



Artificial Health Intelligence-Making sense of health data



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Research Excellence Funds awarded to four UM academics

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Data Science Research Group RESEARCH 08:57.04 Jan



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RESEARCH 10:19, 22 Dec





Global health challenges





Global health challenges

- 1. Obesity and Chronic diseases
- 2. Aging
- 3. Drug resistance, Hospital acquired infections and medical errors
- 4. Global warming and pollution
- 5. Health inequality and healthcare finance
- 6. Infectious and/or zoonotic diseases and viruses
- 7. Stress and sleep aponia
- 8. Relationships and social health



Future healthcare technologies





Future healthcare technologies

- 1. Artificial Intelligence in Healthcare: AI/ML predictive/prescriptive analytics and digital twin
- 2. Sensors: Smart wearables, cyborg, satellites, wireless sensor networks and IoTs
- 3. 5G/6G: Cloud computing, telemedicine and Mobile health
- 4. Robotics (computer vision and natural language processing)
- 5. Gene editing, genomics, epigenomics proteomics and metabolomic

Enabler: Globalization and economic growth

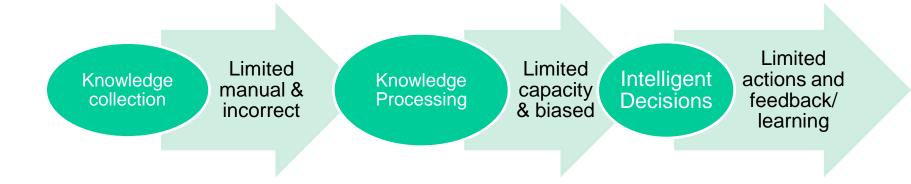


Why Artificial Intelligence in Healthcare?





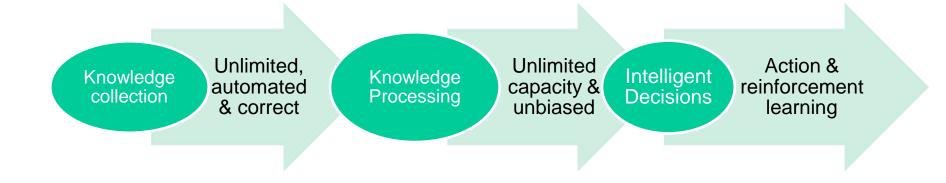
Human Intelligence



https://www.chirpbooks.com/audiobooks/paradise-lost-by-john-milton



Artificial Intelligence



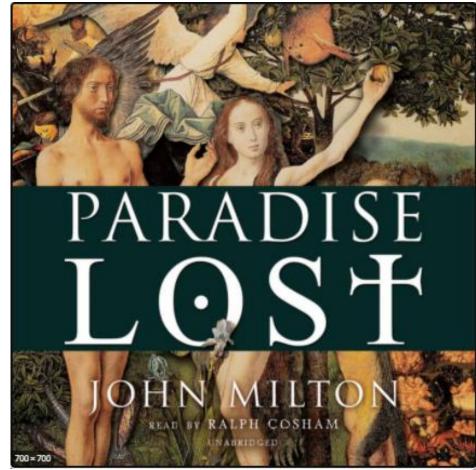
https://www.chirpbooks.com/audiobooks/paradise-lost-by-john-milton



Cognitive biases







https://www.chirpbooks.com/audiobooks/paradise-lost-by-john-milton

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Where ignorance is bliss, 'Tis folly to be wise. Thomas Gray

https://www.brainyquote.com/quotes/thomas_gray_150669







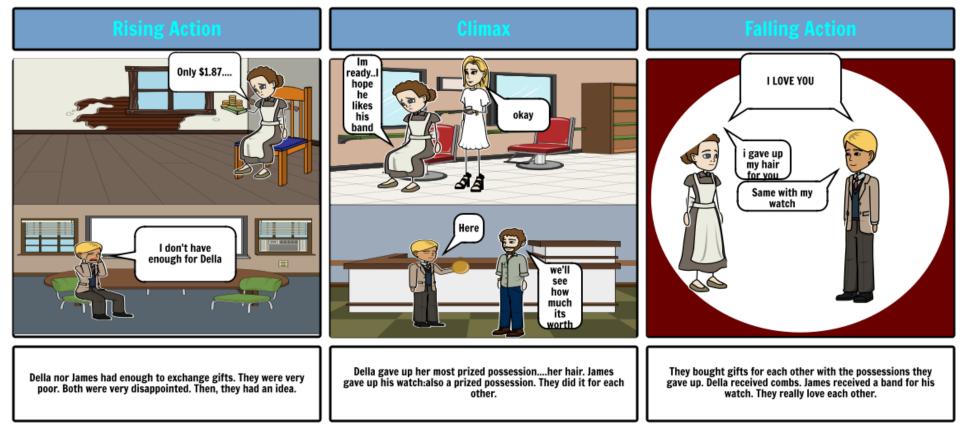
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Requires

1. Human Behavioural Modelling





Requires

- 1. Human Behavioural Modelling
- 2. Modelling the effect of others' Behaviour (using game theory),





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- 1. Human Behavioural Modelling
- 2. Modelling the effect of others' Behaviour (using game theory),
- 3. Modelling of cultural, social, economical, financial and environmental effects (Big data analytics),





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- 4. Most difficult: modelling spontaneous (uncorrelated) changes in sentiments,





Requires

- 1. Human Behavioural Modelling
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- 4. Most difficult: modelling spontaneous (uncorrelated) changes in sentiments,

Reality vs perception.



Reality vs Perception







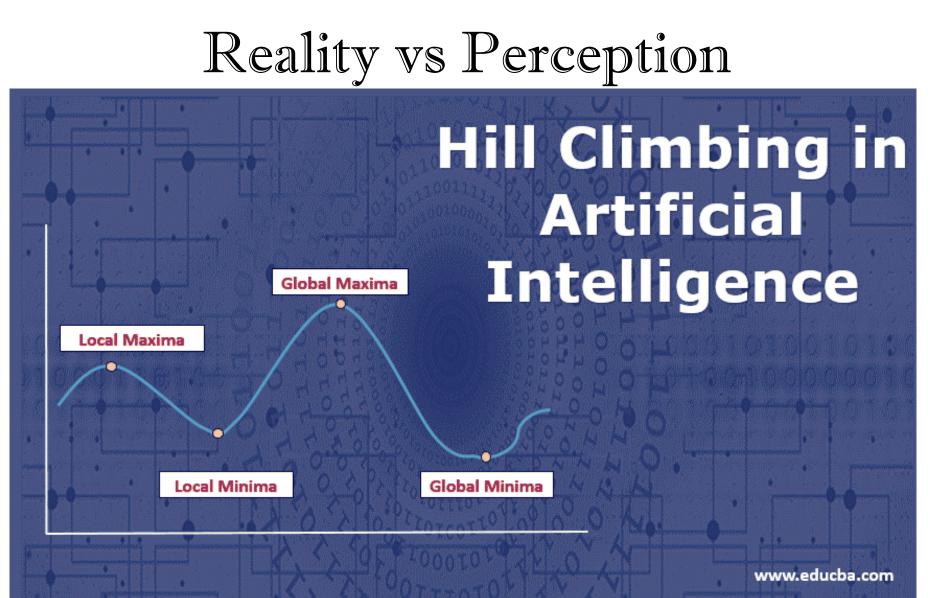
Reality vs Perception

Karmanye Vadhikaraste. Ma Phalesha Kadachana..

WWW.SWAMIRARA.COM



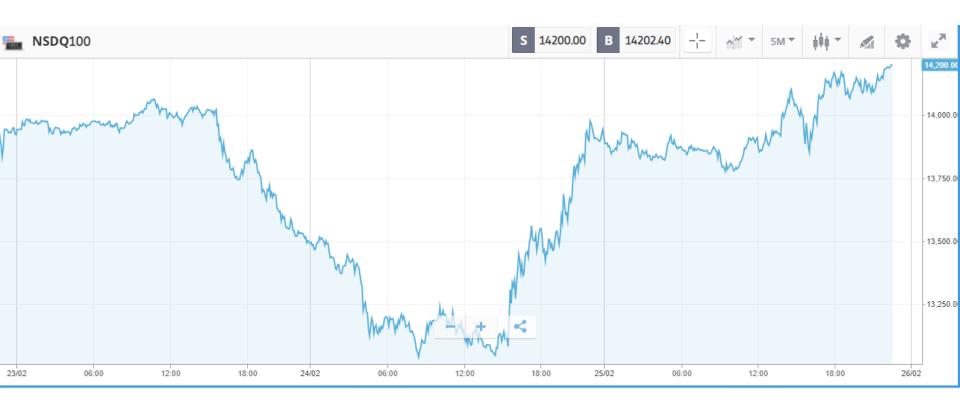








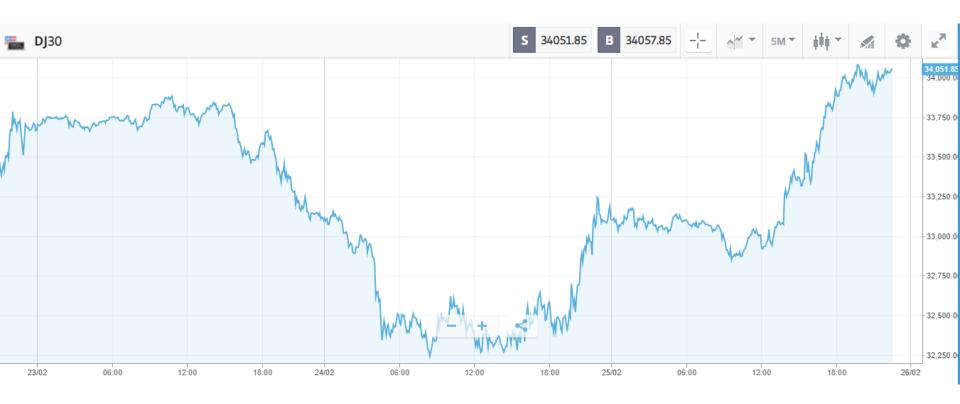
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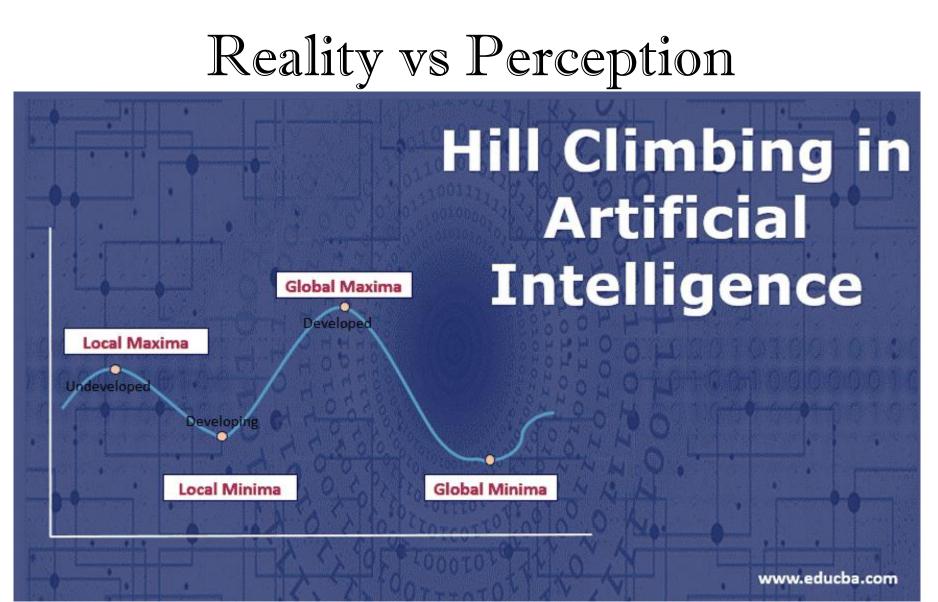


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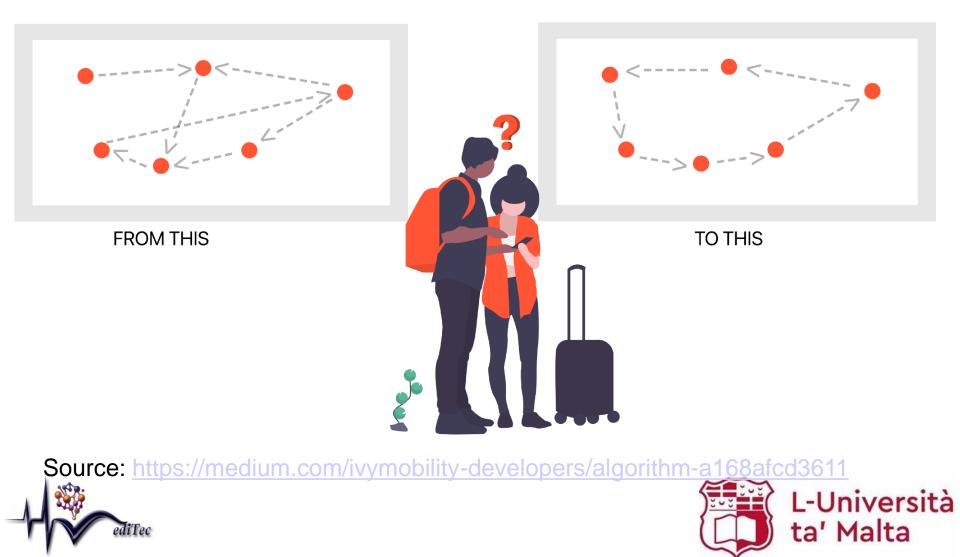






Reality vs Perception

Greedy Algorithm



Optimism bias in COVID-19



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23% migrant workers walked back to villages during coronavirus lockdown

Despite the hardships some optimism remains as 33 per cent respondents said they wanted to return to the city they worked in once the lockdown is lifted

BusinessToday.In | August 11, 2020 | Updated 08:41 IST







Coronavirus lockdown: The Indian migrants dying to get home

By Vikas Pandey BBC News, Delhi

3 20 May 2020

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Coronavirus pandemic





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Coronavirus: India's pandemic lockdown turns into a human tragedy



30 March 2020





Coronavirus: Heartbreaking scenes as India lockdown sparks mass migration



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Covid-19 2.0: Migrant crisis returns! Indian Railways steps up special services

By: FE Bureau | April 22, 2021 2:30 AM

At present, the Indian Railways is running 1,512 mail/express and festival specials, on an average per day, up from the 1,490 such services last week.



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Other biases





- Confirmation bias
- Anchoring bias
- Bandwagon effect
- Halo effect
- Availability bias/heuristic
- Ostrich effect
- Recency/serial position effect
- Choice-supportive bias





- Fundamental attribution error
- Outcome bias
- Illusory correlation bias
- Dunning Kruger effect
- Exponential-growth bias
- Magical Beliefs
- Conspiracy Theory Beliefs
- Overconfidence





- Conformity bias
- Authority bias
- Loss-aversion bias
- False causality bias
- Action bias
- Self-serving bias
- Framing bias
- Ambiguity bias





- Strategic misrepresentation
- Projection bias
- Pro-innovation bias
- Status-quo bias
- Feature positive effect
- Bounded Rationality
- Certainty Effect
- Cognitive Dissonance





- Commitment
- Decision Fatigue
- Decoy Effect
- Time Discounting / Present Bias
- Diversification Bias
- Ego Depletion
- Elimination-By-Aspects
- Hot-Cold Empathy Gap





- Commitment
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- Ego Depletion
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- Endowment Effect
- Fear of Missing Out (FOMO)
- Gambler's Fallacy (Monte Carlo Fallacy)
- Habit
- Hedonic Adaptation
- Herd Behaviour
- Hindsight Bias (Knew-It-All-Along Effect)
- IKEA Effect





- Less-Is-Better Effect
- Licensing Effect
- Mental Accounting
- Naive Diversification
- Over justification Effect
- Pain of paying
- Partitioning
- Peak-End Rule





- Priming
- Procrastination
- Ratio bias
- Reciprocity
- Regret aversion
- Representativeness heuristic
- Scarcity
- Social proof





- Sunk Cost Fallacy
- Zero Price Effect









Our Economy: A Complex System More demand then supply = More profit





More demand then supply = More profit More profit = More attractive industry





More demand then supply = More profit More profit = More attractive industry = More players





More demand then supply = More profit

- = More players
- = More supply than demand





More demand then supply = More profit

- = More players
- = More supply than demand
- = Less price = Less profit





More demand then supply = More profit

- = More players
- = More supply than demand
- = Less price = Less profit
- = Some will leave the market with loss





More demand then supply = More profit

- = More players
- = More supply than demand
- = Less price = Less profit
- = Some will leave the market with loss
- = More demand than supply























Private healthcare:

• Some patients want cheap healthcare





- Some patients want cheap healthcare
- Some patients want best (luxurious) healthcare





- Some patients want cheap healthcare
- Some patients want best (luxurious) healthcare
- Health providers want maximum profit





- Some patients want cheap healthcare
- Some patients want best (luxurious) healthcare
- Health providers want maximum profit
- maximum profit = maximum hospital visits





Private healthcare:

- Some patients want cheap healthcare
- Some patients want best (luxurious) healthcare
- Health providers want maximum profit
- maximum profit = maximum hospital visits

= maximum readmissions

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+ maximum hospital duration of stay

Private healthcare:

• At the time of COVID-19 pandemic:





- At the time of COVID-19 pandemic:
- Should they make more hospitals and employ more health professionals?





- At the time of COVID-19 pandemic:
- Should they make more hospitals and employ more health professionals?
- Can they make more hospitals and employ more health professionals instantly?











Public healthcare:

• Everyone gets the same healthcare





- Everyone gets the same healthcare
- Health providers want minimum cost





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = least duration in hospitals + minimum admissions





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = least duration in hospitals + waiting list





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = more readmissions + waiting list





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = more readmissions + waiting list
 - = longer waiting list







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- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = longer waiting list
 - = Poor healthcare





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = longer waiting list
 - = Poor healthcare
 - = Public outcry
 - = Preference





- Everyone gets the same healthcare
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = longer waiting list
 - = Poor healthcare
 - = Public outcry
 - = Preference





- Corruption
- Health providers want minimum cost
- Minimum cost
 - = Limited resources
 - = longer waiting list
 - = Poor healthcare
 - = Public outcry
 - = Preference





- Public healthcare:
- More resources





Public healthcare:

• More resources = More cost





- Public healthcare:
- More resources = short waiting lists





- More resources = short waiting lists
- Short waiting list = longer hospital stay





- More resources = short waiting lists
- Short waiting list = longer hospital stay minimum readmissions





- More resources = short waiting lists
- Short waiting list = longer hospital stay minimum readmissions more patients





- Even more resources = no waiting lists
- Short waiting list = longer hospital stay minimum readmissions more patients underutilization





Public healthcare:

- Even more resources = no waiting lists
- Short waiting list = longer hospital stay minimum readmissions
 more patients underutilization

misuse





Public healthcare:

- Even more resources = no waiting lists
- Short waiting list = longer hospital stay

minimum readmissions

more patients

underutilization

misuse

more cost





Public healthcare:

- Even more resources = no waiting lists
- Short waiting list = longer hospital stay

minimum readmissions

more patients

underutilization

misuse

more cost

Some waiting lists

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Public healthcare:

• Optimum resources = optimum waiting time





- Optimum resources = optimum waiting time
- = Optimum hospital stay





- Optimum resources = optimum waiting time
 - = Optimum hospital stay= minimum readmissions





- Optimum resources = optimum waiting time
 - = Optimum hospital stay
 - = minimum readmissions
 - = optimum patients' number





- Optimum resources = optimum waiting time
 - = Optimum hospital stay
 - = minimum readmissions
 - = optimum patients' number
 optimum utilization





- Optimum resources = optimum waiting time
 - = Optimum hospital stay
 - = minimum readmissions
 - optimum patients' number
 optimum utilization
 minimum misuse





- Optimum resources = optimum waiting time
 - = Optimum hospital stay
 - = minimum readmissions
 - = optimum patients' number optimum utilization minimum misuse optimum cost





Public healthcare:

• Optimum resources = optimum waiting time

= Optimum hospital stay

= minimum readmissions

= optimum patients' number optimum utilization minimum misuse optimum cost

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Some waiting lists



Public healthcare:

• Optimum resources = Proper planning

= Continuously adding resources (if population is increasing/changing)

= Resource requirement forecasting





Public healthcare:

• Optimum resources = Proper planning

= Continuously adding resources (if population is increasing/changing)

- = Resource requirement forecasting
- = Admission rate estimation
- = Length of stay estimation





- At the time of COVID-19 pandemic:
- Should they make more hospitals and employ more health professionals?
- Can they make more hospitals and employ more health professionals instantly?

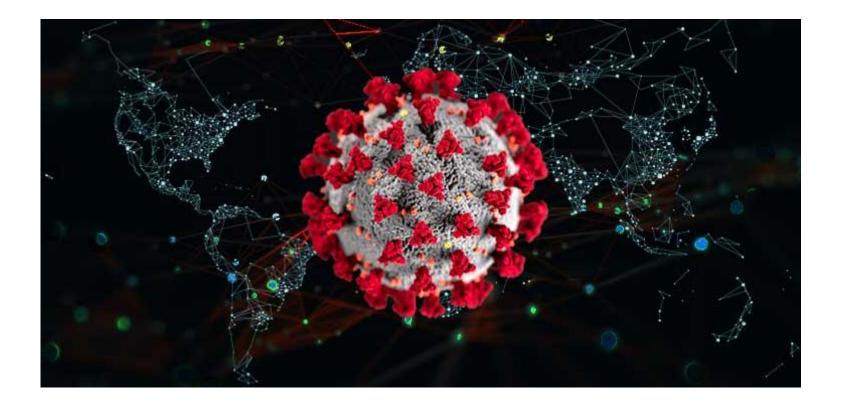




- Do we prepare ourselves for such a pandemic?
- Do we keep a hospital bed booked for us?
- Do we keep resources for homecare?

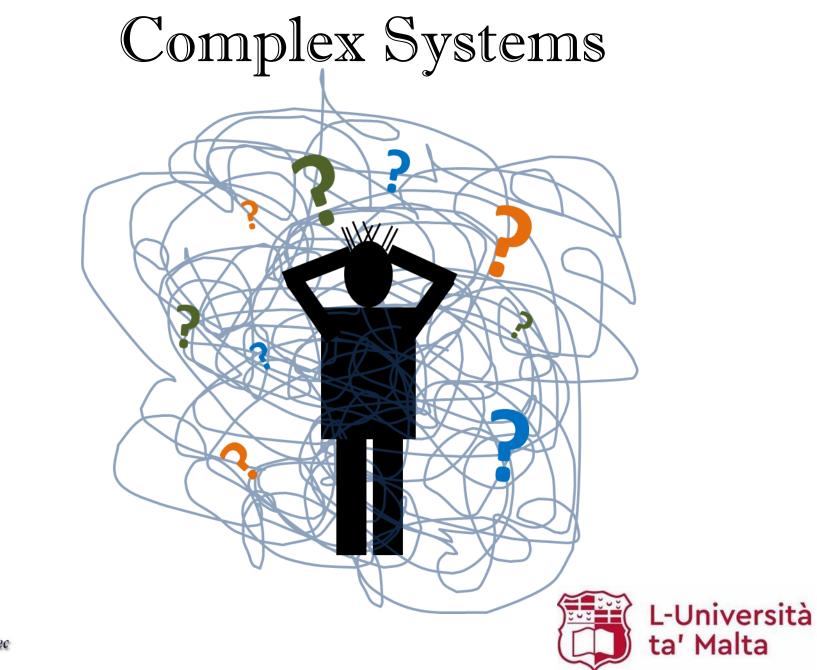


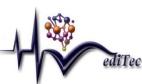












We need an efficient mechanism/system for Acquiring/collecting, analysing, and sharing information from different domains of the complex system for decision making to fight COVID-19 and monitor the progress. I.e. we need Artificial Intelligence in Healthcare.





Covid-19 is not just a medical problem but It is also a

- 1. Social problem
- 2. Political problem
- 3. Cultural problem
- 4. Community problem
- 5. Communication problem
 - 5. Transportation problem



- 7. Management problem
- 8. Supply chain problem
- 9. Administration problem
- 10.Education problem
- 11. Financial problem
- 12. Economical problem
- 13.Behavioural/ psychological problem
 - 14.Geological problem



Some COVID-19 Statistics

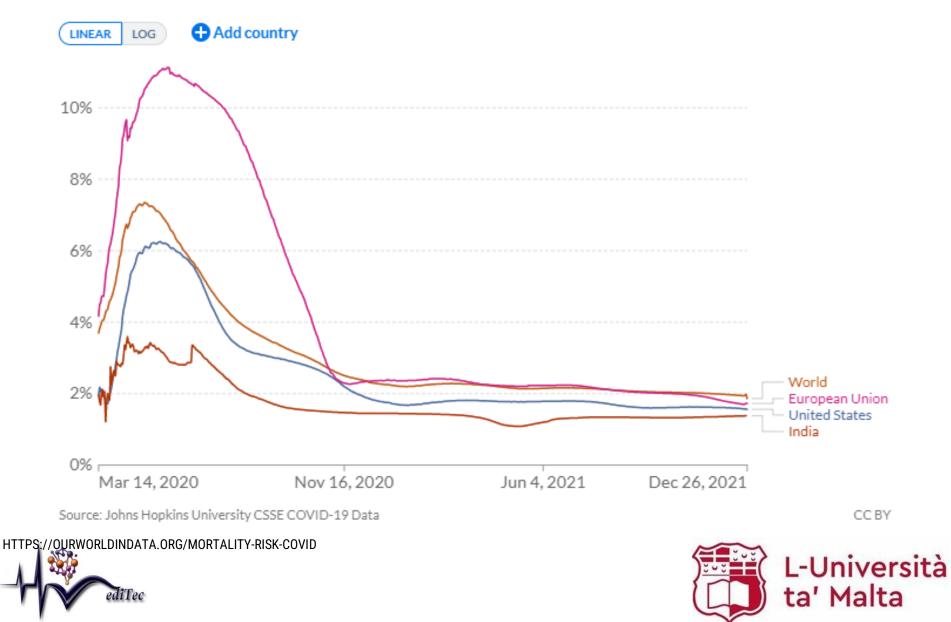




Case fatality rate of COVID-19



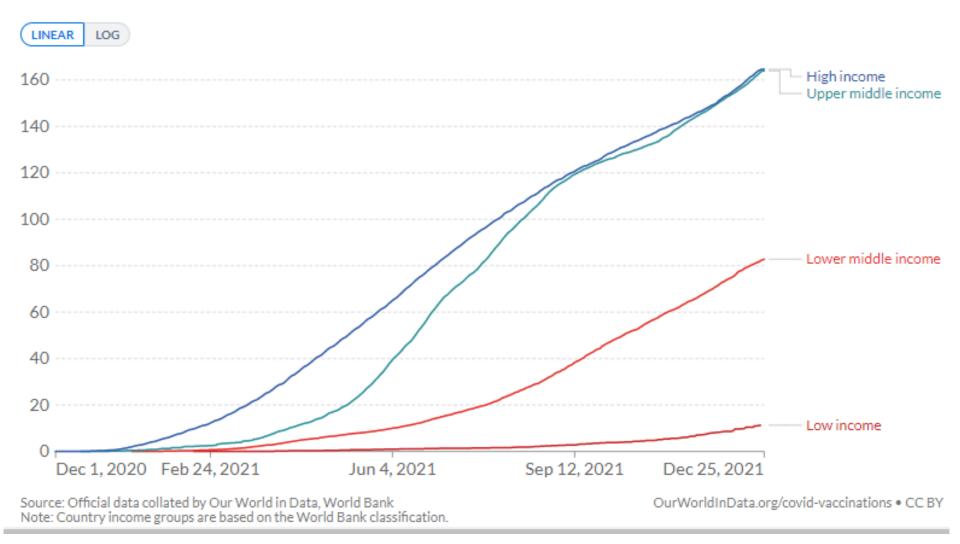
The case fatality rate (CFR) is the ratio between confirmed deaths and confirmed cases. The CFR can be a poor measure of the mortality risk of the disease. We explain this in detail at OurWorldInData.org/mortality-risk-covid



COVID-19 vaccine doses administered per 100 people, by income group



All doses, including boosters, are counted individually. As the same person may receive more than one dose, the number of doses can be higher than the number of people in the population.

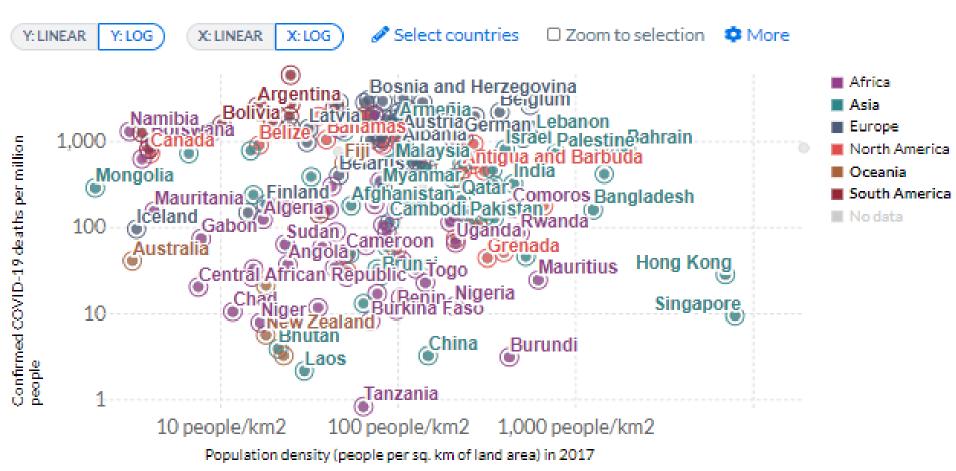






COVID-19 death rate vs. Population density, Sep 4, 2021

The death rate is the number of total confirmed deaths due to COVID-19 per million people.



Source: Johns Hopkins University CSSE COVID-19 Data – Last updated 5 September, 09:03 (London time), World Bank OurWorldInData.org/coronavirus • CC BY

TOTAL CONFIRMED COVID-19 DEATHS PER MILLION VS GDP PER CAPITA, JAN 6, 2021 HTTPS://OURWORLDINDATA.ORG/GRAPHER/TOTAL-CONFIRMED-DEATHS-OF-COVID-19-PER-MILLION-PEOPLE-VS-GDP-PER-CAPITA?YSCALE=LINEAR&TIME=LATEST





Our World in Data

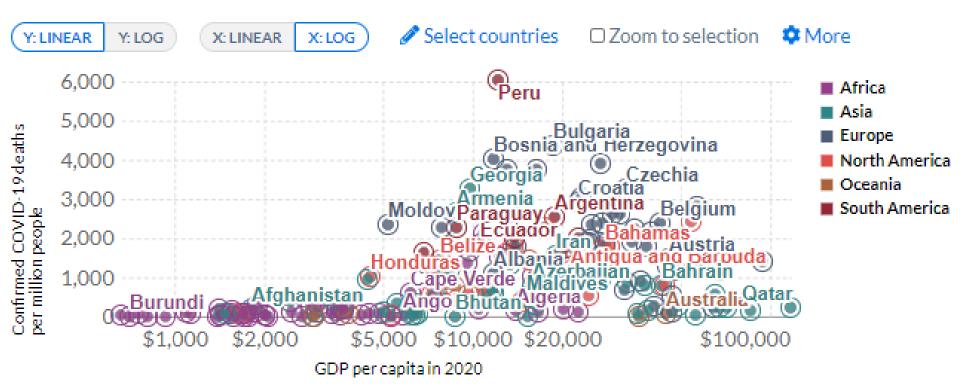
Total confirmed COVID-19 deaths per million vs GDP per capita, Dec 17, 2021

Due to limited testing and challenges in the attribution of the cause of death, confirmed deaths can be lower than the true number of deaths. GDP per capita is adjusted for price differences between countries (it is expressed in international dollars).

Our World in Data

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Source: Johns Hopkins University CSSE COVID-19 Data – Last updated 18 December, 05:05 (London time), World Bank OurWorldInData.org/coronavirus • CC BY

TOTAL CONFIRMED COVID-19 DEATHS PER MILLION VS GDP PER CAPITA, JAN 6, 2021 HTTPS://OURWORLDINDATA.ORG/GRAPHER/TOTAL-CONFIRMED-DEATHS-OF-COVID-19-PER-MILLION-PEOPLE-VS-GDP-PER-CAPITA?YSCALE=LINEAR&TIME=LATEST



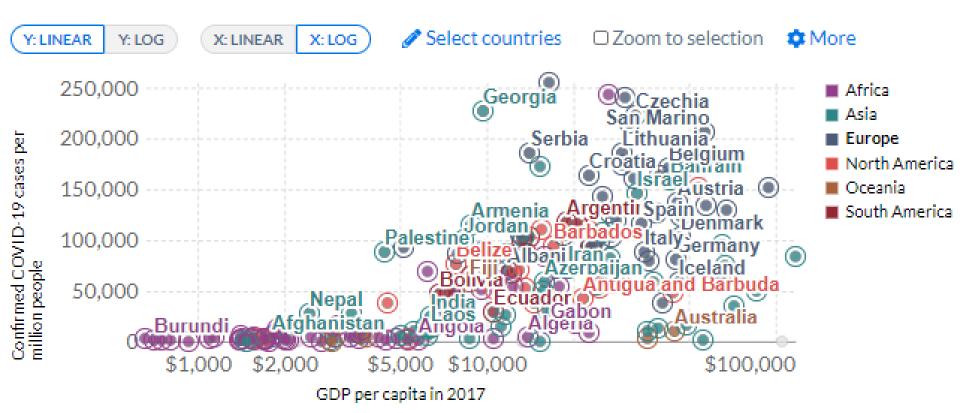
Cumulative confirmed COVID-19 cases per million vs. GDP per capita, Dec 17, 2021

Due to limited testing, the number of confirmed cases is lower than the true number of infections. GDP per capita is adjusted for price differences between countries (it is expressed in international dollars).

Our World in Data

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Source: Johns Hopkins University CSSE COVID-19 Data – Last updated 18 December, 05:05 (London time), World Bank OurWorldInData.org/coronavirus • CC BY

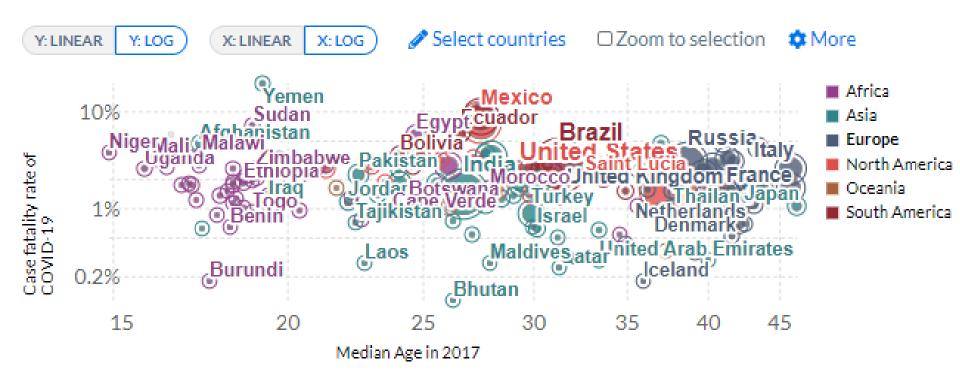
CUMULATIVE CONFIRMED COVID-19 CASES PER MILLION VS. GDP PER CAPITA, JAN 6, 2021: REF: HTTPS://OURWORLDINDATA.ORG/GRAPHER/TOTAL-CONFIRMED-CASES-OF-COVID-19-PER-MILLION-PEOPLE-VS-GDP-PER-CAPITA



<u>Case fatality rate of COVID-19 vs. Median age of the</u> <u>population</u>



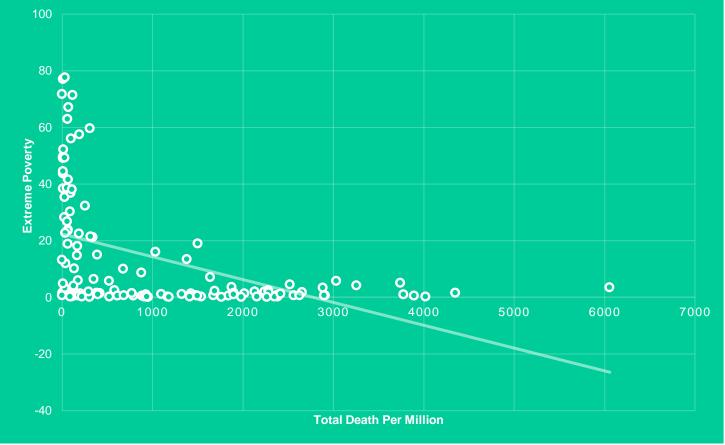
The Case Fatality Rate (CFR) is the ratio between confirmed deaths and confirmed cases. During an outbreak of a pandemic the CFR is a poor measure of the mortality risk of the disease. We explain this in detail at OurWorldInData.org/coronavirus



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: REF: HTTPS://OURWORLDINDATA.ORG/GRAPHER/CASE-FATALITY-RATE-OF-COVID-19-VS-MEDIAN-AGE?XSCALE=LOG&VSCALE=LOG&COUNTRY=AFG~ALB~DZA~AGO~ARG~ARM~ATG~AUS~AUT~BHS~AZE~BHR~BGD~BRB~BLR~BEL~BLZ~LVA~LBN~LSO~LBR~LBY~LTU~LU X~MDG~MWI~MYS~MDV~MLI~MLT~MRT~MUS~MEX~BEN~BTN~BOL~BIH~BWA~BRA~BRN~BGR~BFA~BDI~KHM~CMR~CAN~CPV~CAF~TCD~CHL~CHN~COL~COM~COG~ CRI~CIV~HRV~CYP~CZE~COD~DNK~DMA~DOM~ECU~EGY~SLV~GNQ~EST~SWZ~ETH~FJI~FIN~FRA~GAB~GMB~GEO~DEU~GHA~GRC~GRD~GTM~GIN~GNB~GUY~HTI~HN D~HKG~HUN~ISL~IND~IDN~IRN~IRQ~IRL~ISR~ITA~JAM~JPN~JOR~KAZ~KEN~OWID_KOS~KWT~KGZ~LAO~MDA~MNG~MNE~MOZ~MAR~MMR~NAM~NPL~NLD~NZL~NIC ~NER~NGA~MKD~NOR~OMN~PAK~PSE~PAN~PNG~PRY~PER~PHL~POL~PRT~QAT~ROU~RUS~RWA~KNA~LCA~VCT~SMR~STP~SAU~SEN~SRB~SYC~SLE~SGP~SVK~SVN ~ZAF~KO~SD~ESP~LKA~SDN~SUR~SWE~CHE~TJK~TZA~THA~TLS~TGO~TTO~TUN~TUR~UGA~UKR~ARE~GBR~USA~URY~VZB~VUT~VNM~YEM~ZMB~ZWE addited

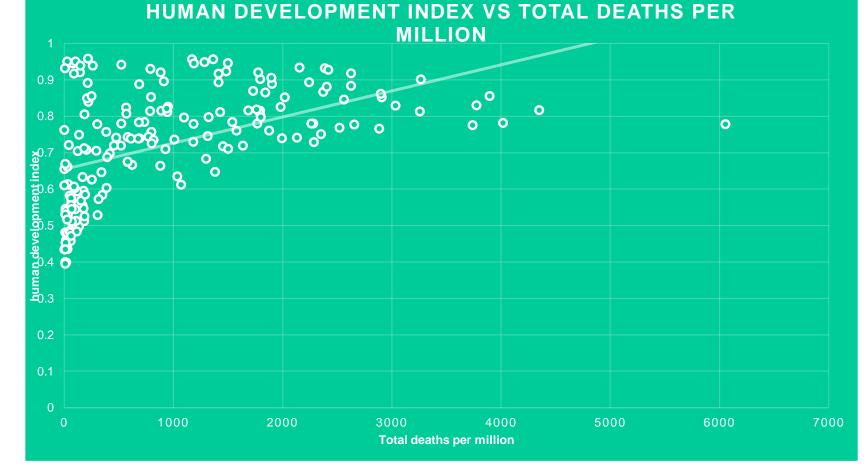
COVID-19: Deaths per Million vs. Extreme Poverty







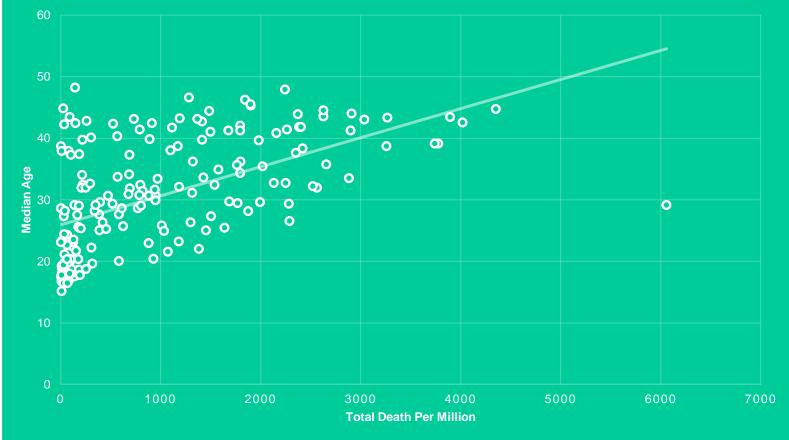
COVID-19: Deaths per Million vs. Human Development Index





COVID-19: Median Age Vs Total Deaths Per Million

MEDIAN AGE VS TOTAL DEATH PER MILLION





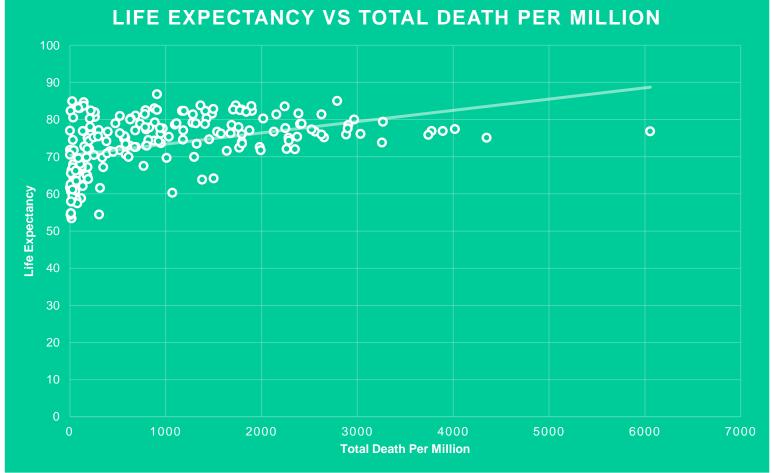
COVID-19: Percentage of 65+ Vs Total Deaths Per Million

PERCENTAGE OF 65+ VS TOTAL DEATH PER MILLION



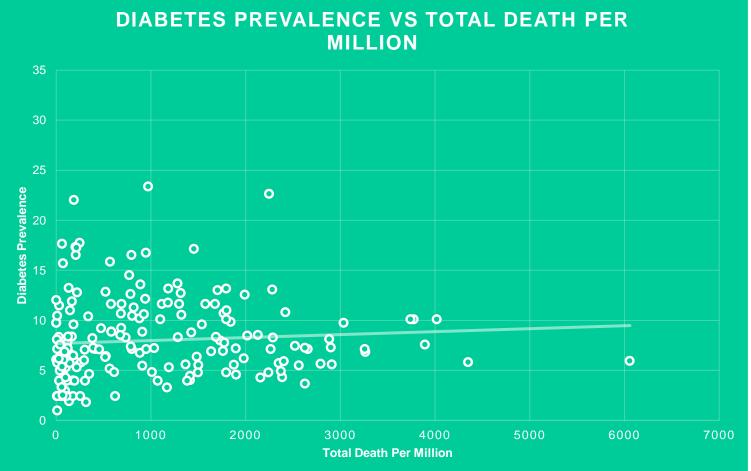


COVID-19: Life expectancy Vs Total Deaths Per Million





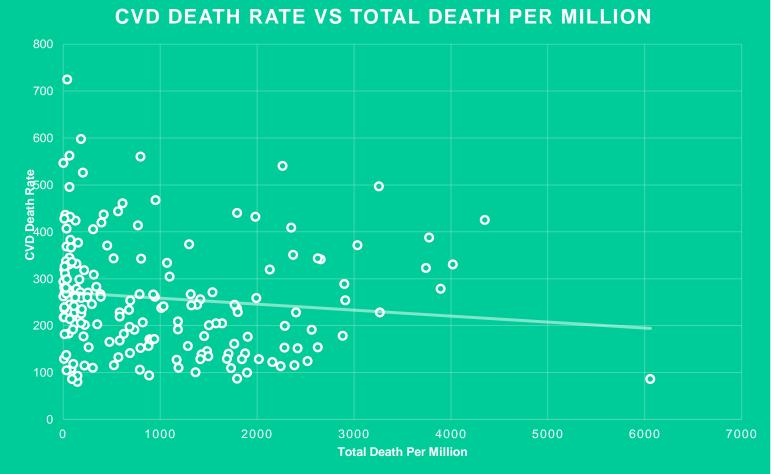
COVID-19: Diabetes Prevalence Vs Total Deaths Per Million



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



COVID-19: Cardiovascular Death Rate Vs Total Deaths Per Million

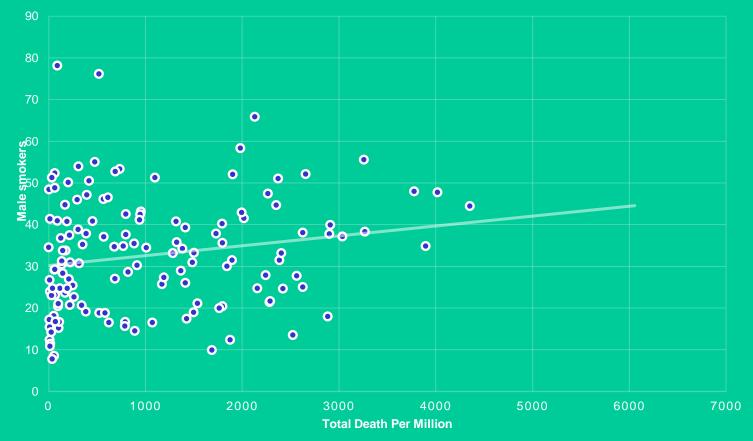


CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



COVID-19: Male Smokers Vs Total Deaths Per Million

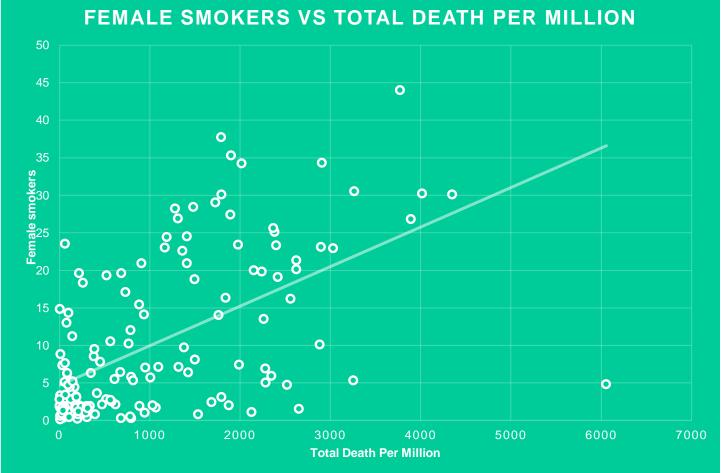
MALE SMOKERS VS TOTAL DEATH PER MILLION



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



COVID-19: Female Smokers Vs Total Deaths Per Million

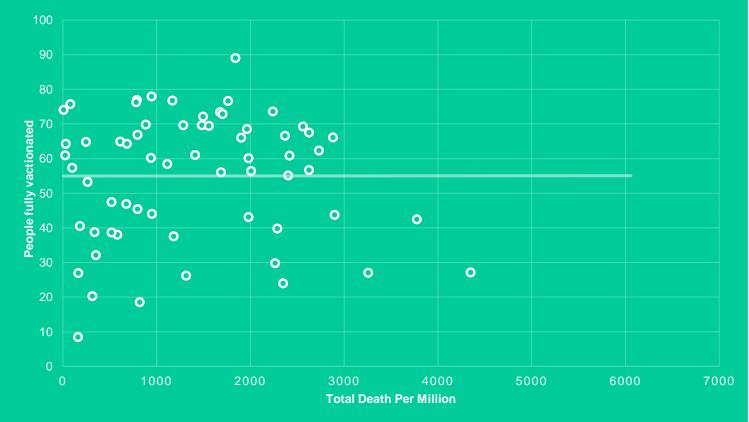


CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



COVID-19: Vaccination rate Vs Total Deaths Per Million

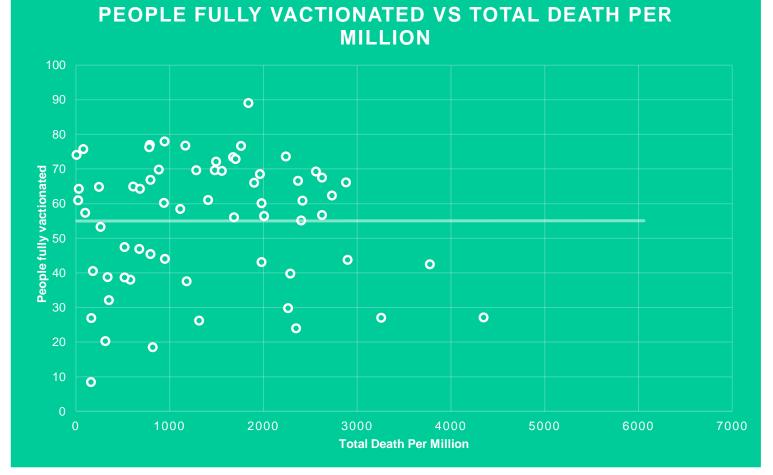
PEOPLE FULLY VACTIONATED VS TOTAL DEATH PER MILLION



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



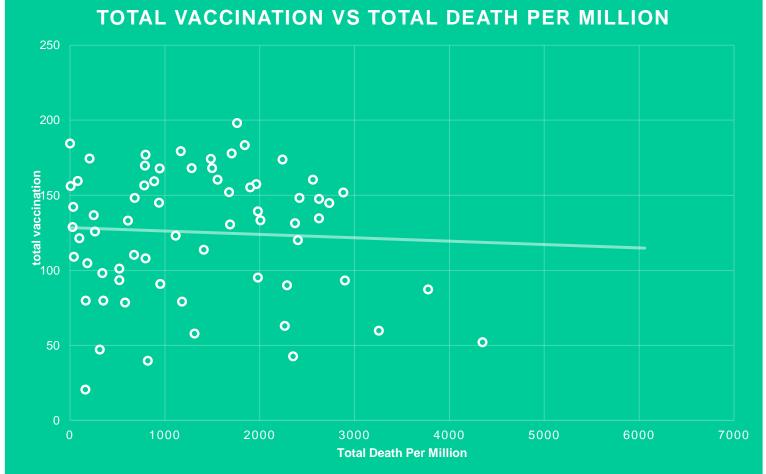
COVID-19: Fully Vaccination rate Vs Total Deaths Per Million



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



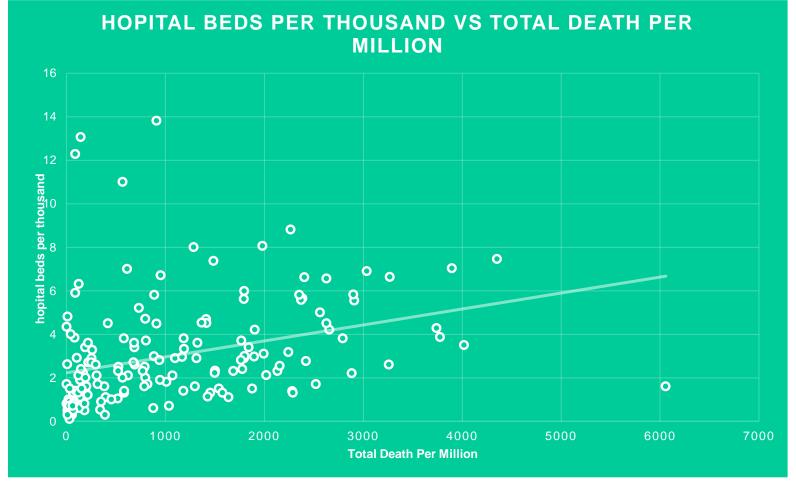
COVID-19: Total Vaccination rate Vs Total Deaths Per Million



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/



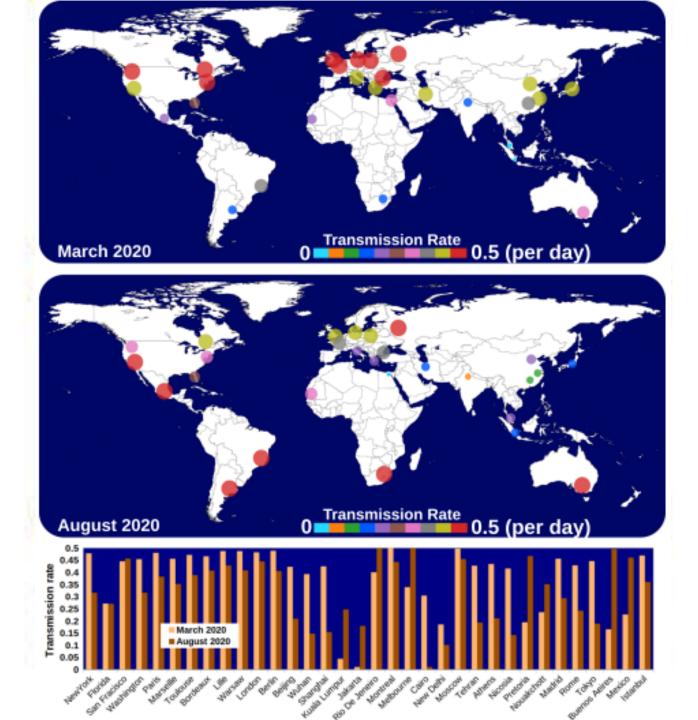
COVID-19: Hospital beds per thousand Vs Total Deaths Per Million



CASE FATALITY RATE OF COVID-19 VS. MEDIAN AGE OF THE POPULATION: SOURCE: HTTPS://OURWORLDINDATA.ORG/

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Why COVID-19 is dangerous

- 1. Low IFR (<1%) and CFR (<2%)
- 2. High incubation period (up to 14 days)
- 3. High number of asymptomatic cases
- 4. Spread by droplets (or airborne)
- 5. Global supply chain
- 6. The Aviation sector growth

Global Covid-19 Case Fatality Rates <u>HTTPS://WWW.CEBM.NET/COVID-19/GLOBAL-COVID-19-CASE-FATALITY-RATES/</u> WHO (2020):Transmission of SARS-CoV-2: implications for infection prevention precautions <u>https://www.who.int/news-room/commentaries/detail/transmission-of-sars-cov-2-implications-for-infection-precautions-precautionsta' Malta</u>

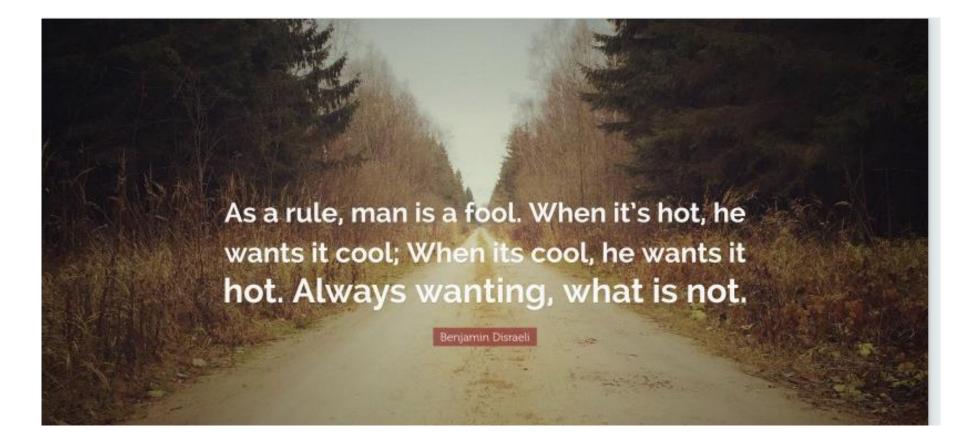
Economic impact and suicide

- 1. 5.2% contraction in global GDP in 2020.
- Might result into 10-15% increase in depression, anxiety disorder and suicide rates
- Might result in 2-5 million extra suicides in 2021
- 4. More than 10 million suicides in future due to COVID-19





The problem







Who is spreading COVID-19

- 1. Those who do not know COVID-19 exists
- 2. Those who know but do not believe
- 3. Those who unknow about protection measures
- 4. Those who know but do not

Global Covid-19 Case Fatality Rates <u>HTTPS://WWW.CEBM.NET/COVID-19/GLOBAL-COVID-19-CASE-FATALITY-RATES/</u> WHO (2020):Transmission of SARS-CoV-2: implications for infection prevention precautions <u>https://www.who.int/news-room/commentaries/detail/transmission-of-sars-cov-2-implications-for-infection-precautions-precautionsunderside the second sec</u>

Hospital bed occupancy and requirements forecasting



Intelligent Patient Management and Resource Planning for Complex, Heterogeneous, and Stochastic Healthcare Systems

Lalit Garg, Member; IEEE, Sally I. McClean, Maria Barton, Brian J. Meenan, and Ken Fullerton

Abstract-Effective resource requirement forecasting is necessary to reduce the escalating cost of care by ensuring optimum utilization and availability of scarce health resources. Patient hospital length of stay (LOS) and thus resource requirements depend on many factors including covariates representing patient characteristics such as age, gender, and diagnosis. We therefore propose the use of such covariates for better hospital capacity planning. Likewise, estimation of the patient's expected destination after discharge will help in allocating scarce community resources. Also, probable discharge destination may well affect a patient's LOS in hospital. For instance, it might be required to delay the discharge of a patient so as to make appropriate care provision in the community. A number of deterministic models such as ratio-based methods have failed to address inherent variability in complex health processes. To address such complexity, various stochastic models have therefore been proposed. However, such models fail to consider inherent heterogeneity in patient behavior. Therefore, we here use a phase-type survival tree for groups of patients that are homogeneous with respect to LOS distribution, on the basis of covariates such as time of admission, gender, and disease diagnosed; these homogeneous groups of patients can then model patient flow through a care system following stochastic pathways that are characterized by the covariates. Our phase-type model is then extended by further growing the survival tree based on covariates

provide a stochastic approach to capacity planning across complex heterogeneous care systems. The approach is illustrated using a five year retrospective data of patients admitted to the stroke unit of the Belfast City Hospital.

Index Terms—Capacity planning, cost, decision-making, forecasting, health information management, medical information systems, operations research, optimal control, prognostics and health management, stochastic systems.

I. INTRODUCTION

E FFECTIVE resource requirement forecasting is necessary to minimize the escalating cost of care by ensuring optimum utilization and availability of scarce health resources [1]. Patient hospital length of stay (LOS) and thus resource requirements depend on many factors including covariates representing patient characteristics such as age, gender, and diagnosis [2]. It is therefore necessary to consider the effect of such covariates for better capacity planning. Information about the patient demography helps in making better allocation of scarce resources. Predicting different treatment outcome,

A non-homogeneous discrete time Markov model for admission scheduling and resource planning in a cost or capacity constrained healthcare system

Lalit Garg · Sally McClean · Brian Meenan · Peter Millard

Received: 5 March 2009 / Accepted: 23 October 2009 © Springer Science+Business Media, LLC 2009

Abstract Healthcare resource planners need to develop policies that ensure optimal allocation of scarce healthcare resources. This goal can be achieved by forecasting daily resource requirements for a given admission policy. If resources are limited, admission should be scheduled according to the resource availability. Such resource availability or demand can change with time. We here model patient flow through the care system as a discrete time Markov chain. In order to have a more realistic representation, a nonhomogeneous model is developed which incorporates timedependent covariates, namely a patient's present age and the present calendar user. The model presented in this paper can using a historical dataset from the geriatric department of a London hospital.

Keywords Resource management · Admission scheduling · Non-homogeneous Markov model · Stochastic optimal control

1 Introduction

Admission scheduling [1, 2] and resource planning [3] are fundamental problems which require complex strategies to effectively manage care services ensuring entimum utilize INFORMATICA, 2011, Vol. 22, No. 1, 57–72 © 2011 Vilnius University

Phase-Type Survival Trees and Mixed Distribution Survival Trees for Clustering Patients' Hospital Length of Stay

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Received: October 2009; accepted: January 2011

Abstract. Clinical investigators, health professionals and managers are often interested in developing criteria for clustering patients into clinically meaningful groups according to their expected length of stay. In this paper, we propose two novel types of survival trees; phase-type survival trees and mixed distribution survival trees, which extend previous work on exponential survival trees. The trees are used to cluster the patients with respect to length of stay where partitioning is based on covariates such as conder, are at the time of admission and primary diagnosis code. Likelihood

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IMA Journal of Management Mathematics (2009) 20, 327–344 doi:10.1093/imaman/dpn030 Advance Access publication on November 20, 2008

Non-homogeneous Markov models for sequential pattern mining of healthcare data

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BRIAN MEENAN§

School of Engineering, University of Ulster, Jordanstown Campus, Newtownabbey, Co. Antrim, BT37 0QB, UK

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[Received May 2007; accepted May 2008]

Sequential pattern mining has been a popular data mining technique for extracting useful information from large databases and has successfully been used for numerous industrial and commercial problems. This paper presents a new mathematical modelling application to healthcare, providing important information to health service managers and policy makers to help them identify sequential patterns which require attention for efficiently managing scarce healthcare resources and developing effective healthcare management policies. In healthcare, these sequential patterns are analogous to the patient pathways. We present a non-homogeneous Markov model for identifying not only patient pathways which have high probability but also for identifying pathways which incur high cost or time. In order to have a more

Hospital bed occupancy and requirements forecasting

•**Collaborative partners:** Nanyang Technological University and Tan Tock Seng Hospital, Singapore.

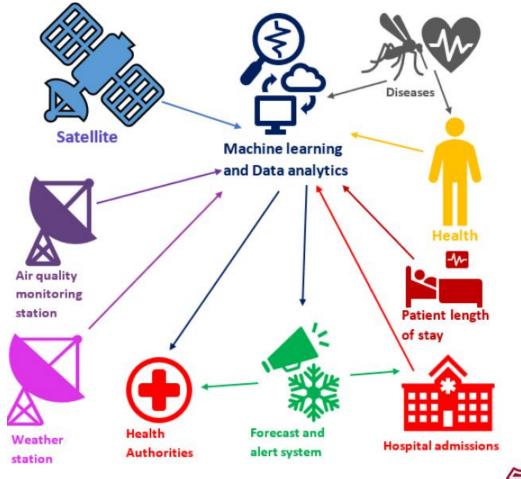
•Approach: Markov modelling, reinforcement learning

•Data: Tan Tock Seng Hospital, Singapore.





Hospital bed requirements forecasting using satellite, weather & air quality data





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Introduction

Life expectancy has increased with improvement in health services and standard of living.



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Introduction

- Life expectancy has increased with improvement in health services and standard of living.
- Higher demand to the healthcare resources



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Introduction

- Life expectancy has increased with improvement in health services and standard of living.
- Higher demand to the healthcare resources
- Healthcare challenge is to continue providing the same quality of care



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• Healthcare system facing major problems





- Healthcare system facing major problems
 - Lack of beds in hospitals





- Healthcare system facing major problems
 - Lack of beds in hospitals and
 - Lack of other hospital resources.







• To work with these problems the healthcare system needs :



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- To work with these problems the healthcare system needs :
 - An efficient way to forecast the resources required





- To work with these problems the healthcare system needs :
 - An efficient way to forecast the resources required

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• To minimize the cost of care while maintaining the quality of care.

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• When modelling the healthcare system it would help:



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- When modelling the healthcare system it would help:
 - To better understand the process for the design of polices that can improve the quality of care





• When modelling the healthcare system it would help:

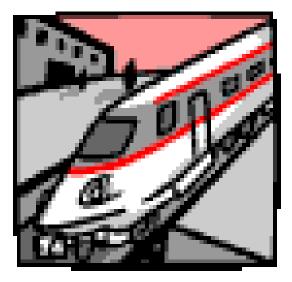
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- To better understand the process for the design of polices that can improve the quality of care
- To ensure the optimal utilization of the available resources

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• Among popular choices to fit spell length of stay data.







- Among popular choices to fit spell length of stay data.
- Provide a simple interpretation of fit for the length of stay data.



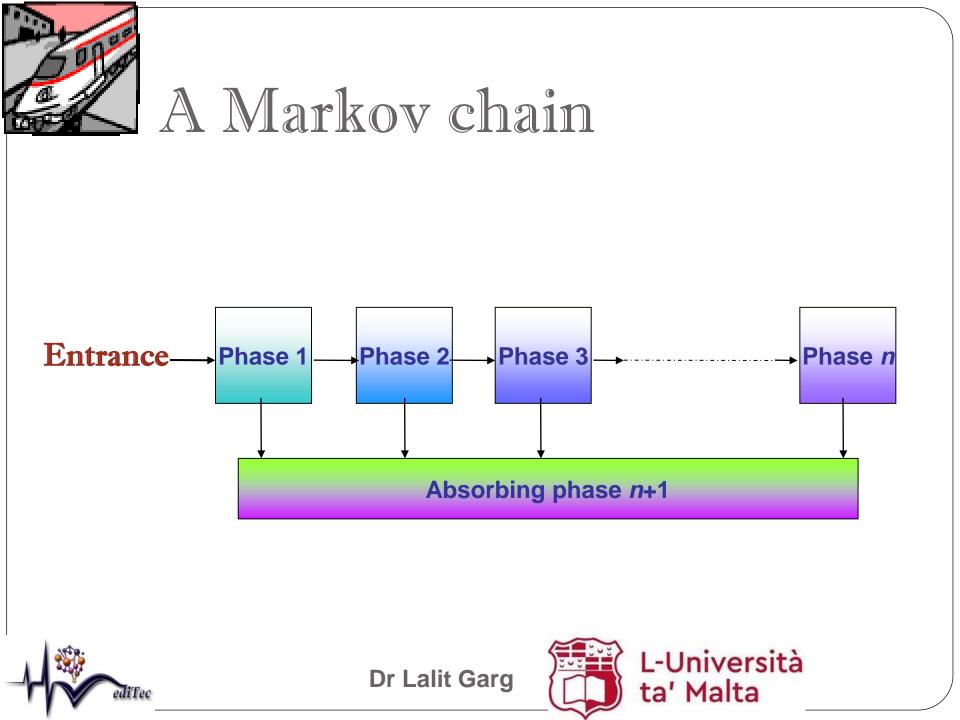




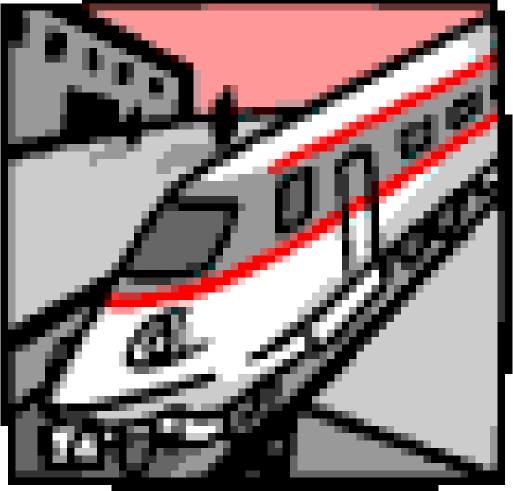
- Among popular choices to fit spell length of stay data.
- Provide a simple interpretation of fit for the length of stay data.
- Parameter estimation is easier than other phase type distributions.







A Markov chain



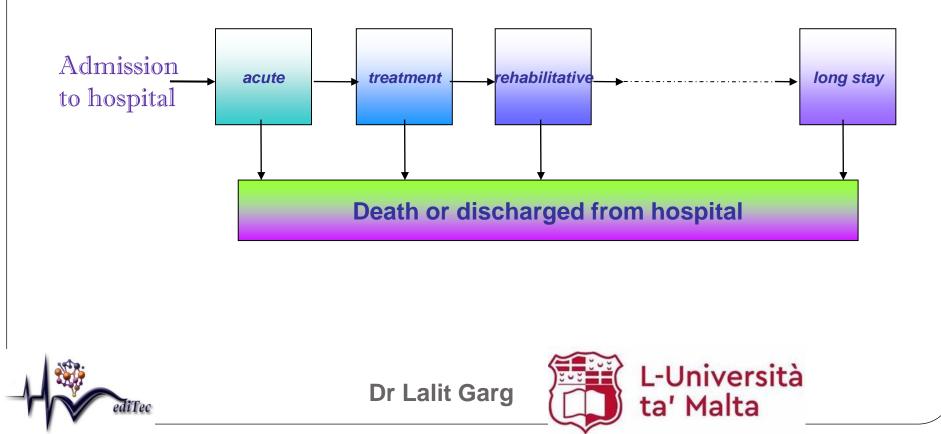


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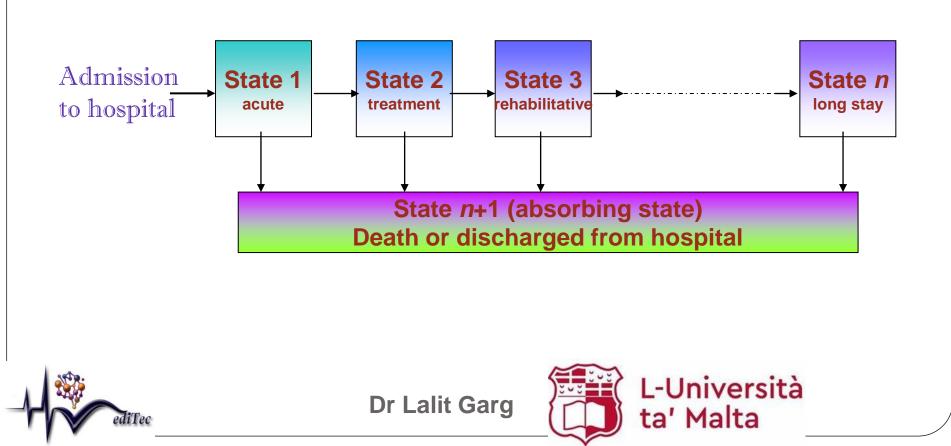


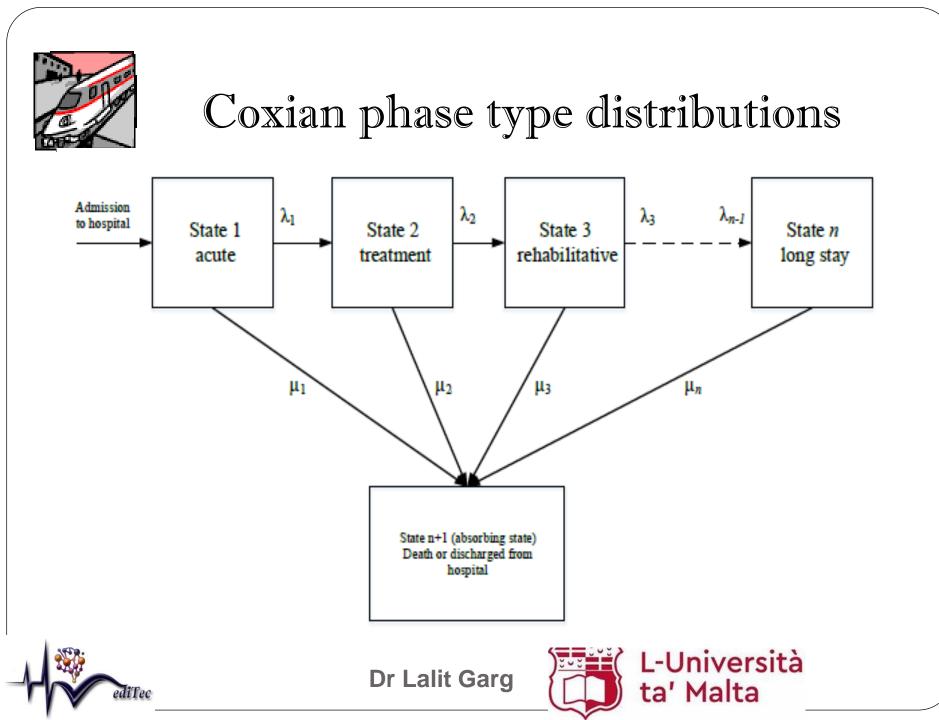
Hospital care system as a Markov chain

Patient flow in the stroke care system can be modelled as an *n* state Markov process with Coxian phase type distributions











- A process can start only in the first state (state 1).
- Sequential transition rate is λ_k .
- Also transition rate from any state k to the absorbing state n+1 is μ_k .





• The PDF for the duration before absorption:

$$f(t) = \mathbf{p} \exp(\mathbf{Q}t)\mathbf{q}$$

• where the initial state probability distribution

p=(1 0 0 ... 0 0)

• absorption probabilities

$$\mathbf{q} = \begin{pmatrix} \mu_1 & \mu_2 & \dots & \mu_{n-2} & \mu_n \end{pmatrix}^{\mathrm{T}}$$



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• And the transition matrix

$$\mathbf{Q} = \begin{pmatrix} -(\lambda_{1} + \mu_{1}) & \lambda_{1} & 0 & \cdots & 0 & 0 \\ 0 & -(\lambda_{2} + \mu_{2}) & \lambda_{2} & \cdots & 0 & 0 \\ \vdots & \vdots & \vdots & \cdots & 0 & 0 \\ 0 & 0 & 0 & 0 & -(\lambda_{n-1} + \mu_{n-1}) \lambda_{n-1} \\ 0 & 0 & 0 & \cdots & 0 & -\mu_{n} \end{pmatrix}$$

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• The likelihood function:

$$l = \prod_{i=1}^{N} \left(\mathbf{p} \exp\left\{\mathbf{Q}t_{i}\right\} \mathbf{q} \right)$$

• where *N* is the total number of patients in the care system.







• The loglikelihood function

$$L = \sum_{i=1}^{N} \left(\log \left(\mathbf{p} \exp \left\{ \mathbf{Q} t_i \right\} \mathbf{q} \right) \right).$$

• Or

$$L = \sum_{i=1}^{N} f(t_i)$$

$$f(t_i) = \log(\mathbf{p}\exp{\{\mathbf{Q}t_i\}}\mathbf{q})$$



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Weighted-Average Information Criterion

• WIC (Weighted-Average Information Criterion) is a weighted average of the Bayesian information criterion and the Akaike information criterion with a small sample size correction.





Weighted-Average Information Criterion

- WIC (Weighted-Average Information Criterion) is a weighted average of the Bayesian information criterion and the Akaike information criterion with a small sample size correction.
- The splitting criteria based on the WIC combines the strengths of both the AIC and the BIC it works well with small and large sample sizes and in the situation where sample size is not known.





Weighted-Average Information Criterion

• The performance of WIC was compared with several other popular criteria in the study and the results showed that WIC is very reliable.

$$WIC = -2L + d + \frac{d(((\log(N) - 1) \log(N))(N - (d - 1))^{2} + 2N(N + (d + 1)))}{(2N + (\log(N)(N - (d + 1)))(N - (d + 1)))}$$

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Survival tree

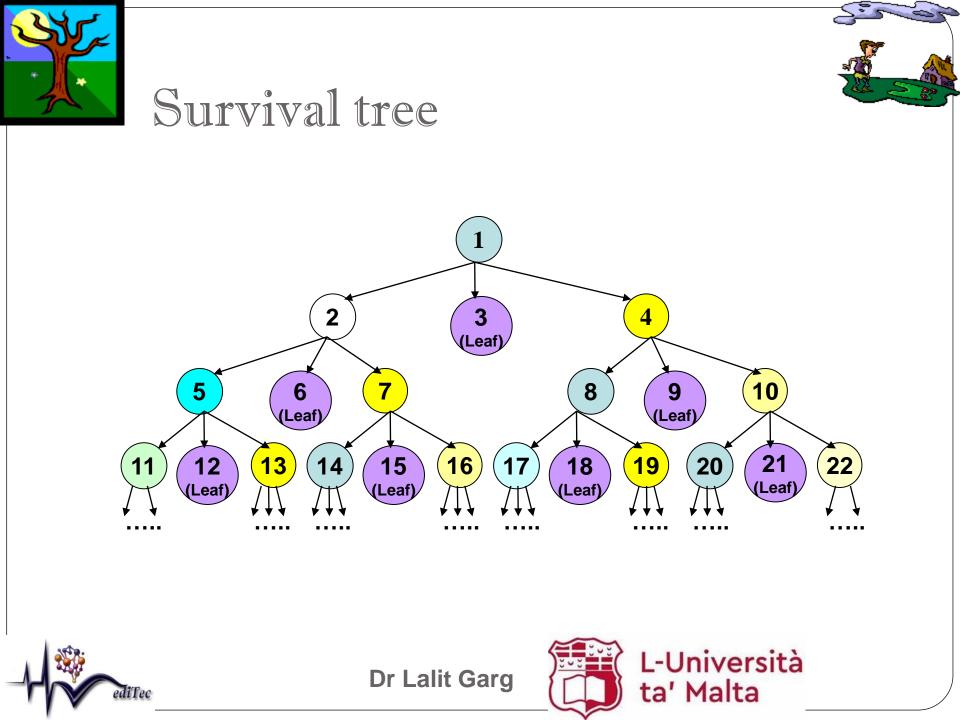






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Survival trees

• Decision trees in survival analysis





Survival trees

- Decision trees in survival analysis
- A type of classification and regression trees



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Survival trees

- Decision trees in survival analysis
- A type of classification and regression trees
- Constructed by recursively partitioning the given dataset in to subsets based on some splitting and selection criteria.





Phase type survival tree





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Phase type survival trees

• A powerful non-parametric method of clustering survival data for prognostication





Phase type survival trees

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- A powerful non-parametric method of clustering survival data for prognostication
 - To determine importance and effect of various covariates (such as patient's characteristics)

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Phase type survival trees

- A powerful non-parametric method of clustering survival data for prognostication
 - To determine importance and effect of various covariates (such as patient's characteristics)
 - Their interrelation on patient's survival, treatment outcome, disease risk, disease progress or hospital length of stay







Phase type survival tree

• Each node of *the survival tree* is separately modeled by *phase type distributions*





Phase type survival tree

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- Each node of *the survival tree* is separately modeled by *phase type distributions*
- It combines the merits of both phase type distributions and survival trees.

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Phase type survival tree

- Each node of *the survival tree* is separately modeled by *phase type distributions*
- It combines the merits of both phase type distributions and survival trees.
- Reduces the dimensionality of data and explains the variations in the data.











• Two steps

• Growing: splitting a node into child nodes











- Two steps
 - Growing: splitting a node into child nodes
 - Selection: determining if a node is terminal node. If it is not then selecting the best possible partition by exploring all possible splits.













- Growing: by recursively partitioning into sub groups by the covariates based on some splitting criteria.
 - At each node apply one covariate at a time and repeat this with other covariates.











Tree growing



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- **Splitting criteria**: maximizing either within node homogeneity or between node separation.
 - We used splitting criteria to maximize within node homogeneity based on improvement of WIC functions



Tree growing

diTec

• A covariate *a* can have any of the *l* values such that

• The loglikelihood of node *a* is
$${}^{l}_{N} = N_{a1} + N_{a2} + \ldots + N_{al} = \sum_{i=1}^{l} N_{ai}$$

• Or

$$L = \sum_{j=1}^{l} \sum_{i=1}^{N_{aj}} f(t_{iaj}) = \sum_{i=1}^{N_{a1}} f(t_{ia1}) + \sum_{i=1}^{N_{a2}} f(t_{ia2}) + \dots + \sum_{i=1}^{N_{al}} f(t_{ial})$$

$$L = L_{a1} + L_{a2} + \dots + L_{al} = \sum_{i=1}^{l} L_{ai}$$

$$L = U_{a1} + U_{a2} + \dots + U_{al} = \sum_{i=1}^{l} L_{ai}$$

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Tree growing

• Similarly, WIC of node *a* is

$$WIC = WIC_{a1} + WIC_{a2} + \ldots + WIC_{al} = \sum_{i=1}^{l} WIC_{ai}$$
.







Node selection



- For each possible split of a node, record the total WIC after the split.
- The split which maximizes the total WIC of sub-groups is determined as follows:

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$$WIC_{\max} = \max(WIC_a, WIC_b, \dots, WIC_l)$$

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Node selection

• If WIC_{max} is greater than WIC of the node before the split, select the split with WIC equal to WIC_{max} else record the node as a terminal node.



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Node selection

- If WIC_{max} is greater than WIC of the node before the split, select the split with WIC equal to WIC_{max} else record the node as a terminal node.
- **Terminal node:** A terminal node is the node at which within node homogeneity cannot significantly be improved by any possible split.

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• To evaluate the model we used the discharge dataset from the Emergency department at the Mater Dei Hospital Malta of all patients discharged in 2011-2012.





- We used covariates that represent the patient characteristics:
 - Age
 - Gender
 - District
 - Source of Admissions





- For the length of stay :
 - The continuous covariate was the patient's age
 - Three categorical covariates Gender, District and Source of Admission.





- Categorical covariate data was divide in three groups.
- The cut points of the age are:
 - 1 to 40,
 - 41 to 70 and
 - 71 and over.
 - Patients with 0 age at admission were omitted from the data.







• The gender covariate has two different values that are Female and Male.





- The gender covariate has two different values that are Female and Male.
- The district covariate has six different values that are the geographical districts of Malta.



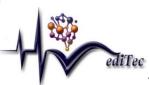


- The gender covariate has two different values that are Female and Male.
- The district covariate has six different values that are the geographical districts of Malta.
- Source of admission is from where the patient was admitted and has five different covariates.





- The gender covariate has two different values that are Female and Male.
- The district covariate has six different values that are the geographical districts of Malta.
- Source of admission is from where the patient was admitted and has five different covariates.
- Each cluster was given a group number for running the Coxian Phase fittings.



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- For the admissions:
 - The categorical covariate was the district of the patient and







- For the admissions:
 - The categorical covariate was the district of the patient and
 - The categorical covariates are the age and the gender.





- For the admissions:
 - The categorical covariate was the district of the patient and
 - The categorical covariates are the age and the gender.
- Each value in the covariate is given a group number to run the Coxian phase fittings for each group.



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• Daily and Monthly Admission Values:

	Sun.	Mon.	Tues.	Wed.	Thurs.	Fri.	Sat.	Total
Jan.	1135	2423	1897	1837	1630	1650	1250	11822
Feb.	989	1942	1663	2013	1585	1518	1202	10912
Mar.	917	1855	1799	2010	1941	1851	1258	11631
Apr.	999	2179	1634	1783	1580	1555	1305	11035
May.	999	2064	1941	1994	1780	1621	1128	11527
Jun.	867	1835	1528	1888	1629	1731	1252	10730
Jul.	1113	2174	1873	1745	1528	1793	1222	11448
Aug.	849	2042	1779	2097	1802	1815	1112	11496
Sept.	934	1874	1623	1668	1756	1666	1189	10710
Oct.	973	2402	1947	2026	1683	1783	1394	12208
Nov.	946	1942	1971	2091	1891	1865	1278	11984
Dec.	894	1671	1404	1572	1469	1729	1262	10001
Tot.	11651	24403	21059	22724	20274	20577	14852	135504





Node	Covariate	Covariate Value	Total Number of Patients	WIC	Mean LOS	Number of phases	Total WIC	Gain in WIC	
	Level 1								
	All	Root Node	64439	351604.66	6.8411	б	351604.66	-	
		1 to 40	20631	87222.35	4.1304	6			
	Age	41 to 70	22600	122877.8	6.7443	5	341295.6	10309.1	
		71 +	21208	131195.4	9.5813	5	 351604.66 341295.6 351775.15 351078.46 351637.51 		
		South	22237	121077.72	6.756	5			
		Central	19480	107177.13	6.9864	4		-170.49	
	District	West	8423	46460.1	7.0515	5	351775.15		
		North	13542	72716.7	6.6032	4			
1		Gozo	539	3227.25	8.3358	5	351604.66 341295.6 351775.15 351078.46 351637.51		
Root		Unknown	218	1116.25	5.5	4			
Node		Elderly Home	1925	11775.05	9.4732	6			
		Home	61356	332501.72	6.7339	6			
		Labour Ward	2	32.84	4.5	of phasesTotal WICW16 351604.66 463535554455248544269626351078.46524226351637.51-33			
	Source	Other (Gov Hospital,					351078.46	526.2	
		Private, Mental and	1060	6297.08	8.4632	6			
		Abroad)							
		Police Custody	96	471.77	4.7604	2			
	Gender	Female	32886	179393.48	6.8672		351637 51	-32.85	
		Male	31553	172244.02	6.814			-52.05	
1	eurrec			L					

Node	Covariate	Covariate Value	Total Number of Patients	WIC	Mean LOS	Number of phases	Total WIC	Gain in WIC	
	Level 3								
	All	Age 41 to 70, Female	9088	49410.24	6.817	4	49410.24	-	
		41 to 70, South, F	3164	17051	6.8587	6			
		41 to 70, Central, F	2782	15094.21	6.8724	5			
	District	41 to 70, West, F	1123	6118.53	6.9154	5	10140 24	261.0	
	District	41 to 70, North, F	1933	10357.31	6.5525	3	49148.34	261.9	
		41 to 70, Gozo, F	55	366.03	9.9454	1			
10 (Age		41 to 70, Unknown, F	31	161.25	4.9678	3			
41 to 70,		41 to 70, Elderly Home, F	81	561.03	12.4445	3			
Female)		41 to 70, Home, F	8835	47791.27	6.7268	4			
	Source of	41 to 70, Labour Ward, F	1	3.89	7	1	10206 16	12 70	
	Admission	41 to 70, Other (Gov Hospital, Private, Mental and Abroad), F	170	1038.88	8.8529	4	49396.46	13.78	
		41 to 70, Police Custody, F	1	1.39	2	1			

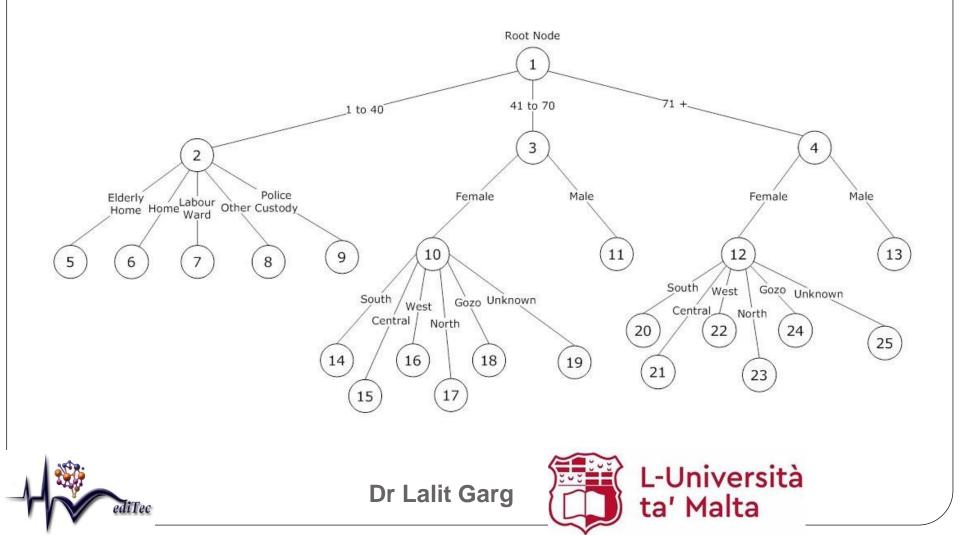




Node	Covariate	Covariate Value	Total Number of Patients	WIC	Mean LOS	Number of phases	Total WIC	Gain in WIC
			Level 3					
	All	Age 71 +, Female	11578	72543.24	9.9719	5	72543.24	-
		71 +, South, F	3663	22859.81	9.8444	6		
	District	71 +, Central, F	3880	24104.55	9.8023	6		
		71 +, West, F	1736	11040.79	10.4919	4	72219.66	323.58
12 ()		71 +, North, F	2242	13837.2	10.0589	6	12219.00	343.30
12 (Age		71 +, Gozo, F	40	287.23	12.825	1		
71 +,		71 +, Unknown, F	17	90.08	4.8235	1		
Female)		71 +, Elderly Home, F	1257	7655.41	9.4121	4		
	Source of	71 +, Home, F	10093	63415.04	10.0396	6		
	Admission	71 +, Other (Gov Hospital, Private, Mental and Abroad), F	228	1462.44	10.057	4	72532.89	10.35







Admissions Phase-Type Survival Tree Construction

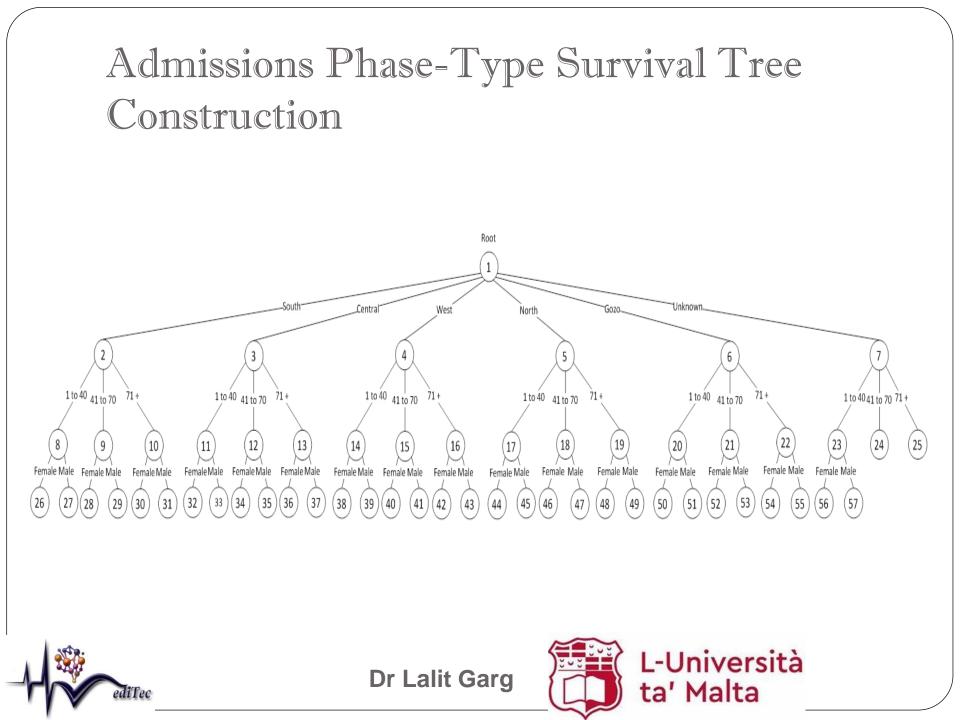
Node	Covariate	Covariate Value	Total Admissio ns	WIC	Mean	Number of Phases	Average WIC	Total WIC	Gain in WIC		
	Level 1										
	All	Root Node	32277	3171.43	89.43	22	3171.43	3171.43	-		
		1 to 40	10386	2561.57	29.45	10	853.86				
	Age	41 to 70	11244	2590.39	31.81	10	863.46	2576.47	594.96		
		71 +	10647	2577.45	30.17	10	859.15				
	Gender	Female	16510	2793.52	44.2	10	1396.76	2811.39	360.04		
1 (Root		Male	15767	2829.26	46.23	10	1414.63				
Node)		South	11211	2581.18	31.72	10	430.2				
		Central	9690	2491.79	27.55	10	415.3				
		West	4270	2051.09	12.7	10	341.85	1756 20	1415 04		
	District	North	6774	2289.19	19.56	10	381.53	1756.39	1415.04		
		Gozo	289	895.58	1.79	6	149.26				
		Unknown	43	229.51	1.12	10	38.25				
+	Dr Lalit Garg ta' Malta										

Admissions Phase-Type Survival Tree Construction

Node	Covariat e	Covariat e Value	Total Admissio ns	WIC	Mean	Number of Phases	Average WIC	Total WIC	Gain in WIC	
	Level 3									
8 (South,	Gender	Female	2263	1817.71	7.2	5	50.49	94.97	17.71	
1 to 40)	Genuer	Male	1518	1601.38	5.16	5	44.48	94.97	1/./1	
9 (South,	Gender	Female	1602	1617.75	5.39	5	44.94	94.31	18.11	
41 to 70)	Genuer	Male	2413	1777.52	7.61	7	49.38	94.31	10.11	
10 (South,	Gender	Female	1804	1680.7	5.94	5	46.69	91.78	17.34	
71 +)	Genuer	Male	1611	1623.45	5.41	5	45.1	91.70		
11		Female	1761	1719.87	5.82	5	47.77			
(Central, 1 to 40)	Gender	Male	1191	1496.32	4.26	5	41.56	89.34	16.18	
12		Female	1325	1565.73	4.63	5	43.49			
(Central, 41 to 70)	Gender	Male	1942	1716.2	6.32	6	47.67	91.16	17.21	
13		Female	1934	1725.28	6.3	5	47.92			
(Central, 71 +)	Gender	Male	1537	1599.83	5.21	5	44.44	92.36	18.68	
14 (West,	Gender	Female	820	1357.36	3.25	4	37.7	60.44	10 /0	
1 to 40)		Male	506	1142.3	2.39	4	31.73	69.44	18.49	
15 (West,	Condor	Female	565	1200.36	2.55	4	33.34	70 41	16 41	
41 to 70)	Gender	Male	840	1334.26	3.3	4	37.06	70.41	16.41	
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Admissions Phase-Type Survival Tree Construction

Node	Covariat e	Covariat e Value	Total Admissio ns	WIC	Mean	Number of Phases	Average WIC	Total WIC	Gain in WIC		
	Level 3										
16 (West,	Gender	Female	908	1387.43	3.49	4	38.54	71.95	18.17		
71 +)	Genuer	Male	631	1202.62	2.73	4	33.41	/1.75	10.17		
17 (North,	Gender	Female	1304	1563.15	4.57	4	43.42	81.14	15.86		
1 to 40)	Genuer	Male	882	1357.83	3.42	4	37.72	01.14	13.00		
18 (North,	Gender	Female	959	1411.44	3.63	4	39.21	84.06	17.54		
41 to 70)	Genuer	Male	1469	1614.66	5.02	5	44.85	84.00	17.54		
19 (North,	Gender	Female	1125	1488.1	4.08	4	41.34	81.41	17.05		
71 +)		Male	1035	1442.69	3.84	4	40.07				
20 (Gozo,	Condor	Female	64	323.82	1.18	10	8.99	16.17	12.16		
1 to 40)	Gender	Male	50	258.44	1.14	10	7.18				
21 (Gozo,	Condor	Female	64	323.82	1.18	10	8.99	20.15	9.06		
41 to 70)	Gender	Male	82	401.76	1.23	10	11.16	20.15			
22 (Gozo,	Condon	Female	24	100.2	1.07	10	2.78	7 77	0.26		
71 +)	Gender	Male	35	161.34	1.1	10	4.48	7.27	9.26		
23		Female	13	22.86	1.04	10	0.64				
(Unknow n, 1 to 40)	Gender	Male	14	45.21	1.04	10	1.26	1.89	5.29		
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Phase-Type Survival Tree Construction

• The Length of Stay phase-type survival tree has 19 leaf nodes and has a total Gain in WIC of 12619.16.





Phase-Type Survival Tree Construction

- The Length of Stay phase-type survival tree has 19 leaf nodes and has a total Gain in WIC of 12619.16.
- The Admissions phase-type survival tree has 34 leaf nodes and a total Gain in WIC of 2111.41.





Prognostication

- Both phase-type survival trees are showing
 - Analysis of the determined patient groups from our dataset.







Prognostication

• Predictions can be made from the data used to construct the Phase-type survival tree

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• For the number of admissions by the patient grouping and

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Prognostication

- Predictions can be made from the data used to construct the Phase-type survival tree
 - For the number of admissions by the patient grouping and
 - We can predict the LOS of a patient by his/her characteristics.





LOS-Prediction

Gender	Age	District	Source	Admission Date	Discharge Date	Actual LOS	Predicted LOS
М	1	South	Home	15/12/2012	19/12/2012	5	4.122102
М	67	Central	Home	21/12/2012	31/12/2012	11	6.744455
F	86	South	Home	18/12/2012	24/12/2012	7	9.960199
F	24	West	Home	22/12/2012	24/12/2012	3	4.122102
М	64	South	Home	15/12/2012	18/12/2012	4	6.744455
М	77	West	Elderly Home	26/12/2012	31/12/2012	6	9.189538
М	16	North	Home	20/12/2012	20/12/2012	1	4.122102
F	94	South	Home	18/12/2012	20/12/2012	3	9.960199
М	57	Central	Home	15/12/2012	19/12/2012	5	6.744455
F	49	Central	Home	20/12/2012	21/12/2012	2	6.916771





Admission Predictions

Admissions Date	Group	Actual Admissions	Predicted Admissions
31/12/2011	41 to 70 Unknown	0	0.04
28/12/2011	1 to 40, South, Male	3	4.15
24/12/2011	1 to 40, Central Males	2	3.28
28/12/2011	1 to 40, West, Males	1	1.39
26/12/2011	1 to 40, North, Males	1	2.45
27/12/2011	1 to 40, Gozo, Males	0	0.14
27/12/2011	1 to 40, Unknown, Males	0	0.04
19/12/2011	1 to 40, South, Females	7	6.30
29/12/2011	1 to 40, Central, Females	3	4.87
30/12/2011	1 to 40, West, Females	2	2.28
28/12/2011	1 to 40, North, Females	5	3.59
24/12/2011	1 to 40, Gozo, Females	0	0.18
24/12/2011	1 to 40, Unknown, Females	0	0.03
28/12/2011	41 to 70, South, Males	12	6.54
19/12/2011	41 to 70, Central, Males	7	5.30
26/12/2011	41 to 70, West, Males	5	2.27
15/12/2011	41 to 70, North, Males	6	2.61
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Admission Predictions

Admissions Date	Group	Actual Admissions	Predicted Admissions
29/12/2011	41 to 70, Gozo, Males	0	0.22
23/12/2011	41 to 70, South, Females	7	4.39
29/12/2011	41 to 70, Central, Females	3	3.63
20/12/2011	41 to 70, West, Females	0	1.59
25/12/2011	41 to 70, North, Females	4	4.02
28/12/2011	41 to 70, Gozo, Females	0	0.18
24/12/2011	71 +, South, Males	8	4.41
30/12/2011	71 +, Central, Males	4	4.16
31/12/2011	71 +, West, Males	1	1.71
17/12/2011	71 +, North, Males	3	2.80
26/12/2011	71 +, Gozo, Males	1	0.10
17/12/2011	71 +, South, Females	4	4.87
16/12/2011	71 +, Central, Females	6	5.15
30/12/2011	71 +, West, Females	3	2.47
16/12/2011	71 +, North, Females	3	3.06
31/12/2011	71 +, Gozo, Females	0	0.07





Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	Root	66166	6.88	361646.80	361646.80	
	MinTemp	0°C-10°C (1)	16465	7.19	91916.01		
		$11^{\circ}C-20^{\circ}C$ (2)	33516	6.76	181607.62	361631.50	15.30
		$21^{\circ}C-30^{\circ}C$ (3)	16185	6.83	88107.87		
	MaxTemp	$0^{\circ}\text{C-}10^{\circ}\text{C}(1)$	303	8.13	1786.56		
		$11^{\circ}C-20^{\circ}C$ (2)	28333	6.95	143924.01	349779.14	11867.67
		$21^{\circ}C-30^{\circ}C$ (3)	25205	6.83	137012.30	545775.14	11007.07
ot)		$31 + ^{\circ}C(4)$	12325	6.82	67056.27		
(Root)	AvgTemp	$0^{\circ}C-10^{\circ}C(1)$	4834	7.23	26828.01		
1 (11°C-20°C (2)	34493	6.87	188586.75	361381.17	265.63
		$21^{\circ}C-30^{\circ}C$ (3)	26090	6.83	141956.96	301301.17	205.05
		$31 + ^{\circ}C(4)$	749	6.88	4009.44		
	MaxVar	$x < -2^{\circ}C(1)$	4032	7.02	22086.49		
		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	18199	6.78	99118.57		
		$0^{\circ}C(3)$	19042	6.79	103741.30	361419.43	227.37
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	21365	7.02	117284.96		
		$x > 2^{\circ}C(5)$	3528	6.88	19188.12		
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Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	$0^{\circ} C$ -1 $0^{\circ} C$ (1)	303	8.13	1786.56	1786.56	
	MinTemp	0°C-10°C (1)	303	8.13	1786.56		
		11°C-20 (2)	0	0.00	0.00	1786.56	0.00
		21°C-30°C (3)	0	0.00	0.00		
Max)	AvgTemp	0°C-10°C (1)	303	8.13	1786.56		
		11°C-20°C (2)	0	0.00	0.00	1790 50	0.00
10°		21°C-30°C (3)	0	0.00	0.00	1786.56	
(0°C-10°C		31+°C (4)	0	0.00	0.00		
0)	MaxVar	x<-2°C (1)	104	9.50	619.01		
7		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	97	7.59	584.81		
		$0^{\circ}C(3)$	102	7.25	605.99	1809.80	-23.24
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	0	0.00	0.00		
		$x > 2^{\circ}C(5)$	0	0.00	0.00		





Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	$11^{\circ}C-20^{\circ}C(2)$	28333	6.83	143924.01	143924.01	
	MinTemp	0°C-10°C (1)	15983	7.19	88145.39		
		11°C-20°C (2)	12350	6.63	66639.23	154784.63	-10860.62
		$21^{\circ}C-30^{\circ}C$ (3)	0	0.00	0.00		
	AvgTemp	$0^{\circ}C^{\circ}C-10^{\circ}C(1)$	4531	7.17	25082.63		
$\widehat{\mathbf{v}}$		11°C-20°C (2)	23802	6.90	130527.73	155010.90	-11686.34
Mao		21°C-30°C (3)	0	0.00	0.00	155610.36	
(11°C-20°C Max)		31+°C (4)	0	0.00	0.00		
20°	MaxVar	x<-2°C (1)	1818	6.98	10045.36		
°,		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	8495	6.78	45964.04		
.11		$0^{\circ}C(3)$	8287	6.72	44646.01	154715.89	-10791.88
5			·	I	,	,	
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	8551	7.23	47346.72		
		$x > 2^{\circ}C(5)$	1182	7.59	6713.75		
	ł	ł	ł	i	ł	ŀ	





Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	21° C-30° C (3)	25205	6.83	137012.30	137012.30	
	MinTemp	$0^{\circ}C-10^{\circ}C(1)$	179	6.13	967.79		
		$11^{\circ}C-20^{\circ}C$ (2)	20347	6.83	110539.41	136794.76	217.54
		$21^{\circ}C-30^{\circ}C$ (3)	4679	6.84	25287.56		
ax)	AvgTemp	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00		
Μ		$11^{\circ}C-20^{\circ}C$ (2)	10691	6.81	57269.63	190905 41	746.88
0.0		$21^{\circ}C-30^{\circ}C$ (3)	14514	6.84	78995.78	136265.41	
2(21°C-30°C Max)		$31+^{\circ}C(4)$	0	0.00	0.00		
1°C	MaxVar	x<-2°C (1)	1203	6.92	6576.60		
2(2)		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	6861	6.77	36677.42		
		$0^{\circ}C(3)$	7826	6.89	42694.92	136579.50	432.80
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	8472	6.88	46191.76		
		$x > 2^{\circ}C(5)$	843	6.04	4438.80		

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Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	$11^{\circ}C-20^{\circ}C(2)$	10691	6.81	57269.63	57269.63	
. 5	MinTemp	$0^{\circ}C-10^{\circ}C(1)$	179	6.13	961.29		
Max, vg)		$11^{\circ}C-20^{\circ}C$ (2)	10512	6.82	57122.54	58083.83	-814.20
C Ma Avg)		$21^{\circ}C-30^{\circ}C$ (3)	0	0.00	0.00		
3(21°C-30°C 11°C-20°C A	MaxVar	x<-2°C (1)	397	5.66	2036.82		
-50 C-0		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	2405	6.95	13061.90		
3(21°C-30° 11°C-20°C		$0^{\circ}C(3)$	2666	6.88	14272.56	57493.12	-223.49
3($1^{\circ}C \le x \le 2^{\circ}C$ (4)	4736	6.87	25550.60		
		$x > 2^{\circ}C(5)$	487	6.00	2571.25		
	All	$21^{\circ}C-30^{\circ}C(3)$	14514	6.84	78995.78	78995.78	
	MinTemp	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00		
dax g)		$11^{\circ}C-20^{\circ}C$ (2)	9835	6.85	52787.28	78354.56	641.22
3(21° C-30° C Max, 21° C-30° C Avg)		$21^\circ ext{C-}30^\circ ext{C}(3)$	4679	6.84	25567.28		
3(21°C-30° 21°C-30°C	MaxVar	$x < -2^{\circ}C(1)$	806	7.54	4555.23		
90 (5		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	4456	6.68	23671.22		
21°		$0^{\circ}C(3)$	5160	6.90	28290.46	78571.35	424.43
$\frac{3}{21}$		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	3736	6.88	20163.21		
		$x > 2^{\circ}C(5)$	356	6.10	1891.23		
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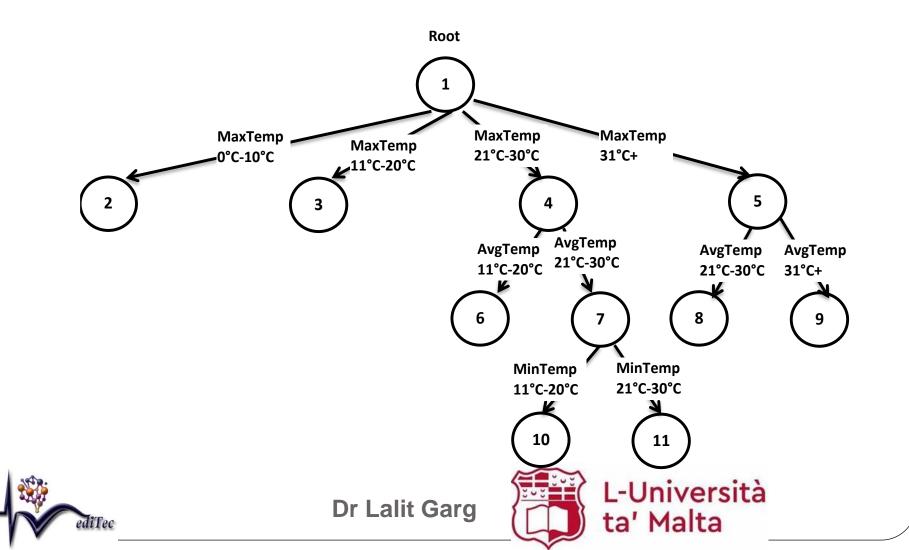
Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	31+°C (4)	12325	6.82	67056.27	67056.27	
	MinTemp	0°C-10°C (1)	0	0.00	0.00		
		$11^{\circ}C-20^{\circ}C$ (2)	820	6.70	4466.70	67053.54	2.73
		21°C-30°C (3)	11505	6.83	62586.84		
(xi	AvgTemp	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00		
Max)		$11^{\circ}C-20^{\circ}C$ (2)	0	0.00	0.00	66238.27	818.00
°C		$21^{\circ}C-30^{\circ}C$ (3)	11576	6.82	62203.93	00230.27	818.00
2(31+°C		$31 + ^{\circ}C$ (4)	749	6.88	4034.35		
2(3	MaxVar	x<-2°C (1)	907	6.96	4967.36		
		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	2746	6.79	14698.72		
		$0^{\circ}C(3)$	2827	6.70	15140.27	66443.03	613.24
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	4342	6.90	23535.51		
		$x > 2^{\circ}C(5)$	1503	6.80	8101.17		

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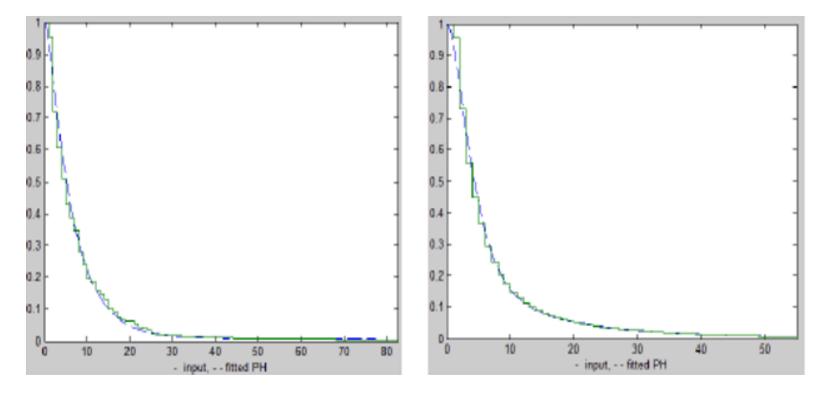


Level	Covariate	Covariate	No. of	Mean	WIC	Total	WIC
		Group	Patients	LOS		WIC	Gain
	All	$21^{\circ}C-30^{\circ}C(3)$	11576	6.82	62203.93	62203.93	
g)	MinTemp	0°C-10°C (1)	0	0.00	0.00		
		$11^{\circ}C-20^{\circ}C$ (2)	820	6.70	4466.70	63023.76	-819.83
Max, C Avg)		$21^{\circ}C-30^{\circ}C$ (3)	10756	6.83	58557.06		
C N	MaxVar	x<-2°C (1)	820	7.04	4489.42		
-30.		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	2546	6.80	13659.83		
3(31+°C M 21°C-30°C .						62465.66	-261.73
3(0°C (3)	2827	6.70	15140.27		
		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	4243	6.90	23001.58		
		$x > 2^{\circ}C(5)$	1140	6.68	6174.55		
		$31 + ^{\circ}C(4)$	749	6.88	4034.35	4034.35	
	MinTemp	0°C-10°C (1)	0	0.00	0.00		
		$11^{\circ}C-20^{\circ}C$ (2)	0	0.00	0.00	4061.80	-27.45
Max, vg)		$21^{\circ}C-30^{\circ}C$ (3)	749	6.88	4061.80		
C Ma Avg)	MaxVar	x<-2°C (1)	87	6.20	485.00		
C / C		$-2^{\circ}C \le x \le -1^{\circ}C$ (2)	200	6.62	1074.44		
3(31+°C 31+°C A		$0^{\circ}C(3)$	0	0.00	0.00	4082.57	-48.23
3(31		$1^{\circ}C \le x \le 2^{\circ}C$ (4)	99	6.93	554.99		
		$x > 2^{\circ}C(5)$	363	7.17	1968.14		
				Y			



• Node 2

Node 3

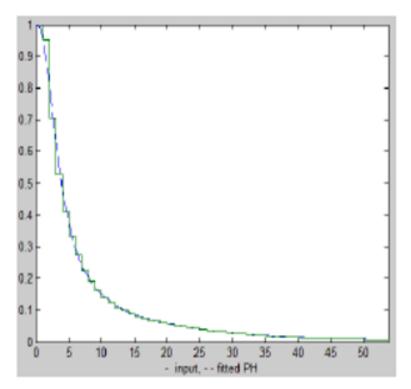






• Node 8

Node 9

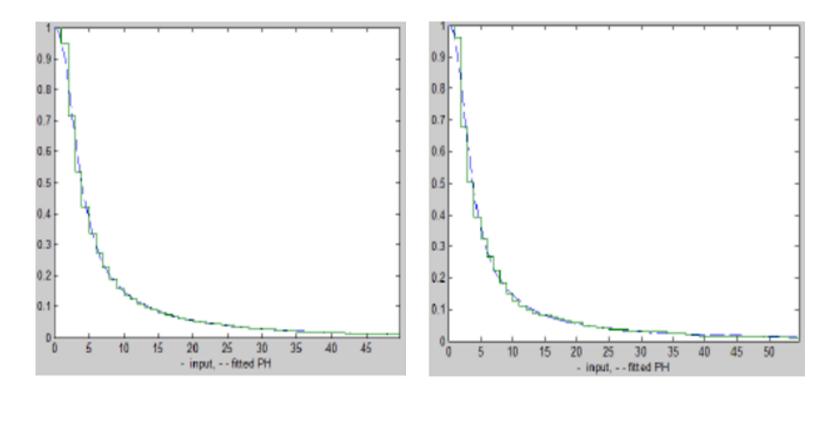






• Node 8

Node 9

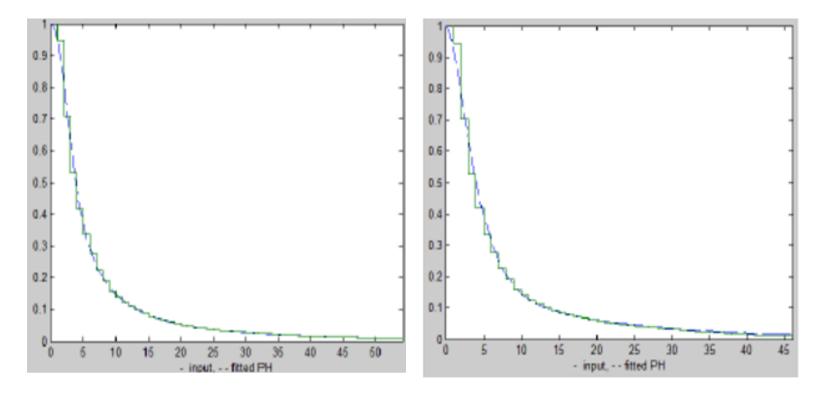






• Node 10

Node 11







• Most significant prognostic factor affecting the patients' length of stay (LOS) is the maximum temperature.



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- Most significant prognostic factor affecting the patients' length of stay (LOS) is the maximum temperature.
- The average temperature affects the patients' length of stay only when the maximum temperature rises beyond 20°C.







• The minimum temperature does not significantly affect the patients' length of stay.





- The minimum temperature does not significantly affect the patients' length of stay.
- Also, the maximum variability in the average temperature between one day and the next does not affect patients' length of stay as patients usually stay inside.

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• These results might be different for different geographic regions due to different weather conditions and different genetic profile of inhabitants there.





• Predictions and Accuracy Tests

Group	No. of				Squared	Absolute	Percentage
	Patients	Mean LOS	Mean LOS	Error	Error	Error	Error (%)
$MaxTemp(0^{\circ}C-10^{\circ}C)$	0	-	8.13	-	-	-	-
$MaxTemp(11^{\circ}C-20^{\circ}C)$	13406	7.19	6.83	-0.36	0.13	0.36	5.01
$\begin{array}{l} \text{MaxTemp}(21^{\circ}\text{C-}30^{\circ}\text{C}), \\ \text{AvgTemp}(11^{\circ}\text{C-}20^{\circ}\text{C}) \end{array}$	6003	7.01	6.81	-0.20	0.04	0.20	2.85
MaxTemp(21°C-°C30),AvgTemp(21°C-30°C), MinTemp(11°C-20°C)	5850	6.78	6.85	0.07	0.00	0.07	1.03
$MaxTemp(21^{\circ}C-30^{\circ}C), AvgTemp(21^{\circ}C-30^{\circ}C), MinTemp(21^{\circ}C-30^{\circ}C)$	4520	6.47	6.84	0.37	0.14	0.37	5.72
$\begin{array}{c} \text{MaxTemp}(31+^{\circ}\text{C}),\\ \text{AvgTemp}(21^{\circ}\text{C}\text{-}30^{\circ}\text{C}) \end{array}$	0	-	6.82	-	-	-	-
$\begin{array}{c} \text{MaxTemp}(31+^{\circ}\text{C}), \\ \text{AvgTemp}(31+^{\circ}\text{C}) \end{array}$	4471	6.72	6.88	0.16	0.03	0.16	2.38





Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	Records	Admissions		WIC	Average WIC	Gain
	ALL	Root	721	91.04	6522.86	6522.86	<u>6522.86</u>	
	Min	$0^{\circ}C-10^{\circ}C(1)$	174	94.63	1653.37	551.12		
		$11^{\circ}C-20^{\circ}C(2)$	376	89.14	3421.33	1140.44	2249.21	4273.65
		$21^{\circ}C-30^{\circ}C$ (3)	181	89.41	1672.91	557.64		
	Max	$0^{\circ}C-10^{\circ}C(1)$	3	101.00	38.93	9.73		
		$11^{\circ}C-20^{\circ}C$ (2)	306	92.59	2848.41	712.10	1600.81	4832.05
		$21^{\circ}C-30^{\circ}C$ (3)	283	89.07	2580.33	645.08	1690.81	
(Root)		$31 + ^{\circ}C(4)$	139	88.67	1295.56	323.89		
Ro	Avg	0°C-10°C (1)	49	98.65	495.38	123.84		
1 (11°C-20°C (2)	379	91.01	3490.43	872.61	1000 54	1000.00
		$21^{\circ}C-30^{\circ}C$ (3)	295	88.44	2685.31	671.33	1690.54	4832.32
		$31 + ^{\circ}C(4)$	8	93.62	91.04	22.76		
	MaxVar	$x < -2^{\circ}C(1)$	45	89.60	449.32	89.86		
		$-2^{\circ}C \leq x \leq -1^{\circ}C$ (2)	200	91.00	1867.44	373.49		
		$0^{\circ}C(3)$	212	89.82	1956.63	391.33	1369.18	5153.68
		$1^{\circ}C \leq x \leq 2^{\circ}C$ (4)	236	90.53	2186.84	437.37		
		${ m x}>2^{\circ}{ m C}~(5)$	38	92.87	385.68	77.14		





Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
Level	Covariate							
		Group	Records	Admissions		WIC	Average WIC	Gain
	All	$x < -2^{\circ} C (1)$	45	89.60	449.32	89.86	89.86	
	Min	$0^{\circ}C-10^{\circ}C(1)$	11	91.64	121.58	40.53		
(1))		$11^{\circ}C-20^{\circ}C$ (2)	19	87.37	200.39	66.80	161.37	-71.51
		$21^{\circ}C-30^{\circ}C$ (3)	15	90.93	162.15	54.05		
2°(Max	$0^{\circ}C-10^{\circ}C(1)$	1	104.00	7.07	1.77		
Š.		$11^{\circ}C-20^{\circ}C$ (2)	20	90.90	211.72	52.93	120.11	-30.25
r., 5		$21^{\circ}C-30^{\circ}C$ (3)	14	85.93	150.35	37.59	120.11	
2(MaxVar, x<-2°C		$31 + ^{\circ}C(4)$		90.70	111.31	27.83		
Max	Avg	0°C-10°C (1)	6	91.83	68.79	17.20		
2(]		$11^{\circ}C-20^{\circ}C$ (2)	20	88.40	210.66	52.66	119.49	-29.63
		$21^{\circ}C-30^{\circ}C$ (3)	18	90.33	191.81	47.95	119.49	-29.03
		$31 + ^{\circ}C(4)$	1	87.00	6.71	1.68		
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Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	Records	Admissions		WIC	Average WIC	Gain
	All	$-2^{\circ}C \leq x \leq -1^{\circ}C(2)$	200	91.00	1867.44	373.49	373.49	
(2))	Min	$0^{\circ}C-10^{\circ}C(1)$	44	96.32	454.40	151.47		
		$11^{\circ}C-20^{\circ}C$ (2)	106	89.50	1003.42	334.47	650.97	-277.48
-1°C		$21^{\circ}C-30^{\circ}C$ (3)	50	89.48	495.09	165.03		
VI	Max	$0^{\circ}C-10^{\circ}C(1)$	1	97.00	6.93	1.73		
×		$11^{\circ}C-20^{\circ}C$ (2)	92	92.34	896.50	224.12	487.78	-114.29
		$21^{\circ}C-30^{\circ}C$ (3)	76	90.28	730.09	182.52	401.10	-114.29
-2°		$31 + ^{\circ}C(4)$	31	88.58	317.61	79.40		
Var	Avg	0°C-10°C (1)	1	99.63	175.32	43.83		
ax		$11^{\circ}C-20^{\circ}C$ (2)	103	91.29	992.95	248.24	484.07	111 49
2(MaxVar,-2°C		$21^{\circ}C-30^{\circ}C$ (3)	79	88.63	761.01	190.25	484.97	-111.48
CN.		$31 + ^{\circ}C(4)$	2	100.00	10.59	2.65		
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Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	Records	Admissions		WIC	Average WIC	Gain
	All	$0^\circ C$ (3)	212	89.82	1956.63	391.33	391.33	
	Min	$0^{\circ}C-10^{\circ}C(1)$	60	93.87	593.16	197.72		
		$11^{\circ}C-20^{\circ}C$ (2)	109	88.20	1025.83	341.94	682.65	-291.33
ÌÌ		21°C-30°C (3)	43	88.28	428.98	142.99		
(3))	Max	0°C-10°C (1)	1	102.00	7.17	1.79		
		$11^{\circ}C-20^{\circ}C$ (2)	90	92.08	864.84	216.21	510.66	-119.34
0°C		21°C-30°C (3)	89	87.93	844.45	211.11		
		$31 + ^{\circ}C(4)$	32	88.34	326.19	81.55		
XVi	Avg	$0^{\circ}C-10^{\circ}C(1)$	14	100.79	154.78	38.70		
2(MaxVar,		$11^{\circ}C-20^{\circ}C$ (2)	108	89.30	1021.80	255.45	507.59	-116.27
2(21°C-30°C (3)	90	88.74	853.79	213.45	507.59	-110.27
		31+°C (4)	0	0.00	0.00	0.00		
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Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	$\mathbf{Records}$	Admissions		WIC	Average WIC	Gain
	All	$1^{\circ}C \le x \le 2^{\circ}C(4)$	236	90.53	2186.84	437.37	437.37	
(4))	Min	$0^{\circ}C-10^{\circ}C(1)$	50	93.86	507.91	169.30		
		$11^{\circ}C-20^{\circ}C$ (2)	128	89.68	1208.86	402.95	761.50	-324.13
2°C		$21^{\circ}C-30^{\circ}C$ (3)	58	89.56	567.72	189.24		
∨I ×	Max	$0^{\circ}C-10^{\circ}C$ (1)	0	0.00	0.00	0.00		
vi –		$11^{\circ}C-20^{\circ}C$ (2)	92	92.95	896.15	23.24	260.2E	68.02
1°C		$21^{\circ}C-30^{\circ}C$ (3)	95	89.18	899.29	224.82	369.35	
		$31 + ^{\circ}C$ (4)	49	88.61	485.15	121.29		
2(MaxVar,	Avg	$0^{\circ}C-10^{\circ}C(1)$	10	99.00	113.07	28.27		
Iax		$11^{\circ}C-20^{\circ}C$ (2)	134	91.77	1271.13	317.78	563.66	196.90
2(N		$21^{\circ}C-30^{\circ}C(3)$	91	87.68	863.45	215.86	003.00	-126.29
		$31 + ^{\circ}C(4)$	1	99.00	6.97	1.74		
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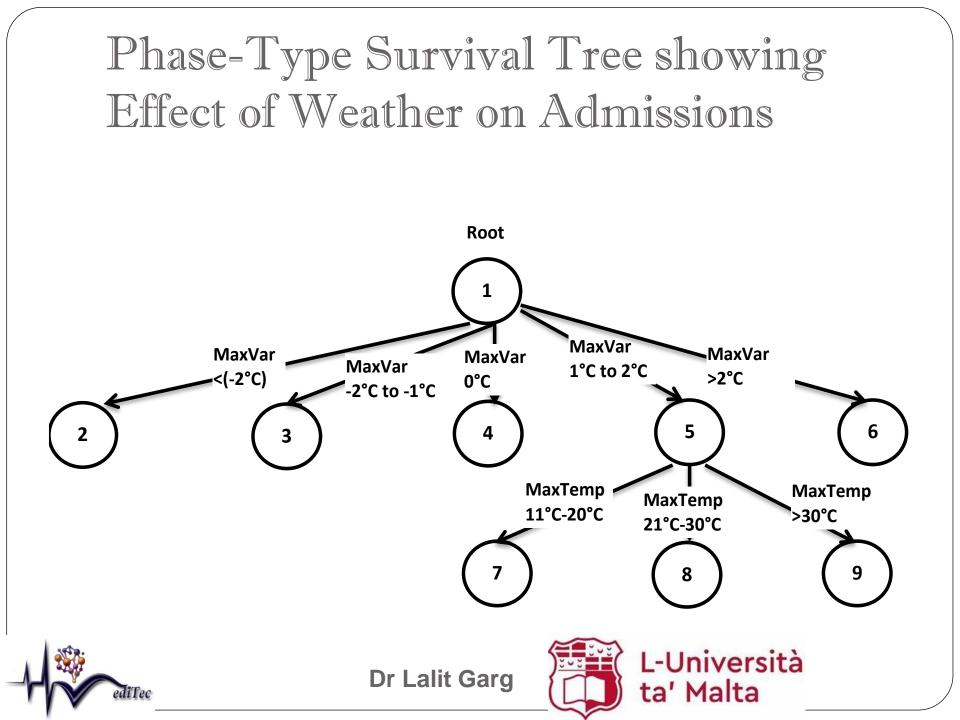


Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	$\mathbf{Records}$	Admissions		WIC	Average WIC	Gain
(2))	All	11-20 (2)	92	92.95	896.15	23.24	23.24	
	Min	$0^{\circ}C-10^{\circ}C(1)$	49	93.78	498.37	124.59		
(4), Max		$11^{\circ}C-20^{\circ}C$ (2)	43	92.00	434.78	108.69	233.29	-210.05
4),		$21^{\circ}C-30^{\circ}C$ (3)	0	0.00	0.00	0.00		
r (4	Avg	0°C-10°C (1)	49	93.78	498.37	124.59		
3(MaxVar		$11^{\circ}C-20^{\circ}C$ (2)	43	92.00	434.78	108.69	233.29	-210.05
Max		$21^{\circ}C-30^{\circ}C$ (3)	0	0.00	0.00	0.00	233.29	-210.05
3(1		$31 + ^{\circ}C(4)$	0	0.00	0.00	0.00		
(3))	All	21-30 (3)	95	89.18	899.29	224.82	224.82	
x (;	Min	$0^{\circ}C-10^{\circ}C(1)$	1		BAD WIC			
Max		$11^{\circ}C-20^{\circ}C$ (2)	81	89.05	773.46	257.82	BAD WIC	BAD WI
(I),]		$21^{\circ}C-30^{\circ}C$ (3)	13	89.31	141.15	47.05		
r (4	Avg	0°C-10°C (1)	0	0.00	0.00	0.00		
3(MaxVar (4), Max		$11^{\circ}C-20^{\circ}C$ (2)	52	91.08	512.81	128.20	235.24	10.41
		$21^{\circ}C-30^{\circ}C$ (3)	43	86.88	428.13	107.03	230.24	-10.41
3(1		31+°C (4)	0	0.00	0.00	0.00		
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Level	Covariate	Covariate	No. of	Mean	WIC	Average	Total	WIC
		Group	Records	Admissions		WIC	Average WIC	Gain
4))	All	31+(4)	49	88.61	485.15	121.29	121.29	
× (*	Min	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00	0.00		
Ma		$11^{\circ}C-20^{\circ}C$ (2)	4	77.50	46.13	15.38	164.93	-43.64
4),		$21^{\circ}C-30^{\circ}C$ (3)	45	89.60	448.66	149.55		
r (Avg	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00	0.00		
кVа		$11^{\circ}C-20^{\circ}C$ (2)	0	0.00	0.00	0.00		BAD WI
Maz		$21^{\circ}C-30^{\circ}C$ (3)	48	88.40	476.47	119.12	BAD WIC	DAD WI
(5)) 3(MaxVar (4), Max (4))		$31 + ^{\circ}C(4)$	1		BAD WIC			
(5))	All	$x>2^\circ C$ (5)	38	92.87	385.68	77.14	77.14	
õ	Min	$0^{\circ}C-10^{\circ}C(1)$	9	99.33	103.12	34.37		
>2°C		$11^{\circ}C-20^{\circ}C$ (2)	14	91.29	151.95	50.65	138.99	-61.85
		$21^{\circ}C-30^{\circ}C$ (3)	15	90.47	161.90	53.97		
ar,	Max	$0^{\circ}C-10^{\circ}C(1)$	0	0.00	0.00	0.00		
Vxe		$11^{\circ}C-20^{\circ}C$ (2)	12	98.50	133.30	33.32	104.15	-27.01
2(MaxVar, x		$21^{\circ}C-30^{\circ}C$ (3)	9	93.78	102.12	25.53	104.15	-27.01
0		31+°C (4)	17	88.41	181.18	45.29		
	Avg	0°C-10°C (1)	3	96.00	38.63	9.66		
	Ŭ	11°C-20°C (2)	14	98.64	154.06	38.52	405.00	
		21°C-30°C (3)	17	88.06	181.07	45.27	105.29	-28.15
		31+°C (4)	4	90.75	47.39	11.85		
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Most significant prognostic factor affecting the number of admissions is the maximum variability in the average temperature between one day and the next.



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- Most significant prognostic factor affecting the number of admissions is the maximum variability in the average temperature between one day and the next.
- The maximum temperature affects the number of admissions only when the average temperature increases by 1°C-2°C than the previous day.

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• The minimum temperature and average temperature do not affect number of admissions.



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- The minimum temperature and average temperature do not affect number of admissions.
- These results might be different for different geographic regions due to different weather conditions and different genetic profile of inhabitants there.





• Predictions and Accuracy Tests

Group	No. of	Actual	Predicted	1	Squared	Absolute	Percentage
	Records	Mean Adm.	Mean Adm.	Error	Error	Error	Error (%)
$MaxVar(x < -2^{\circ}C)$	31	92.13	89.60	-2.53	6.40	2.53	2.75
$MaxVar(-2^{\circ}C \leq x \leq -1^{\circ}C)$	99	92.34	91.00	-1.34	1.80	1.34	1.45
$MaxVar(x = 0^{\circ}C)$	93	92.77	89.82	-2.95	8.70	2.95	3.18
$MaxVar(x>2^{\circ}C)$	19	97.63	92.87	-4.76	22.66	4.76	4.88
$\begin{array}{c} MaxVar(1^{\circ}C \leq x \leq 2^{\circ}C), \\ MaxTemp \ (11^{\circ}C - 20^{\circ}C) \end{array}$	42	100.95	92.95	-8.00	64.00	8.00	7.92
$\begin{array}{l} MaxVar(1^{\circ}C \leq x \leq 2^{\circ}C), \\ MaxTemp \ (21^{\circ}C\text{-}30^{\circ}C) \end{array}$	54	91.63	89.18	-2.45	6.00	2.45	2.67
$\begin{array}{c} \operatorname{MaxVar}(1^{\circ}\mathrm{C} \leq x \leq 2^{\circ}\mathrm{C}), \\ \operatorname{MaxTemp} (31 + ^{\circ}\mathrm{C}) \end{array}$	27	95.48	88.61	-6.87	47.20	6.87	7.20





Accuracy test for all predictions

		MSE	RMSE	MAD	BIAS
LOS	Weather	0.08	0.28	0.26	-0.09
105	Personal Characteristics	1.15	1.07	0.74	-0.69
Admissions	Weather	16.17	4.02	3.37	-3.37
Admissions	Personal Characteristics	1.38	1.17	0.96	-0.82

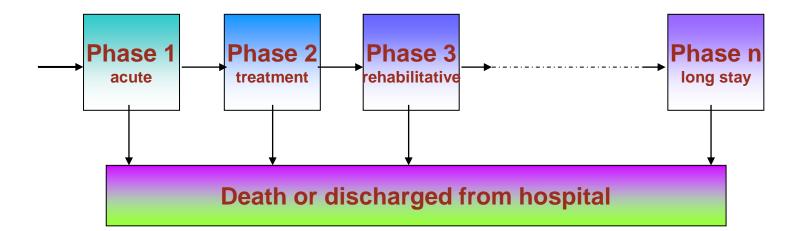
MSE: Mean Square Error, RMSE: Root Mean Square Error, MAD: Mean Absolute Deviation BIAS: Bias





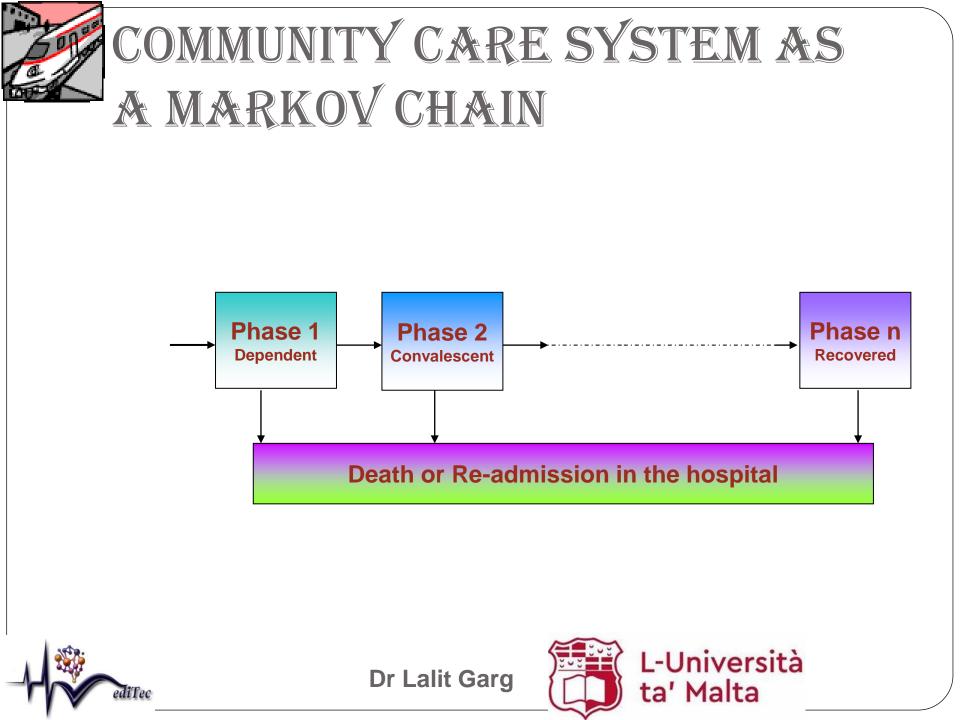


HOSPITAL CARE SYSTEM AS A MARKOV CHAIN

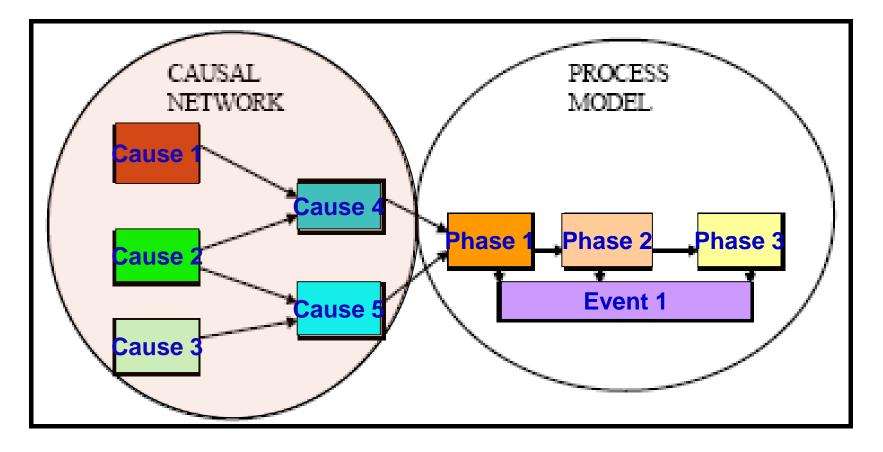








NDITIONAL PHASE TYPE STRIBUTION



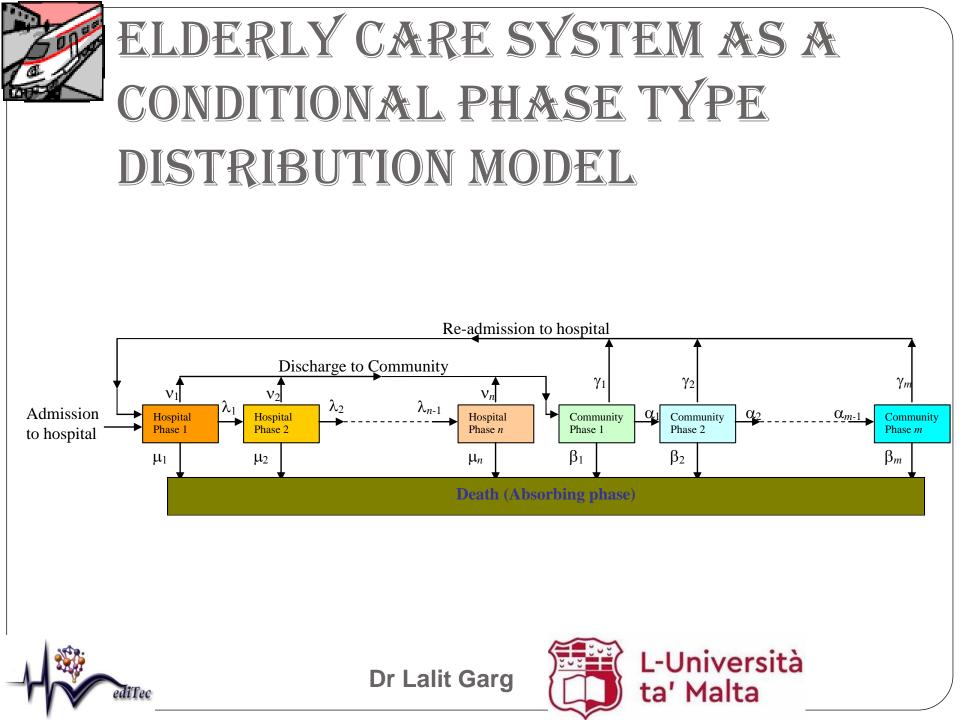
Conditional phase type distribution (taken from Marshall et al., 2000a)



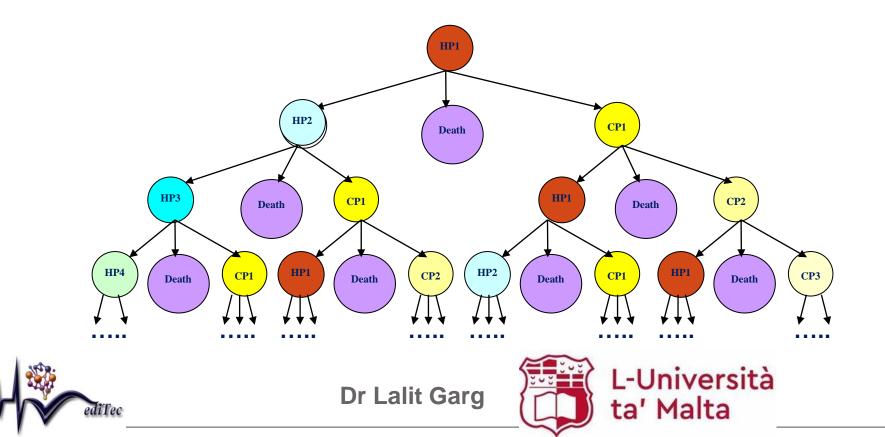
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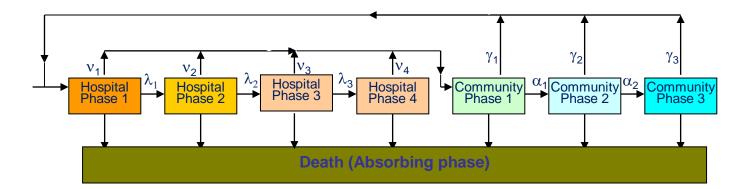


Sequential pattern mining: Patient pathways



8-PHASE MARKOV MODEL Acute Dependent **Treatment** 2 Convalescent 6 Rehabilitative 3 Recovered 7 Long stay Δ Death L-Università **Dr Lalit Garg** ta' Malta diTec

An Example: Eight-phase model

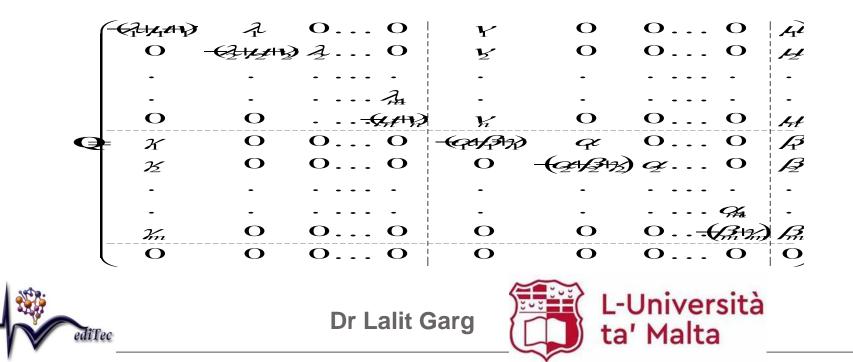






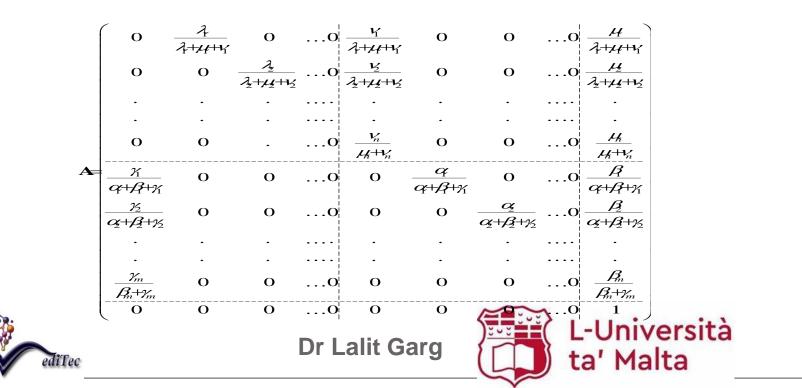
Transition matrix

Transition matrix $Q = \{q_{ij}\}$ =Transition rate (next transition is to phase *j* | currently in phase *i* }



Transient probability matrix

Associated transient probability matrix $A = \{a_{ij}\}$, where $a_{ij} =$ Probability {next transition is to phase *j* | currently in phase *i* } and,



Expected time

The expected time spent in a phase *i* (when next transition is to phase *j* / given that the patient has entered phase *i*)





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Expected cost

The expected cost of care in phase *i* (when next transition is to phase *j* / given that the patient has entered phase *i*)





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Probability of occurrence of a patient pathway



where, pathway r is having in total k transitions among various hospital and community phases and finally death in phase l

 p_h is the transient probability for the h^{th} transition. (If the h^{th} transition is from phase *i* to phase *j* then $p_h = a_{ij}$),

 $a_{l,(m+n+1)}$ is the probability of death in phase l





Expected duration of a patient pathway



where,

 t_h is the expected time spent in the phase before the h^{th} transition. If the h^{th} transition is from phase *i* to phase *j* then $t_h = t_{ij} =$ time spent in the phase *l*

 $t_{l,(m+n+1)}$ is the time spent in phase *l* before death.

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L-Università ta' Malta Expected total cost of a patient pathway



where,

 c_h is the expected cost of stay in the phase before the h^{th} transition. If the h^{th} transition is from phase *i* to phase *j* then $c_h = c_{ij} =$ time spent in the phase *I*

 $c_{l,(m+n+1)}$ is the cost of stay in phase *l* before death.

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Daily resource requirements

On day *d*:

Number of patients in phase i =

 η_i = (number of patients on the day *d*-1) + (number of patients entered in phase *I* on day *d*) – (number of patients left phase *i* on day *d*)





Daily resource requirements

In fact, we will calculate the number (η_i) of patients in phase *i* after every transition event.

For a transition from phase *i* to phase *j* with probability a_{ij}

 $\eta_i = (1 - a_{ij})^*$ (the number of patients in phase *i* before the transition)

 η_j = (the number of patients in phase *j* before the transition) + (a_{ij}) *(the number of patients in phase *i* before the transition)





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Daily resource requirements

Total daily resource requirement for the given initial population of patients in various phases



where, $\eta_i(t_0) =$ Given initial number of patients in phase *i*.

 $\eta_i(t_{given}) = \text{daily resource requirement at time } t_{given}$ for the care system starting with one patient in phase *i* (no new admissions).



Incorporating new admissions

Number of patients in phase 1 =

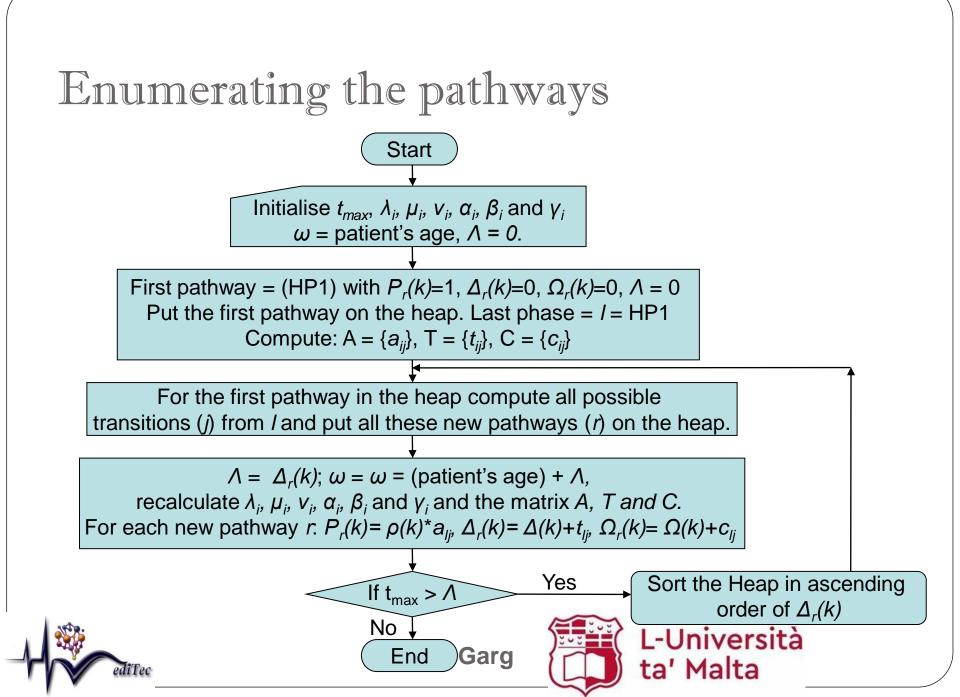
 $\eta_1 = 1 + (\text{the number of patients in the hospital phase 1 before the new admission})$

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Number of admissions required For a new hospital

Where $B(t_{given})$ = number of beds available at time t_{given}

assuming that there are no patients in the hospital at t = 0.



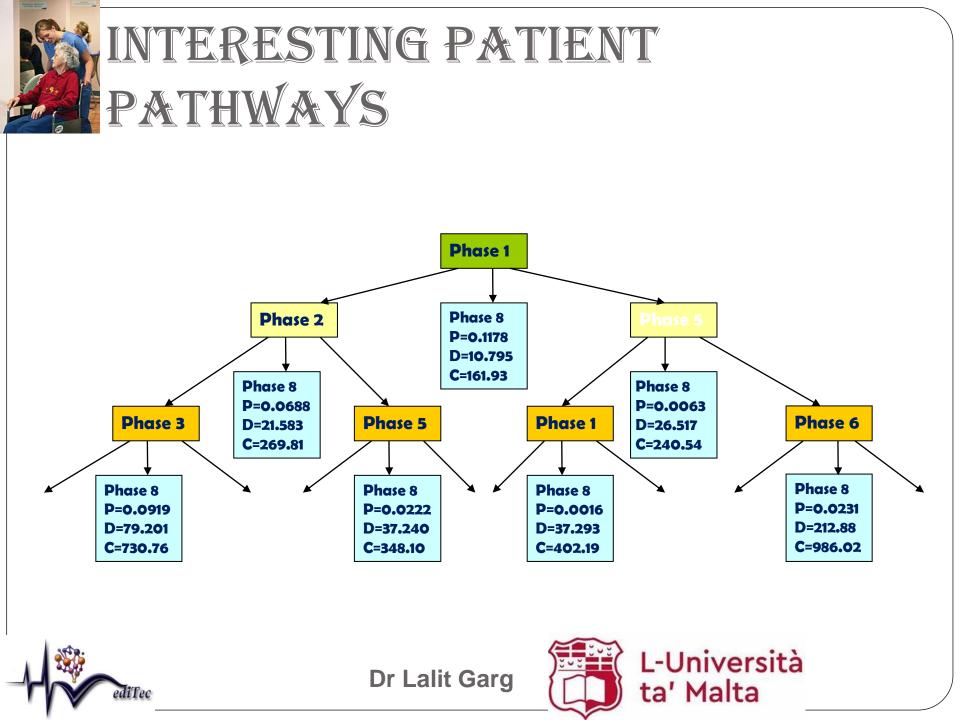




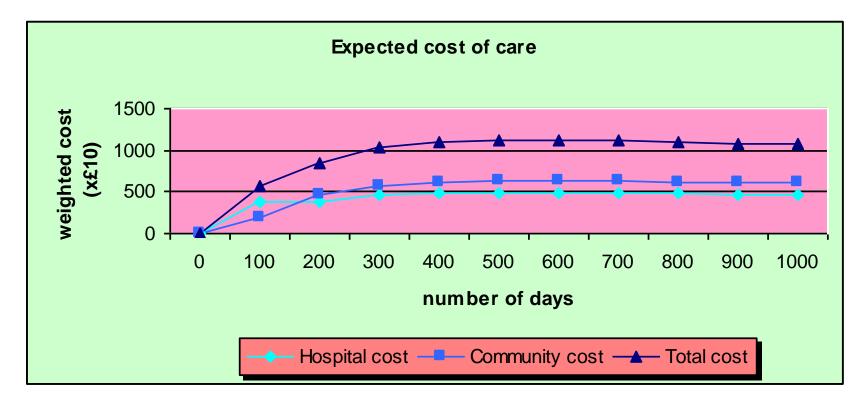
- where, $\eta(t_{given})$ = the total number of patients still remaining at t_{given} for the initial population of patients in various phases (without any new admissions).
- $\eta'(t_{given})$ = the total number of patients remaining at t_{given} having been admitted before t_{given} (with one admission per day and no initial population)







Expected cost of care

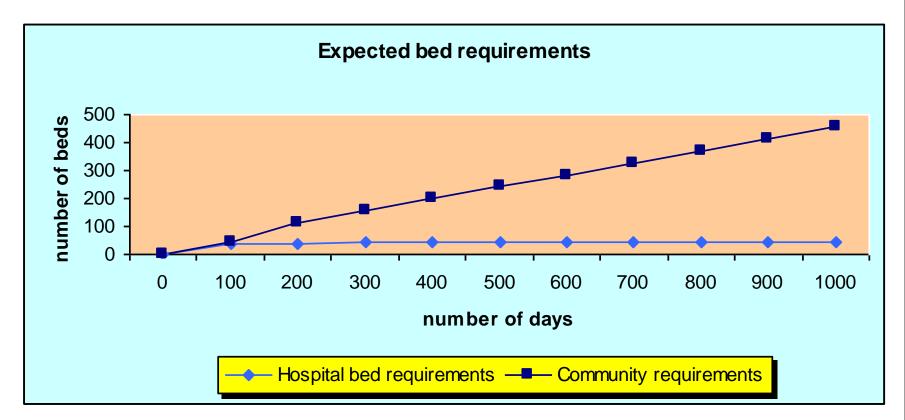


Data source: S.I. McClean and P.H. Millard, "Patterns of length of stay after admission in geriatric medicine: an event history approach", *The Statistician*, 42(3), 1993, pp. 263–274





Expected bed requirements

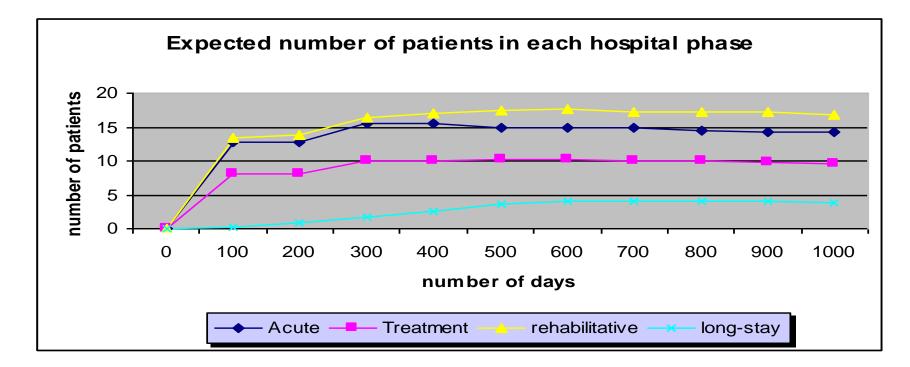


Data source: S.I. McClean and P.H. Millard, "Patterns of length of stay after admission in geriatric medicine: an event history approach", *The Statistician*, 42(3), 1993, pp. 263–274

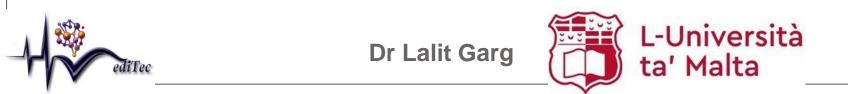




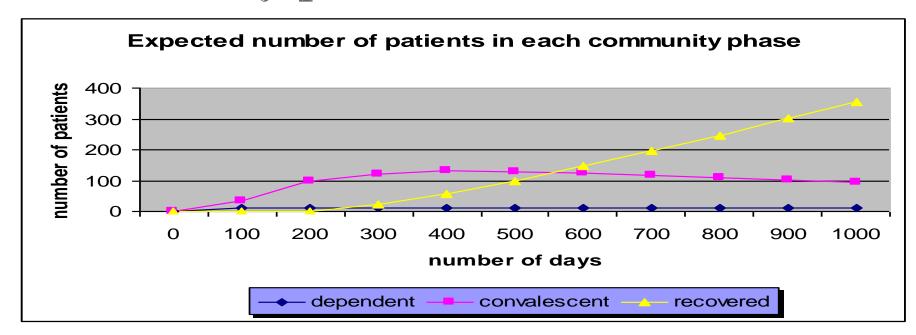
Expected number of patients in each hospital phase



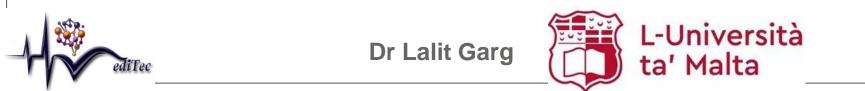
Data source: S.I. McClean and P.H. Millard, "Patterns of length of stay after admission in geriatric medicine: an event history approach", *The Statistician*, 42(3), 1993, pp. 263–274

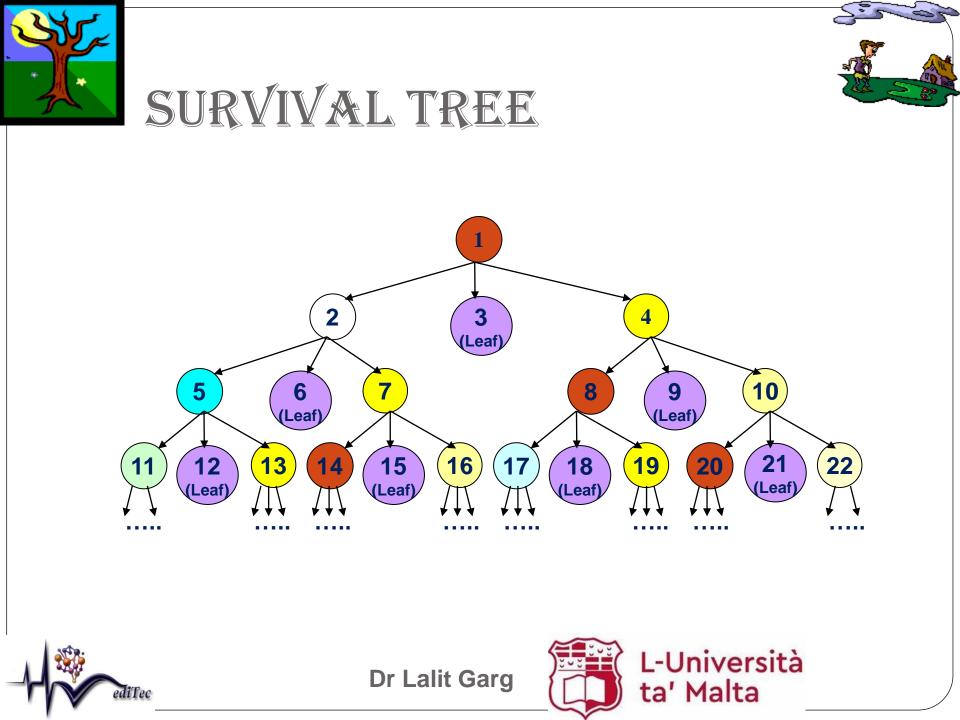


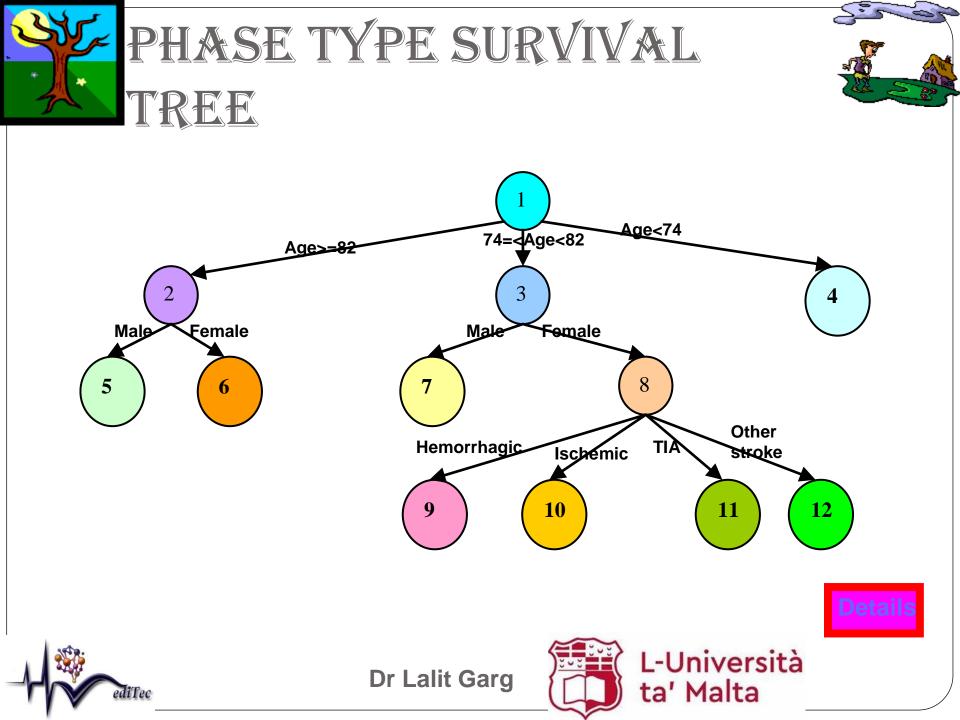
Expected number of patients in each community phase



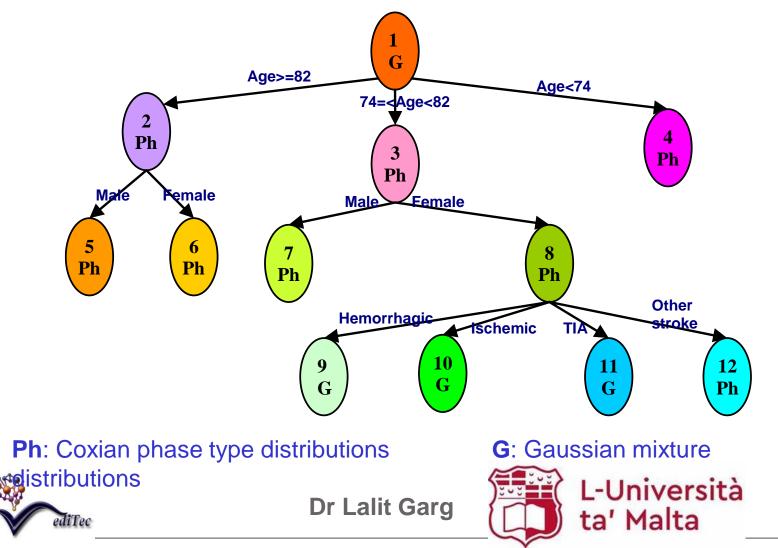
Data source: S.I. McClean and P.H. Millard, "Patterns of length of stay after admission in geriatric medicine: an event history approach", *The Statistician*, 42(3), 1993, pp. 263–274







MIXTURE DISTRIBUTION SURVIVAL TREE





Splitting criteria used

- MLIC (Maximum likelihood ratio criterion)
- AIC (Akaike Information Criterion)
- AICc (Corrected AIC)
- BIC (Bayesian Information Criterion)
- BICc (Corrected BIC)
- HQIC (Hannan and Quinn Criterion)
- WIC (The Weighted-Average Information Criterion)

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Maximum likelihood ratiobased Splitting criteria

Maximum likelihood ratio criterion

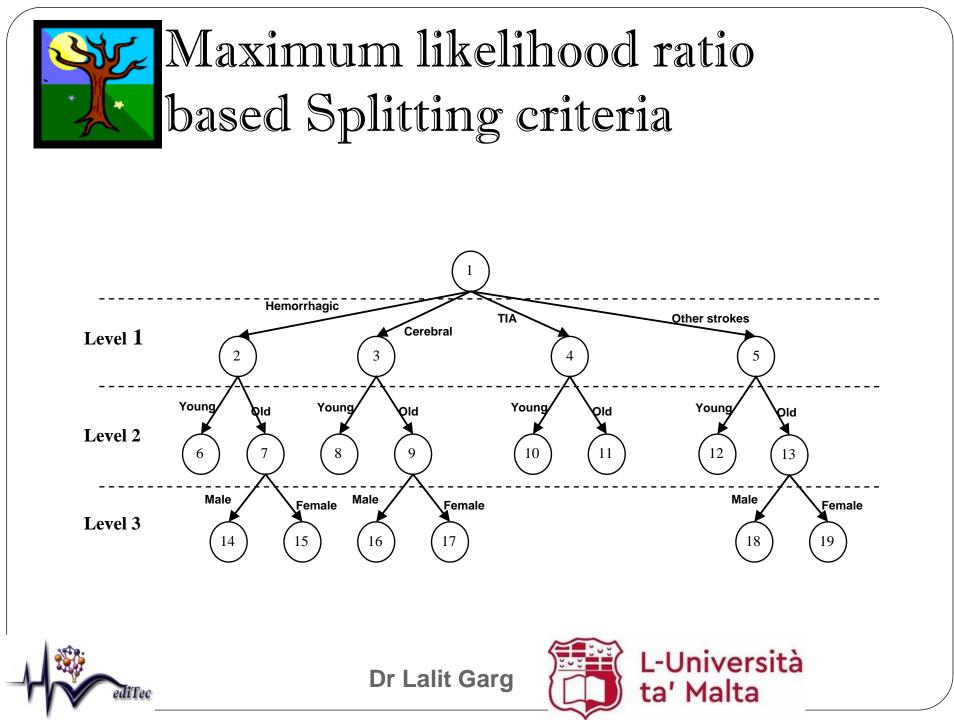
MLIC(df) = -2*Log likelihood.

$$\text{MLIC}(df_1) - \text{MLIC}(df_2) \sim \chi^2_{df_1 - df_2} (p < \alpha)$$

• **df*: number of free parameters required to be estimated









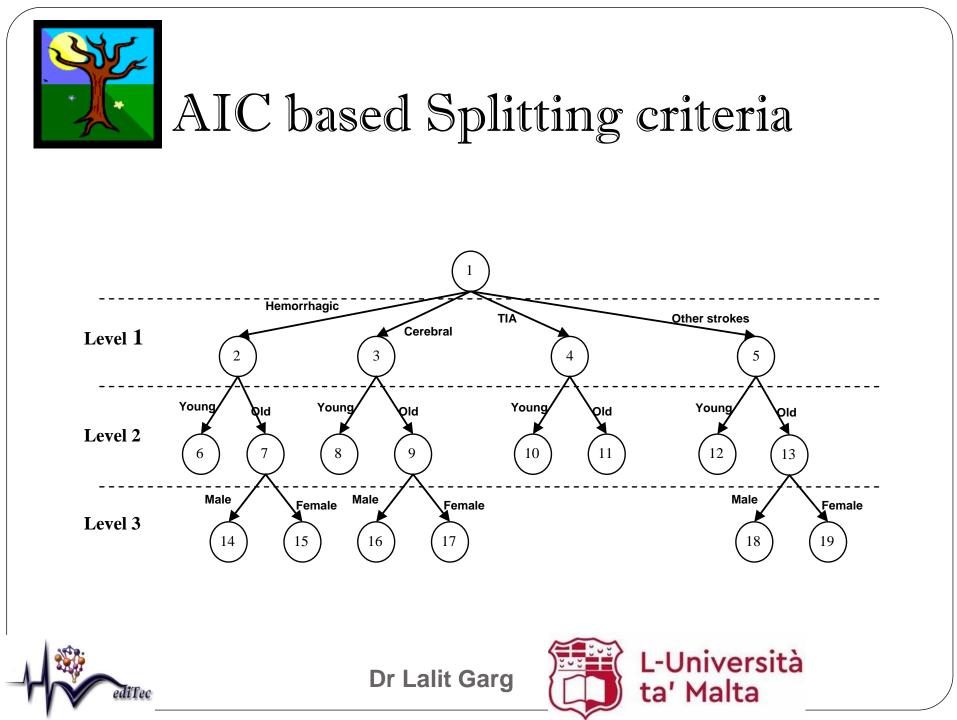
AIC based Splitting criteria

• Akaike Information Criterion

AIC(df) = -2*Log likelihood + 2*df.









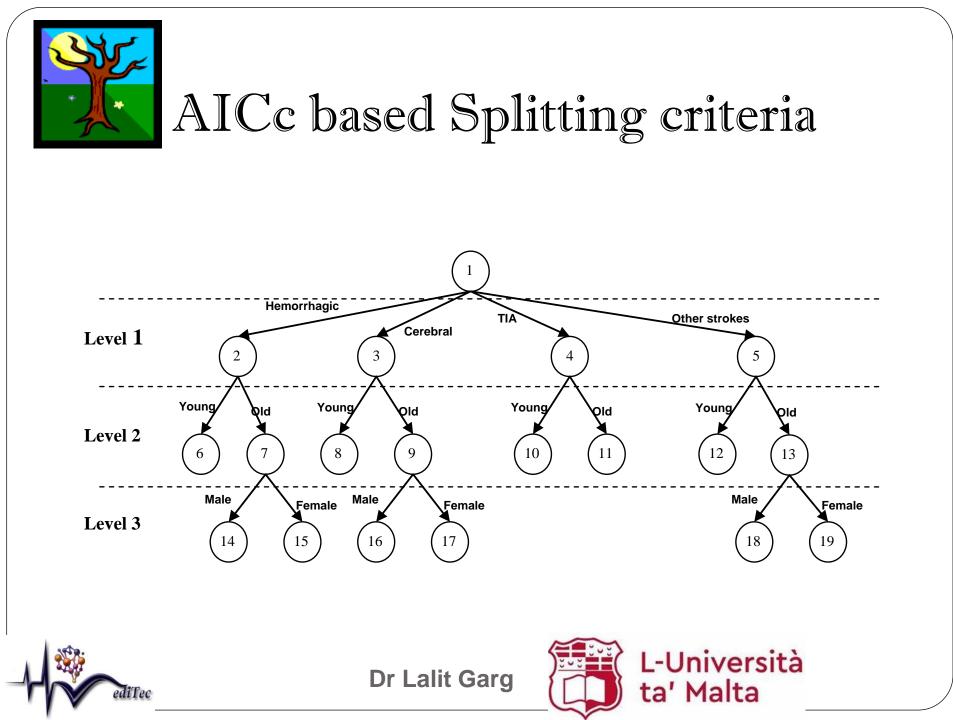
AICc based Splitting criteria

• Corrected AIC

AICc(df) = -2*Log likelihood $+2*df + \frac{2*df*(df+1)}{(n-(df+1))}$







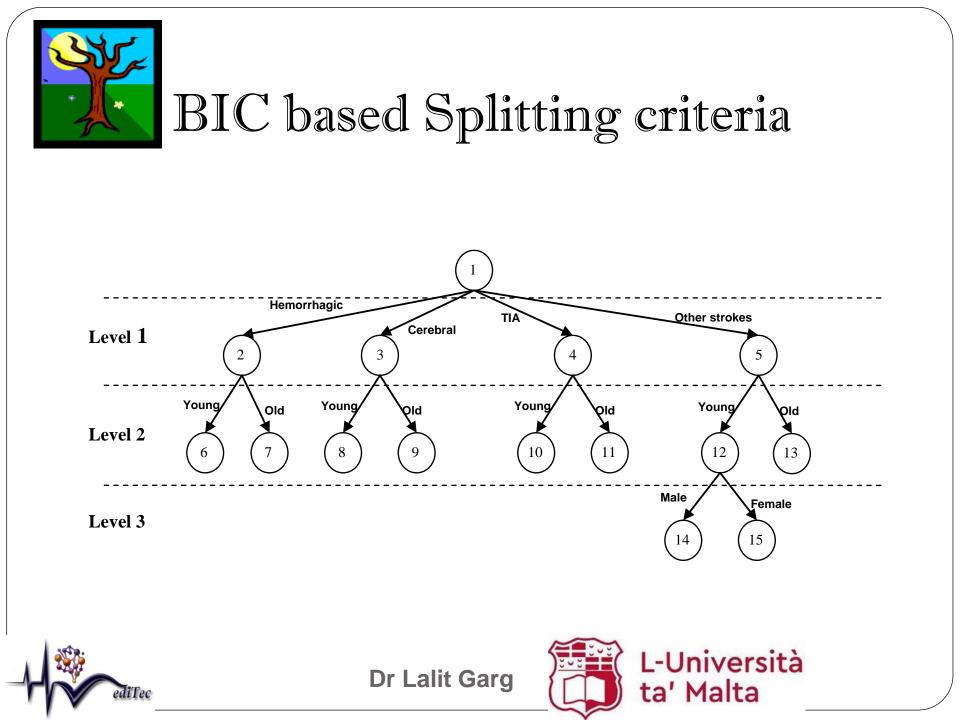


• Bayesian Information Criterion

BIC(df) = -2*Log likelihood+df *log(n)







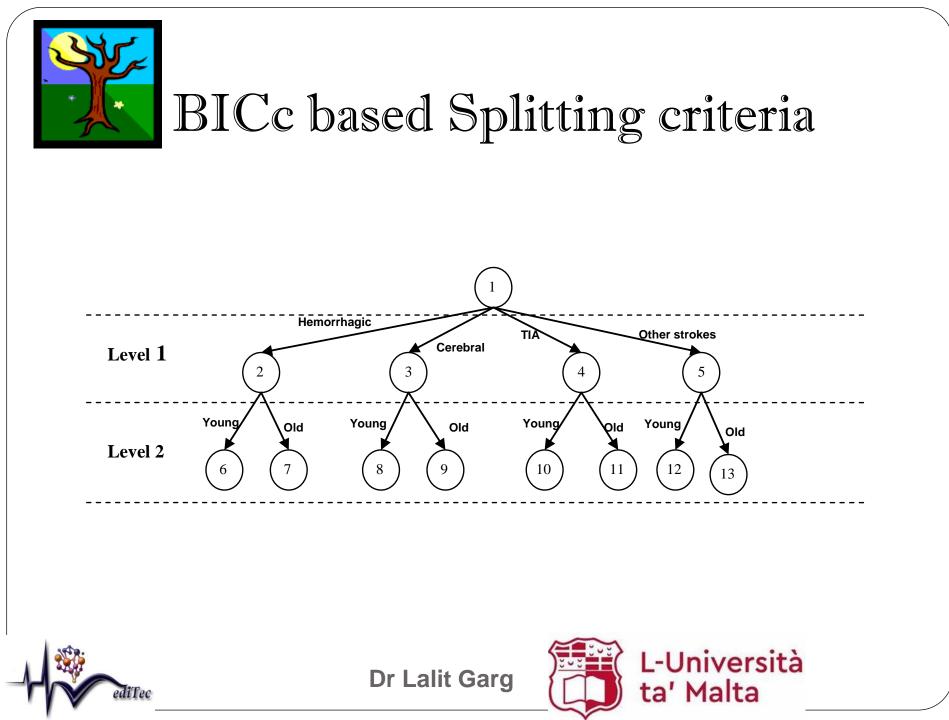


• Corrected BIC

$BICc(df) = -2*Log \ likelihood + df * log(n) + \frac{2df * (df + 1)}{(n - (df + 1))}$







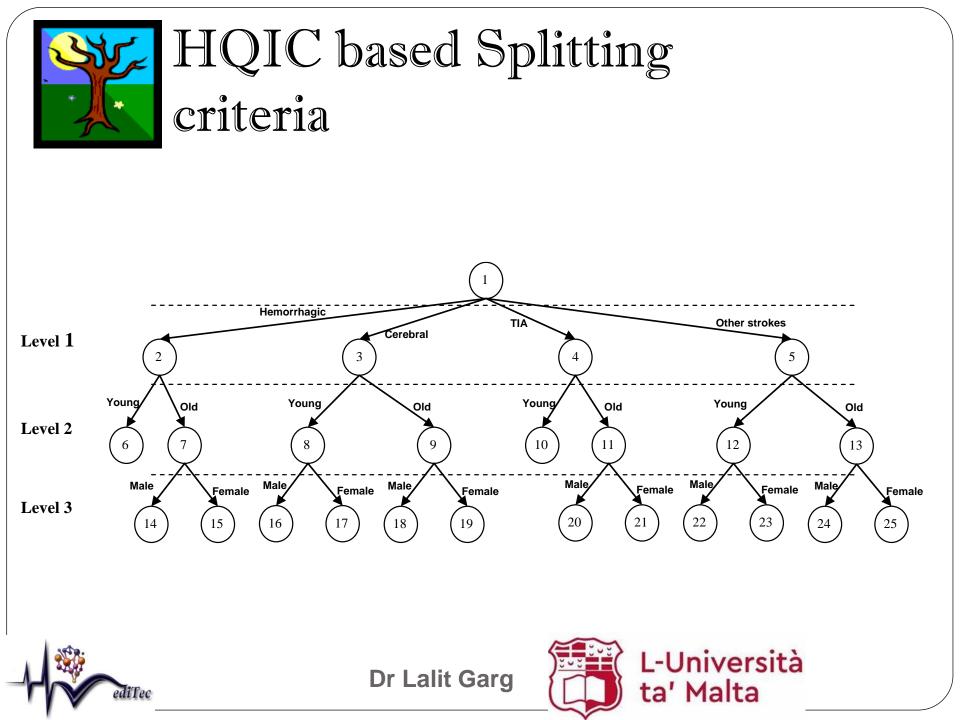


• Hannan and Quinn Criterion

HQIC(df) = -2*Log likelihood+df * log(log(n))









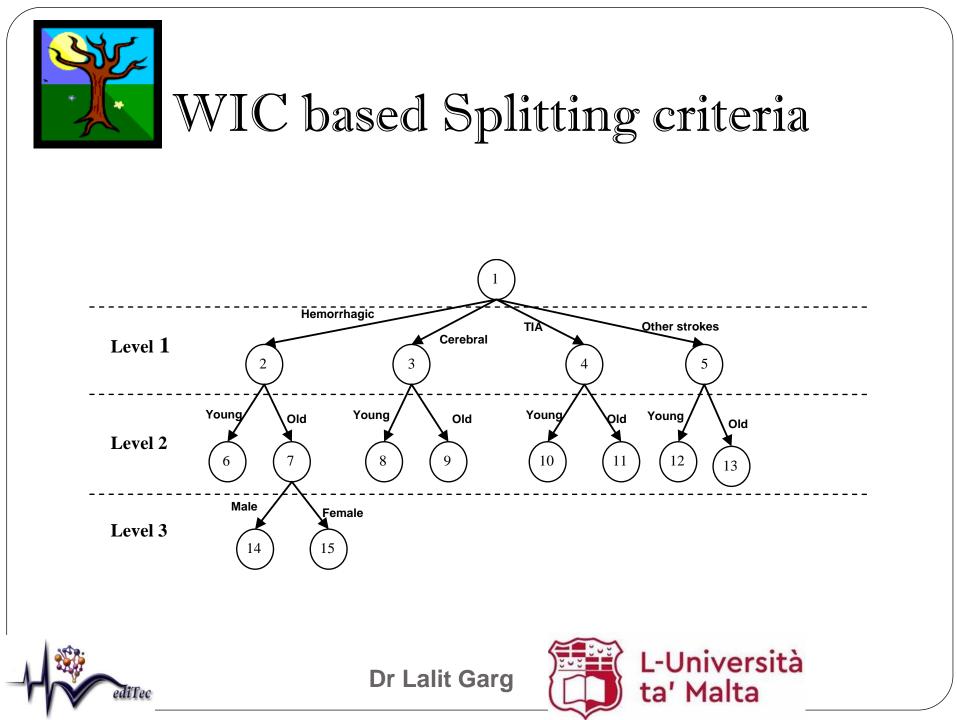
WIC based Splitting criteria

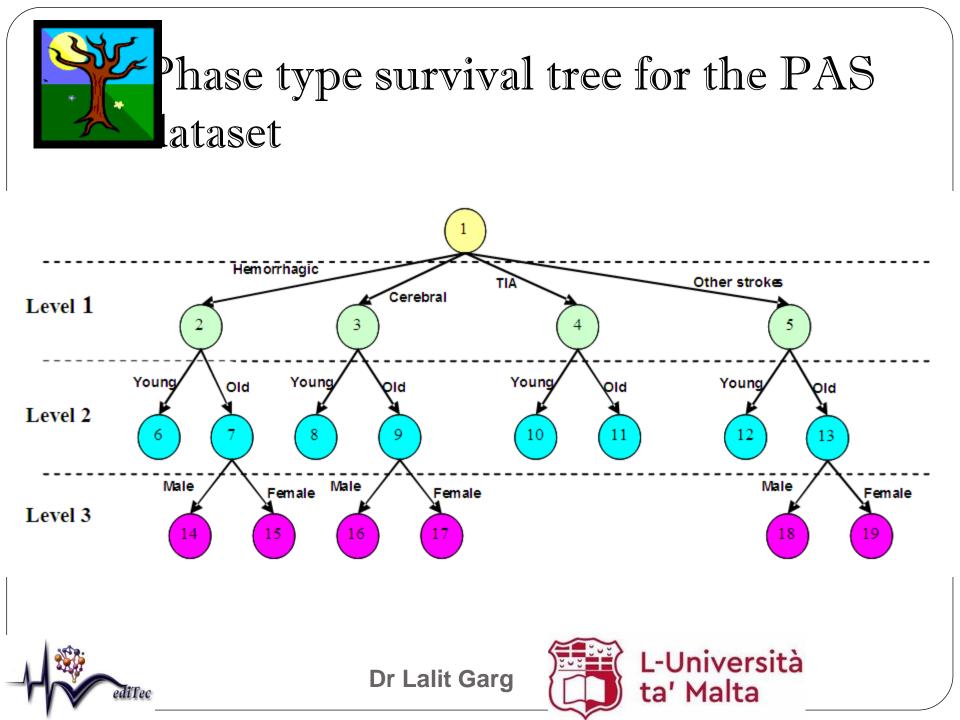
• The Weighted-Average Information Criterion:

WIC(df) =
$$\left(\frac{2*n}{2*n + (\log(n)*(n - (df + 1)))}\right) * AICc + \left(\frac{\log(n)*(n - (df + 1))}{2*n + (\log(n)*(n - (df + 1)))}\right) * BIC$$





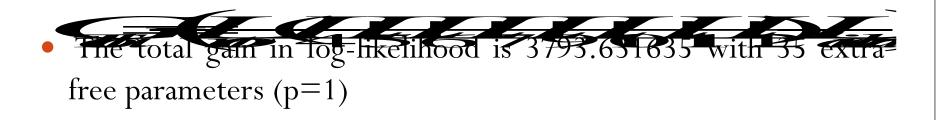






Phase type survival tree for the PAS dataset

• The total gain in the with in node homogeneity







Extended Phase type survival tree

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• The phase type survival tree approach can be extended by further growing the survival tree by partitioning the terminal nodes into subgroups with more homogeneous patient pathways based on covariates representing outcome measures such as discharge destination.



Extended Phase type survival tree

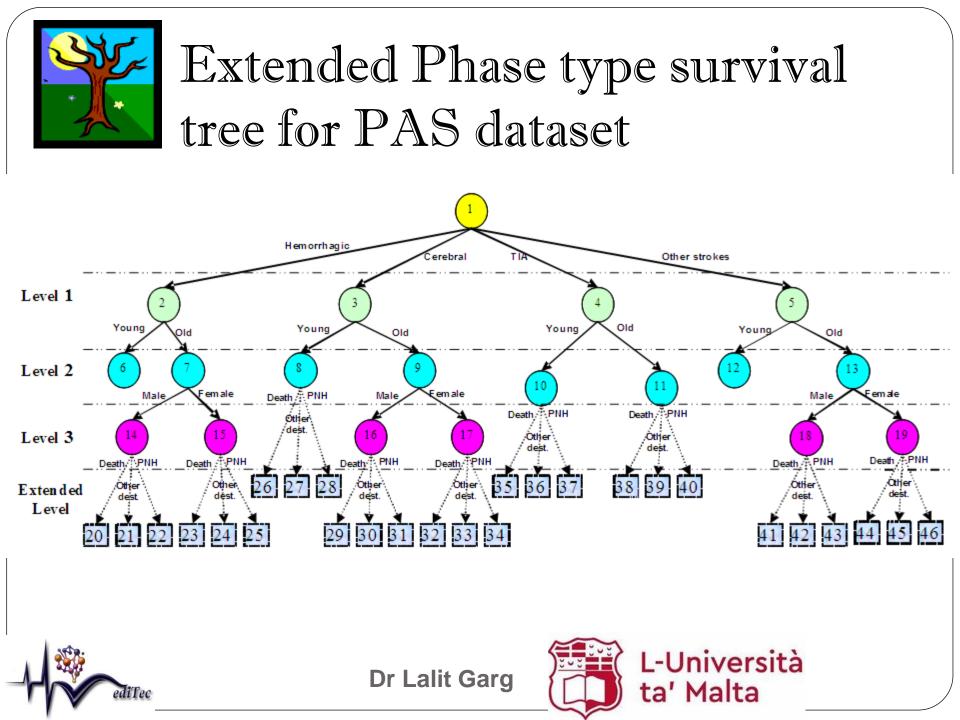
Although the information about the discharge destination is not available at the time of admission, the probability of each discharge destination can be assigned using cohort analysis.







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Extended Phase type survival tree for PAS dataset

$G_{Total} = 163.53$ at the cost of 16 additional free parameters (p<0.000001).





DECISION SUPPORT SÝSTEM

This DSS model has been further enhanced and now it can answer the questions like

- ✓ Bed occupancy and resource allocation: Resource requirements in various care units at various times
- ✓ Survival Analysis: Possibility of a patient to survive after a particular duration
- ✓ Budgetary requirements: The expected cost of care after a particular duration
- ✓ What-if analysis: Forecasting effects of various policies
 This model has been presented in CBMS-2008.









The transition probability matrix •





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• The initial distribution of patients (at t = 0)



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• If there are no patient admissions, then the expected distribution of patients (\mathbf{s}_t)

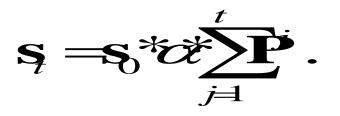
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$\mathbf{s}_t = \mathbf{s}_0 * \mathbf{P} \, .$



• If patient admissions (or arrivals) are modeled using a Poisson process with a mean arrival rate α .





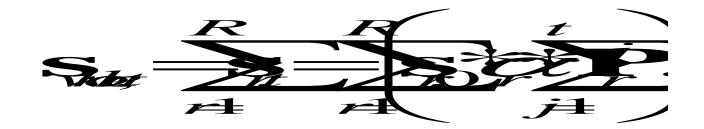
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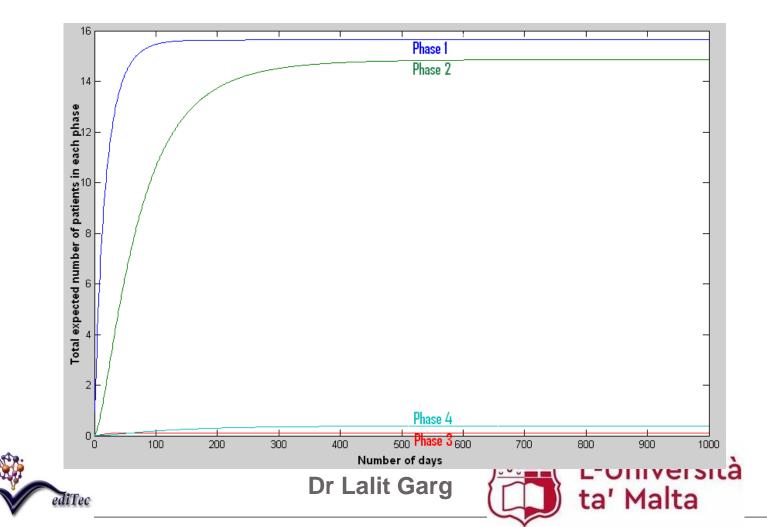
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• For the whole care system with *R* clusters, patient distribution:

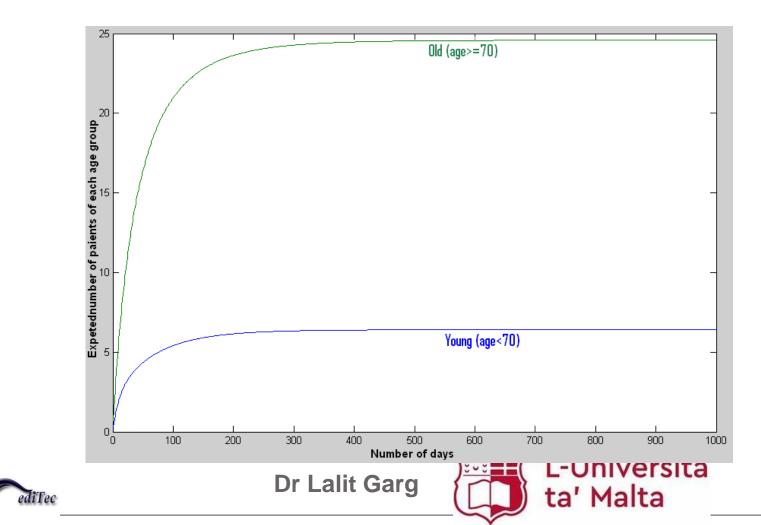




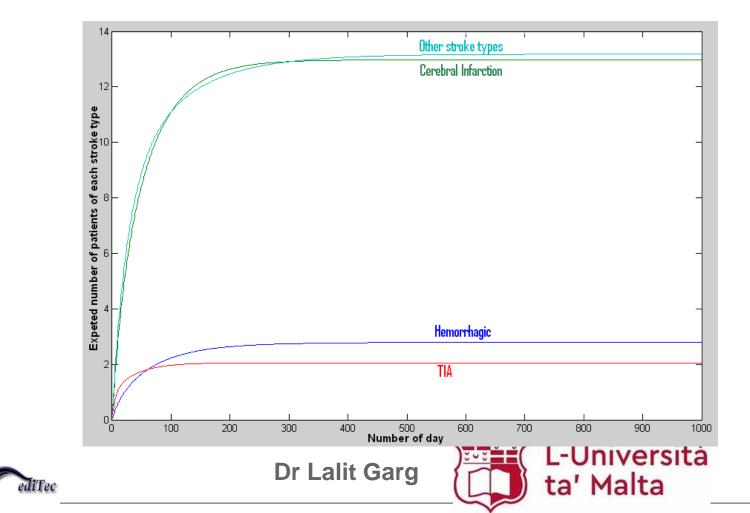
Total expected number of patients in each phase:



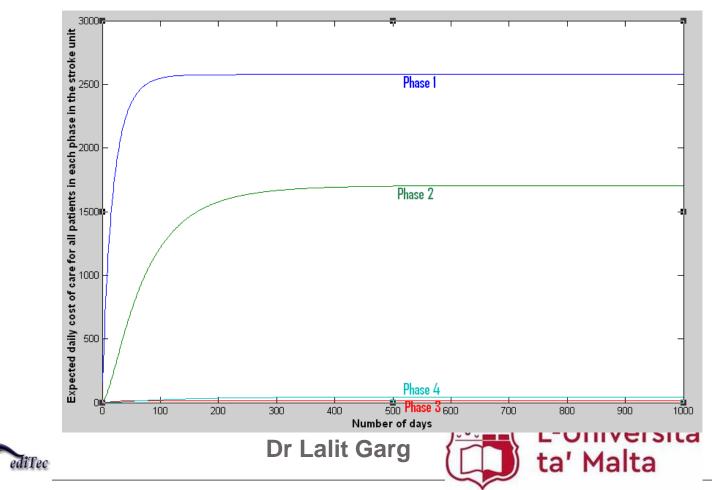
Expected number of patients of each age group



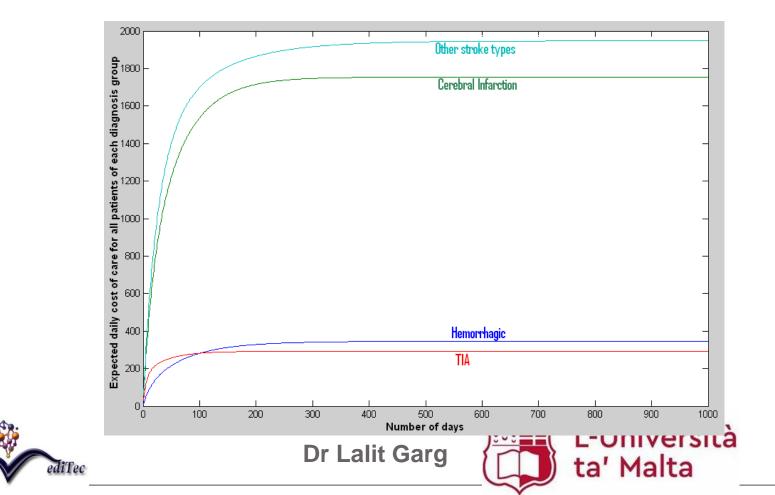
Expected number of patients of each stroke type



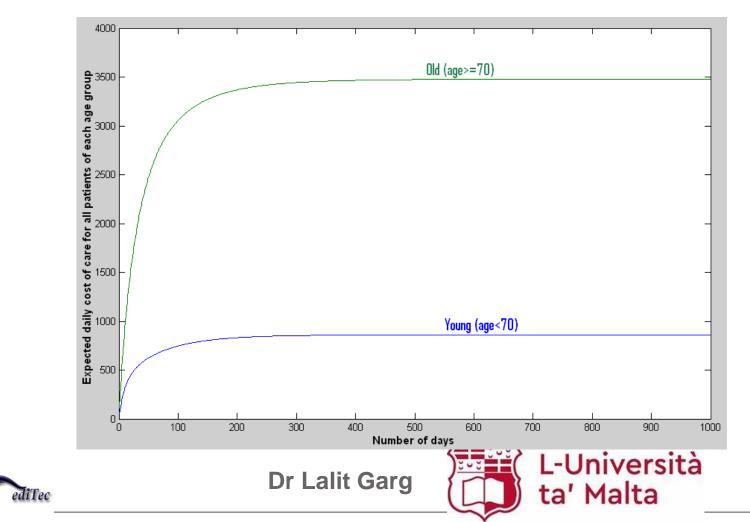
Expected daily cost of care (in £s) for all patients in each phase of the stroke unit



Expected daily cost of care (in £s) for all patients of each diagnosis group



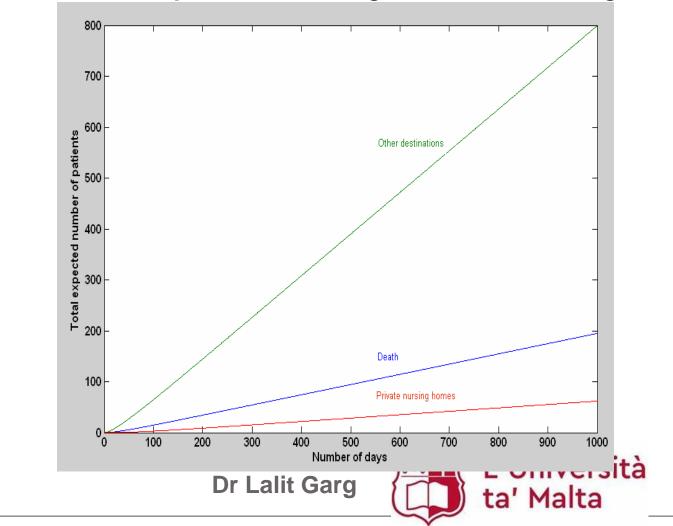
Expected daily cost of care (in £s) for all patients of each age group



Hospital Capacity Planning

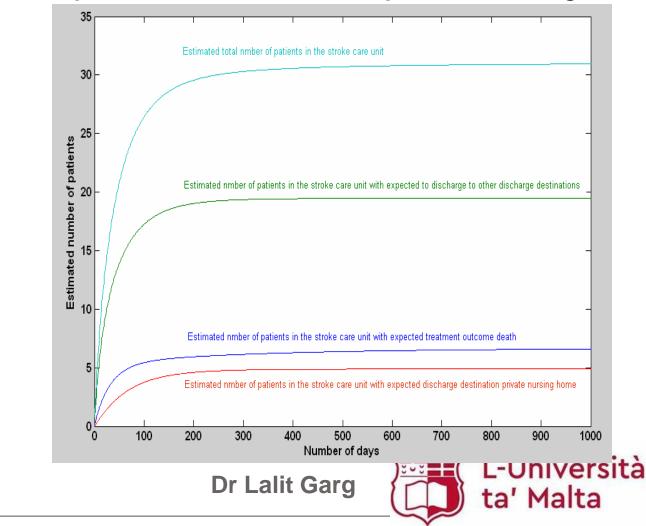
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Total expected number of patients discharged to each discharge destinations



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Distribution of patients based on their expected discharge destinations





• The total expected daily cost after k days:

$$\boldsymbol{\Omega}_{k} = \mathbf{s}_{whole,k} * \mathbf{c}.$$

• Where cost vector

$$\mathbf{c} = \{c_1, c_2, c_3, \dots, c_n, c_{n+1}, c_{n+2}, \dots, c_{n+m}\}^T$$

• where c_i is the daily cost of care in state *i*.



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- We attach unit costs* of £164.80 per day for stay in acute care (phase 1) and £114.80 per day for stay in rehabilitative care or long stay care (phase 2, phase 3 and phase 4).
- *using estimates from Saka et al. (2009) which is adjusted from 2005.
- Saka O, McGuire A, Wolfe C (2009). Cost of stroke in the United Kingdom. Age and Aging. 38: 27-32.

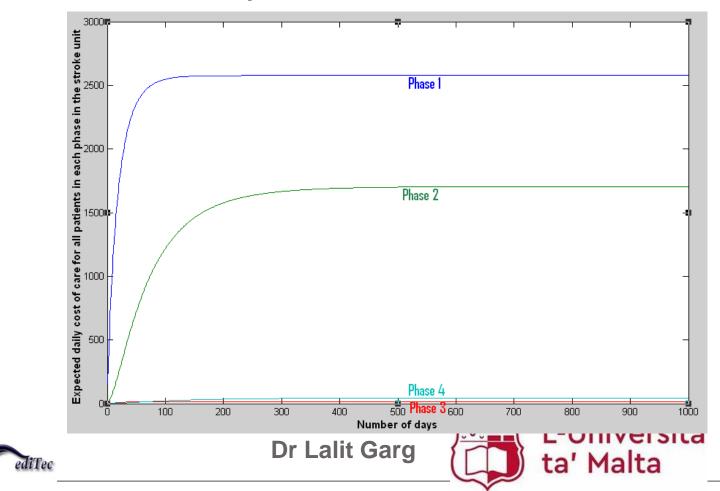




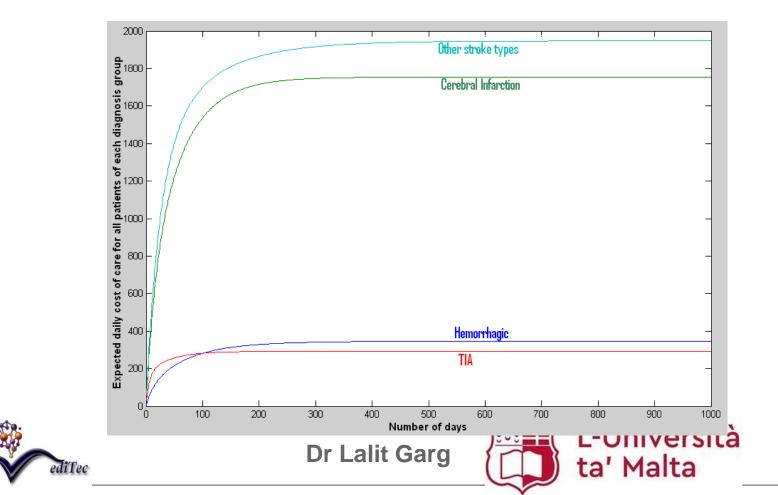




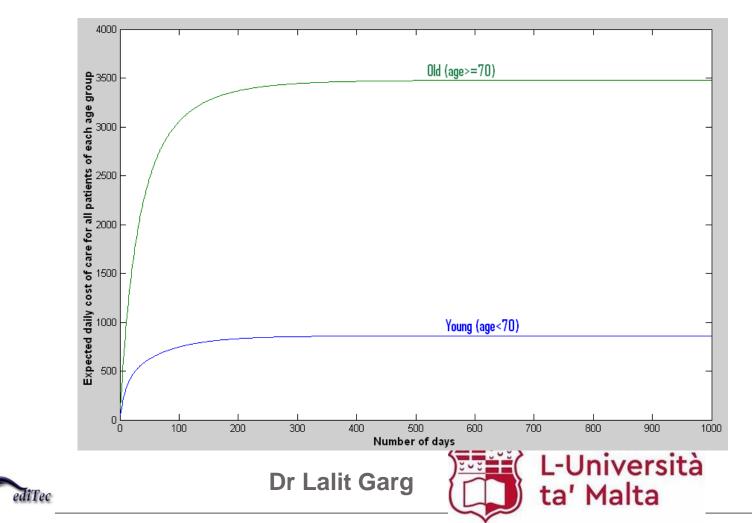
Expected daily cost of care (in £s) for all patients in each phase of the stroke unit



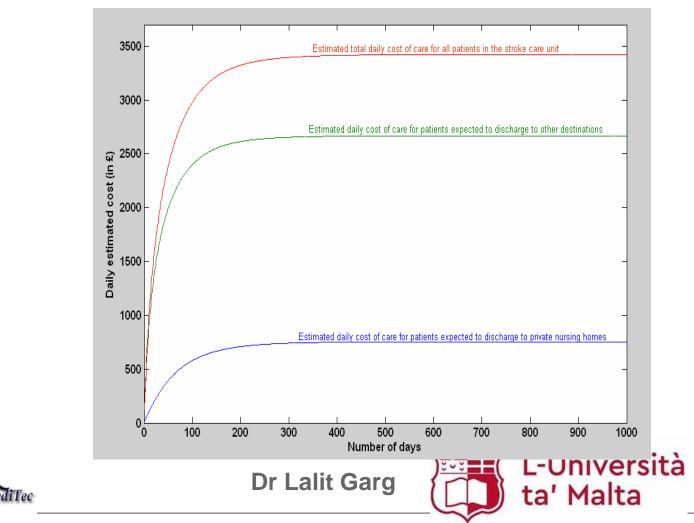
Expected daily cost of care (in £s) for all patients of each diagnosis group



Expected daily cost of care (in £s) for all patients of each age group



Estimated daily cost (in £s) of care of patients for each discharge destination



Opportunities unlimited

- Discharge delay modelling
- Activity mining in sensor network
- Disease progression modelling (HIV)
- Mental state detection (using fMRI and EEG neuroimages)
- Behavioral analysis





Discharge delay modelling

- A challenge for healthcare managers and policy makers
- Negatively affects the hospital performance metrics

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• Has other serious consequences for the healthcare system such as affecting patients' health

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Discharge delay modelling

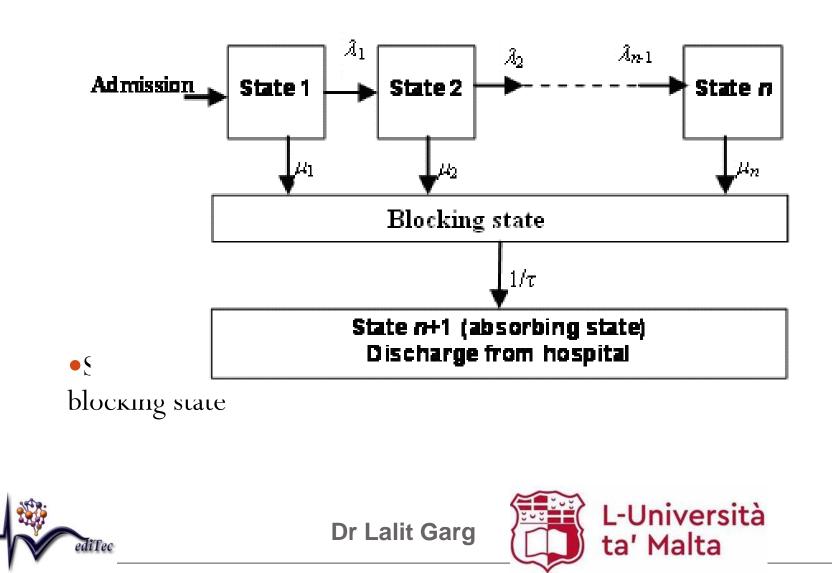
- Phase type distribution survival tree based clustering can be used for modelling delayed discharge and its effects
- Delayed discharge patients waiting for discharge can be modeled as a special state in the Markov chain called 'blocking state'
- A model can be developed to recognize association between demographic factors and discharge delays and its effects, and to identify groups of patients who require attention in order to resolve the most common delays and prevent them from happening again.

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Discharge delay modelling



- A sensor network is deployed in the smart home environment to aid assistive living in self care of patients with Dementia (Alzheimer's disease).
- Sensors in the sensor network are placed such that each sensor detects and records (time and duration of) a particular activity each time it is performed by the user.
- In order to carry out a task, a user performs combination of these activities in a particular sequence.

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- Activity mining or exception mining from the sensor network data might help in monitoring patient condition, recognising alarming events, determining care needs, treatment effects and progress etc.
- The time spent in each activity can separately be modelled by Coxian Phase Type Distribution.
- A sequential pattern can be defined as a sequence of activities followed by a user in order to perform a task.

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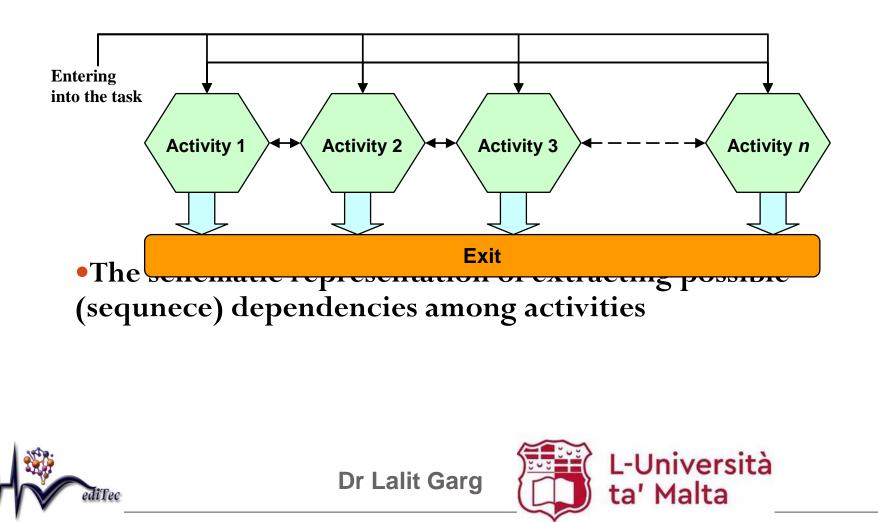


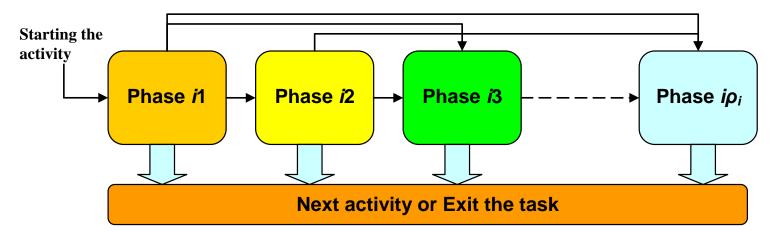
- The values of fitting parameters and dependency parameters can be estimated using semi-supervised learning.
- Frequent sequential patters satisfying given criteria of interestingness can be enumerated using an algorithm based on global optimization (Falk and Soland, 1969, Garg et al. 2009a, Lawler and Wood 1966) or some other suitable algorithm based on the given criteria.
- We can use the model to recognise unexpected patterns based on given criteria such as patterns having likelihood less than the given threshold or pattern with duration more than the given threshold duration.



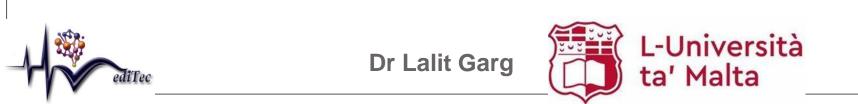


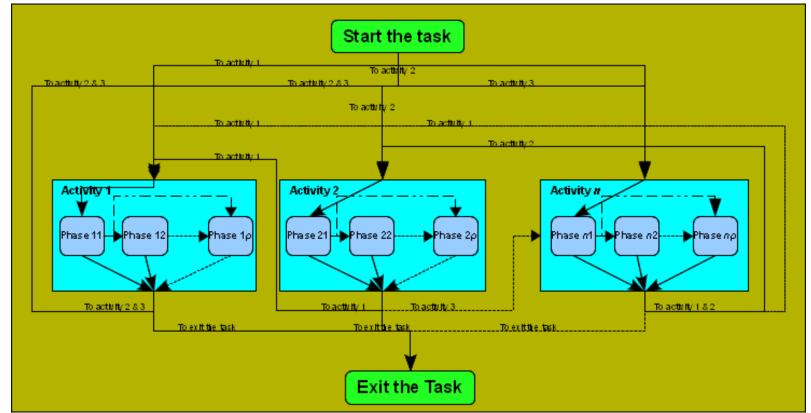






•The schematic representation of an activity as a Markov chain





separately modelled by Coxian-Phase type distribution.



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Disease progression modelling (HIV)

- Disease progression models can be useful tools for gaining a systems understanding of the transitions to disease states, clustering patients based on their disease progression rates and characterizing the relationship between disease progress and factors affecting it such as patients profile, treatment, stage at which disease was diagnosed or stage at which patient was first institutionalized.
- WHO classifies the progression of HIV disease as a 4 stage bidirectional process in which a patient's disease progression stage is determined by his/her absolute peripheral blood CD4+ T-lymphocyte count.





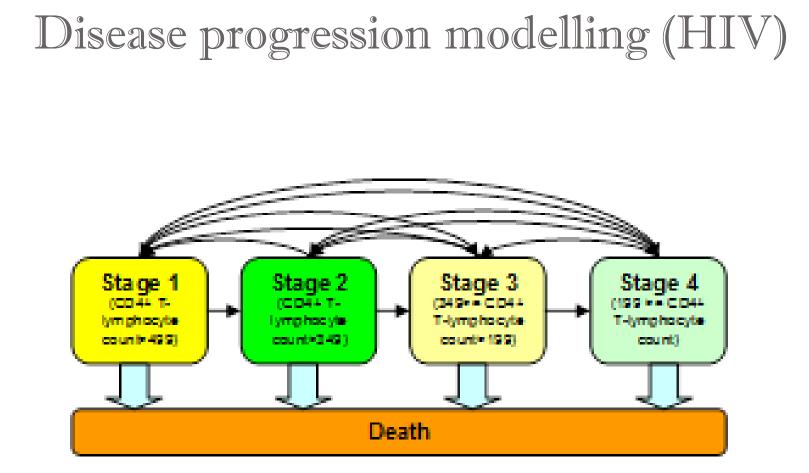
Disease progression modelling (HIV)

- The patient's immunological status can not only progress sequentially from stage 1 to stage 4 but also regress or jump from one stage to the another stage.
- We are developing a novel approach of modelling progression of HIV disease using phase type distributions.
- Model can then be extended to illustrate how it can be used to model effects of the affecting factors such as stage at which disease was diagnosed or stage at which the patient was first institutionalized.





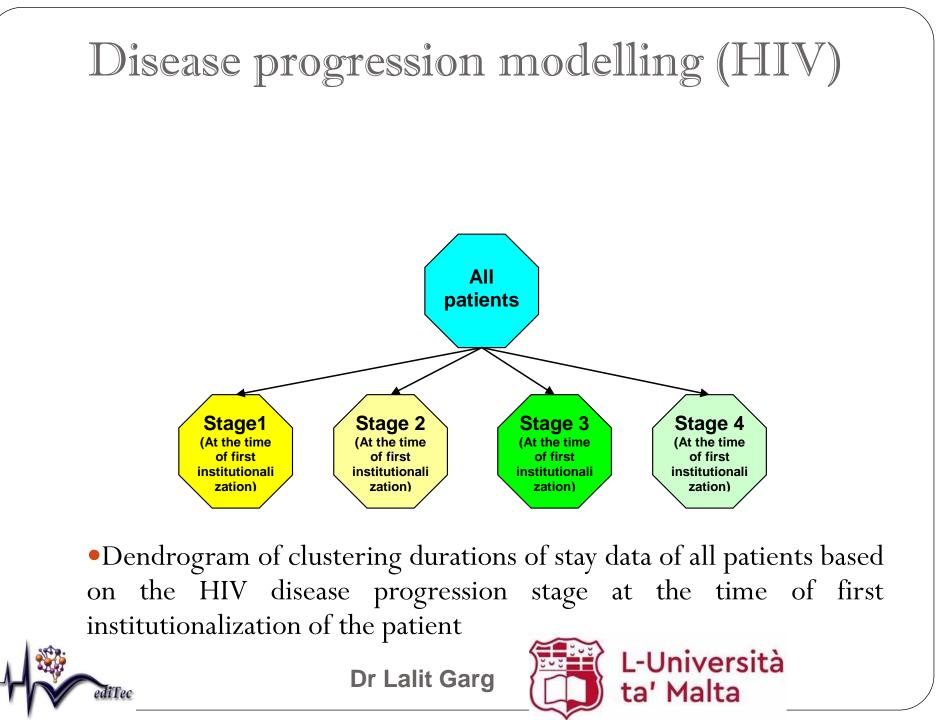




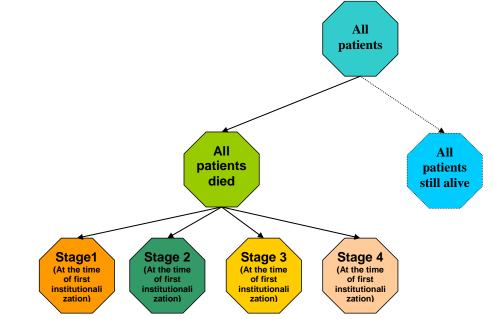
•Stages of HIV progression







Disease progression modelling (HIV)

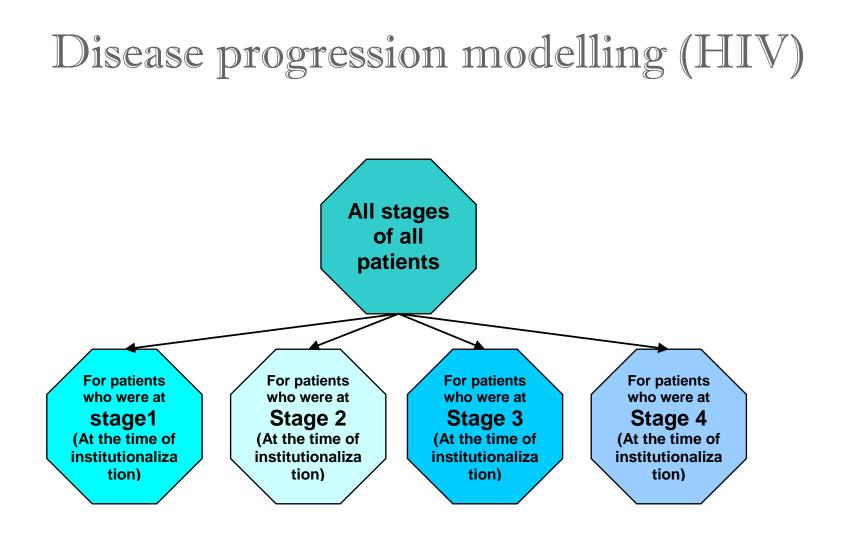


•Dendrogram of clustering durations of stay data of all died patients based on the HIV disease progression stage at the time of first institutionalization of the patient

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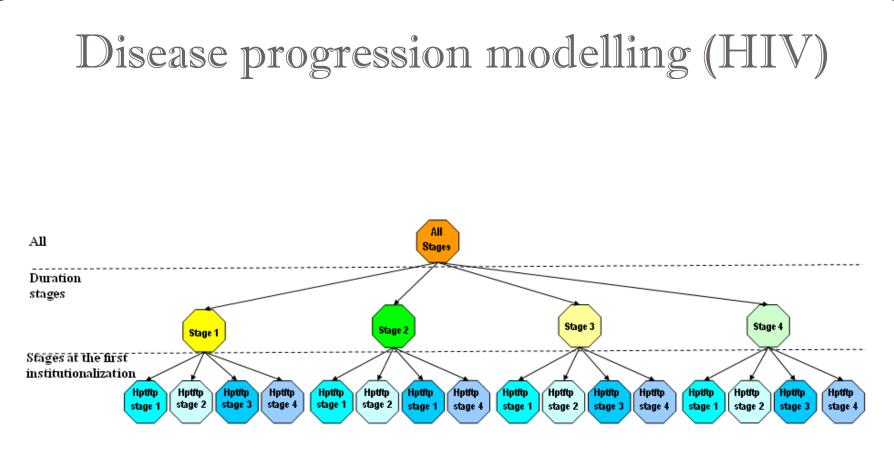




•Dendrogram of clustering durations of stay in each stage data of all died patients based on the HIV disease progression stage at the time of first institutionalization of the patient







•Dendrogram of clustering time spent in each stage data of all patients first based on the HIV progression stage in which time is spent and then each such cluster in further sub-clustered based on the HIV disease progression stage at the time of first institutionalization of the patient





- Mental state detection (using fMRI and EEG neuroimages)
- To identify the activation regions in fMRI data
- To model brain response to different activities using fMRI data





- Repeating these with EEG data
- Characterising similar and complimentary information in fMRI and EEG data.
- To develop an integrated model for mental state detection using both fMRI and EEG data.





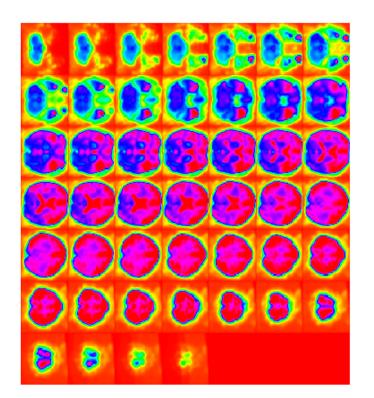
- fMRI provides Blood Oxygen Level Dependent (BOLD) responses in brain.
- Clustering (pattern recognition) whole fMRI data using Gaussian Mixture Distributions
- Developing Gaussian Mixture Distribution tree using covariates such as subjects, Time, Slices and Regions



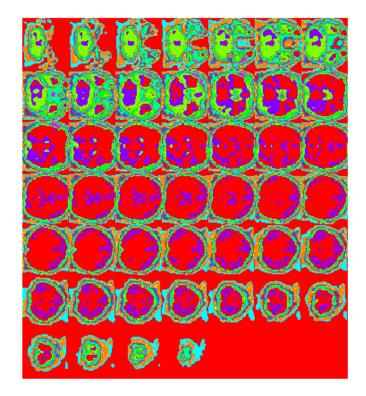
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Different slices of a single scan before clustering



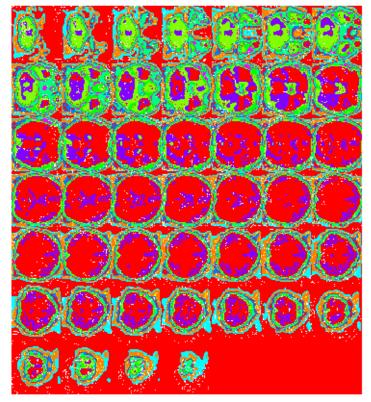
Different slices of a single scan after clustering



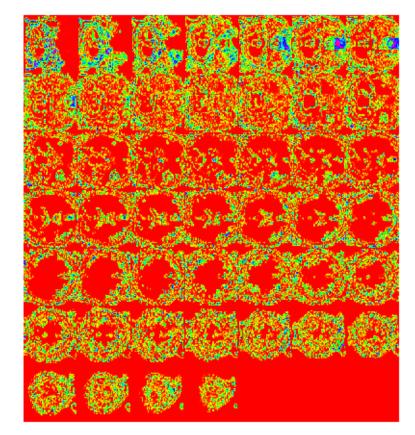
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Different slices of a single scan after clustering



Changes in time domain in different slices



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- Again Clustering whole fMRI data using Phase Type Distributions, which gives the phase changes in data on the time axis.
- Developing Phase Type Survival tree using covariates such as subjects, Slices and Regions





- Dirichlet Mixture Model based analysis of probability of shift in the given cluster.
- Complete brain model for changes in Blood Oxygen Level Dependent (BOLD) activity.

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- Dirichlet Mixture Model based analysis of probability of shift in the given cluster.
- Complete brain model for changes in Blood Oxygen Level Dependent (BOLD) activity.

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- We can use phase-type survival tree analysis to
 - Effectively prognosticate survival data and





- We can use phase-type survival tree analysis to
 - Effectively prognosticate survival data and

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• Cluster survival data into groups of patients following homogeneous patient pathways.

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• Our models can be used to forecast bed occupancy and the requirements.



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- Our models can be used to forecast bed occupancy and the requirements.
- The LOS can be predicted at admission by the use of this model.







- Our models can be used to forecast bed occupancy and the requirements.
- The LOS can be predicted at admission by the use of this model.
- The number of admissions can be forecasted by the patients' characteristics.





• These models can also be used to characterize the effect of weather on LOS and admissions.



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- These models can also be used to characterize the effect of weather on LOS and admissions.
- We can also use these models to predict effect of other factors affecting LOS and admissions.





• These forecasts can help us better designing policies to ensure optimal utilization of scarce health resources.





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