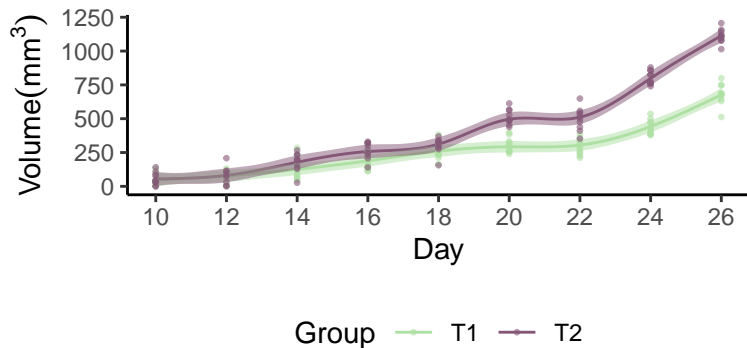


Figure 8: GAM fitted to simulated data

- The model captures the trend of the data

Fitting a GAM

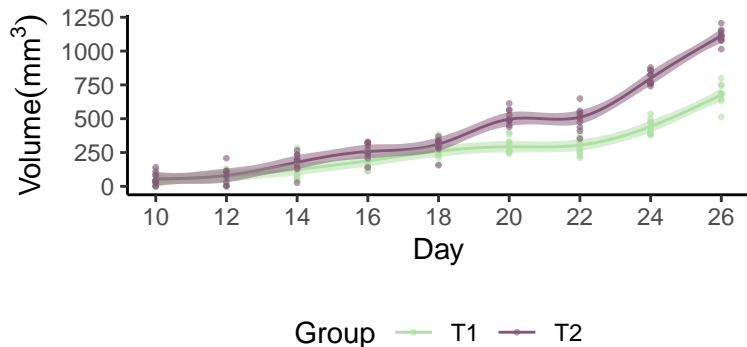


Figure 8: GAM fitted to simulated data

- The model captures the trend of the data
- We can furthermore compare the trends.

Differences

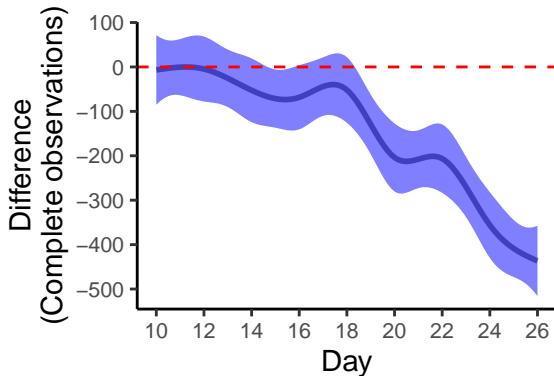


Figure 9: Pairwise comparisons between smooths

- We can compare the smooths for each group. Here, we see that T2 is significantly higher after day 18.

Differences

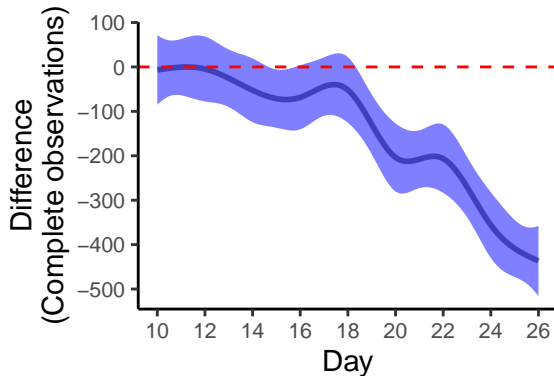


Figure 9: Pairwise comparisons between smooths

- We can compare the smooths for each group. Here, we see that T2 is significantly higher after day 18.
- This can give an idea of further explorations of biological

Conclusions

- GAMs are useful to analyze longitudinal data because they provide:

Conclusions

- GAMs are useful to analyze longitudinal data because they provide:
 - A model that captures non-linear trends in the data

Conclusions

- GAMs are useful to analyze longitudinal data because they provide:
 - A model that captures non-linear trends in the data
 - This allows to examine specific time points that might be of interest, where metabolic, or physiological relevant changes might be occurring

Conclusions

- GAMs are useful to analyze longitudinal data because they provide:
 - A model that captures non-linear trends in the data
 - This allows to examine specific time points that might be of interest, where metabolic, or physiological relevant changes might be occurring
 - Lets the data speak for itself

Addressing Reproducibility

- There is an ongoing need of making papers reproducible.

Addressing Reproducibility

- There is an ongoing need of making papers reproducible.
- This is specially important in the case of data/methods of health research.

Addressing Reproducibility

- There is an ongoing need of making papers reproducible.
- This is specially important in the case of data/methods of health research.
 - Otherwise, tools cannot be used by others.

Addressing Reproducibility

- There is an ongoing need of making papers reproducible.
- This is specially important in the case of data/methods of health research.
 - Otherwise, tools cannot be used by others.
- How are we addressing this in our research?

Addressing Reproducibility

- Using GitHub to share:

Addressing Reproducibility

- Using GitHub to share:
 - Data: Making publicly available the datasets used

Addressing Reproducibility

- Using GitHub to share:
 - Data: Making publicly available the datasets used
 - Methods: Sharing the code used for statistical analyses

Addressing Reproducibility

- Using GitHub to share:
 - Data: Making publicly available the datasets used
 - Methods: Sharing the code used for statistical analyses
- In synthesis, sharing all the information used to create a paper such that anyone can re-create the analysis, results, and the paper itself from the files provided.

Addressing Reproducibility

- For GAMs

<https://github.com/aimundo/GAMs-biomedical-research>

Addressing Reproducibility

- For GAMs
<https://github.com/aimundo/GAMs-biomedical-research>
- COVID-19: Work is ongoing, but repository will be ready when paper is submitted

Conclusion

- There is an ongoing need of analyzing public health data to address important disparities in areas such as vaccination.

Conclusion

- There is an ongoing need of analyzing public health data to address important disparities in areas such as vaccination.
- Semi-parametric statistical to analyze biomedical/public health longitudinal data, such as GAMs can provide better insight on periods where important biological changes might occur.

Acknowledgements

- The Nasri Lab (Université de Montréal)
 - Bouchra Nasri, PhD (PI)
 - Idriss Sekkak, PhD
 - Rado Ramasy
 - Fatima El-Mousawi
 - Rawda Berkat
- The Muldoon Lab (University of Arkansas)
 - Timothy J. Muldoon (PI)
- John R. Tipton (Los Alamos National Laboratory)



FIELDS INSTITUTE
FOR RESEARCH IN MATHEMATICAL SCIENCES

Arkansas
BIOSCIENCES
INSTITUTE



-  Beck, Nathaniel and Simon Jackman (Apr. 1998). “Beyond Linearity by Default: Generalized Additive Models”. In: *American Journal of Political Science* 42.2, p. 596. DOI: 10.2307/2991772. URL: <https://doi.org/10.2307/2991772>.
-  Gerretsen, Philip et al. (Nov. 2021). “Individual determinants of COVID-19 vaccine hesitancy”. In: *PLOS ONE* 16.11. Ed. by Leeberk Raja Inbaraj, e0258462. DOI: 10.1371/journal.pone.0258462. URL: <https://doi.org/10.1371/journal.pone.0258462>.
-  Hawkins, Devan (June 2020). “Differential occupational risk for COVID-19 and other infection exposure according to race and ethnicity”. In: *American Journal of Industrial Medicine* 63.9, pp. 817–820. DOI: 10.1002/ajim.23145. URL: <https://doi.org/10.1002/ajim.23145>.

-  Nafilyan, Vahe et al. (July 2021). “Sociodemographic inequality in COVID-19 vaccination coverage among elderly adults in England: a national linked data study”. In: *BMJ Open* 11.7, e053402. DOI: 10.1136/bmjopen-2021-053402. URL: <https://doi.org/10.1136/bmjopen-2021-053402>.
-  Sen, Arindam et al. (May 2011). “Mild Elevation of Body Temperature Reduces Tumor Interstitial Fluid Pressure and Hypoxia and Enhances Efficacy of Radiotherapy in Murine Tumor Models”. In: *Cancer Research* 71.11, pp. 3872–3880. DOI: 10.1158/0008-5472.can-10-4482. URL: <https://doi.org/10.1158/0008-5472.can-10-4482>.
-  Tsasis, Peter, Jenna M. Evans, and Susan Owen (Sept. 2012). “Reframing the challenges to integrated care: a complex-adaptive systems perspective”. In: *International Journal of Integrated Care* 12.5. DOI: 10.5334/ijic.843. URL: <https://doi.org/10.5334/ijic.843>.