

**New York City Department of Health & Mental Hygiene
Bureau of Communicable Disease
&**

**New York City Department of Environmental Protection
Bureau of Water Supply**

**Waterborne Disease Risk Assessment Program
2019 Annual Report**

March 2020

*Prepared in accordance with Section 8.1 of the NYSDOH
2017 NYC Filtration Avoidance Determination*



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LIST OF ACRONYMS

| Acronym | Description |
|----------------|---|
| ADM | Anti-diarrheal medication |
| BCD | Bureau of Communicable Disease |
| CGAP | <i>Cryptosporidium</i> and <i>Giardia</i> Action Plan |
| CIDT | Culture independent diagnostic test |
| CUSUM | Cumulative sums |
| DEP | Department of Environmental Protection |
| DOHMH | Department of Health and Mental Hygiene |
| ED | Emergency Department |
| GI | Gastrointestinal |
| NYC | New York City |
| NYSDOH | New York State Department of Health |
| O&P | Ova and parasite test |
| OTC | Over the counter medication |
| PCR | Polymerase chain reaction |
| UHF | United Hospital Fund |
| WDRAP | Waterborne Disease Risk Assessment Program |

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EXECUTIVE SUMMARY

NYC's Waterborne Disease Risk Assessment Program (WDRAP) helps assure the microbial safety of the municipal water supply, and it is a component of NYC's Filtration Avoidance Program. The primary objectives of WDRAP are to: (a) obtain data on the rates of giardiasis and cryptosporidiosis, along with demographic and risk factor information on patients; and (b) provide a system to track gastrointestinal illness (diarrhea or vomiting) to ensure rapid detection of any outbreaks. The program began in 1993, and is jointly administered by two NYC agencies, the Department of Health and Mental Hygiene (DOHMH) and the Department of Environmental Protection (DEP). This report provides an overview of program activities and data collected during 2019.

DISEASE SURVEILLANCE

Active disease surveillance for giardiasis and cryptosporidiosis began in July 1993 and November 1994, respectively, and continued through 2010 when it was replaced by an electronic reporting system. This report presents the number of cases and case rates for giardiasis and cryptosporidiosis in 2019 and includes data from past years for context. Demographic information for cases of giardiasis and cryptosporidiosis diagnosed in 2019 is also summarized in this report. Telephone interviews of cryptosporidiosis patients were conducted to gather potential risk exposure information, and selected results are presented.

Giardiasis and cryptosporidiosis rates declined over the first twenty years of this surveillance program. However, the introduction of new and more sensitive diagnostic assays has led to an increase in parasitic disease rates, particularly cryptosporidiosis, since 2015. In 2019, there were 1,205 reported cases of giardiasis, compared to 1,112 in 2018. The rate of giardiasis per 100,000 population increased from 12.9 in 2018 to 14.3 in 2019, which exceeded the range of observed rates over the last decade (rate range 2009–2018: 9.2–12.9, median: 10.6). In 2019, there were 395 reported cases of cryptosporidiosis, compared to 250 in 2018. The rate of cryptosporidiosis per 100,000 population increased from 2.9 in 2018 to 4.7 in 2019, which again exceeded the range of observed rates over the last decade (rate range 2009–2018: 1.0–2.9, median: 1.4).

In 2015, the introduction of a new type of diagnostic test coincided with an increasing trend in observed cases of giardiasis and cryptosporidiosis. These assays, known as syndromic multiplex polymerase chain reaction (PCR) panels, can test for the presence of a wide range of enteric organisms including *Cryptosporidium* and *Giardia*. Prior to the availability of these new tests, physicians would have to specifically request testing for *Cryptosporidium spp.* The poor sensitivity of traditional diagnostics in addition to specific testing requirements likely contributed to under-reporting of cryptosporidiosis. However, since 2015, physicians at an increasing number of hospitals and laboratories across NYC can order a single test for a patient with diarrheal disease and evaluate the presence of approximately 20 different pathogens. The increased number of cases of parasitic disease observed since 2015 are hypothesized to reflect an increase in the detection of cases and not a true increase in disease for both cryptosporidiosis as well as giardiasis. This trend has also been observed across multiple jurisdictions in the United States.

An outbreak of cryptosporidiosis was detected by routine DOHMH cluster detection algorithms in September 2019. After investigation, it was determined that the outbreak was related to person-to-person transmission among the Orthodox Jewish community in Brooklyn. In total, 47 cases diagnosed between August and November 2019 were linked to this outbreak. There was no evidence to suggest this outbreak was related to the NYC water supply.

Additionally, work by DOHMH suggests that both giardiasis and cryptosporidiosis infections are commonly sexually transmissible enteric infections among men who have sex with men in NYC. In addition, the data suggests that international travel is a likely risk factor for infection for some residents of New York City.

SYNDROMIC SURVEILLANCE AND OUTBREAK DETECTION

The tracking of sentinel populations (e.g., nursing homes) or surrogate indicators of disease (e.g., drug sales) through “syndromic surveillance” can be useful in assessing gastrointestinal (GI) disease trends in the general population. Such tracking programs provide greater assurance against the possibility that a citywide outbreak would remain undetected. In addition, such programs can play a role in limiting the extent of an outbreak by providing an early indication of an outbreak so that control measures are rapidly implemented.

DOHMH maintains four distinct and complementary outbreak detection systems: one system involves the tracking of chief complaints from hospital emergency department (ED) databases; a second system involves the monitoring of sales of over-the-counter (non-prescription) anti-diarrheal medications; a third system tracks the number of stool specimens submitted to a clinical laboratory for microbiological testing; the fourth system involves DOHMH monitoring and assisting in the investigation of GI outbreaks in eight sentinel nursing homes.

A summary of syndromic surveillance findings for 2019 pertaining to GI illness is presented. Citywide trends and signals observed in the ED system were generally consistent with GI viral trends. There was no evidence of a drinking water-related outbreak in NYC in 2019.

INFORMATION SHARING AND RESPONSE PLANNING

Information on *Cryptosporidium* and *Giardia*, WDRAP, and related topics, is available on the websites of NYC’s DEP and DOHMH as listed in Section 4 of this report. Included are annual reports on program activities, fact sheets on giardiasis and cryptosporidiosis, and results from the DEP’s source water protozoa monitoring program. An updated version of NYC’s “*Hillview Reservoir Cryptosporidium and Giardia Action Plan*” (CGAP) was issued in 2019, per annual requirement. Finally, a manuscript detailing the epidemiology of cryptosporidiosis from 1995–2018 was written by DOHMH, in collaboration with DEP, and was accepted for publication by the journal *Emerging Infectious Diseases*. It was published in volume 26/number 3 of the journal in March 2020 (Alleyne, Fitzhenry et al. 2020).

1. INTRODUCTION

The Waterborne Disease Risk Assessment Program (WDRAP) is a multi-faceted public health assessment program that provides enhanced assurance of the microbial safety of New York City's (NYC) drinking water supply. This program is a critical element of NYC's Filtration Avoidance Program, which was developed in response to US Environmental Protection Agency's Surface Water Treatment Rule regulations. WDRAP is a joint agency program involving the NYC Department of Health & Mental Hygiene (DOHMH) and NYC Department of Environmental Protection (DEP). This partnership was originally established in 1993, under a joint-agency (DEP-DOHMH) Memorandum of Understanding. The intra-agency agreement between DEP and DOHMH for continuation of WDRAP was updated and signed in 2017, laying out the organizational & funding foundation for WDRAP until 2022.

The ongoing primary objectives of WDRAP are to:

- Obtain data on the rates of giardiasis and cryptosporidiosis, along with demographic and risk factor information on patients; and
- Provide a system to track gastrointestinal illness (diarrhea and vomiting) to ensure rapid detection of any waterborne disease outbreaks.

This report provides a summary of WDRAP highlights and data for the year 2019.

2. DISEASE SURVEILLANCE

2.1 *Giardiasis*

Giardiasis is a notifiable disease in NYC, per the NYC Health Code. From 1993 through 2010 active laboratory surveillance – involving visits or calls to labs by DOHMH staff – was conducted under WDRAP to ensure complete reporting of laboratory diagnosed cases of giardiasis. Since 2011, *Giardia* positive laboratory results have been reported to DOHMH via an electronic laboratory reporting system.

During 2019, a total of 1,205 cases of giardiasis were reported to DOHMH resulting in an annual case rate of 14.3 per 100,000 ([Table 1](#)). The annual case count increased 8% from 2018 to 2019. After a steep decline in giardiasis rates from 1994–2004 (rate range: 13.4–32.4 per 100,000, median 22.9 per 100,000, decline 59%) giardiasis rates remained relatively constant during 2005–2016 (rate range: 9.2–11.4 per 100,000, median: 10.5 per 100,000), as shown in [Figure 1A](#). In 2018, the giardiasis rate was 12.9 per 100,000 and rose to 14.3 per 100,000 in 2019 ([Figure 1B](#)). The introduction of new syndromic multiplex panels in clinical practice in 2015 has likely impacted the incidence of giardiasis (see further discussion later in this report).

Table 1: Giardiasis, the number of cases and case rates, New York City, 1994–2019.

| Year | Number of Cases | Case Rate per 100,000 |
|------|-----------------|-----------------------|
| 1994 | 2,457 | 32.3 |
| 1995 | 2,484 | 32.4 |
| 1996 | 2,288 | 29.6 |
| 1997 | 1,787 | 22.9 |
| 1998 | 1,959 | 24.9 |
| 1999 | 1,896 | 23.9 |
| 2000 | 1,771 | 22.1 |
| 2001 | 1,530 | 19 |
| 2002 | 1,423 | 17.6 |
| 2003 | 1,214 | 15 |
| 2004 | 1,088 | 13.4 |
| 2005 | 875 | 10.7 |
| 2006 | 938 | 11.4 |
| 2007 | 852 | 10.3 |
| 2008 | 840 | 10 |
| 2009 | 844 | 10.1 |
| 2010 | 923 | 11.3 |
| 2011 | 918 | 11.2 |
| 2012 | 872 | 10.7 |
| 2013 | 767 | 9.2 |
| 2014 | 864 | 10.4 |
| 2015 | 869 | 10.2 |
| 2016 | 899 | 10.5 |
| 2017 | 975 | 11.4 |
| 2018 | 1,112 | 12.9 |
| 2019 | 1,205 | 14.3 |

Note: Active disease surveillance for giardiasis began in July 1993. Starting January 2011, active laboratory surveillance was replaced by an electronic reporting system. Case numbers in this table conform to the case numbers as they appear in the NYC Department of Health and Mental Hygiene Bureau of Communicable Disease surveillance database for the years 1994–2019, and rates have been accordingly adjusted. Minor variations in the data may be related to reporting delays, corrections, the removal of duplicate reports, and other data processing refinements. Yearly case numbers and rates in this table may therefore differ from case numbers and rates that appeared in prior WDRAP reports.

Since 1995, case investigations for giardiasis have been conducted only for patients who are known or suspected to be in a secondary transmission risk category (e.g., food handler, health care worker, child attending child care, or child care worker), or when giardiasis clusters or outbreaks are suspected. In 2019, five patients diagnosed with giardiasis were excluded from work or school to reduce the risk of secondary transmission: two cases were children attending child care, two worked as healthcare professionals and one patient worked as a food handler. No cases were associated with outbreaks.

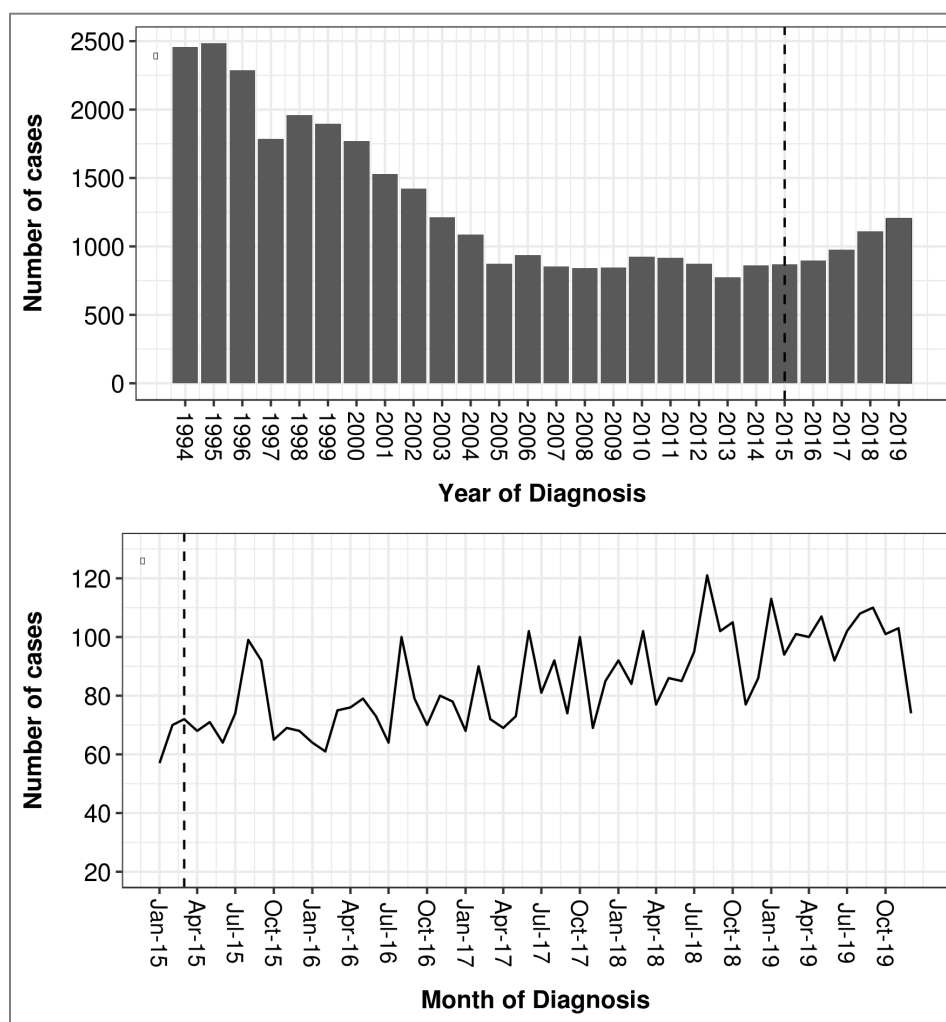


Figure 1: Annual **giardiasis** counts for all years in (A) and monthly counts for the last five years (B). The vertical dotted lines show the date when the first NYC laboratory reported results from using syndromic multiplex panels for enteric diseases.

The following provides highlights from the surveillance data for giardiasis among NYC residents diagnosed from January 1 through December 31, 2019. Data are presented in Figures 1 and 2 and Tables 1–6.

2.1.1 Borough of Patient Residence

Borough of patient residence was known for all 1,205 cases of giardiasis patients who resided in NYC. Manhattan had the highest borough-specific annual case rate (27.8 cases per 100,000) ([Table 3](#)). The highest United Hospital Fund (UHF) neighborhood-specific case rate was found in the Chelsea-Clinton neighborhood in Manhattan (67.6 cases per 100,000) ([Figure 2](#) and [Table 4](#)).

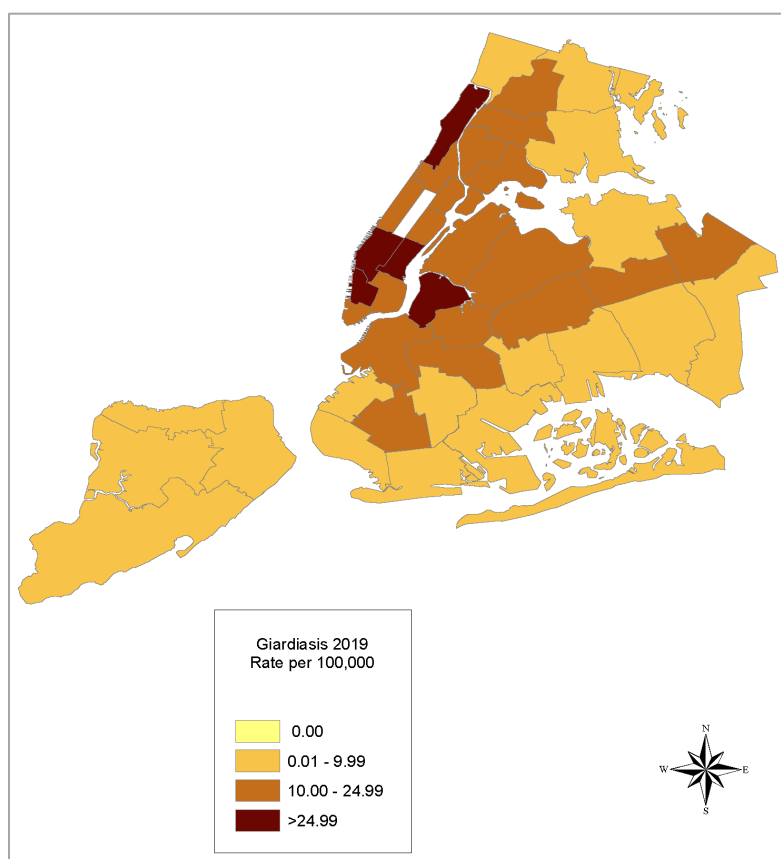


Figure 2: Map of giardiasis annual case rate per 100,000 population by United Hospital Fund Neighborhood, NYC, 2019.

2.1.2 Sex

Information regarding patient sex was available for all cases. The count and rate of giardiasis cases were higher in males than females, with 865 males (21.6 per 100,000) and 340 females (7.7 cases per 100,000) reported ([Table 3](#)). The highest sex- and borough-specific case rate was observed among males residing in Manhattan (46.3 cases per 100,000) ([Table 3](#)).

2.1.3 Age

Information regarding patient age was available for all cases. The highest age group-specific case rate was among persons aged 20–44 years (19.0 cases per 100,000). The highest age group and sex-specific case rate was among males aged 20–44 years (30.2 cases per 100,000) ([Table 5](#)). The two highest age-group and borough-specific case rates were in persons aged 45–59 years in Manhattan (36.7 cases per 100,000), followed by persons aged 20–44 years in Manhattan (35.7 cases per 100,000) ([Table 6](#)).

2.1.4 Race/Ethnicity

Information regarding patient race/ethnicity was available for only 292 of 1,205 (24%) cases. Ascertainment of race/ethnicity status for patients with giardiasis was poor. As mentioned, giardiasis patients are not routinely interviewed unless they are in occupations or settings that put them at increased risk for secondary transmission or if they are part of a suspected cluster or outbreak. Race/ethnicity information among giardiasis patients should be interpreted with caution as it may be based on the impressions of health care providers and may not reflect the patient's self-reported identity. For this reason, and because race/ethnicity information was missing for the majority of giardiasis disease reports, race/ethnicity findings pertaining to giardiasis patients diagnosed in 2019 are not presented in this report.

2.1.5 Census Tract Poverty Level

Age-adjusted case rates for giardiasis among four levels of census tract poverty, with levels encompassing low poverty to very high poverty, ranged from 14.6 to 22.1 cases per 100,000 population, with the lowest rate occurring in census tracts with very high poverty levels, and the highest rates occurring in census tracts with medium poverty levels ([Table 7](#)). Based on data from earlier WDRAP reports and from previous analyses (Greene, Levin-Rector et al. 2015), giardiasis is not typically associated with a high neighborhood poverty level in NYC. However, because giardiasis patients are not routinely interviewed, specific risk factors for giardiasis (e.g. international travel) in areas of low poverty versus high poverty are not known (see [APPENDIX A](#) for poverty definition).

2.1.6 Laboratory Diagnosis Trends

Syndromic multiplex panels are highly sensitive and specific in the detection of a large variety of enteric pathogens, including giardiasis (Navidad, Griswold et al. 2013, Madison-Antenucci, Relich et al. 2016). These panels are also a quick and less expensive method to screen for a large number (>20) of enteric pathogens, and their use has increased in recent years. In 2015, the proportion of giardiasis patients diagnosed exclusively by a syndromic multiplex panel at a hospital or commercial laboratory was 5%. This proportion grew to 12% in 2016, to 16% in 2017 and to 32% in 2018. By 2019, close to half (n=551, 46%) of all cases of giardiasis were exclusively diagnosed by a syndromic multiplex panel at a commercial or hospital laboratory. A

variety of laboratories began using syndromic multiplex panels to test for giardiasis in 2019. There are now 27 known laboratories reporting to NYC DOHMH using syndromic multiplex panels for enteric diagnoses, including 11 large hospitals, eight high volume commercial laboratories, six small volume commercial laboratories and two small clinics. There were approximately 21 known laboratories reporting to NYC DOHMH before 2019. Laboratories with syndromic multiplex panels are now used in all five boroughs. The proportion of giardiasis cases diagnosed in NYC exclusively by syndromic multiplex panels was less than that of cryptosporidiosis, as discussed below. This may potentially be related to the higher sensitivity of traditional diagnostics like an ova and parasite exam for giardiasis compared to cryptosporidiosis. It may be that reported giardiasis incidence prior to 2015 was closer to the true burden of disease than was the reported incidence of cryptosporidiosis, given the relatively robust sensitivity of traditional diagnostic assays for giardiasis, and the fact that the use of syndromic multiplex panels is having a smaller impact on reported giardiasis incidence in NYC.

2.1.7 Giardiasis as a Sexually Transmissible Enteric Infection

As in previous years, the age/sex demographic group with the largest number of diagnosed giardiasis cases in 2019 was adult men aged 20–44 years (39%, 468/1205) followed by adult men aged 45–59 years (15%, 175/1205). Adult men have been consistently over-represented in surveillance data since the WDRAP began. Giardiasis rates have historically and consistently been elevated in Chelsea-Clinton, a neighborhood in Manhattan with a higher prevalence of men who have sex with men compared to the rest of NYC (Bureau of Epidemiology Services New York City Department of Health and Mental Hygiene 2017). It is hypothesized that giardiasis is a sexually transmissible enteric infection among men who have sex with men in NYC and accounts for a considerable burden of reported disease.

Giardiasis is known to be a sexually transmissible enteric infection among men who have sex with men (Mitchell and Hughes 2018). Studies from several decades in NYC demonstrated that giardiasis and amebiasis were commonly detected in this population (Kean, William et al. 1979, Phillips, Mildvan et al. 1981). The authors of one study hypothesized that enteric parasitic infections are hyperendemic in men who have sex with men because of three factors: a high prevalence in the population, the prevalence of sexual behavior that facilitates transmission, and the frequency of exposure to infected persons (Phillips, Mildvan et al. 1981). However, information on exposures such as sexual behavior is not routinely collected for giardiasis patients in NYC so it is not possible to determine how prevalent sexual behavior with increased risk of fecal/oral contact is among reported giardiasis patients.

2.2 Cryptosporidiosis

Cryptosporidiosis was added to the list of reportable diseases in the NYC Health Code in January 1994. Active disease surveillance for cryptosporidiosis involving lab visits and calls began in November 1994 and continued through 2010. Starting in 2011, active surveillance was replaced

by electronic laboratory reporting. Patient interviews for demographic and risk factor data were initiated in 1995 and are ongoing.

Table 2: Cryptosporidiosis, number of cases and case rates, New York City, 1994–2019

| Year | Number of Cases | Case Rate per 100,000 |
|------|-----------------|-----------------------|
| 1994 | 288 | 3.8 |
| 1995 | 471 | 6.1 |
| 1996 | 334 | 4.3 |
| 1997 | 172 | 2.2 |
| 1998 | 207 | 2.6 |
| 1999 | 261 | 3.3 |
| 2000 | 172 | 2.1 |
| 2001 | 122 | 1.5 |
| 2002 | 148 | 1.8 |
| 2003 | 126 | 1.6 |
| 2004 | 138 | 1.7 |
| 2005 | 148 | 1.8 |
| 2006 | 155 | 1.9 |
| 2007 | 105 | 1.3 |
| 2008 | 107 | 1.3 |
| 2009 | 81 | 1 |
| 2010 | 107 | 1.3 |
| 2011 | 86 | 1.1 |
| 2012 | 125 | 1.5 |
| 2013 | 80 | 1 |
| 2014 | 102 | 1.2 |
| 2015 | 133 | 1.6 |
| 2016 | 192 | 2.2 |
| 2017 | 163 | 1.9 |
| 2018 | 250 | 2.9 |
| 2019 | 395 | 4.7 |

Note: Case numbers in this table conform to the case numbers as they appear in the NYC Department of Health and Mental Hygiene Bureau of Communicable Disease surveillance database for the years 1994–2019, and rates have been accordingly adjusted. Minor variations in the data may be related to reporting delays, corrections, the removal of duplicate reports, and other data processing refinements. Yearly case numbers and rates in this table may therefore differ from case numbers and rates that have appeared in prior WDRAP reports.

During 2019, a total of 395 cases of cryptosporidiosis were reported to DOHMH, all of which met the case definition for confirmed cryptosporidiosis (see [APPENDIX A](#) for case definition description). The 2019 annual case rate was 4.7 per 100,000 ([Table 2](#)). The annual case count

increased 58% from 2018 to 2019. After a sharp decline in cryptosporidiosis rates from 1995–2006 (rate range: 1.5–6.1 per 100,000, median 2.1 per 100,000, decline 75%), cryptosporidiosis rates remained relatively constant during 2007–2014 (rate range: 1.0–1.5 per 100,000, median: 1.3 per 100,000) as shown in [Figure 3A](#). Cryptosporidiosis rates started to increase in 2015, rising from 1.6 per 100,000 to 4.7 per 100,000 in 2019.

In 2019, 34 patients diagnosed with cryptosporidiosis were excluded from work or school to reduce the risk of secondary transmission. The majority of exclusions were children aged <5 years in child care or preschool (n=14), followed by food handlers (n=9), individuals who worked with children in child care facilities or preschools (n=7), healthcare workers (n=3), and a child in a camp (n=1). Six excluded patients were associated with an outbreak among the Orthodox Jewish community ([section 2.2.10](#)).

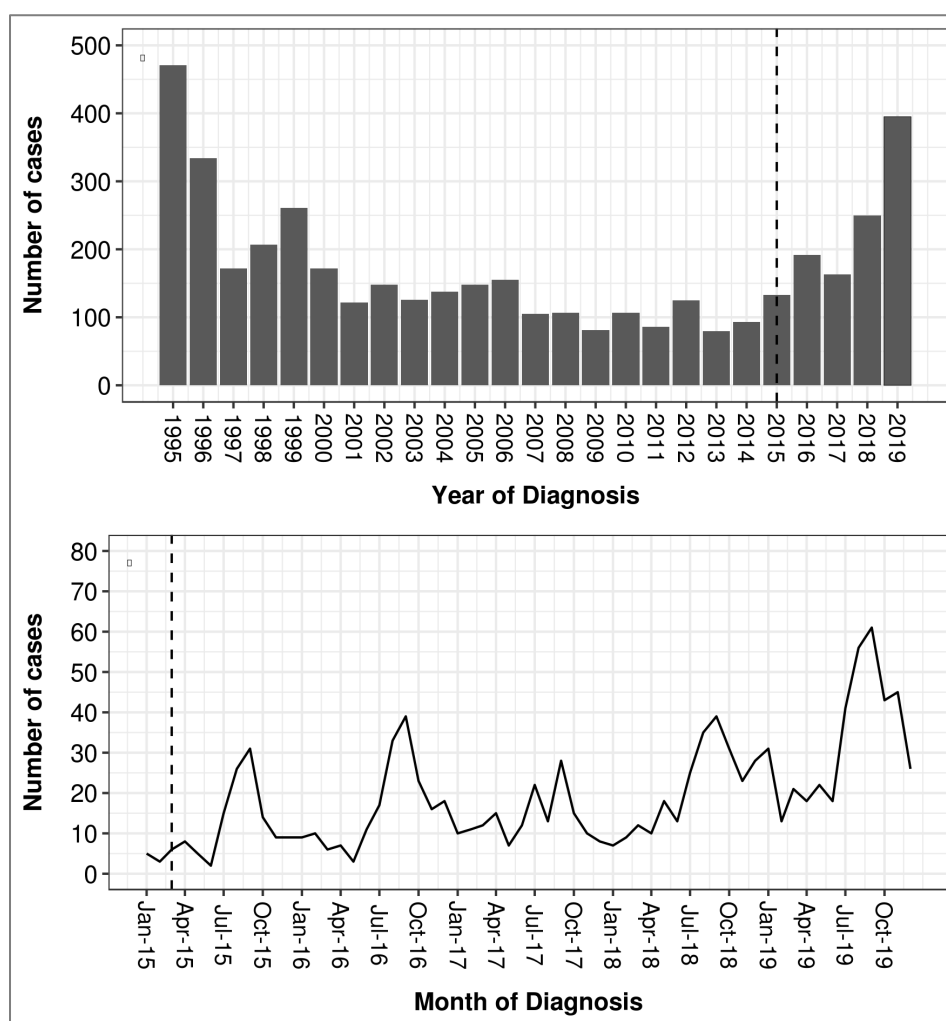


Figure 3: Annual **cryptosporidiosis** counts for all years in (A) and monthly counts for the last five years (B). The vertical dotted lines show the date when the first laboratory NYC reported results from syndromic multiplex panels for enteric diseases.

Cryptosporidiosis is highly seasonal in NYC, as shown in [Figure 3B](#). In 2019, cryptosporidiosis patients were most often diagnosed in August (n=56, 14%) or September (n=61, 15%). Because diagnosis may occur sometime after onset, information is collected in the interview regarding date of symptom onset. The date of onset can be used more accurately than date of diagnosis to estimate when patients were likely exposed to *Cryptosporidium* and is used to determine the risk exposure period.

The following provides highlights from the surveillance data for cryptosporidiosis among NYC residents from January 1 through December 31, 2019. Data are presented in Figures 3–5 and Tables 8–18.

2.2.1 Borough of patient residence

Information on borough of residence was available for all 395 cases of cryptosporidiosis. Manhattan had the highest borough-specific annual case rate (8.0 cases per 100,000) ([Table 8](#)). The highest UHF neighborhood-specific case rate was in the Greenpoint neighborhood in Brooklyn (16.9 cases per 100,000), followed by Upper East Side in Manhattan (12.8 cases per 100,000), and Chelsea-Clinton (12.7 cases per 100,000) ([Figure 4](#) and [Table 9](#)).

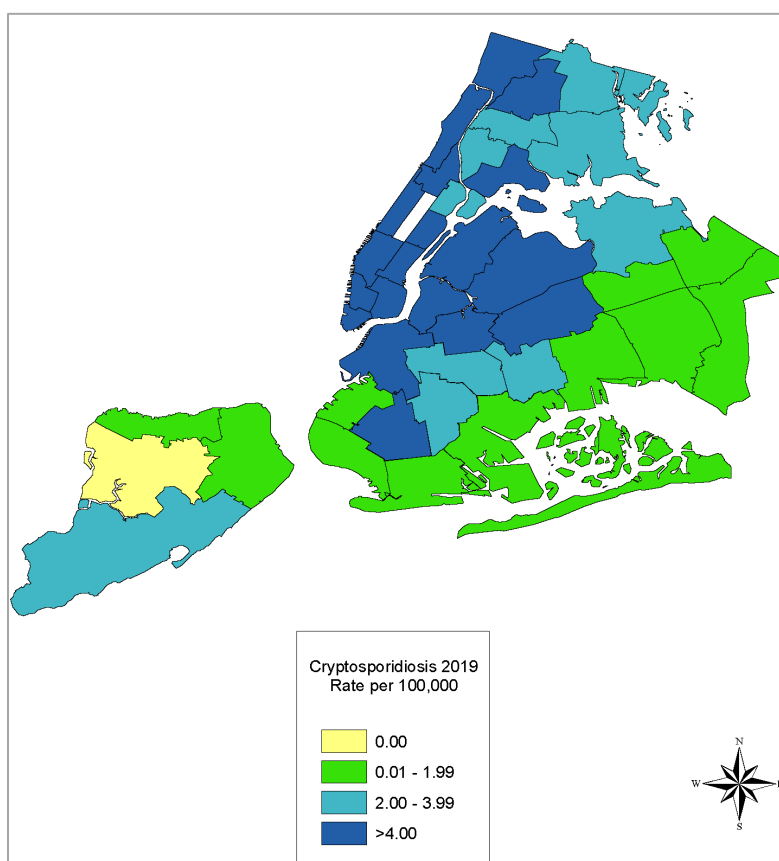


Figure 4: Map of cryptosporidiosis annual case rate per 100,000 population by United Hospital Fund neighborhood, NYC, 2019.

2.2.2 Sex

Information regarding patient sex was available for all cases. The count and rate of cryptosporidiosis cases was higher in males than females, with 221 males (5.5 cases per 100,000), and 174 females (4.0 cases per 100,000) ([Table 8](#)). The borough- and sex-specific case rate was highest for males in Manhattan (10.2 cases per 100,000) ([Table 8](#)).

2.2.3 Age

Information regarding patient age was available for all cases. The highest age group-specific case rates were in children aged <5 years (8.8 cases per 100,000), followed by persons aged 20–44 years (6.6 cases per 100,000). The highest age group- and sex-specific case rates were in males aged <5 years (9.1 cases per 100,000), followed by females aged <5 years (8.4 cases per 100,000) ([Table 10](#)). The highest age group and borough-specific case rates occurred in persons aged <5 years in Brooklyn (13.9 cases per 100,000), followed by persons aged 20–44 years in Manhattan (12.3 cases per 100,000) ([Table 11](#)). The high case rate among children in Brooklyn was related to the outbreak among the Orthodox Jewish population, please see [section 2.2.10](#).

2.2.4 Race/Ethnicity

Patient race/ethnicity information was available for 366 of 395 cases (92.6%). Among the major racial/ethnic groups, White, non-Hispanic persons had the highest cryptosporidiosis rate (6.5 per 100,000) followed by Hispanic persons (4.2 per 100,000) and Black/African American persons (3.3 per 100,000) ([Table 12](#)). Cryptosporidiosis rates were highest among White, non-Hispanic persons in Manhattan (9.7 per 100,000), highest among Hispanic persons in Manhattan (5.7 per 100,000) and Queens (5.6 per 100,000), and highest among Black/African Americans in the Bronx (5.5 per 100,000) ([Table 12](#)). By age group, rates were highest among white, non-Hispanic children aged <5 years (18.2/100,000). Among Hispanic persons, rates were evenly distributed across age groups. Among Black/African American persons, rates were highest among persons aged 20–44 years (5.8 per 100,000) ([Table 13](#)). This paragraph does not describe some race/ethnic groups due to relatively small number of people in those groups.

2.2.5 Census Tract Poverty Level

Age-adjusted case rates for cryptosporidiosis among four levels of census tract poverty ranged from 4.4 to 7.9 cases per 100,000 population, with no clear pattern between age-adjusted rate and increasing or decreasing census tract poverty level in 2019 ([Table 14](#)).

2.2.6 Laboratory Diagnosis Trend

Similar to the trends in giardiasis testing, a number of large healthcare facilities in NYC began to report cryptosporidiosis diagnosed by syndromic multiplex panels to DOHMH during 2015–2019. Notably, Columbia University Medical Center began using a syndromic multiplex

panel in 2015; and in 2019, they published a manuscript detailing the increased sensitivity they found with these panels in comparison with the traditional microscopy assay. The authors found that traditional testing identified a pathogen in 4% of samples from 2012–2015 compared to 29% of samples positive for a pathogen using syndromic multiplex panel on samples from 2015–2017 (Axelrad, Freedberg et al. 2019).

In 2015, the proportion of NYC cryptosporidiosis patients diagnosed exclusively by a syndromic multiplex panel at a hospital or commercial laboratory was 20%. This proportion grew to 34% in 2016, 48% in 2017 and by 2018 was 75%. This proportion of all cases of cryptosporidiosis diagnosed by a syndromic multiplex panel held steady in 2019 (74%, n=282). This trend has been mirrored across a number of different jurisdictions in the United States (Huang, Henao et al. 2016, Marder, Cieslak et al. 2017). The sensitivity and specificity of these panels for the detection of cryptosporidiosis over traditional microscopic diagnostic techniques is described by others as well (Navidad, Griswold et al. 2013, Buss, Leber et al. 2015). And, importantly, the panels are considerably less expensive. Of note, as indicated in Section 2.1.6, the number of laboratories reporting to NYC DOHMH using syndromic multiplex panels for enteric diseases was higher in 2019 than in prior years.

In the manuscript prepared by the NYCDOHMH-based team in 2019, which details the descriptive epidemiology of cryptosporidiosis in NYC between 1995–2018, we noted that the reported incidence of cryptosporidiosis increased significantly after the introduction of syndromic multiplex panels in 2015, and that the demographic profile of patients changed (Alleyne, Fitzhenry et al. 2020). The median age-adjusted annual incidence increased from 1.46/100,000 in 2000–2014 to 2.11/100,000 during 2015–2018, following the introduction of syndromic multiplex panels (incidence rate ratio: 1.49, 95% CI: 1.17–1.91). After these new tests were adopted, cryptosporidiosis patients were more likely to be aged 10-19 years of age, HIV negative, and non-Hispanic White, compared with the period prior to syndromic multiplex availability. Given the dramatic consequences of cryptosporidiosis amongst people living with HIV/AIDS, clinicians treating this population would have likely been more aware of the need to specifically request testing for this parasite in the pre-syndromic multiplex panel era. The increase in HIV-negative patients ([Figure 5](#)) likely reflects increased case finding among the general population. A change in the racial profiles of patients might reflect both the populations residing in the specific catchment areas of the laboratories using syndromic multiplex panels, as well as disparities in health care access by race/ethnicity.

The increased range of hospitals and laboratories using the syndromic multiplex panels is leading to an increase in reported incidence of cryptosporidiosis across a range of neighborhoods in NYC. Importantly, DOHMH has also observed substantial increases in reported incidence of a range of additional enteric infections included on syndromic multiplex panels across NYC. Some infections with increasing incidence because of the use of syndromic multiplex panels, such as norovirus, are transmitted predominately by person-to-person contact or fecal-oral contact and are not normally related to waterborne transmission.

2.2.7 Cryptosporidiosis and Immune Status

Trends observed over the years in reported numbers of cryptosporidiosis cases have differed between persons living with HIV/AIDS and those who are immunocompetent. Reported cryptosporidiosis cases among persons living with HIV/AIDS declined dramatically during 1995–1997, corresponding with the introduction of highly active antiretroviral therapy for HIV/AIDS. The count of cryptosporidiosis cases among persons living with HIV/AIDS has continued to decline since then, with only 60 cases reported in 2019 (representing 15.2% of all cases). The count of cryptosporidiosis cases among immunocompetent patients has increased since 2015, however, rising from 78 to 313 in 2019 (300% increase) (Figure 5). This trend is also coincident with the introduction of syndromic multiplex panels in 2015 as mentioned in [section 2.2.6](#). As cryptosporidiosis infection can be particularly severe among people living with HIV/AIDS (Blanshard, Jackson et al. 1992, Poznansky, Coker et al. 1995, Rashmi and Kumar 2013), physicians were historically more likely to consider cryptosporidiosis in their differential diagnosis of diarrheal disease among persons living with HIV/AIDS than in a person without HIV/AIDS. However, now that syndromic multiplex panels can be ordered for diagnosis of any diarrheal infection in hospitals and clinics that have adopted these assays, cryptosporidiosis is more frequently identified in immunocompetent patients who likely would not have been tested for cryptosporidiosis before 2015.

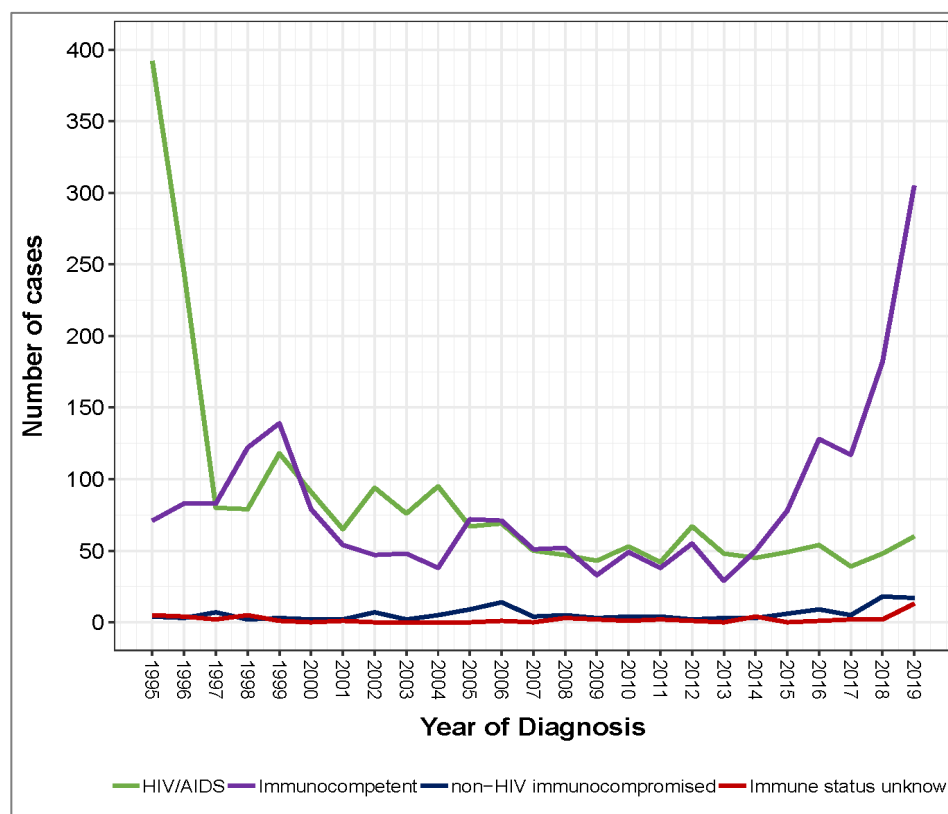


Figure 5: Cryptosporidiosis, number of cases by year of diagnosis and immune status, New York City, 1995–2019.

2.2.8 Cryptosporidiosis and Potential Risk Exposures

Of the 395 cryptosporidiosis cases diagnosed among NYC residents in 2019, questionnaires concerning potential exposures were completed for 314 (79.5%) patients. For patients with missing interview data, investigators were either unable to locate the patient (44 cases, 11.1%) or the patient refused interview (37 cases, 9.4%). Of the immunocompetent patients, interviews were completed for 264 patients (84.3%). Among persons with HIV/AIDS, interviews were completed for 38 patients (63.3%), and interviews were completed for 12 patients (85.7%) who were immunocompromised for reasons other than HIV/AIDS. Summary data for 1995 through 2019 on commonly reported potential risk exposures, obtained from patient interviews of persons with HIV/AIDS and from interviews of persons who are immunocompetent, are presented in [Table 15](#) and [Table 16](#), respectively. Information has also been collected regarding type of tap water consumption, and is presented in [Table 17](#) and [Table 18](#). Compared to previous years, reported use of filtered water increased among both persons living with HIV/AIDS as well as immunocompetent patients. It is unclear why this change was observed.

Tables 15–18 indicate the percentage of patients who reported engaging in each of the listed potential risk exposures for cryptosporidiosis before disease onset. However, it must be noted that the determination of an association between exposure to possible risk factors for cryptosporidiosis and acquisition of cryptosporidiosis cannot be made without reference to a suitable control population (i.e., non-*Cryptosporidium*-infected controls). As exposure data for a control population are not available, such determinations of association cannot be made.

Though no conclusions about association can be reached, in an attempt to assess if there are any patterns of interest, data have been compared between patients who are immunocompromised because of HIV/AIDS and patients who are immunocompetent. In 2019, interviewed patients who were immunocompetent were significantly more likely to report international travel (39% compared to patients with HIV/AIDS (11%) ($p=0.0004$, Fishers exact test). Additionally, interviewed immunocompetent patients were also more likely to report exposure to recreational water (30%) compared to patients with HIV/AIDS (5%) ($p=0.0007$, Fishers exact test). There were no significant differences in reported contact with an animal between the two groups (33% and 32%, respectively, $p=1.0$, chi-square test). Finally, interviewed patients with HIV/AIDS were more likely to report high-risk sexual activity (39%) compared to immunocompetent patients (7%) ($p<0.0001$, chi-square test). It should be noted that high-risk sex in this context refers to having a penis, finger or tongue in a partner's anus. Information about sexual practices is gathered via phone interview and may not be reliable. Overall, these data indicate that, for most years, immunocompetent patients were more likely to travel internationally and have greater recreational water exposure than immunocompromised patients. International travel and exposure to recreational water may be more likely risk factors for the acquisition of cryptosporidiosis in the immunocompetent group. However, as noted above, the extent to which these risk factors may have been associated with cryptosporidiosis cannot be determined without comparison to a control population.

2.2.9 Cryptosporidiosis as a Sexually Transmissible Enteric Infection

As in previous years, and similar to giardiasis, the age/sex demographic group with the largest number of diagnosed cryptosporidiosis cases in 2019 was adult men aged 20–44 years (29%, 116/395). Adult men aged 45–59 years accounted for an additional 8% of all people diagnosed with cryptosporidiosis in 2019. This demographic group has been consistently over-represented in surveillance data since the WDRAP began, again similar to the profile of giardiasis. Furthermore, cryptosporidiosis rates have historically and consistently been elevated in Chelsea-Clinton, a neighborhood in Manhattan with a higher prevalence of men who have sex with men compared to the rest of NYC (Bureau of Epidemiology Services New York City Department of Health and Mental Hygiene 2017). Therefore, it is hypothesized that cryptosporidiosis is often an infection among men who have sex with men in NYC.

Men who have sex with men are historically at greater risk for cryptosporidiosis, not only because of a higher prevalence of AIDS in this population (Centers for Disease Control and Prevention 2006), but also because of higher risk sexual practices related to oral/anal contact that entail a low risk for HIV transmission but increase the risk for fecal contact (Hellard, Hocking et al. 2003). In 2019, there were a total of 94 adult men aged 20–59 years who answered questions related to sexual behavior in their cryptosporidiosis incubation period. There were a total of 126 other adults (men aged 18 and 19 years and men aged >59 years as well as all women ≥ 18 years) who answered the sexual behavior questions during interview. Among men diagnosed with cryptosporidiosis aged 20–59 years, 35% (33/94) reported high-risk sexual practices, compared to 6% (8/126) of all other adult cryptosporidiosis patients ($p < 0.001$, Fishers exact test). There are considerable limitations with large amounts of missing data in the sexual behavior questions. However, the data suggest that adult men diagnosed with cryptosporidiosis are likely to report engaging in sexual behaviors that increase the risk of fecal/oral contact.

2.2.10 Cryptosporidiosis outbreak among the Orthodox Jewish population of Brooklyn

In 2019, routine DOHMH cluster detection algorithms detected an increase of cryptosporidiosis in Brooklyn at the end of September. Initial investigations suggested the infections were concentrated in the Orthodox Jewish communities of Borough Park and Williamsburg in Brooklyn. In total, there were 47 cases diagnosed between August–November 2019 that were linked to this outbreak. Diagnoses peaked the week of October 6, 2019 ($n=11$), and onset dates peaked the week of August 25, 2019 ($n=8$). The median patient age was 13 years, ranging from <1–46 years. The proportion of young children aged <5 years (36%) was similar to adults aged >18 years (43%) and cases were equally as likely to be male (49%) or female (51%).

A supplemental questionnaire was developed to ask patients about recent exposures related to water and contact with other cases in the community. Twelve patients (26%) reported traveling to upstate NY during the summer months, the majority to Sullivan ($n=8$) or Orange Counties ($n=2$). Many patients with upstate travel reported spending the summer months in Orthodox Jewish communities or camps, returning home for the start of the school year on September 1,

2019. Patients with reported travel to upstate NY had onset dates at the start of the outbreak (August 2–September 1, 2019).

Seventeen patients (36%) reported attending (n=16) or working in (n=1) child care centers or preschools in the Orthodox community. Six patients were excluded from attending child care or preschool while infectious as per the NYC Health Code (New York City Health Code 2019). Additionally, there were eight likely instances of intrahousehold transmission, all related to an initial infection in a child or worker at a school or child care center. Two patients (4%) reported water exposure (aside from drinking or bathing) in NYC: both patients reported attending mikva, which is a ceremonial bath in water at a temple. The patients did not attend the same temple.

DOHMH conducted substantial outreach to the Orthodox community. Letters in English and Yiddish were sent to child care centers and schools in the community, informing principals, teachers, and parents about the outbreak. The letters included details on cryptosporidiosis symptoms and transmission, and instructed staff and children to wash hands frequently with soap and water. A public messaging campaign was introduced into a popular messaging forum to alert the Orthodox community to wash hands with soap and instructed people to stay home from work or school if ill. Finally, an article was published in a local newspaper read widely among the Orthodox community (Borchardt October 30, 2019).

Data gathered by NYCDOHMH indicated that this outbreak was related to person-to-person transmission within the Orthodox Jewish community in Borough Park and Williamsburg, Brooklyn. Similar outbreaks among the Orthodox community have been documented for the bacterial diarrheal infection shigellosis, including a documented risk for intrahousehold transmission and transmission in child care centers (Pilon, Camara et al. 2016, Cohen, Korin et al. 2019). This outbreak was not related to the water supply of NYC. A manuscript describing this outbreak will be submitted for publication in 2020 to inform clinicians treating this community in NYC.

3. SYNDROMIC SURVEILLANCE AND OUTBREAK DETECTION

The tracking of sentinel populations or surrogate indicators of disease (“syndromic surveillance”) can be useful in assessing gastrointestinal (GI) disease trends in the general population. Such tracking programs provide greater assurance against the possibility that a citywide outbreak would remain undetected. In addition, such programs can play a role in limiting the extent of an outbreak by providing an early indication of a problem so that control measures can be rapidly implemented. Beginning in the 1990s, NYC established and has maintained a number of distinct and complementary outbreak detection systems. One system utilizes hospital emergency department (ED) chief complaint logs to monitor for outbreaks. The ED system is relied upon most heavily for monitoring the burden of diarrheal illness in NYC. A second system monitors sales of anti-diarrheal medications: the Anti-Diarrheal Monitoring System (ADM)/over-the-counter medication (OTC) system. A third system monitors the number

of stool specimens submitted to a participating clinical laboratory for microbiological testing. Finally, the fourth system monitors for GI outbreaks in sentinel nursing homes and DOHMH staff assist in the investigation of any identified outbreaks. A full description of each system can be found in [APPENDIX B](#).

Other than the ED system, which is mandated under the NYC Health Code, all systems rely upon the voluntary participation of the organizations providing the syndromic data. A summary of syndromic surveillance findings pertaining to GI illness for 2019 is provided in the final section of Section 3.1 and in [Figure 6](#), [Figure 7](#), and [Figure 8](#).

Throughout 2019, DOHMH received electronic data from all of NYC's 53 EDs, which reported approximately 11,500 visits per day. Additionally, data were received daily from approximately 515 pharmacies in 2019 as part of the ADM/OTC system. Finally, WDRAP team members made site visits to seven of eight nursing homes participating in the Nursing Home Sentinel Surveillance system in 2019. The remaining nursing home was visited in February 2020.

3.1 Findings: Summary of Syndromic Surveillance Signals

Syndromic surveillance signals alone cannot be used to determine etiologic diagnoses. Also, experience has shown that most signals, especially localized spatial signals in the emergency department system or signals in the laboratory or ADM monitoring systems, may be statistical aberrations and not related to public health events. The systems are therefore used in concert. A signal in one system is compared to other systems to evaluate the presence of concurrent signals. In this report, Figures 6–8 summarize GI disease signals from NYC's syndromic surveillance systems. [Figure 6](#) and [Figure 7](#) summarize signals from the Emergency Department system only. [Figure 8](#) summarizes signal results from all syndromic surveillance systems operated by DOHMH during 2019.

Of note, DOHMH saw a significant increase in norovirus reports through routine surveillance activities during November 2018–April 2019 and from mid-November 2019–December 2019. There were approximately 100 reports of norovirus per week from mid-January through mid-March 2019. For context, the maximum weekly report case count in 2018 was <60. Additionally, there was an increase in rotavirus counts in 2019 compared with 2018, with weekly reported rotavirus cases peaking from the start of February to the start of May 2019. These increases were likely related to the introduction of syndromic multiplex panels. For the most up to date data on all communicable diseases from DOHMH, please see the Epiquery webpage (New York City Department of Health and Mental Hygiene 2020).

[Figure 6](#) shows the ratio of daily ED visits for the diarrhea syndrome to all other daily ED visits for syndromes not tracked by ED syndromic surveillance (“other visits”) from January 1 to December 31, 2019. The graph also indicates the occurrence of citywide signals and of the spatial residential zip code and hospital signals. There were no citywide diarrheal ED signals in 2019. The zip code ED signals in July 2019 were related to an outbreak of shigellosis traced back to a contaminated, recirculating fountain in Flushing Meadows/Corona Park. There were over 30

cases of shigellosis identified, most of them children, who had exposure to recreational water in the park. DOHMH worked together with the Department of Parks and Recreation to ensure cleaning and disinfection several splash pads and the fountain in question in late July 2019.

The diarrheal ED signals in June 2019 occurred at two hospitals in Manhattan. WDRAP staff reviewed the signals and found that there was no unusual spatial distribution of cases. WDRAP staff also reviewed the DEP water contamination dashboard and did not identify any water testing parameters that exceeded normal limits. The diarrheal ED signals in November 2019 occurred in a neighborhood of Queens. WDRAP staff reviewed syndromic systems and identified that the signals were likely driven by children aged 0–4 years. The hospital ED was contacted and per the attending physician, there had been a recent increase in the number of children coming to the pediatric ED with diarrhea and vomiting. The children were seen and discharged to home as the symptoms were mild. No stool samples were taken at the discretion of the attending physicians. Neither set of the ED signals (June and November) were determined to be related to a waterborne disease outbreak.

[Figure 7](#) shows the ratio of daily ED visits for the vomiting syndrome compared to all other daily ED visits for syndromes not tracked by ED syndromic surveillance for 2019. There were no citywide ED signals for vomiting in 2019. There were single-day spatial-hospital signals in January, June, and November and one multi-day spatial-hospital signal in December. Given the signals were localized, were short duration, and lacked corresponding signals in the other monitoring systems, these were not determined to be related to a waterborne disease outbreak.

[Figure 8](#) shows the timing of signals from all four surveillance systems in 2019. There were no citywide signals in either diarrhea or vomiting from the ED signals in 2019. However, there were a large number of OTC/ADM signals throughout the year, concentrating specifically in April and July. The signals in April likely relate to coincident increase in both norovirus and rotavirus at that time. The signals in July were driven by a new store that opened in mid-July in zip code 10455. The majority of the OTC/ADM signals were found to be related to promotional sales at the pharmacy chains, specifically for Pepto Bismol®/Bismuth sales. There was no evidence to suggest that the OTC/ADM signals were related to a waterborne disease outbreak. Additionally, there were eight signals in the Clinical Laboratory surveillance system throughout the year. There was a [large outbreak](#) of foodborne *Cyclospora* identified in July 2019, which may account for some of the signaling in July as *Cyclospora*, *Cryptosporidium*, and *Giardia* are all detected by similar microscopy assays at the Clinical Laboratory. The mean length of these signals was 1.5 days (range: 1–3 days). The longest signal was 3 days, March 21–March 23, 2019. During this week, there were no positive stool specimens for *Cryptosporidium* in a NYC resident from the clinical laboratory.

There was one GI outbreak in a sentinel nursing home in 2019. The sentinel nursing home GI outbreak occurred in a facility in Brooklyn, beginning on November 20, 2019. Six patients on two units and one staff member were reported to be ill. The symptoms were vomiting and diarrhea. There were no deaths or hospitalizations. The facility sent three stool specimens from three residents to the Public Health Laboratory for viral pathogen testing on November 21, 2019. All three specimens were positive for norovirus genogroup II RNA. The facility also sent three

specimens from three patients for bacterial and *Clostridium difficile* testing to a private laboratory. These tests were negative. Based on the positive norovirus test and the citywide increase in norovirus starting in mid-November 2019, DOHMH is confident this nursing home outbreak was related to norovirus.

In summary, there were no citywide ED signals for GI illness in 2019. The ED system was robust enough to signal in response to a localized shigellosis outbreak in Queens. The OTC/ADM signals were related primarily to promotional sales, and a period of signaling coincided with the incidence of viral GI illness in 2019, which was elevated compared to 2018

In conclusion, during 2019, there were no signals consistent with a waterborne disease outbreak from the four syndromic surveillance systems set up to detect an outbreak related to the water supply. This finding is consistent with all prior years of WDRAP surveillance.

4. INFORMATION SHARING AND RESPONSE PLANNING

In 2019, DOHMH authored a manuscript in collaboration with DEP detailing the epidemiology of cryptosporidiosis in NYC from 1995–2018 (Alleyne, Fitzhenry et al. 2020) as mentioned in [section 2.2.6](#). The paper appeared in the March 2020 edition of the Journal of Emerging Infectious Diseases and is expected to reach a large audience of public health practitioners and infectious disease clinicians, both in NYC and elsewhere.

World Pride 2019 occurred in NYC. World Pride is a global celebration of the LGBTQ community and NYC DOHMH updated the enterics among MSM [postcard](#) and handed out them out at Pride activities throughout the City in June 2019.

Additionally DOHMH conducted extensive outreach to the Orthodox Jewish community in Brooklyn alerting them to the existence of an outbreak of cryptosporidiosis, as described in [section 2.2.10](#). This outbreak was not related to the NYC water supply.

Information pertaining to NYC’s Waterborne Disease Risk Assessment Program and related issues are available on both the DEP and DOHMH websites, including results from the City’s source water protozoa monitoring program. Documents on the websites include:

DOHMH Webpages:

[The Giardiasis and Cryptosporidiosis Fact Sheets were modified by DOHMH in 2019]

- *Giardiasis fact sheet*
<https://www1.nyc.gov/site/doh/health/health-topics/giardiasis.page>
- *Cryptosporidiosis fact sheet*
<https://www1.nyc.gov/site/doh/health/health-topics/cryptosporidiosis.page>
- Communicable Disease Surveillance Data
<https://a816-healthpsi.nyc.gov/epiquery/CDSS/index.html>

- Diarrheal Infections in Gay Men and Other Men Who Have Sex with Men
<https://www1.nyc.gov/site/doh/health/health-topics/diarrheal-infections.page>

DEP Webpages:

- ***Waterborne Disease Risk Assessment Program's Annual Reports***
 - For the latest WDRAP annual report posted:
<https://www1.nyc.gov/site/dep/water/waterborne-disease-risk-assessment.page>
 - For WDRAP Annual reports going back to 1997:
<https://www1.nyc.gov/site/dep/about/document-portal.page>
- ***New York City Drinking Water Supply and Quality Statement (for latest posted report):***
<https://www1.nyc.gov/site/dep/about/drinking-water-supply-quality-report.page>
- ***DEP Water Supply Testing Results for Giardia and Cryptosporidium***
(Data are collected and entered on the website each week; historical data are also included).
<https://data.cityofnewyork.us/Environment/DEP-Cryptosporidium-And-Giardia-Data-Set/x2s6-6d2j>

With regard to response planning, NYC has developed an action plan for responding to elevations in levels of either *Giardia* cysts or *Cryptosporidium* oocysts at a key water supply monitoring location. The initial response plan was developed in 2001. The plan in its current form is known as, NYC's "*Hillview Reservoir Cryptosporidium and Giardia Action Plan*" (CGAP), and the plan is reviewed & updated on an annual basis; it was updated in 2019.

5. REFERENCES

- Alleyne, L., R. Fitzhenry, K. A. Mergen, N. Espina, E. Amoroso, D. Cimini, S. Balter, A. M. Fireteanu, A. Seeley, L. Janus, B. Gutelius, S. Madison-Antenucci and C. N. Thompson (2020). "Epidemiology of cryptosporidiosis in New York City, New York, USA, 1995-2018." Emerging Infectious Disease **26**(3).
- Axelrad, J. E., D. E. Freedberg, S. Whittier, W. Greendyke, B. Lebwohl and D. A. Green (2019). "Impact of Gastrointestinal Panel Implementation on Health Care Utilization and Outcomes." Journal of Clinical Microbiology **57**(3): e01775-01718.
- Blanshard, C., A. M. Jackson, D. C. Shanson, N. Francis and B. G. Gazzard (1992). "Cryptosporidiosis in HIV-seropositive patients." The Quarterly Journal of Medicine **82**(307-308): 813-823.
- Borchardt, R. (October 30, 2019). Outbreak of Parasitic Infection in Boro Park and Wililamsburg. Hamodia. <https://hamodia.com/2019/10/30/outbreak-parasitic-infection-boro-park-williamsburg>.
- Bureau of Epidemiology Services New York City Department of Health and Mental Hygiene (2017). Prevalence of Men Who Had Sex with Men in the past 12 months in NYC by United Hospital Fund Neighborhood, Community Health Survey, 2012-2016; <http://www1.nyc.gov/site/doh/data/data-sets/community-health-survey-public-use-data.page>.
- Buss, S., A. Leber, K. Chapin, P. Fey, M. Bankowski, M. Jones, M. Roqatcheva, K. Kanack and K. Bourzac (2015). "Multicenter evaluation of the BioFire FilmArray gastrointestinal panel for etiologic diagnosis of infectious gastroenteritis." Journal of Clinical Microbiology **53**(3): 915-925.
- Centers for Disease Control and Prevention (2006). "Epidemiology of HIV/AIDS -- United States, 1981-2005." Morbidity and Mortality Weekly Report **55**(21): 589-592.
- Cohen, D., H. Korin, R. Bassal, M. P. Markovich, Y. Sivan, S. Goren and K. Muhsen (2019). "Burden and risk factors of *Shigella sonnei* shigellosis among children aged 0-59 months in hyperendemic communities in Israel." International Journal of Infectious Diseases **82**: 117-123.
- Greene, S. K., A. Levin-Rector, J. L. Hadler and A. D. Fine (2015). "Disparities in Reportable Communicable Disease Incidence by Census Tract-Level Poverty, New York City, 2006--2013." American Journal of Public Health **105**(9): e27-e34.
- Heffernan, R., F. Mostashari, D. Das, A. Karpati, M. Kulldorf and D. Weiss (2004). "Syndromic Surveillance in Public Health Practice, New York City." Emerging Infectious Disease **10**(5): 858 -- 864.
- Hellard, M., J. Hocking, J. Willis, G. Dore and C. Fairley (2003). "Risk factors leading to *Cryptosporidium* infection in men who have sex with men." Sexually Transmitted Infections **79**: 412-414.

Huang, J. Y., O. L. Henao, P. M. Griffin, D. J. Vugia, A. B. Cronquist, S. Hurd, M. Tobin-D'Angelo, P. Ryan, K. Smith, S. Lathrop, S. Zansky, P. R. Cislak, J. Dunn, K. G. Holt, B. J. Wolpert and M. E. Patrick (2016). "Infection with Pathogens Transmitted Commonly Through Food and the Effect of Increasing Use of Culture-Independent Diagnostic Tests on Surveillance - Foodborne Diseases Active Surveillance Network, 10 U.S. Sites, 2012-2015." Morbidity and Mortality Weekly Report **65**(14): 368-371.

Hutwagner, L., E. Maloney, N. Bean, L. Slutsker and S. Martin (1997). "Using Laboratory-Based Surveillance Data for Prevention: An Algorithm for Detecting *Salmonella* Outbreaks." Emerging Infectious Disease **3**(3): 395-400.

Kean, B. H., D. C. William and S. K. Luminais (1979). "Epidemic of amoebiasis and giardiasis in a biased population." British Journal of Venereal Diseases **55**(5): 375-378.

Klein, R. J. and C. A. Schoenborn (2001). Age Adjustment Using the 2000 Projected U.S. Population. Hyattsville, Maryland, Centers for Disease Control and Prevention, National Center for Health Statistics.

Madison-Antenucci, S., R. F. Relich, L. Doyle, N. Espina, D. Fuller, T. Karchmer, A. Lainesse, J. E. Mortensen, P. Pancholi, W. Veros and S. M. Harrington (2016). "Multicenter Evaluation of BD Max Enteric Parasite Real-Time PCR Assay for Detection of *Giardia duodenalis*, *Cryptosporidium hominis*, *Cryptosporidium parvum*, and *Entamoeba histolytica*." Journal of Clinical Microbiology **54**(11): 2681-2688.

Marder, E. P., P. R. Cieslak, A. B. Coquist, J. Dunn, S. Lathrop, T. Rabatsky-Ehr, P. Ryan, K. Smith, M. Tobin-D'Angelo, D. J. Vugia, S. Zansky, K. G. Holt, B. J. Wolpert, M. Lynch, R. Tauxe and A. L. Geissler (2017). "Incidence and Trends of Infections with Pathogens Transmitted Commonly Through Food and the Effect of Increasing Use of Culture-Independent Diagnostic Tests on Surveillance -- Foodborne Diseases Active Surveillance Network, 10 U.S. Sites, 2013-2016." Morbidity and Mortality Weekly Report **66**(15).

Mitchell, H. and G. Hughes (2018). "Recent Epidemiology of Sexually Transmissible Enteric Infections in Men Who Have Sex With Men." Current Opinion in Infectious Diseases **31**(1): 50-56.

Navidad, J. F., D. J. Griswold, M. S. Gradus and S. Bhattacharyya (2013). "Evaluation of Luminex xTAG Gastrointestinal Pathogen Analyte-Specific Reagents for High-Throughput Simultaneous Detection of Bacteria, Viruses, and Parasites of Clinical and Public Health Importance." Journal of Clinical Microbiology **51**(9): 3018-3024.

New York City Department of City Planning (2010). Decennial Census - Census 2010; <https://www1.nyc.gov/site/planning/data-maps/nyc-population/census-2010.page>.

New York City Department of Health and Mental Hygiene. (2020). "Communicable Disease Surveillance Data; <https://a816-health.nyc.gov/hdi/epiquery/>."

New York City Health Code (2019). Article 11: Reportable Diseases and Conditions; <https://www1.nyc.gov/assets/doh/downloads/pdf/about/healthcode/health-code-article11.pdf>.

Peterson, E. R., A. M. Fireteanu and S. K. Greene (2018). Adapting Reportable Disease Cluster Detection Methods for Increased Use of Culture-Independent Diagnostic Testing;

<https://cste.confex.com/cste/2018/meetingapp.cgi/Paper/9135>. Council of State and Territorial Epidemiologists. West Palm Beach, Florida.

Phillips, S. C., D. Mildvan, D. C. William, A. M. Gelb and M. C. White (1981). "Sexual Transmission of Enteric Protozoa and Helminths in a Venereal-Disease-Clinic Population." The New England Journal of Medicine **305**(11): 603-606.

Pilon, P., B. Camara and S. Bekal (2016). "Outbreak of *Shigella sonnei* in Montreal's ultra-Orthodox Jewish community, 2015." Canadian Communicable Disease Reports **42**: 86-90.

Poznansky, M. C., R. Coker, C. Skinner, A. Hill, S. Bailey, L. Whitaker, A. Renton and J. Weber (1995). "HIV positive patients first presenting with an AIDS defining illness: characteristics and survival." British Medical Journal **311**(6998): 156-158.

Rashmi, K. S. and K. L. R. Kumar (2013). "Intestinal Cryptosporidiosis and the Profile of the CD4 Counts in a Cohort of HIV Infected Patients." Journal of Clinical and Diagnostic Research **7**(6): 1016-1020.

6. TABLES AND FIGURES

Table 3: Giardiasis, number of cases and annual case rate per 100,000 population (in parentheses) by sex and borough of residence, New York City, 2019.

| Sex | Borough of residence | | | | | |
|--------|----------------------|---------------|---------------|---------------|---------------|---------------|
| | Citywide | Manhattan | Bronx | Brooklyn | Queens | Staten Island |
| Male | 865 (21.6) | 357 (46.3) | 114 (16.9) | 213 (17.4) | 160 (14.5) | 21 (9.1) |
| Female | 340 (7.7) | 95 (11.1) | 50 (6.6) | 109 (8.0) | 76 (6.5) | 10 (4.1) |
| Total | 1,205 (14.3) | 452 (27.8) | 164 (11.5) | 322 (12.5) | 236 (10.4) | 31 (6.5) |

Table 4: Giardiasis, number of cases and annual case rate per 100,000 by United Hospital Fund neighborhood of residence, New York City, 2019.

| United Hospital Fund Neighborhood | Borough | Number Of Cases | Population | Case Rate |
|-----------------------------------|---------------|-----------------|------------|-----------|
| Chelsea-Clinton | Manhattan | 101 | 149438 | 67.6 |
| Greenwich Village-Soho | Manhattan | 27 | 84055 | 32.1 |
| Gramercy Park-Murray Hill | Manhattan | 40 | 131358 | 30.5 |
| Greenpoint | Brooklyn | 40 | 135944 | 29.4 |
| Washington Heights-Inwood | Manhattan | 79 | 269142 | 29.4 |
| Union Sq-Lower East Side | Manhattan | 43 | 189305 | 22.7 |
| Upper West Side | Manhattan | 49 | 216284 | 22.7 |
| Upper East Side | Manhattan | 47 | 219079 | 21.5 |
| Lower Manhattan | Manhattan | 13 | 60936 | 21.3 |
| Long Island City-Astoria | Queens | 43 | 208828 | 20.6 |
| C Harlem-Morningside Hgts | Manhattan | 34 | 179078 | 19.0 |
| Downtown Heights-Slope | Brooklyn | 49 | 259325 | 18.9 |
| Williamsburg-Bushwick | Brooklyn | 39 | 217865 | 17.9 |
| Hunts Point-Mott Haven | Bronx | 25 | 141190 | 17.7 |
| East Harlem | Manhattan | 19 | 113268 | 16.8 |
| Borough Park | Brooklyn | 48 | 337153 | 14.2 |
| Bedford Stuyvesant-Crown Hgts | Brooklyn | 45 | 325715 | 13.8 |
| West Queens | Queens | 59 | 457316 | 12.9 |
| Fordham-Bronx Park | Bronx | 32 | 261273 | 12.2 |
| High Bridge-Morrisania | Bronx | 26 | 218196 | 11.9 |
| Bayside-Littleneck | Queens | 10 | 87432 | 11.4 |
| Crotona-Tremont | Bronx | 24 | 215942 | 11.1 |
| Ridgewood-Forest Hills | Queens | 26 | 251726 | 10.3 |
| Fresh Meadows | Queens | 10 | 99867 | 10.0 |
| Pelham-Throgs Neck | Bronx | 30 | 304035 | 9.9 |
| Northeast Bronx | Bronx | 19 | 201522 | 9.4 |
| Bensonhurst-Bay ridge | Brooklyn | 19 | 204630 | 9.3 |
| Coney Island-Sheepshead Bay | Brooklyn | 26 | 288894 | 9.0 |
| Kingsbridge-Riverdale | Bronx | 8 | 92535 | 8.6 |
| Flushing-Clearview | Queens | 21 | 253235 | 8.3 |
| East New York | Brooklyn | 15 | 184764 | 8.1 |
| East Flatbush-Flatbush | Brooklyn | 24 | 297255 | 8.1 |
| Southwest Queens | Queens | 23 | 284964 | 8.1 |
| South Beach-Tottenville | Staten Island | 14 | 191845 | 7.3 |
| Southeast Queens | Queens | 15 | 211955 | 7.1 |
| Jamaica | Queens | 22 | 316627 | 6.9 |
| Willowbrook | Staten Island | 6 | 90834 | 6.6 |
| Stapleton-St.George | Staten Island | 8 | 124623 | 6.4 |
| Canarsie-Flatlands | Brooklyn | 12 | 204565 | 5.9 |
| Rockaway | Queens | 6 | 121153 | 5.0 |
| Port Richmond | Staten Island | 3 | 68877 | 4.4 |
| Sunset Park | Brooklyn | 3 | 126721 | 2.4 |

Note: this table does not include three cases of giardiasis in which UHF neighborhood could not be determined.

Table 5: Giardiasis, number of cases and annual case rate per 100,000 population (in parentheses) by age group and sex, New York City, 2019.

| Age Group | Sex | | |
|--------------|-------------------------|-----------------------|----------------------|
| | Total | Male | Female |
| <5 years | 72 (13.5) | 38 (13.9) | 34 (13.0) |
| 5–9 years | 82 (16.9) | 48 (19.3) | 34 (14.3) |
| 10–19 years | 86 (9.6) | 54 (11.9) | 32 (7.2) |
| 20–44 years | 604 (19.0) | 468 (30.2) | 136 (8.3) |
| 45–59 years | 224 (14.2) | 175 (23.3) | 49 (5.9) |
| ≥ 60 years | 137 (7.9) | 82 (11.2) | 55 (5.5) |
| Total | 1,205 (14.3) | 865 (21.6) | 339 (7.7) |

Table 6: Giardiasis, number of cases and annual case rate per 100,000 population (in parentheses) by age group and borough of residence, New York City, 2019.

| Age Group | Borough of residence | | | | | |
|-------------|----------------------|---------------|---------------|---------------|---------------|---------------|
| | Citywide | Manhattan | Bronx | Brooklyn | Queens | Staten Island |
| <5 years | 72 (13.5) | 9 (11.7) | 18 (17.6) | 26 (13.9) | 18 (12.7) | 1 (3.6) |
| 5–9 years | 82 (16.9) | 14 (21.6) | 26 (25.8) | 25 (15.2) | 16 (12.6) | 1 (3.5) |
| 10–19 years | 86 (9.6) | 16 (12.7) | 18 (9.5) | 20 (7.0) | 29 (12.4) | 3 (5.1) |
| 20–44 years | 604 (19.0) | 253 (35.7) | 61 (11.8) | 171 (17.2) | 105 (13.0) | 14 (9.2) |
| 45–59 years | 224 (14.2) | 108 (36.7) | 28 (10.5) | 47 (10.4) | 35 (7.5) | 6 (6.0) |
| ≥ 60 years | 137 (7.9) | 52 (14.5) | 13 (5.0) | 33 (6.6) | 33 (6.6) | 6 (5.5) |
| Total | 1,205 (14.3) | 452 (27.8) | 164 (11.5) | 322 (12.5) | 236 (10.4) | 31 (6.5) |

Table 7: Giardiasis, number of cases and case rates by census tract poverty level, New York City, 2019.

| Census Tract Poverty Level | Number of cases | Case Rate per 100,000 | Age adjusted rate |
|----------------------------|-----------------|-----------------------|-------------------|
| Low ^a | 321 | 13.8 | 19.1 |
| Medium ^b | 411 | 15.6 | 22.1 |
| High ^c | 267 | 15.4 | 20.5 |
| Very high ^d | 202 | 11.8 | 14.6 |

Poverty levels are defined by the American Community Survey, 2013–2017 and are defined as the proportion of residents that have household incomes below 100% of the federal poverty level: ^aLow poverty: <10%; ^b Medium poverty: 10–19%; ^c High poverty: 20–29%; ^d Very high poverty: ≥30%.

Note: Four cases (0.3%) were excluded from this table because geolocating information for census tract identification was unavailable.

Table 8: Cryptosporidiosis, number of cases and annual case rate per 100,000 population (in parentheses) by sex and borough of residence, New York City, 2019.

| Sex | Borough of residence | | | | | |
|--------|----------------------|--------------|-------------|--------------|-------------|---------------|
| | Citywide | Manhattan | Bronx | Brooklyn | Queens | Staten Island |
| Male | 221 (5.5) | 79 (10.2) | 36 (5.3) | 66 (5.4) | 36 (3.3) | 4 (1.7) |
| Female | 174 (4.0) | 51 (5.9) | 21 (2.8) | 67 (4.9) | 32 (2.7) | 3 (1.2) |
| Total | 395 (4.7) | 130 (8.0) | 57 (4.0) | 133 (5.1) | 68 (3.0) | 7 (1.5) |

Table 9: Cryptosporidiosis, number of cases and annual case rate per 100,000 population by United Hospital Fund neighborhood of residence, New York City, 2019

| United Hospital Fund Neighborhood | Borough | Number Of Cases | Population | Case Rate |
|-----------------------------------|---------------|-----------------|------------|-----------|
| Greenpoint | Brooklyn | 23 | 135944 | 16.9 |
| Upper East Side | Manhattan | 28 | 219079 | 12.8 |
| Chelsea-Clinton | Manhattan | 19 | 149438 | 12.7 |
| Borough Park | Brooklyn | 34 | 337153 | 10.1 |
| Greenwich Village-Soho | Manhattan | 8 | 84055 | 9.5 |
| Downtown Heights-Slope | Brooklyn | 24 | 259325 | 9.3 |
| Lower Manhattan | Manhattan | 5 | 60936 | 8.2 |
| Williamsburg-Bushwick | Brooklyn | 17 | 217865 | 7.8 |
| Gramercy Park-Murray Hill | Manhattan | 10 | 131358 | 7.6 |
| Washington Heights-Inwood | Manhattan | 20 | 269142 | 7.4 |
| Fordham-Bronx Park | Bronx | 17 | 261273 | 6.5 |
| Union Sq-Lower East Side | Manhattan | 12 | 189305 | 6.3 |
| Upper West Side | Manhattan | 13 | 216284 | 6.0 |
| Ridgewood-Forest Hills | Queens | 15 | 251726 | 6.0 |
| CHarlem-Morningside Hgts | Manhattan | 9 | 179078 | 5.0 |
| Hunts Point-Mott Haven | Bronx | 7 | 141190 | 5.0 |
| Long Island City-Astoria | Queens | 10 | 208828 | 4.8 |
| Kingsbridge-Riverdale | Bronx | 4 | 92535 | 4.3 |
| West Queens | Queens | 19 | 457316 | 4.2 |
| Crotona-Tremont | Bronx | 8 | 215942 | 3.7 |
| East Harlem | Manhattan | 4 | 113268 | 3.5 |
| Northeast Bronx | Bronx | 7 | 201522 | 3.5 |
| East Flatbush-Flatbush | Brooklyn | 10 | 297255 | 3.4 |
| Bedford Stuyvesant-Crown Hgts | Brooklyn | 9 | 325715 | 2.8 |
| High Bridge-Morrisania | Bronx | 6 | 218196 | 2.7 |
| East New York | Brooklyn | 5 | 184764 | 2.7 |
| Flushing-Clearview | Queens | 6 | 253235 | 2.4 |
| Pelham-Throgs Neck | Bronx | 7 | 304035 | 2.3 |
| South Beach-Tottenville | Staten Island | 4 | 191845 | 2.1 |
| Canarsie-Flatlands | Brooklyn | 4 | 204565 | 2.0 |
| Southeast Queens | Queens | 4 | 211955 | 1.9 |
| Southwest Queens | Queens | 5 | 294964 | 1.8 |
| Rockaway | Queens | 2 | 121153 | 1.7 |
| Stapleton-St.George | Staten Island | 2 | 124623 | 1.6 |
| Jamaica | Queens | 5 | 316627 | 1.6 |
| Port Richmond | Staten Island | 1 | 68877 | 1.5 |
| Coney Island-Sheepshead Bay | Brooklyn | 4 | 288894 | 1.4 |
| Bayside-Littleneck | Queens | 1 | 87432 | 1.1 |
| Fresh Meadows | Queens | 1 | 99867 | 1.0 |
| Bensonhurst-Bay Ridge | Brooklyn | 2 | 204630 | 1.0 |
| Sunset Park | Brooklyn | 1 | 126721 | 0.8 |

Note: This table does not include three cases of cryptosporidiosis in which UHF neighborhood could not be determined.

Table 10: Cryptosporidiosis, number of cases and annual case rate per 100,000 population (in parentheses) by age group and sex, New York City, 2019.

| Age Group | Sex | | |
|-------------|--------------|--------------|--------------|
| | Total | Male | Female |
| <5 years | 47 (8.8) | 25 (9.1) | 22 (8.4) |
| 5–9 years | 25 (5.1) | 11 (4.4) | 14 (5.9) |
| 10–19 years | 33 (3.7) | 20 (4.4) | 13 (2.9) |
| 20–44 years | 209 (6.6) | 116 (7.5) | 93 (5.7) |
| 45–59 years | 45 (2.9) | 30 (4.0) | 15 (1.8) |
| ≥ 60 years | 36 (2.1) | 19 (2.6) | 17 (1.70) |
| Total | 395 (4.7) | 221 (5.5) | 174 (4.0) |

Table 11: Cryptosporidiosis, number of cases and annual case rate per 100,000 population (in parentheses) by age group and borough, New York City, 2019.

| Age Group | Borough of residence | | | | | |
|-------------|----------------------|--------------|-------------|--------------|-------------|---------------|
| | Citywide | Manhattan | Bronx | Brooklyn | Queens | Staten Island |
| <5 years | 47 (8.8) | 3 (3.9) | 4 (3.9) | 26 (13.9) | 14 (9.9) | 0 |
| 5–9 years | 25 (5.1) | 0 | 6 (5.9) | 15 (9.1) | 4 (3.2) | 0 |
| 10–19 years | 33 (3.7) | 5 (4.0) | 4 (2.1) | 14 (4.9) | 10 (4.3) | 0 |
| 20–44 years | 209 (6.6) | 87 (12.3) | 32 (6.2) | 64 (6.4) | 22 (2.7) | 4 (2.6) |
| 45–59 years | 45 (2.9) | 19 (6.5) | 8 (3.0) | 9 (2.0) | 8 (1.7) | 1 (1.0) |
| ≥ 60 years | 36 (2.1) | 16 (4.5) | 3 (1.2) | 5 (1.0) | 10 (2.0) | 2 (1.8) |
| Total | 395 (4.7) | 130 (8.0) | 57 (4.0) | 133 (5.1) | 68 (3.0) | 7 (1.5) |

Table 12: Cryptosporidiosis, number of cases and annual case rate per 100,000 population (in parentheses) by race/ethnicity and borough of residence, New York City, 2019

| Race/Ethnicity | Borough of residence | | | | | |
|--|----------------------|----------------------|---------------------|----------------------|---------------------|--------------------|
| | Citywide | Manhattan | Bronx | Brooklyn | Queens | Staten Island |
| Hispanic | 103 (4.2) | 24 (5.7) | 24 (3.0) | 19 (3.9) | 36 (5.6) | 0 |
| White, non-Hispanic | 175 (6.5) | 74 (9.7) | 3 (2.3) | 80 (8.5) | 14 (2.5) | 4 (1.4) |
| Black/African American, non-Hispanic | 61 (3.3) | 10 (4.9) | 23 (5.5) | 18 (2.3) | 9 (2.2) | 1 (2.2) |
| Asian, non-Hispanic | 16 (1.3) | 9 (4.4) | 1 (1.7) | 2 (0.6) | 3 (0.5) | 1 (2.1) |
| Pacific Islander, Native Hawaiian, American Indian, non-Hispanic | 0 | 0 | 0 | 0 | 0 | 0 |
| Two or more races, other, non-Hispanic | 11 (7.3) | 2 (6.0) | 3 (21.3) | 4 (8.2) | 1 (2.1) | 1 (14.1) |
| Unknown | 29 | 11 | 3 | 10 | 5 | 0 |
| Total | 395 (4.7) | 130 (8.0) | 57 (4.0) | 133 (5.1) | 68 (3.0) | 7 (1.5) |

Table 13: Cryptosporidiosis, number of cases and annual case rate per 100,000 population (in parentheses) by race/ethnicity and age group, New York City, 2019.

| Race/Ethnicity | Age group | | | | | | |
|--|----------------------|---------------------|---------------------|---------------------|----------------------|---------------------|---------------------|
| | Total | <5 years | 5–9 years | 10–19 years | 20–44 years | 45–59 years | ≥ 60 years |
| Hispanic | 103 (4.2) | 10 (5.4) | 10 (5.7) | 13 (4.1) | 44 (4.7) | 16 (3.6) | 10 (2.6) |
| White, non-Hispanic | 175 (6.5) | 27 (18.2) | 11 (8.7) | 15 (6.7) | 92 (8.9) | 13 (2.7) | 17 (2.5) |
| Black/African American, non-Hispanic | 61 (3.3) | 3 (2.8) | 1 (1.0) | 2 (0.9) | 38 (5.8) | 11 (2.9) | 6 (1.5) |
| Asian, non-Hispanic | 16 (1.3) | 0 | 1 (1.6) | 2 (1.8) | 8 (1.6) | 3 (1.2) | 2 (0.8) |
| Pacific Islander, Native Hawaiian, American Indian, non-Hispanic | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Two or more races, other, non-Hispanic | 11 (7.3) | 1 (4.7) | 1 (5.9) | 0 | 8 (14.6) | 1 (4.9) | 0 |
| Unknown | 29 | 6 | 1 | 1 | 19 | 1 | 1 |
| Total | 395 (4.7) | 47 (8.8) | 25 (5.1) | 33 (3.7) | 209 (6.6) | 45 (2.9) | 36 (2.1) |

Table 14: Cryptosporidiosis, number of cases and case rates by census tract poverty level, New York City, 2019.

| Census Tract Poverty Level | Number of cases | Case Rate per 100,000 | Age adjusted rate |
|----------------------------|-----------------|-----------------------|-------------------|
| Low ^a | 110 | 4.7 | 6.1 |
| Medium ^b | 99 | 3.8 | 4.4 |
| High ^c | 65 | 3.7 | 5.1 |
| Very high ^d | 118 | 6.9 | 7.9 |

Poverty levels are defined by the American Community Survey, 2013–2017 and are defined as the proportion of residents that have household incomes below 100% of the federal poverty level: ^aLow poverty: <10%; ^bMedium poverty: 10–19%; ^cHigh poverty: 20–29%; ^dVery high poverty: ≥30%.

Note: Three cases (0.38%) were excluded from this table because geolocating information for census tract identification was unavailable

Table 15: Percentage of interviewed **cryptosporidiosis** patients reporting selected potential risk exposures before disease onset, persons with HIV/AIDS, New York City 1995–2019, median (range).

| Exposure Type ^a | Persons with HIV/AIDS | | | | | |
|---|-----------------------|------------------|------------------|------------------|------------------|------|
| | 1995–1999 | 2000–2004 | 2005–2009 | 2010–2014 | 2015–2018 | 2019 |
| Contact with an animal ^b | 25% (33%–36%) | 40% (24%–43%) | 38% (31%–44%) | 34% (20%–43%) | 37% (25–45%) | 32% |
| High-risk sexual activity ^c (aged ≥ 18 years) | 20% (9%–22%) | 24% (16–34%) | 31% (21%–39%) | 17% (7%–25%) | 37% (21%–42%) | 39% |
| International travel ^d | 9% (9%–18%) | 13% (10%–15%) | 8% (6%–17%) | 6% (4%–13%) | 10% (7%–13%) | 11% |
| Recreational water contact ^e | 16% (8%–16%) | 13% (8%–21%) | 14% (5%–18%) | 10% (4%–14%) | 10% (5%–13%) | 5% |

Note:

Determination of an association between exposure to possible risk factors for cryptosporidiosis and acquisition of cryptosporidiosis cannot be made without reference to a suitable control population (i.e., non-*Cryptosporidium*-infected controls).

The format of the patient interview form changed in 1997, 2001, 2002 and 2010:

a: From January 1, 1995 to April 25, 2010, patients were asked about potential risk exposures during the month before disease onset. Beginning April 26, 2010, patients were asked about potential risk exposures during the 14 days before onset.

b: Contact with an animal: includes having a pet, or visiting a farm or petting zoo (1995–1996); expanded to include: visiting a pet store, or veterinarian office (1997–2012); or other animal exposure (2019).

c: High-risk sexual activity: includes having a penis, finger or tongue in a sexual partner’s anus (1995–2019)

d: International travel: travel outside of the United States (1995–2019)

e: Recreational water contact: includes swimming in a pool, or swimming in or drinking from a stream, lake, river or spring (1995–1996); expanded to include: swimming in the ocean or visiting a recreational water park (1997–2012); swimming in a hot tub or swimming or drinking water from a pond or body of water (2019).

Table 16: Percentage of interviewed **cryptosporidiosis** patients reporting selected potential risk exposures before disease onset, immunocompetent persons, New York City, 1995–2019, median (range).

| Exposure Type ^a | Immunocompetent persons | | | | | |
|--|-------------------------|------------------|------------------|------------------|------------------|------|
| | 1995–1999 | 2000–2004 | 2005–2009 | 2010–2014 | 2015–2018 | 2019 |
| Contact with an animal ^b | 35% (7%–41%) | 34% (23%–37%) | 36% (28%–40%) | 34% (18%–41%) | 31% (30%–41%) | 33% |
| High-risk sexual activity ^c (aged ≥18 years) | 12% (10%–25%) | 23% (13%–31%) | 17% (7%–19%) | 8% (4%–11%) | 13% (14%–29%) | 10% |
| International travel ^d | 28% (26%–30%) | 45% (33%–47%) | 45% (37%–52%) | 44% (35%–62%) | 43% (41%–45%) | 39% |
| Recreational water contact ^e | 24% (21%–40%) | 34% (32%–35%) | 40% (28%–52%) | 35% (32%–48%) | 34% (26%–39%) | 30% |

Note:

Determination of an association between exposure to possible risk factors for cryptosporidiosis and acquisition of cryptosporidiosis cannot be made without reference to a suitable control population (i.e., non-*Cryptosporidium*-infected controls).

The format of the patient interview form changed in 1997, 2001, 2002 and 2010:

a: From January 1, 1995 to April 25, 2010, patients were asked about potential risk exposures during the month before disease onset. Beginning April 26, 2010, patients were asked about potential risk exposures during the 14 days before onset.

b: Contact with an animal: includes having a pet, or visiting a farm or petting zoo (1995–1996); expanded to include: visiting a pet store, or veterinarian office (1997–2012); or other animal exposure (2019).

c: High-risk sexual activity: includes having a penis, finger or tongue in a sexual partner's anus (1995–2019)

d: International travel: travel outside of the United States (1995–2019)

e: Recreational water contact: includes swimming in a pool, or swimming in or drinking from a stream, lake, river or spring (1995–1996); expanded to include: swimming in the ocean or visiting a recreational water park (1997–2012); swimming in a hot tub or swimming or drinking water from a pond or body of water (2019).

Table 17: Percentage of interviewed **cryptosporidiosis** patients by type of tap water exposure before disease onset, persons with HIV/AIDS, New York City, 1995–2019, median (range).

| Exposure Type ^a | Persons with HIV/AIDS | | | | | |
|--|-----------------------|------------------|------------------|------------------|------------------|------|
| | 1995–1999 | 2000–2004 | 2005–2009 | 2010–2014 | 2015–2018 | 2019 |
| Plain tap ^b | 69% (64%–71%) | 55% (49%–77%) | 67% (58%–76%) | 63% (50%–71%) | 53% (46%–63%) | 47% |
| Filtered tap ^c | 12% (9%–20%) | 20% (13%–22%) | 14% (7%–18%) | 11% (8%–25%) | 11% (8%–15%) | 24% |
| Boiled tap ^d | 5% (3%–7%) | 6% (0%–6%) | 7% (0%–11%) | 4% (2%–11%) | 2% (0%–8%) | 3% |
| Incidental plain tap only ^e | 15% (8%–16%) | 15% (4%–19%) | 10% (4%–17%) | 18% (8%–20%) | 24% (13%–24%) | 24% |
| No tap ^f | 2% (0%–5%) | 4% (2%–6%) | 2% (0%–6%) | 4% (0%–4%) | 5% (0%–13%) | 3% |

Note:

Determination of an association between exposure to possible risk factors for cryptosporidiosis and acquisition of cryptosporidiosis cannot be made without reference to a suitable control population (i.e., non-*Cryptosporidium*-infected controls).

The format of the patient interview form changed in 1997, 2001, 2002 and 2010:

a: From January 1, 1995 to April 25, 2010, patients were asked about tap water exposure during the month before disease onset. Beginning April 26, 2010, patients were asked about tap water exposure during the 14 days before onset.

b: Plain tap: drank unboiled/unfiltered NYC tap water (1995–5/10/2001) or drank greater than 0 cups of unboiled/unfiltered NYC tap water (5/11/2001–12/31/2013).

c: Filtered tap: drank filtered NYC tap water (1995–5/10/2001) or drank greater than 0 cups of filtered NYC tap water, and 0 or more cups of boiled NYC tap water, and no unboiled/unfiltered NYC tap water (5/11/2001–12/13/2019).

d: Boiled tap: drank boiled NYC tap water (1995–5/10/2001) or drank greater than 0 cups of boiled NYC tap water, and no unboiled/unfiltered NYC tap water, and no filtered NYC tap water (5/11/2001–12/31/2019).

e: Incidental plain tap only: did not drink any NYC tap water but did use unboiled/unfiltered NYC tap water to brush teeth, or to wash vegetables/fruits, or to make ice (1995–1996), expanded to include make juice from concentrate (1997–2019).

f: No tap: did not drink any NYC tap water and did not use unboiled/unfiltered NYC tap water to brush teeth, or to wash vegetables/fruits, or to make ice (1995–1996); expanded to include make juice from concentrate (1997–2019).

Table 18: Percentage of interviewed **cryptosporidiosis** patients by type of tap water exposure before disease onset, immunocompetent persons, New York City, 1995–2019, median (range).

| Exposure Type ^a | Immunocompetent persons | | | | | |
|--|-------------------------|------------------|------------------|------------------|------------------|------|
| | 1995–1999 | 2000–2004 | 2005–2009 | 2010–2014 | 2015–2018 | 2019 |
| Plain tap ^b | 58% (56%–67%) | 36% (27%–56%) | 30% (27%–47%) | 33% (28%–49%) | 39% (38%–47%) | 31% |
| Filtered tap ^c | 21% (17%–25%) | 31% (17%–44%) | 23% (20%–30%) | 24% (17%–27%) | 23% (11%–26%) | 30% |
| Boiled tap ^d | 8% (3%–11%) | 2% (0%–7%) | 5% (0%–14%) | 2% (0%–7%) | 4% (2%–6%) | 4% |
| Incidental plain tap only ^e | 9% (7%–12%) | 16% (8%–21%) | 25% (14%–28%) | 15% (11%–22%) | 23% (14%–29%) | 23% |
| No tap ^f | 4% (2%–7%) | 9% (2%–21%) | 14% (3%–27%) | 21% (11%–29%) | 13% (12%–14%) | 11% |

Note:

Determination of an association between exposure to possible risk factors for cryptosporidiosis and acquisition of cryptosporidiosis cannot be made without reference to a suitable control population (i.e., non-*Cryptosporidium*-infected controls).

The format of the patient interview form changed in 1997, 2001, 2002 and 2010:

a: From January 1, 1995 to April 25, 2010, patients were asked about tap water exposure during the month before disease onset. Beginning April 26, 2010, patients were asked about tap water exposure during the 14 days before onset.

b: Plain tap: drank unboiled/unfiltered NYC tap water (1995–5/10/2001) or drank greater than 0 cups of unboiled/unfiltered NYC tap water (5/11/2001–12/31/2013).

c: Filtered tap: drank filtered NYC tap water (1995–5/10/2001) or drank greater than 0 cups of filtered NYC tap water, and 0 or more cups of boiled NYC tap water, and no unboiled/unfiltered NYC tap water (5/11/2001–12/13/2019).

d: Boiled tap: drank boiled NYC tap water (1995–5/10/2001) or drank greater than 0 cups of boiled NYC tap water, and no unboiled/unfiltered NYC tap water, and no filtered NYC tap water (5/11/2001–12/31/2019).

e: Incidental plain tap only: did not drink any NYC tap water but did use unboiled/unfiltered NYC tap water to brush teeth, or to wash vegetables/fruits, or to make ice (1995–1996), expanded to include make juice from concentrate (1997–2019).

f: No tap: did not drink any NYC tap water and did not use unboiled/unfiltered NYC tap water to brush teeth, or to wash vegetables/fruits, or to make ice (1995–1996); expanded to include make juice from concentrate (1997–2019).

Figure 6: Emergency Department Syndromic Surveillance, Trends in visits for the diarrhea syndrome, New York City, January 1, 2017–December 31, 2019.

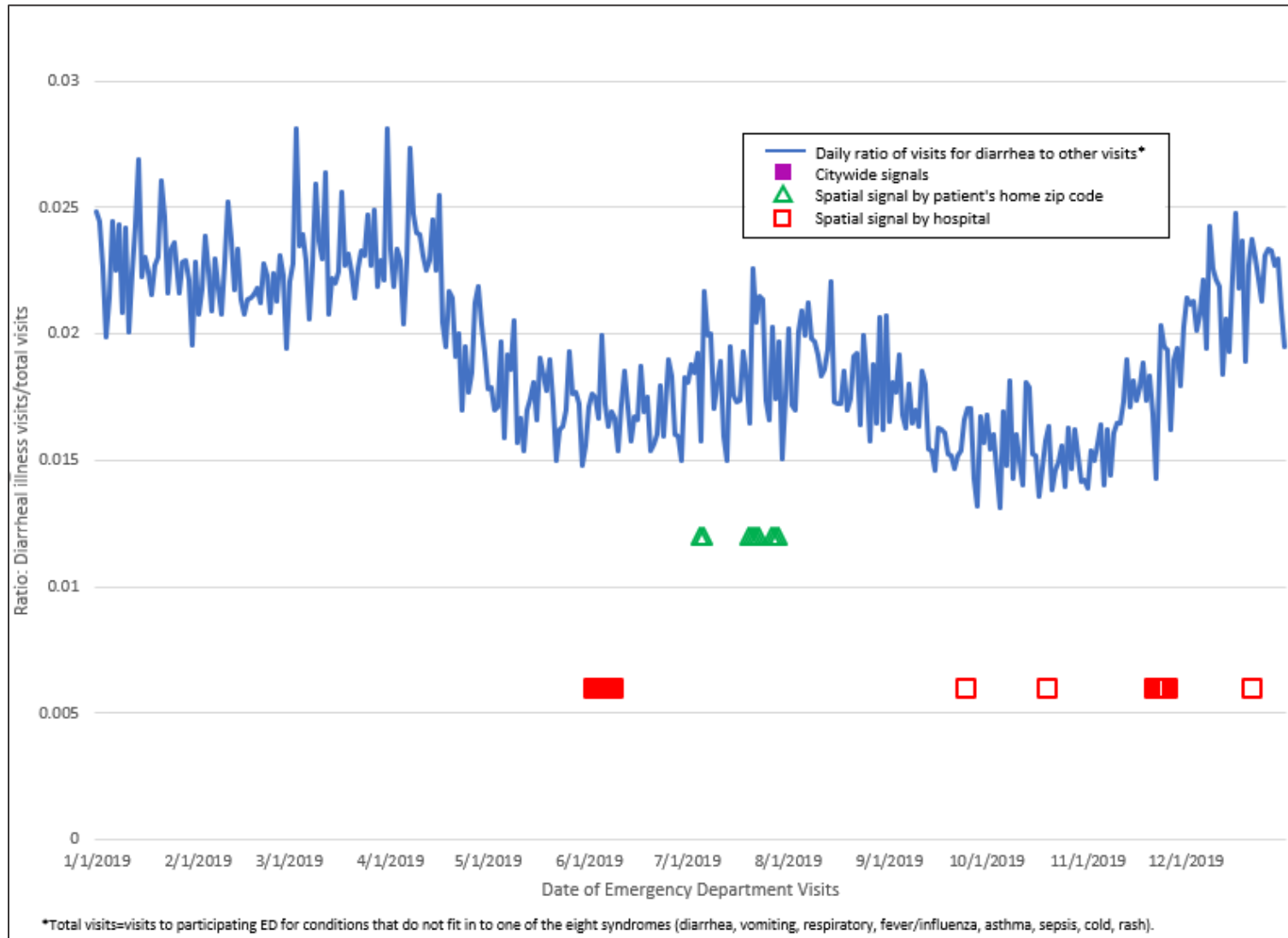


Figure 7: Emergency Department Syndromic Surveillance, Trends in visits for the vomiting syndrome, New York City, January 1, 2017–December 31, 2019.

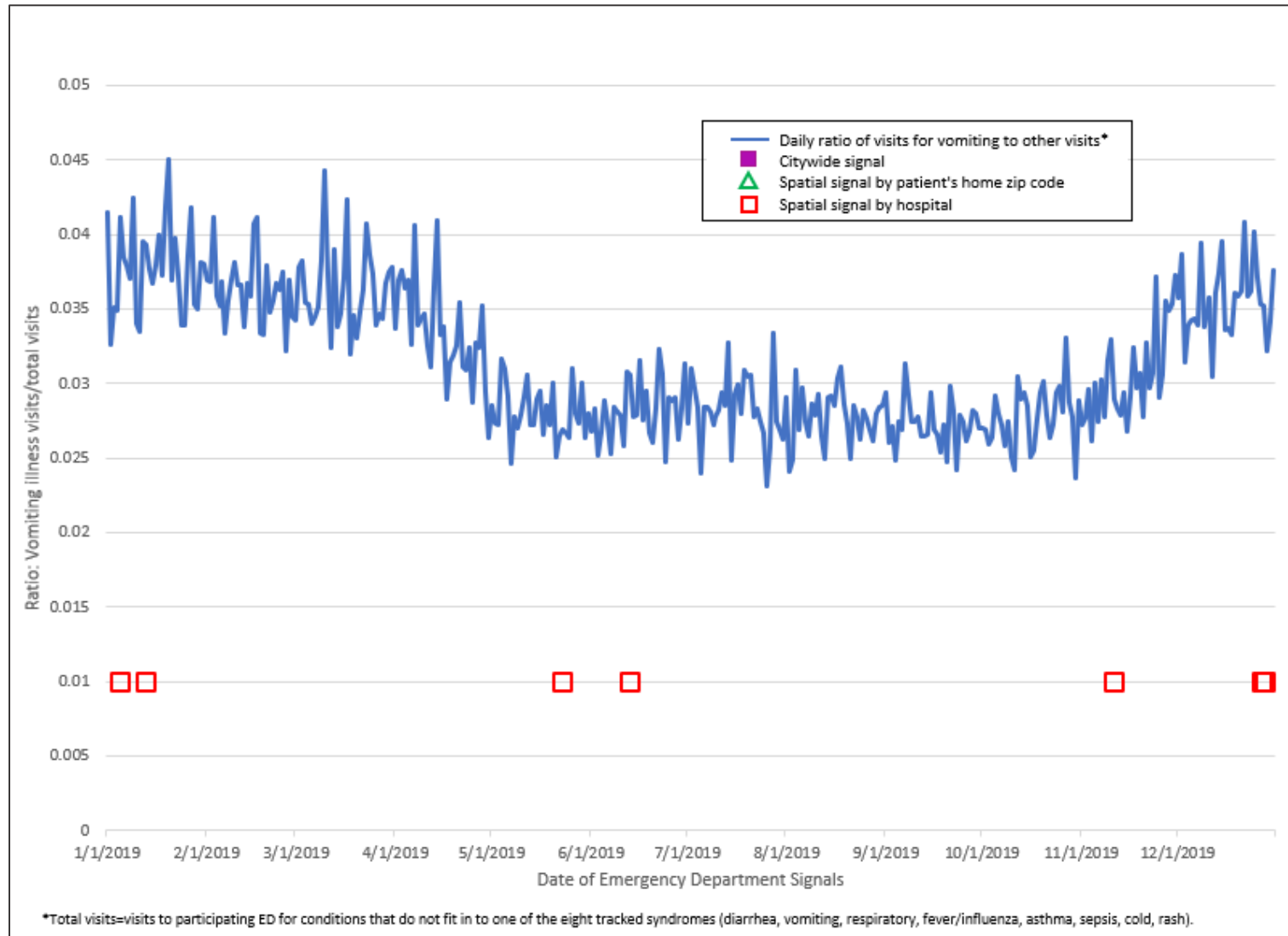
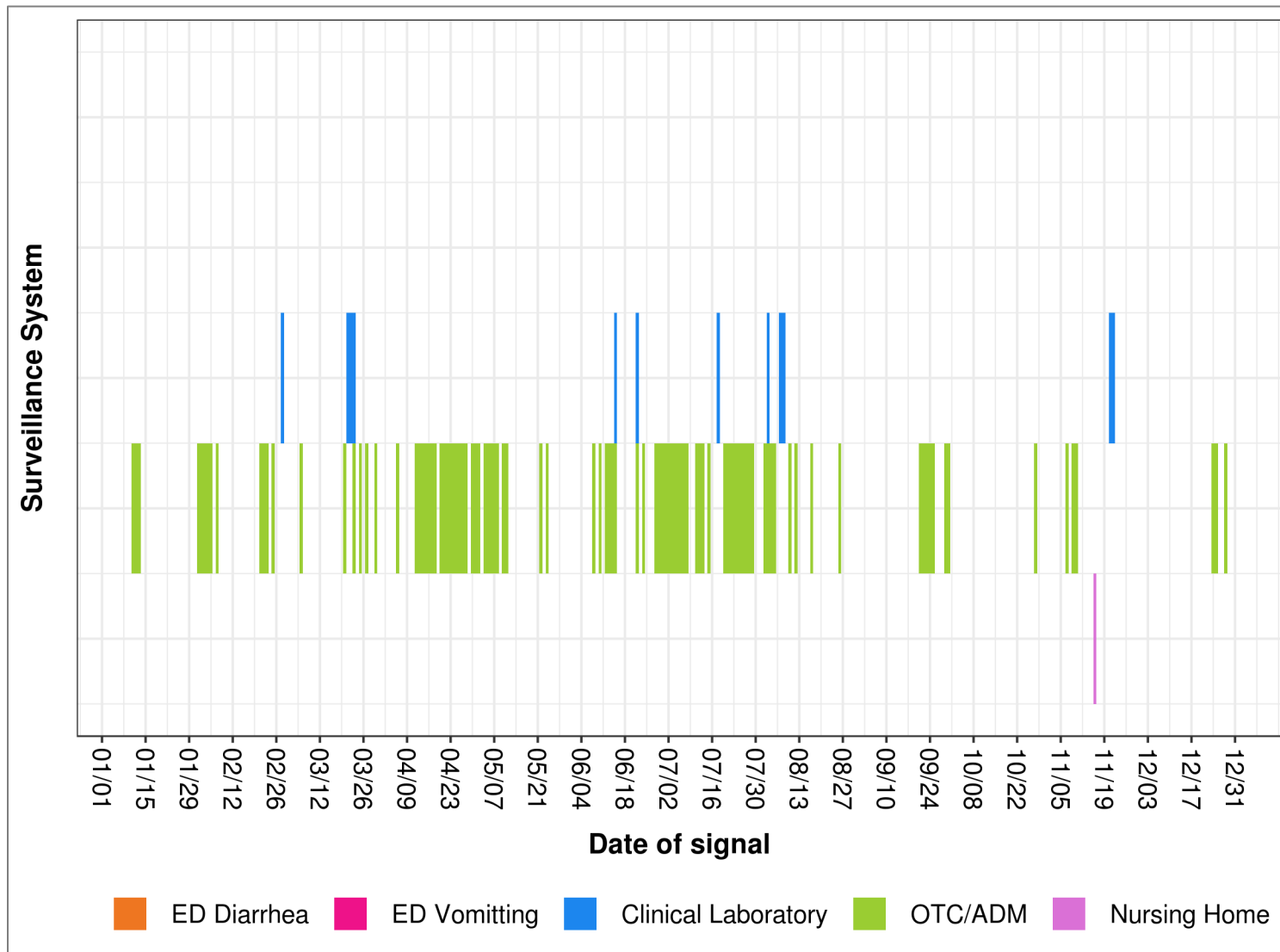


Figure 8: Signals for Gastrointestinal Illness, Syndromic Surveillance Systems, New York City, 2019.



7. APPENDIX A: Information on calculation of rates, case definitions and risk factor collection

Population denominators

The population denominators used to calculate rates were intercensal population estimates for all years except 2000 and 2010 to 2012. For the years 1994 through 1999, intercensal population estimates per year were used based upon linear interpolation between 1990 and 2000 NYC Census. For the years 2001 through 2009 and 2013 through 2019, intercensal population estimates for each year were used from data produced by DOHMH based on the US Census Bureau Population Estimate Program and housing unit data obtained from the NYC Department of City Planning. For 2010 to 2012, the year 2010 NYC Census data were used (New York City Department of City Planning 2010). Because rates for the years 2001 through 2009 and the rates for the years 2014 through 2019 were calculated for this report using intercensal population estimates, they may differ from previously reported rates based on year 2000 and 2010 NY Census data. Other variations in data between this report and previous reports may be because of factors such as disease reporting delays, correction of errors, and refinements in data processing (for example, the removal of duplicate disease reports). All rates in this report are annual rates. Caution must be exercised when interpreting rates based on very small case numbers.

UHF Zones

For mapping purposes, the United Hospital Fund (UHF) neighborhood of patient residence was used. New York City is divided on the basis of zip code into 42 UHF neighborhoods. Maps illustrating annual case rates by UHF neighborhood are included in this report.

Race-Ethnicity Categories

In this report, race/ethnicity-specific case rates for 2019 are based upon intercensal population estimates and include the race/ethnicity categories used by the US Census Bureau Population Estimate Program. Prior to 2011, there was one race/ethnicity category entitled “Asian, Pacific Islander, American Indian, Alaskan Native, non-Hispanic”. Since 2011, separate categories have been used for non-Hispanic Asians, non-Hispanic Pacific Islanders and Native Hawaiians, non-Hispanic American Indian and non-Hispanic of two or more races.

Socioeconomic Status

Beginning with the 2011 WDRAP Annual Report, socioeconomic status (SES) is now included as a measure as part of the demographic description of cases of giardiasis and cryptosporidiosis in NYC. Differences in SES among cases of a disease may indicate economically-related disparities in health. Neighborhood poverty can be used as a proxy for individual SES. The poverty level of the neighborhood of patient resident is measured as the percentage of individuals in the neighborhood who live below the federal poverty level, as reported in census data. Four categories of poverty level were used for the WDRAP analysis (see Tables 7 & 14). Further explanation of how SES designations were made can be found in the 2011–2014 WDRAP Annual reports.

Age-adjusted case rates

Age-adjusted case rates were calculated for each of the four neighborhood poverty levels using direct standardization and weighing by the US 2000 Standard Population. Cases were grouped into three age group categories (aged <24 years, 25–44 years, and ≥45 years) (Klein and Schoenborn 2001).

Confirmed and Probable cases

As was first described in the 2012 Annual Report, confirmed and probable cryptosporidiosis cases are now included in the WDRAP reports. Confirmed cases are those in which the laboratory method used has a high positive predictive value (such as light microscopy of stained slide, enzyme immunoassay, polymerase chain reaction, and direct fluorescent antibody test). Probable cases are those in which the laboratory method used has a low positive predictive value (such as the immunochromatographic card/rapid test) or in which the method used for diagnostic testing was not known. The probable case classification for cryptosporidiosis also includes those cases in which laboratory confirmation was not obtained, but the case was epidemiologically linked to a confirmed case and clinical illness was consistent with cryptosporidiosis. DOHMH BCD reports both confirmed and probable cryptosporidiosis cases to the Centers for Disease Control and Prevention through the National Electronic Telecommunications System for Surveillance. BCD interviews all cases. However, if cases are not confirmed at NYS DOH Wadsworth Center then these patients are not considered to be a case and are not included in the final annual count.

Cryptosporidiosis and Potential Risk Factors

Tables 15, 16, 17, and 18—a change to table format was introduced, starting with the 2015 annual report. This change involves grouping and summarizing data in 5-year sets (e.g., 1995–1999, 2000–2004, etc.). This change was made to continue providing historical data for comparison, and to allow for easier comprehension of trends. Potential risk exposure data for individual years, rather than grouped years, can be viewed in the earlier WDRAP Annual Reports. Only the new data (i.e., the year of the report) is listed independently as a single year.

8. APPENDIX B: Syndromic Surveillance System Descriptions

Hospital Emergency Department (ED) Monitoring

NYC initiated monitoring of hospital ED visits as a public health surveillance system in 2001, and this system has been in operation since that time. Hospitals transmit electronic files each morning containing chief complaint and demographic information for patient visits during the previous 24 hours. Patients are classified into syndrome categories, and daily analyses are conducted to detect any unusual patterns or signals. The two syndromes used to track GI illness are the vomiting syndrome and the diarrhea syndrome. Temporal citywide analyses assess whether the frequency of ED visits for the syndrome has increased in the last one, two, or three days compared to the previous 14 days. Clustering is examined by both hospital location and residential zip code. Statistical significance is based on Monte Carlo probability estimates that adjust for the multiple comparisons inherent in examining many candidate clusters each day. The threshold of significance for citywide and spatial signals was originally set at $p < 0.01$, indicating that less than 1 out of every 100 analyses would generate a cluster due to chance alone. Beginning in 2005, the threshold of significance for spatial signals was changed to $p < 0.005$, while the threshold of significance for citywide signals remained at $p < 0.01$. The system is described further in Heffernan *et al.* (Heffernan, Mostashari et al. 2004).

Anti-Diarrheal Medication Monitoring

NYC began tracking anti-diarrheal drug sales as an indicator of GI illness trends in 1995 via a system operated by DEP. Major modifications and enhancements to NYC's anti-diarrheal medication surveillance program have been made over the years, including: utilization of different data sources, initiation and expansion of DEP's ADM program, initiation of DOHMH's OTC program in 2002, and in 2012, the merger of the ADM and the OTC systems. The ADM and OTC systems were merged to simplify the processing and analysis of pharmacy data, and combine the strengths of the two systems. The combined OTC/ADM system is operated by DOHMH, and the first full year of operation of the merged system was 2013. DOHMH conducted an evaluation of the impact of the merger of the two systems (final report completed in 2014). In 2015, one ADM pharmacy chain data source dropped out of the program, but two additional pharmacy chains were added. Surveillance with both additional pharmacy chains began in 2016.

In summary, the current system involves tracking of sales of over-the-counter, non-bismuth-containing anti-diarrheal medications and of bismuth subsalicylate medications, searching for citywide as well as local signals. DOHMH Bureau of Communicable Disease (BCD) staff review signals on a daily basis to evaluate whether there are any new or sustained signals at citywide and zip-code levels. If there are sustained signals, BCD staff will perform reviews of reportable GI illness, including norovirus and rotavirus, to attempt to rule out a potential waterborne outbreak. Also, information on product promotions (e.g., price discounts) are considered, as these are known to impact on sales volume).

Clinical Laboratory Monitoring System

The number of stool specimens submitted to clinical laboratories for bacterial and parasitic testing also can be a source of information on GI illness trends in the population. The clinical laboratory monitoring system currently collects data from one large laboratory, designated as Laboratory A in this report. The number of participating laboratories has changed over time, as reported in prior WDRAP reports. Laboratory A transmits data by fax to DOHMH BCD 3–4 times per week, indicating the number of stool specimens examined per day for: (a) bacterial culture and sensitivity, (b) ova and parasites, and (c) *Cryptosporidium*.

The Clinical Laboratory Monitoring results are reviewed upon their receipt. Beginning in 2004, DOHMH implemented a model to establish statistical cut-offs for significant increases in clinical laboratory submissions. The model uses the entire historical dataset from November 1995 for Laboratory A. Sundays and holidays are removed because the laboratories do not test specimens on those days. Linear regression is used to adjust for average day-of-week and day-after-holiday effects as certain days routinely have higher volumes than other days. The cumulative sums (CUSUM) method is applied to a two-week baseline to identify statistically significant aberrations (or signals) in submissions for ova and parasites and for bacterial culture and sensitivity. CUSUM is a quality control method that has been adapted for aberration-detection in public health surveillance. CUSUM is described further in Hutwagner, *et al.* (Hutwagner, Maloney et al. 1997).

Nursing Home Sentinel Surveillance

The nursing home surveillance system began in 1997. Under the current protocol, when a participating nursing home documents an outbreak of GI illness that is legally reportable to NYSDOH, the nursing home also notifies the WDRAP team at DOHMH. Such an outbreak is defined as onset of diarrhea and/or vomiting involving three or more patients on a single ward/unit within a seven-day period, or more than expected (baseline) number of cases within a single facility. All participating nursing homes have been provided with stool collection kits in advance. When such an outbreak is noted, specimens are to be collected for testing for bacterial culture and sensitivity, ova and parasites, *Cryptosporidium* spp., viruses, and *Clostridium difficile* toxin. Though *C. difficile* is not a waterborne pathogen, *C. difficile* toxin testing was added in 2010 to address a need expressed by infection control practitioners in the nursing homes, and was intended to help ensure compliance with the sentinel nursing home protocol.

DOHMH BCD staff facilitates transportation of the specimens to the DOHMH Public Health Laboratory, where culture and sensitivity testing is performed. Specimens designated for ova and parasite tests, *Cryptosporidium* as well as for virus and *C. difficile* toxin testing are sent to NYSDOH Wadsworth Center Laboratory. There are currently eight nursing homes participating in the program. Three are in Manhattan, two are in the Bronx, two are in Queens, and one is in Brooklyn. As feedback for their role in outbreak detection, participating nursing homes are provided with copies of the WDRAP annual report.

All participating nursing homes are visited on an annual basis to help ensure compliance with the program protocol. During the site visits, DOHMH staff members reviewed the rationale for the program and program protocol with nursing administration or infection control staff. In addition, the DOHMH staff members verified that the nursing homes had adequate stool collection supplies on hand.