# **Project Tycho**

Preliminary data for the state of Alaska

## Tycho database beta test version

The data presented in this report are of preliminary nature and should not be used for publication or other types of official use

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University of Pittsburgh Graduate School of Public Health



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## **Contact information**

The Tycho database is currently being tested and a login account will be required to use the database and website (www.tycho.pitt.edu). All State Epidemiologists, CDC partners, other federal agencies and our research collaborators will be provided with user accounts. User accounts can also be provided upon specific request. Note that the quality of the data during the testing phase cannot be guaranteed to be sufficient for publication or official use.

#### Point of contact for the Tycho project:

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#### Introduction

**Project Tycho** started in 2009 as part of the Vaccine Modeling Initiative (VMI) at the University of Pittsburgh Graduate School of Public Health, funded by the Bill & Melinda Gates foundation. This project aims to contribute to the availability of public health data for research and policy making. The vision for

this project as described in the next section is a central access point for public health data of a detailed spatial and temporal scale for all countries. Currently, the Tycho database contains 10 million records that each represent a weekly report from a location for a specific disease. These records were extracted from weekly US Nationally Notifiable Disease Surveillance Reports between 1888 and 2009 (6300 reports) using double data entry (200 million keystrokes). The current database includes a total of 90 million reported cases and 4 million reported deaths due to notifiable diseases in the United States for the last 122 years.

Project **Tycho** is named after the Danish nobleman Tycho Brahe (1546 – 1601), who meticulously collected astronomical data. After Brahe's death, his assistant Kepler used these data to develop area laws of planetary motion.

#### **Background**

#### "No health department, State or local, can effectively prevent or control diseases without knowledge of when, where and under what conditions cases are occurring"

(Quote stated above weekly surveillance reports published in the Public Health Reports, 1888-1951)

This quote has accompanied weekly reports on the prevalence of notifiable diseases in the United States as long as these were published in the Public Health Reports, emphasizing the importance of data for disease control policy making. In the current day and age, this statement has lost none of its significance. On the contrary, as infectious diseases continue to pose challenges to the global public health system in an increasingly interconnected global society. The public health system has responded to this challenge by evermore sophisticated disease surveillance systems. The availability of internet and information technology has facilitated collection of detailed data on infectious disease incidence, the spread of pathogens, disease determinants, health behavior, etc. New technologies such as remote sensing and mobile reporting systems have advanced measurement of these factors in both high and low resource settings. The enormous wealth of public health information that is currently being accrued also poses its own challenges for data storage, management, preservation and dissemination. Public health agencies will need to allocate resources and develop capacity to specifically address these issues.

Although the importance of public health data collection has been widely recognized, the dissemination of data has been neglected for many reasons. The above quote does not only imply data collection but also its use and dissemination as knowledge on disease occurrence can only be derived from data after processing and analysis. Not only has disease surveillance become more sophisticated, analytical tools have changed as well and now include computational models to assess disease transmission and to evaluate disease control options. These methods have introduced the opportunity for data intensive studies of fine spatial-temporal patterns using dynamic models. Disease surveillance data at such scale are often not available for analysis for multiple reasons. First, surveillance data are not collected for research purposes but for disease monitoring and planning. Secondly, public health staff often lack time or capacity to prepare these data for analysis. Third, surveillance methodology often lacks transparency and standardization, reducing options for scientific analyses. Fourth, disease surveillance data are often

kept confidential and ownership or data use requirements are often unclear. Finally, archiving and preservation of surveillance data are often not a priority and disaggregated records may get lost, losing the opportunity to ever use these data for analysis.

The Tycho project aims at overcoming some of these challenges by providing open access to large public health datasets of great spatial-temporal detail. Open access to these data will provide great opportunities for scientific analysis and better informed policy making on disease control.

#### The Tycho database and website

As described in the methods section, a large heterogeneity in reporting practice was observed over the past 122 years of weekly disease surveillance and substantial efforts were made to clean and standardize these data. The current online database includes all cleaned and standardized data in a format that allows comparisons across time and space. It does not include parts of the data that require additional cleaning and research such as classification of disease subcategories.

Each record in the database provides a reported number of cases or deaths due to a specific disease for a specific location, for a certain week. The database can be searched online (www.tycho.pitt.edu) by selecting a dataset, an aggregation method and an output format. Data can be viewed in tables, graphics and maps and these can be downloaded for further use.

#### Data availability

The data provided by the Tycho database reflect the weekly US Nationally Notifiable Disease Surveillance System between 1888 and 2009. It includes all diseases that were reported as part of this system, but it does not yet include records from other surveillance systems such as the annual system or disease specific subsystem (such as the influenza surveillance system).

The available data also reflects changes in reporting practices over time. This explains why city or state level reports are only available for a certain time period (1888-1953 for cities and 1927-2009 for states) and why morbidity and mortality reports are available for different time periods. We are currently working on inclusion of additional data as described in the section on collaboration.

#### Testing and release of the Tycho database

The Tycho database is currently in a beta testing phase that will start during the 2011 annual meeting of the Council of State and Territorial Epidemiologists (CSTE) in Pittsburgh June 12-16. This phase will end with a release to the general public in the fall of 2011. During the testing phase, invited users will be provided with login accounts that will enable full use of all features of the website and database. Invited users will include all state epidemiologists, partners in the CDC and other federal agencies as well as research collaborators. During this phase, data cleaning and standardization will continue. The quality of the data will not be sufficient for publication or official use during this phase.

#### **Future developments**

After optimization of the database and website during the testing phase, the first version of the Tycho database will be released to the general public. After that, efforts to clean and standardize the data will continue and more data will be released in subsequent versions. In addition, ongoing and new collaborations for inclusion of new data will allow gradual expansion of the dataset over time and increased opportunities for data driven analysis and policy making.

## Vision and goal

The vision for project Tycho is based on the value of open access to detailed, disaggregated public health data for scientific analysis and policy making. Starting with 122 years of weekly US surveillance data, it is our hope that this example will convince public health authorities worldwide of the value of this resource and will stimulate similar contributions from other countries.

In the past, data sharing and archiving have not been a priority in public health or among health sciences in general. Continuous and diligent public health data collection has led to a large pool of disaggregated data scattered over the world including the internet, ranging from paper archives to basements of local health departments. The need for open access to a central repository of public health data has been recognized and both benefits and challenges of open access to data in general have been well described. Open access to data could lead to increased accountability, transparency, innovation, collaboration, cost-effectiveness, replication of results, the development of new methods and insights, and a narrower data access gap between low- and higher income countries. Ultimately, this would result in the advancement of science and technology to the benefit of all.

Challenges to open access include inconsistent formats, lack of annotation or metadata, lack of information technology (IT) capacity, a lack of incentive, and cultural norms around data ownership. Open access to public health data in particular is confronted by decentralized archiving of disease reports, lack of time and capacity for data management, degraded physical format of records and archives, and governmental level legal constraints. Multiple principles and requirements for a central open access repository for public health data have been described and include (1) capacity building for data management in low- and middle income countries, (2) feedback loops to data contributors, (3) sustainability, (4) common standards, (5) interoperability, and (6) user-friendliness.

The ultimate goal of the Tycho project is to provide a central global public health data access point. Historical as well as current public health data are of great value if archived and accessible for research and analysis. Open access will enable the use of analytical capacity from around the globe which will lead to new discoveries of disease patterns and control policies. Furthermore, a central data access point will facilitate data archiving and preservation into the future which will be an increasing need in a data rich public health environment.

## Collaboration

The application of the surveillance data from the Tycho database as well as the inclusion of new data will require a wide range of collaborations with public health offices in the US and abroad. The Tycho project has currently made the first step of digitizing 122 years of weekly US surveillance reports that had already been published. The next step will be the completion of the existing dataset as well as the inclusion of new data from unpublished sources.

This report provides an overview of data available from the Tycho database for the state of Alaska. This includes all data that has ever been published at state or city level for Alaska in the weekly US Nationally Notifiable Disease Surveillance System. As you will find, the data availability will vary greatly over time, between city and state level reports, between diseases and between morbidity vs. mortality reports.

The Tycho development team would look forward to work with state health departments in the US to:

- 1. Better understand the current Tycho data for each state (eg. by collecting historic documentation)
- 2. Provide better usability and applications of current Tycho data for each state by continued joint development of website and database features.
- 3. Collect and/or digitize official, confirmed data to validate the current (preliminary) weekly data.
- 4. For each state, collect and/or digitize new data that has not yet been included. For example all city level reports discontinued in 1953 and completion of city level data until 2009 would greatly increase opportunities for analysis of disease patterns and trends.
- 5. Provide support to state health offices to manage, preserve and provide access to public health data.

## **Summary of methods**

Detailed documentation on the methods used for the Tycho database has been provided on the website (www.tycho.pitt.edu). This section describes these methods in short.

#### **Data collection**

Weekly reports that contain tables on the occurrence of nationally notifiable diseases have been published since 1888 by public health authorities at the Federal level in various journals. Table 1 provides the list of publications and the responsible agency since 1888. All weekly nationally notifiable disease reports between 1888 and 1951 could be retrieved from the PubMed Central repository of the National Library of Medicine <sup>1</sup>. For 1995 to 2009, these reports could be retrieved from the MMWR digital archive on the CDC website <sup>2</sup>. Most weekly reports between 1952 and 1995 could be retrieved from the HathiTrust Digitial Library <sup>3 4</sup>, but many could not be found and had to be copied from hard copies of MMWR issues in the University of Pittsburgh library.

Table 1, publications and responsible Federal agencies for nationally notifiable disease reports

Time period	Publication title	Responsible federal agency
1888-1889	Weekly Abstract of Sanitary Reports	US Marine Hospital Service
1890-1895	Abstract of Sanitary Reports	US Marine Hospital Service
1896-1901	Public Health Reports	US Marine Hospital Service
1902-1911	Public Health Reports	US Public Health and Marine Hospital
		Service
1912-1951	Public Health Reports	US Public Health Service
1952-1960	Morbidity and Mortality Weekly Report	National Office of Vital Statistics, US Public
		Health Service
1961-1969	Morbidity and Mortality Weekly Report	Communicable Disease Center
1970-1991	Morbidity and Mortality Weekly Report	Center for Disease Control
1992-2009	Morbidity and Mortality Weekly Report	Centers for Disease Control and Prevention

#### Inclusion criteria

Weekly reports of each year were reviewed systematically to assess the diseases reported. We included all tables that provided disease specific information by week for US cities, townships, counties or states. Tables that provided summary or aggregated information by month, year or at the national level were not included. Similarly, tables that did not contain disease specific information (such as all cause mortality) were not included.

#### Data entry

Weekly reports were downloaded or scanned as PDF files and selected tables with notifiable disease reports were entered into computer spreadsheets in a highly standardized fashion using double data entry. During the second round of data entry, operators could not see what had been entered in the first round and could not continue if the system detected a discrepancy between the second and first entry

<sup>&</sup>lt;sup>1</sup> <u>http://www.pubmedcentral.nih.gov/tocrender.fcgi?journal=333&action=archive</u>

<sup>&</sup>lt;sup>2</sup> <u>http://www.cdc.gov/mmwr</u>

<sup>&</sup>lt;sup>3</sup> <u>http://catalog.hathitrust.org/Record/003910026</u>

<sup>&</sup>lt;sup>4</sup> <u>http://catalog.hathitrust.org/Record/003843660</u>

for a specific value. Such discrepancies could only be resolved by checking the PDF file and try again or discussion with the group leader and verification of a value in the source documents.

#### Quality control for data entry

The accuracy of data entry was checked at various levels. First, completeness of data was verified by comparing the content of entered data with PDF sources files. Secondly, accuracy of data entry was verified by multiple rounds of comparing random samples of entered files with PDF source files. Thirdly, data formatting was verified by various checks to ensure appropriate formatting for data loading.

#### Data loading and standardization

All data was entered in Excel spreadsheets and various components of these spreadsheets were loaded in data files. Table titles, column headers, place names and reported numbers were loaded in separate files. These files were used to extract information on each reported number, including:

- 1. the disease reported
- 2. the disease subcategory reported
- 3. cases or deaths reported
- 4. the reporting location (name, state and type of location)
- 5. the time period for which a number was reported
- 6. the date of publication of the original weekly report associated with a reported number

#### Integration

All reported numbers and extracted information was integrated in one database with one record per reported number and associated information.

#### Post-processing quality control

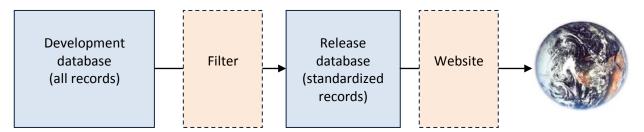
After integration of all data in one database, checks were performed to detect duplicate reports and data inconsistencies. Duplicate records were removed and inconsistencies resolved by verification with original PDF source files.

#### Data filtering

The digitized version of all historical weekly US nationally notifiable disease surveillance records is a very heterogeneous dataset. It took a substantial number of data processing protocols to standardize time and space variables as well as disease names. All standardized records have been separated from non-standard records by a filter. All remaining heterogeneity in non-standard records is inherent to the surveillance system and can only be standardized after further analysis will have been completed (eg. remaining heterogeneity in reporting periods for reports before 1953 and standardization of disease subcategories). The largest proportion of data (>4 million records) has been standardized however and has been made available in the current testing version.

#### **Data visualization**

Figures in this reports were made with the R system, version 2.9.2 and the maps were generated by the GAIA platform developed at the University of Pittsburgh Graduate School of Public Health in collaboration with the Pittsburgh Supercomputing Center (PSC). See http://midas-pitt.psc.edu/gaia for more information.



**Figure 1, Schematic of the Tycho data architecture.** Access to a consistent, standardized subset of the digitized US weekly surveillance reports will be provided for beta testing through a website that will allow querying and downloading of data. Data from the development database will be pushed through a filter to feed the Tycho database.

The current Tycho database will provide the reported number of cases or deaths reported by city or state health authorities to the federal health agency for all weeks between 1888 and 2009. Currently, only standardized, consistent data are being released for testing, according to the following criteria:

- 1. Only weekly reports are provided (reports for other periods such as 10 days, 2 weeks, 1 month, 1 year) are currently not included due to lack of comparability with the weekly reports.
- 2. Only reports that were published in the same year as the reporting period were included (this excludes updates or comparisons that were published more than a year after the original reporting period)
- 3. Only numbers were included for which information about the location, time period and disease could be extracted.
- 4. Only numbers or time periods for which no disease subcategories were reported were included.

Each of the steps described above have been described in detailed protocols that are posted in the documentation part of the Tycho website (www.tycho.pitt.edu).

#### Website

The Tycho website features simple and advanced searches of the data, visualizations and downloading of machine readable files. A dataset can be selected by specifying: 1) the disease of interest, 2) the location(s) of interest and 3) the time period. In a second step, the type of aggregation and length of increments can be specified and in a third step, the type of output (table, graphic or map) can be selected. After output has been generated, data can be downloaded.

As mentioned above, data availability varies largely by disease, time period and location. We aimed to restrict selection options by data availability to avoid "no results available.

Access to the most of the website will be password protected during the testing phase. For invited users after login, full functionality of the database and website will be available. During this phase however, the data quality will not be sufficient for publication or official use.

## Overview of data available for the state of Alaska

In this section, an overview of data available for the state of Alaska will be provided. As described earlier, data availability depends entirely on historical reporting practices in the weekly US Nationally Notifiable Disease Surveillance System between 1888 and 2009. For this report, we only included morbidity reports (cases) to improve the format and limit the size of this report. As mentioned earlier, the analysis and standardization of has not been completed yet for all records in the database and preliminary data are provided here that may not yet be available in the online database.

This section will provide a general overview of data availability. The next sections will provide disease specific data from state and city level reports of Alaska. Data for a maximum of 3 major cities were provided depending on availability (city data was only reported until 1953). Some summary data will be provided at the end of this report, for a subset of diseases for which data was fragmented over time.

Table 1 lists the number of weekly state or city reports that are available for Alaska per each disease and subcategory.

Disease	City	State
Aids	-	896
Brucellosis [undulant fever]	-	96
Chickenpox [varicella]	-	567
Chlamydia	-	663
Cryptosporidiosis	-	200
Diphtheria	-	442
Encephalitis		
Post infectious	-	7
Primary [infectious] including unspecified	-	508
Escherichia coli		
EHEC 0157	-	149
O157:H7 NETSS	-	186
O157:H7 PHLIS	-	86
STEC	-	38
Giardiasis	-	386
Gonorrhea		
Civilian	-	742
Unspecified	-	1054
Haemophilus influenzae		
Age <5 non-serotype B	-	40
Age <5 unknown serotype	-	67
All ages all serotypes	-	568

Disease	City	State
Hepatitis		
Acute type A	-	302
Acute type B	-	345
Acute type C	-	30
All types, <20 years	-	143
All types, >=20 years	-	60
All types, all ages	-	567
Type A [infectious]	-	1229
Туре В [serum]	-	1008
Type NA NB [including C]	-	406
Type unspecified	-	282
Influenza	-	23
Legionellosis	-	206
Leprosy	-	71
Lyme disease	-	342
Malaria		
Civilian	-	1
Military	-	1
Unspecified	-	1085
Measles		
Imported	-	166
Indigenous	-	242
Unspecified	1	1210
Meningitis		
Aseptic	-	536
Meningococcus	-	50
Meningococcal disease		
All serogroups	-	37
Invasive all serogroups	-	165
Invasive serogroup unknown	-	35
Serogroup unspecified	-	1798
Mumps	-	1535
Pneumonia	1	26
Poliomyelitis		
Non paralytic	-	48
Paralytic	-	111
Total	-	267
Rabies in animals	-	1755
Rocky mountain spotted fever	-	28
Rubella	_	781

			,
Table 1, Number of weekly	y state or city reports	per disease and subcategory	i, continued

Disease	City	State
Salmonellosis		
NETSS	-	125
PHLIS	-	104
Unspecified	-	375
Scarlet fever		
Including streptococcal sore throat	-	421
Unspecified	-	69
Shigellosis		
NETSS	-	103
PHLIS	-	82
Unspecified	-	326
Streptococcal disease, invasive group a	-	147
Streptococcal sore throat	-	93
Streptococcus pneumoniae invasive disease	-	94
Syphilis		
Civilian primary and secondary	-	610
Primary and secondary	-	566
Tetanus	-	1
Trichiniasis	-	2
Tuberculosis [phthisis pulmonalis]		
New active	-	96
Unspecified	2	1386
Tularemia	-	264
Typhoid fever [enteric fever]		
Including paratyphoid fever	-	7
Unspecified	-	448
Whooping cough [pertussis]	-	1025

The diseases included in the weekly US Nationally Notifiable Disease Surveillance System varied largely over time and reflected the historical social-political priorities of each time period. Note that the diseases in the weekly system were a subset of all diseases included in the annual Notifiable Disease Surveillance System (that was not entered as part of this project).

The number of locations reporting diseases also varied over time depending on the type of reports submitted. Figure 1 on the next page provides an overview of the weeks for which a report was available per disease at the state or city level. Colors indicate the total number of reporting locations. Figure 2 below displays the number of diseases for which a weekly report was included at the state state level per week between 1888 and 2009.

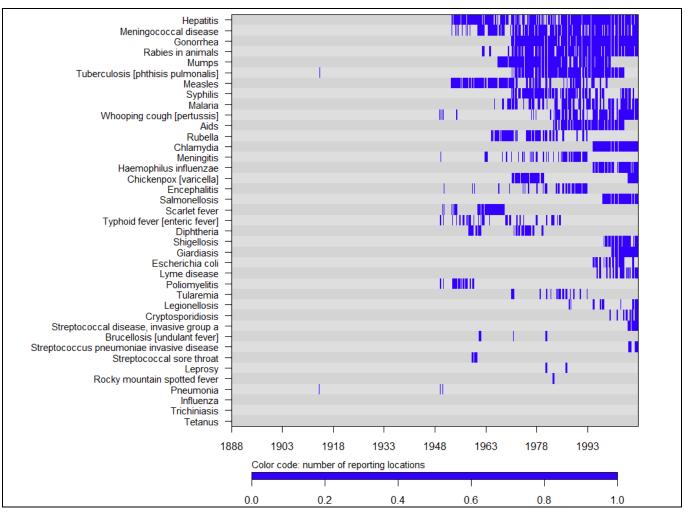
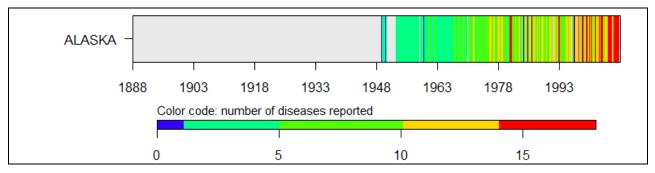


Figure 1, The number of locations (in color code) reporting on each disease per week between 1888 and 2009



**Figure 2,** The number of diseases (in color code) that were reported for the state and city level (state on top row) for each week between 1888 and 2009

## Disease specific data for Alaska

AIDS												
l la cara cifica d												
Unspecified -		4000	4000	4000	4000	4004	4000	4000			0004	_
	1984	1986	1988	1990	1992	1994	1996	1998	2000	2002	2004	

Table D1, Summary information for AIDS					
Indicator	Alaska				
Report period	1984-2005				
Total weeks	896				
Total cases	527				
Max. cases per year	93				
Year (max)	1993				
Max. cases per week	37				
Week (max)	1993, wk 47				
Average cases per year	24				
95%CI	(14-34)				
Average cases per week	1				
95%CI	(1-1)				

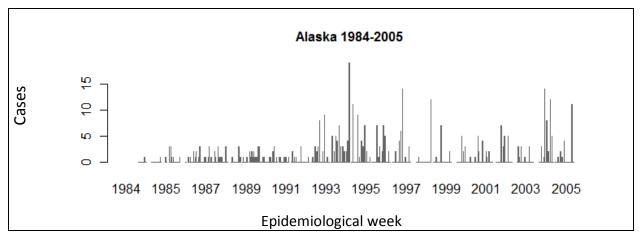


Figure D2, Number of cases reported for AIDS per epidemiological week

Chlamydia							
Unspecified							
	1996	1998	2000	2002	2004	2006	2008

Table D1, Summary information for Chlamydia					
Indicator	Alaska				
Report period	1996-2009				
Total weeks	663				
Total cases	42,878				
Max. cases per year	6,059				
Year (max)	2009				
Max. cases per week	2,588				
Week (max)	2009, wk 29				
Average cases per year	3,063				
95%CI	(2,286-3,840)				
Average cases per week	65				
95%CI	(56-74)				

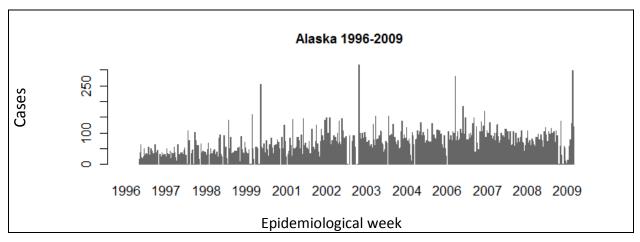


Figure D2, Number of cases reported for Chlamydia per epidemiological week

Giardiasis									
Unspecified									
·	2002	2003	2004	2005	2006	2007	2008	2009	

Table D1, Summary informa	Table D1, Summary information for Giardiasis							
Indicator	Alaska							
Report period	2002-2009							
Total weeks	386							
Total cases	3,518							
Max. cases per year	2,869							
Year (max)	2006							
Max. cases per week	2,784							
Week (max)	2006, wk 35							
Average cases per year	440							
95%CI	(-381-1,261)							
Average cases per week	9							
95%CI	(-5-23)							

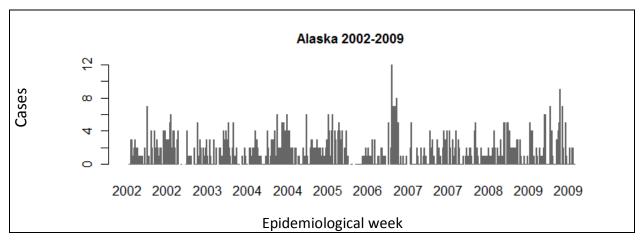


Figure D2, Number of cases reported for Giardiasis per epidemiological week

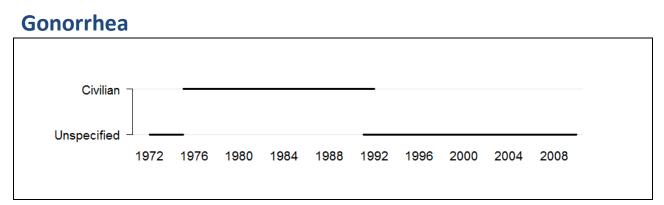


Table D1, Summary information for Gonorrhea (Unspecified and Civilian)

Indicator	Alaska
Report period	1972-2009
Total weeks	1,796
Total cases	70,677
Max. cases per year	5,742
Year (max)	1986
Max. cases per week	4,077
Week (max)	1986, wk 03
Average cases per year	1,860
95%CI	(1,300-2,420)
Average cases per week	39
95%CI	(34-44)

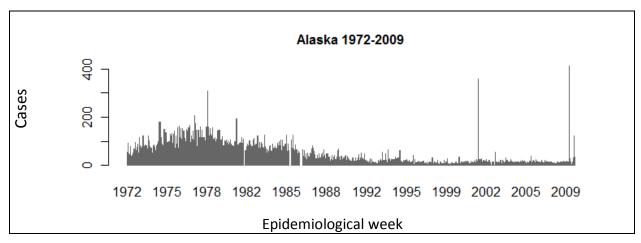
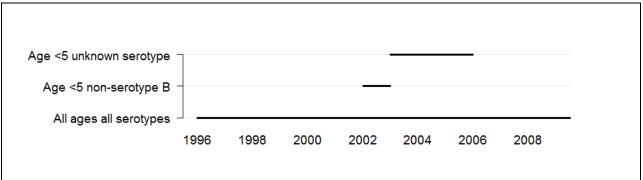


Figure D2, Number of cases reported for Gonorrhea per epidemiological week

## Haemophilus Influenzae



**Table D1,** Summary information for Haemophilus Influenzae (All ages all serotypes)

Indicator	Alaska
Report period	1996-2009
Total weeks	568
Total cases	165
Max. cases per year	44
Year (max)	2000
Max. cases per week	35
Week (max)	2000, wk 44
Average cases per year	12
95%CI	(5-19)
Average cases per week	0
95%CI	(0-0)

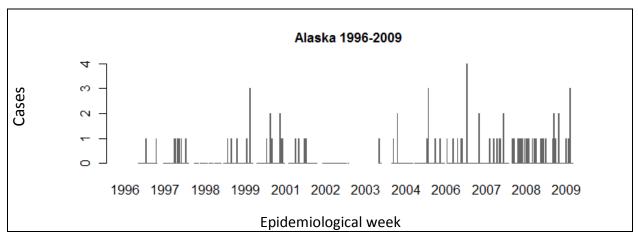
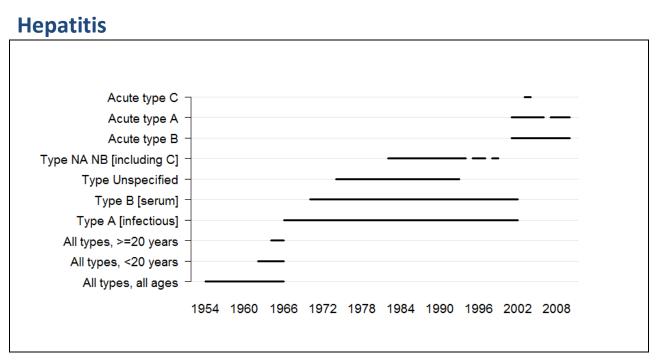


Figure D2, Number of cases reported for Haemophilus Influenzae per epidemiological week



**Figure D1,** Weeks between 1888 and 2009 for which data on the disease and subcategories (if applicable) are available in the Tycho database. If no subcategory was reported, a subcategory of "Unspecified" was assigned.

Indicator	Alaska
Report period	1966-2009
Total weeks	1,744
Total cases	5,051
Max. cases per year	572
Year (max)	1976
Max. cases per week	198
Week (max)	1970, wk 44
Average cases per year	
before 1990	171
95%Cl	(114-228)
after 1990	41
95%CI	(-2-84)
Average cases per week	
before 1990	5
95%CI	(4-6)
after 1990	1
95%CI	(1-1)

**Table D1,** Summary information for Hepatitis (Type A [Infectious], Acute type A, Type B [Serum], and Acute type B)

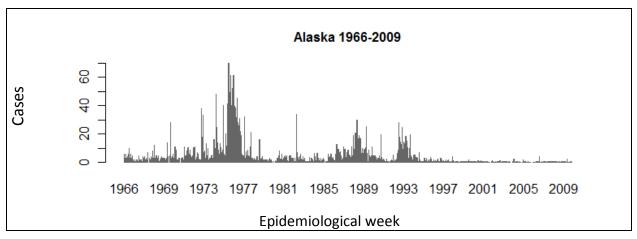


Figure D2, Number of cases reported for Hepatitis per epidemiological week

Lyme Dise	ase						
Unspecified							
·	1997	1999	2001	2003	2005	2007	2009

Table D1, Summary informati	on for Lyme Disease
Indicator	Alaska
Report period	1997-2009
Total weeks	342
Total cases	27
Max. cases per year	9
Year (max)	2007
Max. cases per week	2
Week (max)	2007, wk 43
Average cases per year	2
95%CI	(0-4)
Average cases per week	0
95%CI	(0-0)

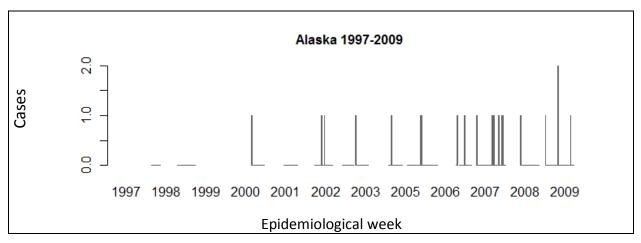
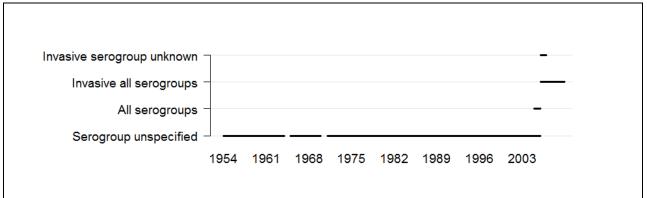


Figure D2, Number of cases reported for Lyme Disease per epidemiological week

## **Meningococcal Disease**



**Figure D1,** Weeks between 1888 and 2009 for which data on the disease and subcategories (if applicable) are available in the Tycho database. If no subcategory was reported, a subcategory of "Unspecified" was assigned.

 Table D1, Summary information for Meningococcal Disease (Serogroup unspecified and Invasive all serogroups)

serogroups)									
Indicator	Alaska								
Report period	1954-2009								
Total weeks	1,963								
Total cases	518								
Max. cases per year	34								
Year (max)	1977								
Max. cases per week	7								
Week (max)	1959, wk 34								
Average cases per year									
before 1980	12								
95%CI	(9-15)								
after 1980	7								
95%CI	(5-9)								

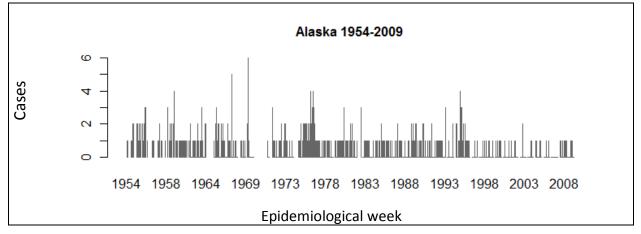


Figure D2, Number of cases reported for Meningococcal Disease per epidemiological week

Mumps										
Unspecified										
	1968	1972	1976	1980	1984	1988	1992	1996	2000	

Indicator	Alaska
Report period	1968-2001
Total weeks	1,535
Total cases	3,635
Max. cases per year	923
Year (max)	1969
Max. cases per week	100
Week (max)	1969, wk 21
Average cases per year	
before 1980	266
95%CI	(61-471)
after 1980	9
95%CI	(6-12)
Average cases per week	
before 1980	5
95%CI	(4-6)

Table D1, Summary information for Mumps

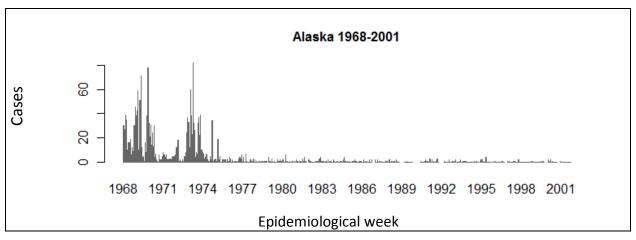


Figure D2, Number of cases reported for Mumps per epidemiological week

## **Rabies in Animals**

Unspecified -	 								
		1970	1975	1980	1985	1990	1995	2000	2005

**Figure D1,** Weeks between 1888 and 2009 for which data on the disease and subcategories (if applicable) are available in the Tycho database. If no subcategory was reported, a subcategory of "Unspecified" was assigned.

Table D1. Summary information for Rabies in Animals

Indicator	Alaska
Report period	1960-2009
Total weeks	1,755
Total cases	895
Max. cases per year	85
Year (max)	1977
Max. cases per week	50
Week (max)	1977, wk 42
Average cases per year	20
95%CI	(14-26)
Average cases per week	1
95%CI	(1-1)

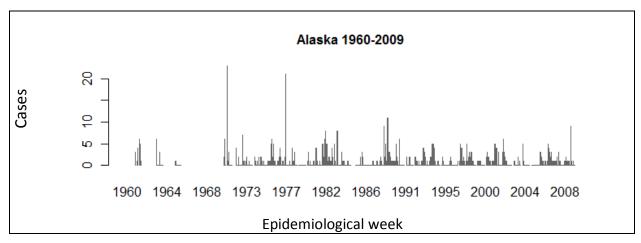


Figure D2, Number of cases reported for Rabies in Animals per epidemiological week

Rubella										
Unspecified										
Unspeched										
	1966	1969	1972	1975	1978	1981	1984	1987	1990	1993

Indicator	Alaska		
Report period	1966-1994		
Total weeks	781		
Total cases	1,407		
Max. cases per year	467		
Year (max)	1969		
Max. cases per week	91		
Week (max)	1969, wk 21		
Average cases per year			
before 1970	252		
95%CI	(54-450)		
after 1970	8		
95%CI	(2-14)		
Average cases per week			
before 1970	6		
95%CI	(5-7)		

Table D1, Summary information for Rubella

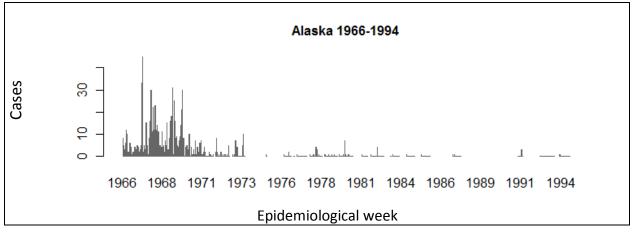


Figure D2, Number of cases reported for Rubella per epidemiological week

# Salmonellosis

Table D1, Summary information for Salmonellosis (PHLIS and Unspecified)

inary information for Samonenosis (Philis an					
Indicator	Alaska				
Report period	1999-2009				
Total weeks	479				
Total cases	641				
Max. cases per year	94				
Year (max)	2003				
Max. cases per week	41				
Week (max)	2009, wk 29				
Average cases per year	58				
95%CI	(40-76)				
Average cases per week	1				
95%CI	(1-1)				

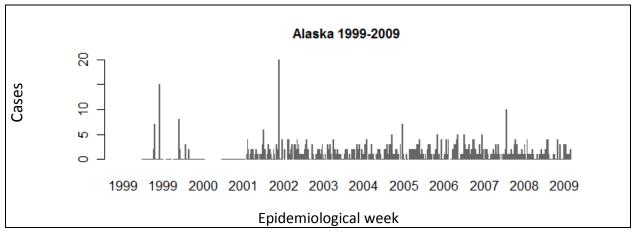


Figure D2, Number of cases reported for Salmonellosis per epidemiological week

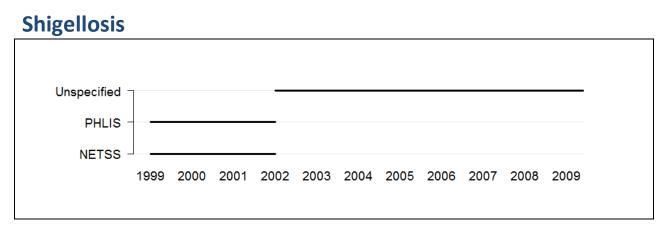


Table D1, Summary information for Shigellosis (Unspecified and PHLIS)

Indicator	Alaska
Report period	1999-2009
Total weeks	408
Total cases	53
Max. cases per year	9
Year (max)	2005
Max. cases per week	5
Week (max)	2001, wk 41
Average cases per year	5
95%CI	(3-7)
Average cases per week	0
95%CI	(0-0)

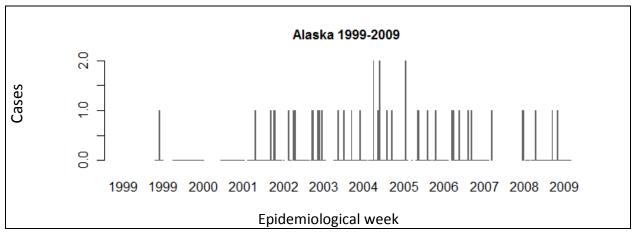


Figure D2, Number of cases reported for Shigellosis per epidemiological week

## **Streptococcal Sore Throat**

Unspecified	1960		1961

**Figure D1,** Weeks between 1888 and 2009 for which data on the disease and subcategories (if applicable) are available in the Tycho database. If no subcategory was reported, a subcategory of "Unspecified" was assigned.

Indicator	Alaska
Report period	1960-1961
Total weeks	93
Total cases	3,747
Max. cases per year	2,021
Year (max)	1960
Max. cases per week	998
Week (max)	1960, wk 50
Average cases per year	1,874
95%CI	(0-3,748)
Average cases per week	40
95%CI	(19-61)

#### Table D1, Summary information for Streptococcal Sore Throat

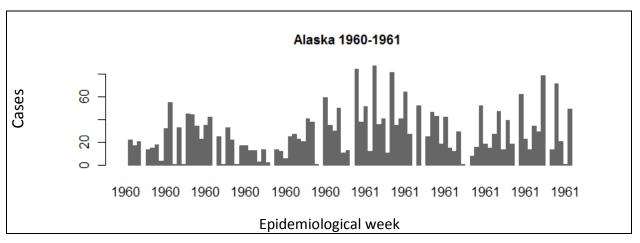


Figure D2, Number of cases reported for Streptococcal Sore Throat per epidemiological week

## **Syphilis**

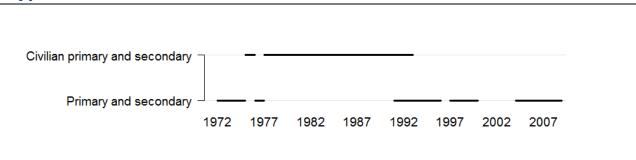


Table D1, Summary information for Syphilis (Primary and secondary and Civilian primary and secondary)

Indicator	Alaska
Report period	1972-2008
Total weeks	1,176
Total cases	440
Max. cases per year	81
Year (max)	1972
Max. cases per week	23
Week (max)	1980, wk 43
Average cases per year	14
95%CI	(8-20)
Average cases per week	0
95%CI	(0-0)

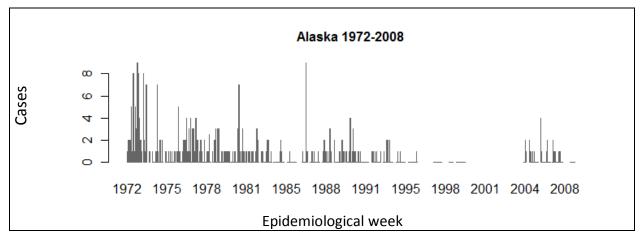
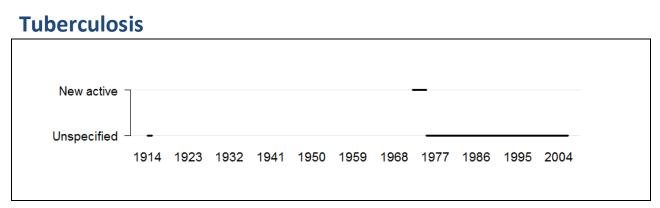


Figure D2, Number of cases reported for Syphilis per epidemiological week



**Figure D1,** Weeks between 1888 and 2009 for which data on the disease and subcategories (if applicable) are available in the Tycho database. If no subcategory was reported, a subcategory of "Unspecified" was assigned.

Indicator	Alaska
Report period	1975-2005
Total weeks	1,386
Total cases	2,093
Max. cases per year	136

1991

80

1991, wk 29

68

(60-76) 2

(2-2)

Year (max)

Week (max)

95%CI

95%CI

Max. cases per week

Average cases per year

Average cases per week

**Table D1,** Summary information for Tuberculosis (Unspecified)

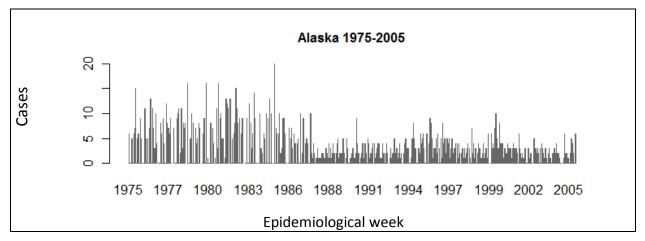


Figure D2, Number of cases reported for Tuberculosis per epidemiological week

#### Diseases with fragmented data over time

For a number of diseases, only fragmented data was available for Alaska. This was due to the inclusion of these diseases in the weekly surveillance system for short periods or widely dispersed periods of time. Due to these wide gaps or limited data, data for these diseases were not presented in separate sections but in the table below. Work will continue to include more data in the Tycho database and complete data sets for each disease where possible. Collaborations with local, state and federal public health agencies will be made to explore data availability and opportunities to include these in the Tycho database.

Table 2, Summary information on the occurrence of diseases with fragmented data in Alaska					
Disease	Report	Report	Number of	Total	
	type	period	reports	cases	
Brucellosis [undulant fever]	State	1958-1982	96	164	
Chickenpox [varicella]	State	1972-2009	567	4453	
Cryptosporidiosis	State	2001-2009	200	19	
Diphtheria	State	1950-1981	442	125	
Encephalitis	State	1950-1994	508	180	
Escherichia coli	State	1996-2008	273	30	
Influenza	State	1950-1951	23	522	
Legionellosis	State	1985-2009	206	8	
Leprosy	State	1982-1988	71	0	
Malaria	State	1967-2009	1085	109	
Measles	City	1914-1914	1	1	
Measles	State	1954-2000	1210	16898	
Meningitis	State	1950-1964	50	13	
Pneumonia	City	1914-1914	1	1	
Pneumonia	State	1950-1951	26	61	
Poliomyelitis	State	1950-1960	267	531	
Rocky mountain spotted fever	State	1983-1984	28	0	
Scarlet fever	State	1950-1969	490	15748	
Streptococcal disease, invasive group a	State	2004-2009	147	123	
Tetanus	State	1972-1972	1	1	
Trichiniasis	State	1954-1955	2	15	
Tularemia	State	1950-1994	264	12	
Typhoid fever [enteric fever]	State	1950-1988	455	44	
Whooping cough [pertussis]	State	1950-2009	1025	1056	

## **Project Tycho**

This report provides preliminary data for the state of Alaska available in the Tycho database. This database is currently being beta tested and these data cannot be used for publication or other official use at this time. An open access release to the general public is planned for later in 2011.

Please visit the Tycho website for more information and to query the database at: www.tycho.pitt.edu. For further information regarding the Tycho project, contact Dr. Wilbert van Panhuis at the University of Pittsburgh Graduate School of Public Health.

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