Letter Health Consultation

PASSYUNK SOIL GAS SITE PHILADELPHIA, PHILADELPHIA COUNTY, PENNSYLVANIA

Evaluation of Residential Indoor Air and Sub-Slab Soil Gas Data

January 09, 2018



Pennsylvania Department of Health Division of Environmental Health Epidemiology

A Disclaimer

"This report was supported in part by funds provided through a cooperative agreement with the Agency for Toxic Substances and Disease Registry (ATSDR), U.S. Department of Health and Human Services. The findings and conclusions in these reports are those of the author(s) and do not necessarily represent the views of the ATSDR or the U.S. Department of Health and Human Services. This document has not been revised or edited to conform to ATSDR standards." Pennsylvania Department of Health Division of Environmental Health Epidemiology Harrisburg, PA 717-787-3500

01/09/2018

Ruth Scharr On Site Coordinator U.S. EPA Region 3 Philadelphia, PA

Re: Review of EPA 2014/2016 indoor air and sub-slab soil gas data at the Passyunk Soil Gas Site

Dear Ms. Scharr,

At the request of the U.S. Environmental Protection Agency (EPA), the Pennsylvania Department of Health (DOH) has prepared this letter health consultation to evaluate potential public health issues and to provide relevant conclusions and recommendations based on data you shared with us in April 2017. DOH worked on this evaluation under a cooperative agreement with the Agency for Toxic Substances and Disease Registry (ATSDR).

SITE BACKGROUND AND STATEMENT OF ISSUES

The Passyunk Soil Gas site ("the site") is located on the west side of South Philadelphia. It is an approximately 3.7-acre residential neighborhood bounded by the Schuylkill River and the Philadelphia Gas Works (PGW) Passyunk facility on the west, the Philadelphia Energy Solutions refining complex (formerly Sunoco, Inc. refinery) to the south, and the Schuylkill Expressway to the east/northeast. The neighborhood is predominately residential with a tavern, a playground, and a Philadelphia Mummers hall. There are no daycares or schools within the neighborhood. (See the Appendix A for a map of the site and surrounding area.)

In 2003, PGW began conducting independent environmental studies on their Passyunk Plant property adjacent to the site. Those studies showed elevated concentrations of benzene in the shallow groundwater aquifer beneath their property, confirmed the transport of benzene towards the adjacent residential neighborhood (the site), and confirmed that vapor intrusion was not a concern on PGW plant property (PGW, 2013). Additional investigations conducted after 2010 confirmed their previous conclusions. In 2012, it was determined that off-property vapor exposure should be investigated and evaluated because of the confirmed off-property transport of benzene (PGW, 2013). The Pennsylvania Department of Environmental Protection (DEP) oversaw PGW's 2012 soil vapor study where soil gas samples were collected along the exterior of residences. The results of the investigation identified chloroform at levels greater than standards established by DEP; however, there was no attributable source of chloroform on PGW property or in the groundwater originating from their property (PGW, 2013). No other compounds were detected in soil gas samples at or above their respective Medium Specific Concentrations – a DEP standard (PGW, 2013).

In 2013, DOH and ATSDR screened the 2012 PGW sampling data using more conservative healthbased comparison values (CVs) – the ATSDR Cancer Risk Evaluation Guide (CREG) values. DOH concluded that maximum concentrations of chloroform and benzene in the soil gas data exceeded their CREG values and required further evaluation; however, it could not be determined if indoor air concentrations of the chemicals were of public health concern in the absence of subslab or indoor air samples from within the residences.

Because chloroform is unrelated to PGW's operations, DOH and ATSDR referred the site, which does not include PGW property, to the EPA removal program to conduct a follow-up residential vapor intrusion investigation. EPA collected 24-hour samples of sub-slab soil gas, indoor air, and outdoor ambient air. Sample collection took place once in January 2014, once in October 2014, and once in March 2016. The samples were collected using SUMMA[®] canisters at residential units at the site and were analyzed in accordance with EPA Toxic Organic Method TO-15 for the full target compound list. Lifestyle products were removed from residences 24 hours prior to sampling to reduce their potential interference (EPA, 2017). Not all sources of indoor air contamination can be eliminated, such as carpeting and building materials.

DOH prepared this letter to review the EPA sampling data and evaluate whether exposure to indoor air contaminants pose a public health threat to residents.

THE VAPOR INSTRUSION EXPOSURE PATHWAY

Vapor intrusion is the general term given to the migration of hazardous vapors from any subsurface contaminant source, such as contaminated soil or groundwater, into a building or structure above (EPA, 2015). (See Appendix B for additional information on vapor intrusion.)

Indoor air in many buildings contain detectable levels of vapor-forming chemicals from a variety of indoor and outdoor sources. If vapor intrusion is occurring, the measured concentration of a contaminant will lessen as it migrates from a subsurface source to sub-slab soil gas and then to indoor air (EPA, 2015), provided there are no other indoor sources of that contaminant.

Exposure to contaminants of concern is determined by examining human exposure pathways. An exposure pathway has five parts:

- 1. A source of contamination (e.g., industrial facilities utilizing hazardous materials);
- 2. An environmental medium that can hold or move the contamination (e.g. water, soil, or air);
- 3. An exposure point at which people could come into contact with a contaminated medium (e.g., private residential well water or a building into which vapors enter);
- 4. An exposure route (e.g., ingestion or inhalation); and
- 5. A population that could come in contact with the contaminants.

For a completed pathway, all five parts must exist and exposure to a contaminant must have occurred, is occurring, or will occur (ATSDR, 2005a). For this investigation, vapor intrusion is the

exposure pathway of concern. The five parts of the vapor intrusion exposure pathway are present or could be present at the site as follows:

- 1. Source: Although the specific source is unknown, the presence of contaminants above their screening levels in the subsurface was confirmed during the 2012 soil vapor study.
- 2. Environmental medium: Groundwater and sub-slab soil are mediums that can hold and transport the contaminants.
- 3. Exposure point: People could become exposed within their residences if the vapors enter their homes, contaminating the indoor air.
- 4. Exposure route: People could inhale the contaminants that may be present in the indoor air.
- 5. Population: The residents living within the site are the potentially exposed population of concern.

This investigation sought to determine if chemical contaminants (particularly benzene and chloroform, which were detected at concentrations above their CREG values in 2012) were present in the indoor air of residences at concentrations that could harm people's health.

DATA EVALUATION

DOH screened the air sampling data against appropriate ATSDR CVs, which are health-based guidelines. Table 1 presents the maximum contaminant concentrations and the corresponding CVs. ATSDR CVs are conservative estimates of contaminant levels below which no health effects would be expected. When an ATSDR CV is not available, DOH uses screening values from other environmental and health agencies such as the EPA or another state agency (ATSDR, 2005a). Concentrations found to be above a CV do not necessarily mean they are harmful but that they require further evaluation to determine if adverse health effects are likely. Contaminants that exceed a CV are further evaluated using other standards and/or scientific studies, where appropriate, to determine whether adverse health effects are likely.

To evaluate non-cancer health effects, the CVs used were ATSDR's chronic and acute Minimal Risk Level (MRL) values, which are estimates of human exposure to a hazardous substance that are unlikely to have an appreciable risk of adverse non-cancer health effects over a specified route and duration of exposure in most people, including sensitive populations. If an MRL was not available, the EPA Reference Concentration (RfC) was used, which is an estimate of a continuous inhalation exposure that is unlikely to have an appreciable risk of harmful health effects during a lifetime, including to members of sensitive populations (EPA, 2017).

To evaluate cancer risk, DOH screened the data using ATSDR's CREG values, which are mediaspecific CVs used to identify concentrations of cancer-causing substances that are unlikely to result in an increase of cancer rates in an exposed population. If the air concentration of a specific contaminant exceeded the ATSDR CREG, DOH reviewed the toxicological information for that chemical and calculated an estimated excess cancer risk for residents using the following equation, which uses default values for an adult's exposure years living at one residence in their lifetime (ATSDR, 2016): Estimated cancer risk =

Exposure Concentration x EPA Inhalation Unit Risk (IUR) x 33 Exposure Years 78 Years.

To compare chemical concentrations in the sub-slab and indoor air across all units, average chemical concentrations were calculated using the results from all three sampling periods. Non-detect values were estimated and were not treated as zero. The assumption that undetected contaminants are absent from samples is often unduly optimistic as some carcinogens can pose potential health risks at levels below detection limits (EPA, 1991b). The non-detect values were estimated by dividing the Method Detection Limit (MDL) by two (EPA, 1991b). The laboratory also reported lab-estimated concentrations if a compound exceeded the MDL and was less than the reporting limit (Table 2, green-highlighted cells).

PUBLIC HEALTH IMPLICATIONS

Non-cancer health effects

With one exception—a single detection of benzene at 14.2 μ g/m³ (Table 1), to be discussed—all contaminants detected in the basement and first-floor indoor air were below their non-cancer health-based CVs; therefore, exposure to the contaminants at the levels detected is not expected to cause harmful non-cancer health effects.

Cancer risk evaluation

There were seven contaminants detected at the site that exceeded their CREG value at least once, requiring further evaluation for cancer risk: benzene; chloroform; 1,2-dicholoroethane; 1,3-butadiene; carbon tetrachloride; tetrachloroethylene; and trichloroethylene. Sub-slab, basement indoor air, and first-floor indoor air sample data for those seven contaminants are presented in Tables 2a–2g.

Benzene

Benzene was historically associated with the site, having been detected above its CVs in the groundwater aquifer below the nearby PGW facility with confirmed off-site transport towards the residential units.

During this investigation, benzene was detected at a maximum concentration of $14.2 \ \mu g/m^3$ in the first floor of unit 175 during October 2014 (Table 2a). The benzene concentration in the basement indoor air was 2.24 $\mu g/m^3$ and was undetected in the sub-slab during the same sampling period – October 2014. The average benzene concentration in the sub-slab, basement indoor air, and first-floor indoor air across all units and sampling periods was 0.368 $\mu g/m^3$, 1.191 $\mu g/m^3$, and 1.878 $\mu g/m^3$, respectively. Benzene was undetected in nine of the 18 sub-slab samples. These values, where the average chemical concentrations are greater in the indoor air, do not indicate a subsurface source of contamination.

A National Human Exposure Assessment Survey (NHEXAS) study of six states in the Great Lakes region found benzene in 100% of indoor air samples with concentrations averaging 7.19

 μ g/m³ [Clayton et al., 1999]. People living in cities or industrial areas are generally exposed to higher levels of benzene in air than those living in rural areas. The major sources of benzene exposure are exhaust from motor vehicles, automobile service stations, industrial emissions, and tobacco smoke. Vapors from products that contain benzene, such as glues, paints, furniture wax, and detergents, can also be a source of exposure. People may be exposed to higher levels of benzene in air by living near hazardous waste sites, petroleum refining operations, petrochemical manufacturing sites, or gas stations. (ATSDR, 2007)

The U.S. Department of Health and Human Services (HHS) has classified benzene as a human carcinogen (NTP, 2016). The International Agency for Research on Cancer (IARC) has also classified benzene as a human carcinogen (IARC, 2017).

The estimated increased cancer risk using the maximum concentration of 14.2 μ g/m³ (detected in the first-floor indoor air of unit 175 during October 2014) was 4.69 x 10⁻⁵ (Table 2a), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶ (EPA, 1991). There is no apparent increased cancer risk from exposure to benzene at the levels detected. Additionally, the estimated increased cancer risk from exposure to the average background level of 7.19 μ g/m³ that the NHEXAS study identified is 2.37 x 10⁻⁵, which is greater than the risk estimated for exposure at the site.

Chloroform

Chloroform is the other chemical historically associated with the site, detected above its CVs, though the chemical is not associated with PGW facility operations.

During this investigation, the maximum *exposure* concentration of chloroform (from an indoor air detection to which residents could be exposed, as opposed to a sub-slab detection) was $9.26 \,\mu\text{g/m}^3$ in the first-floor indoor air of unit 175 during October 2014 (Table 2b). Chloroform was detected at an overall maximum concentration of $67 \,\mu\text{g/m}^3$ in the sub-slab of unit 175 during March 2016. The basement indoor air chloroform concentration in March 2016 was $0.78 \,\mu\text{g/m}^3$. The average chloroform concentration in the sub-slab, basement indoor air, and first-floor indoor air all units and sampling periods was $5.610 \,\mu\text{g/m}^3$, $0.652 \,\mu\text{g/m}^3$, and $0.945 \,\mu\text{g/m}^3$, respectively. Chloroform was undetected in six of the 18 sub-slab samples. The higher average concentration in the sub-slab indicates a potential subsurface source of contamination, though the first-floor average is higher than the basement, indicating an alternate source of contamination.

Typical median indoor air concentrations of chloroform range from 0.98 to 19 μ g/m³. Chloroform has been found in the air from all areas of the United States and in nearly all public drinking water supplies; it is formed in small amounts when chlorine is added to water. Chloroform was once widely used as an anesthetic during surgery in humans. (ATSDR, 1997)

HHS has classified chloroform as *reasonably anticipated to be a human carcinogen* based on sufficient evidence of carcinogenicity from studies in experimental animals (NTP, 2016). IARC has classified chloroform as *possibly* carcinogenic to humans (IARC, 2017).

The estimated increased cancer risk using the maximum *exposure* concentration of 9.26 μ g/m³ (detected in the first-floor indoor air of unit 175 during October 2014) was 9.01 x 10⁻⁵ (Table 2b), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent increased cancer risk from exposure to chloroform at the levels detected. Additionally, the estimated increased cancer risk using the higher background concentration of 19 μ g/m³ (ATSDR, 1997) is 1.85 x 10⁻⁴, which is greater than the risk estimated for exposure at the site.

1,2-Dicholoroethane

During this investigation, 1,2-dicholoroethane was detected at a maximum concentration of 1.58 μ g/m³ in the basement indoor air of unit 70 during October 2014 (Table 2c). The average 1,2-dicholoroethane concentrations in the basement indoor air and first-floor indoor air across all units and sampling periods were 0.247 μ g/m³ and 0.342 μ g/m³, respectively. 1,2-Dicholoroethane was undetected in all sub-slab samples. These values, where the average chemical concentrations are greater in the indoor air, do not indicate a subsurface source of contamination.

1,2-Dicholoroethane is a manufactured chemical that is not found naturally in the environment. It is used in the production of vinyl chloride and is added to leaded gasoline to remove lead. It is a probable human carcinogen. In a 1996 survey of New Jersey and Pennsylvania residences by Heavner et al., 1,2-dichloroethane was detected in the homes of nonsmokers at a mean concentration of 0.03 μ g/m³ and in the homes of smokers at a mean concentration of 0.32 μ g/m³. The maximum concentration of 1,2-dichloroethane reported in nonsmoking households was 0.54 μ g/m³, while the maximum concentration in households where at least one family member smoked was 9.72 μ g/m³. (ATSDR, 2001)

HHS has classified 1,2-dicholoroethane as *reasonably anticipated to be a human carcinogen* based on sufficient evidence of carcinogenicity from studies in experimental animals (NTP, 2016). IARC has classified 1,2-dicholoroethane as *possibly* carcinogenic to humans (IARC, 2017).

The estimated increased cancer risk using the maximum exposure concentration of 1.58 μ g/m³ (detected in the basement indoor air of unit 70 during October 2014) was 1.74 x 10⁻⁵ (Table 2c), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent increased cancer risk from exposure to 1,2-dicholoroethane at the levels detected. Additionally, the estimated increased cancer risk using the maximum background concentration of 0.54 μ g/m³ identified in a 1996 study (ATSDR, 2001) is 5.94 x 10⁻⁶, which is less than the risk estimated for exposure at the site.

1,3-Butadiene

During this investigation, the maximum *exposure* concentration of 1,3-butadiene (from an indoor air detection to which residents could be exposed, as opposed to a sub-slab detection) was 0.23 μ g/m³ in the first-floor indoor air of unit 50 during January 2014 (Table 2d). 1,3-Butadiene was detected at an overall maximum concentration of 0.5 μ g/m³ in the sub-slab of unit 116 during January 2014. The basement indoor air and first-floor indoor air concentration in January 2014 was 0.150 μ g/m³ and 0.190 μ g/m³, respectively. 1,3-Butadiene was undetected in all other sub-slab samples. The average 1,3-butadiene concentration in the sub-slab, basement indoor air, and first-

floor indoor air across all units and sampling periods was 0.035 μ g/m³, 0.047 μ g/m³, and 0.063 μ g/m³, respectively. These values, where the average chemical concentrations are greater in the indoor air, do not indicate a subsurface source of contamination.

1,3-Butadiene is a chemical made from the processing of petroleum. It evaporates easily and as such, it is not expected to be found in water or soil. Automobile exhaust and cigarette smoke are common sources of 1,3-butadiene release into the air. A 2004 Sax et al. study measured 1,3-butadiene in indoor air of homes in New York. The mean concentrations were 1.0 μ g/m³ during the winter and 1.2 μ g/m³ during the summer (ATSDR, 2012).

HHS has classified 1,3-butadiene as a human carcinogen (NTP, 2016). IARC has also classified 1,3-butadiene as a human carcinogen (IARC, 2017).

The estimated increased cancer risk using the maximum exposure concentration of 0.23 μ g/m³ (detected in the first-floor indoor air of unit 50 during January 2014) was 2.92 x 10⁻⁶ (Table 2d), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent increased cancer risk from exposure to 1,3-butadiene at the levels detected. Additionally, the estimated increased cancer risk using the higher background concentration of 1.2 μ g/m³ (ATSDR, 2012) is 1.52 x 10⁻⁵, which is greater than the risk estimated for exposure at the site.

Carbon tetrachloride

During this investigation, the maximum *exposure* concentration of carbon tetrachloride (from an indoor air detection to which residents could be exposed, as opposed to a sub-slab detection) was 0.72 μ g/m³ in the first-floor indoor air of unit 34 during January 2014 (Table 2e). Carbon tetrachloride was detected at an overall maximum concentration of 135 μ g/m³ in the sub-slab of unit 175 during October of 2014. The concentration in the basement indoor air was 0.467 μ g/m³ during the same sampling period – October 2014. The average carbon tetrachloride concentration in the sub-slab, basement indoor air, and first-floor indoor air across all units and sampling periods was 7.642 μ g/m³ in the sub-slab, the average sub-slab concentration is 0.143 μ g/m³). Carbon tetrachloride was undetected in 12 of the 18 sub-slab samples. These values, where the average chemical concentrations are greater in the indoor air, do not indicate a subsurface source of contamination.

Carbon tetrachloride is a manufactured chemical that is not found naturally in the environment. Typical indoor concentrations in homes in several U.S. cities were around 1.0 μ g/m³, with some values up to 9 μ g/m³. (ATSDR, 2005b)

HHS has classified carbon tetrachloride as *reasonably anticipated to be a human carcinogen* based on sufficient evidence of carcinogenicity from studies in experimental animals (NTP, 2016). IARC has classified carbon tetrachloride as *possibly* carcinogenic to humans (IARC, 2017).

The estimated increased cancer risk using the maximum *exposure* concentration of 0.72 μ g/m³ (detected in the first-floor indoor air of unit 34 during January 2014) was 1.83 x 10⁻⁶ (Table 2e), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent

increased cancer risk from exposure to carbon tetrachloride at the levels detected. Additionally, the estimated increased cancer risk using the higher background concentration of 9 μ g/m³ (ATSDR, 2005b) is 2.28 x 10⁻⁵, which is greater than the risk estimated for exposure at the site.

Tetrachloroethylene

During this investigation, the maximum *exposure* concentration of tetrachloroethylene (from an indoor air detection to which residents could be exposed, as opposed to a sub-slab detection) was $6.81 \ \mu g/m^3$ in the first-floor indoor air of unit 34 during March 2016 (Table 2f). Tetrachloroethylene was detected at an overall maximum concentration of 27.1 $\mu g/m^3$ in the sub-slab of unit 116 during October 2014. The tetrachloroethylene concentration in the basement indoor air was $0.526 \ \mu g/m^3$ during the same sampling period – October 2014. The average tetrachloroethylene concentration in the sub-slab, basement indoor air, and first-floor indoor air across all units and sampling periods was $3.856 \ \mu g/m^3$, $0.315 \ \mu g/m^3$, and $0.537 \ \mu g/m^3$, respectively. Tetrachloroethylene was undetected in four of the 18 sub-slab samples. The higher average concentration in the sub-slab indicates a potential subsurface source of contamination, though the first-floor average is higher than the basement, indicating an alternate source of contamination.

Tetrachloroethylene is one of the most commonly detected chemicals in background indoor sources. The median value for indoor air in the United States, from 2,195 entries in the EPA's database of volatile organic contaminants (VOC-AMBI), was approximately 4.9 μ g/m³, with an average value of 20.7 μ g/m³. The use of tetrachloroethylene as a dry-cleaning agent, chemical intermediate, and metal degreasing solvent has led to its release to the environment. It is also used in some consumer products. (ATSDR, 2014a)

HHS has classified tetrachloroethylene as *reasonably anticipated to be a human carcinogen* based on sufficient evidence of carcinogenicity from studies in experimental animals (NTP, 2016). IARC has classified tetrachloroethylene as *probably* carcinogenic to humans (IARC, 2017).

The estimated increased cancer risk using the maximum *exposure* concentration of 6.81 μ g/m³ (detected in the first-floor indoor air of unit 34 during March 2016) was 7.49 x 10⁻⁷ (Table 2f), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent increased cancer risk from exposure to tetrachloroethylene at the levels detected. Additionally, the estimated increased cancer risk using average background concentration of 20.7 μ g/m³ (ATSDR, 2014a) is 2.28 x 10⁻⁶, which is greater than the risk estimated for exposure at the site.

Trichloroethylene

During this investigation, the maximum *exposure* concentration of trichloroethylene was 0.627 μ g/m³ in the first-floor indoor air of unit 175 during October 2014 (Table 2g). Trichloroethylene was detected at an overall maximum concentration of 8.4 μ g/m³ in the sub-slab of unit 175 during January 2014. Trichloroethylene was undetected in the basement indoor air during that sampling period. The average trichloroethylene concentration in the sub-slab, basement indoor air, and first-floor indoor air across all units and sampling periods was 0.591 μ g/m³, 0.135 μ g/m³, and 0.095 μ g/m³, respectively. Trichloroethylene was undetected in 15 of the 18 sub-slab samples. The higher average concentration in the sub-slab indicates a potential subsurface source of contamination.

However, individual basement indoor air detections do not correspond to elevated concentrations in the sub-slab for the same residential unit during the same sampling period, indicating an alternate source of contamination.

Major uses of trichloroethylene are as a solvent to remove grease from metal parts and as an intermediate chemical in the multi-step process of manufacturing of refrigerant chemicals. Exposure most often occurs by drinking water that has been contaminated by trichloroethylene or by breathing trichloroethylene released to the air from contaminated water. Arithmetic mean trichloroethylene concentrations measured in air at locations across the United States are generally between 0.05 μ g/m³ and 1.61 μ g/m³, although mean levels as high as 18.27 μ g/m³ have been reported. (ATSDR, 2014b)

HHS has classified trichloroethylene as *known to be a human carcinogen* (NTP, 2016). IARC has also classified trichloroethylene as a human carcinogen (IARC, 2017).

The estimated increased cancer risk using the maximum *exposure* concentration of 0.627 μ g/m³ (detected in the first-floor indoor air of unit 175 during October 2014) was 1.09 x 10⁻⁶ (Table 2g), which falls within EPA's target cancer risk range of 1 x 10⁻⁴ to 1 x 10⁻⁶. There is no apparent increased cancer risk from exposure to trichloroethylene at the levels detected. Additionally, the estimated increased cancer risk using the higher background concentration of 1.61 μ g/m³ (ATSDR, 2014b) is 2.79 x 10⁻⁶, which is greater than the risk estimated for exposure at the site.

Cumulative cancer risk health effects

For this health evaluation, cumulative cancer risk was estimated using the summation of each individual chemical's cancer risk estimate. A cumulative cancer risk estimate was calculated separately for the first floor and basement of each residential unit for each sampling period. The cumulative cancer risk estimates in the basement indoor air samples ranged from 6.18×10^{-6} to 3.27×10^{-5} and from 1.81×10^{-6} to 1.42×10^{-4} in the first-floor indoor air (Table 3).

The highest estimate of 1.42×10^{-4} (from the first-floor indoor air of unit 175 during October 2014) falls outside of EPA's target cancer risk range of 1×10^{-6} to 1×10^{-4} and was evaluated further. The elevated risk in that location is attributable to benzene and chloroform outliers, which are likely due to a source or sources other than vapor intrusion. While long-term exposure to benzene and chloroform can cause certain cancers, short-term exposure to the levels detected at the site during October 2014 is not expected to increase cancer risk.

Based on our analysis, there is no apparent increased cancer risk from exposure to the contaminants at the levels detected in the indoor air of residential units at the site.

LIMITATIONS

- There is no data to assess indoor air contamination during the summer months. The effects of temperature and other weather variations should be considered when evaluating the concentration of volatile organic compounds in the air.
- Background sources of contamination such as furniture and flooring inside the home, or automobile exhaust and industrial emissions entering the home from outside can affect environmental evaluations where vapor intrusion is the source of concern. Particularly in urban areas, contaminants from ambient sources can be present in indoor air (DEP, 2002).

CONCLUSIONS AND RECOMMENDATIONS

- Exposures to indoor air contaminants at the levels detected are not expected to result in adverse health effects.
- Most indoor air detections did not exceed background levels common in urban environments.
- The data do not indicate that the contaminants detected in the indoor air originated entirely from the sub-slab soil gas. Elevated detections of first-floor and basement indoor air contaminants did not correspond to elevated levels of that chemical in the sub-slab in the same unit during the same sampling period. Additionally, when elevated levels of contaminants were detected in the sub-slab, there was not a corresponding and proportionally elevated detection in the basement indoor air.
- Contaminants detected in the indoor air likely originated from sources of contamination other than vapor intrusion. To protect the current and future health of individuals, DOH recommends that EPA provide health education to these residents to limit their chemical exposure from indoor sources.

Please contact me with any questions you may have regarding this letter health consultation.

Sincerely,

Bevin Durant, MPH Epidemiology Research Associate Pennsylvania Department of Health bdurant@pa.gov

Cc: Farhad Ahmed, MBBS, MPH; DOH Anil Nair, PhD, MPH; DOH

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TABLES

Contominonto	A a b b b b b b b b b b	ATSDR CREG	ATSDR Chronic MRL	ATSDR Acute MRL	EPA RfC
Contaminants	Maximum Indoor Air µg/m	µg/m³	μg/m ³	μg/m ³	µg/m³
Acetone	260	—	31000	62000	_
Benzene	14.2	0.13	9.6	29	30
Bromoform	0.327	0.91	—	—	_
1,3-Butadiene	0.23	0.033	—	_	2
2-Butanone	16	—	_	_	5000
Carbon Tetrachloride	0.72	0.17	190	_	100
Chlorobenzene	0.0955	—	—	—	-
Chloroethane	0.331	—	—	40000	10000
Chloroform	9.26	0.043	98	490	_
Chloromethane	2.41	—	100	1000	90
Cyclohexane	3.06	—	—	—	6000
Dibromochloromethane	2.35	—	—	—	—
1,4-Dichlorobenzene	2.83	—	60	12000	800
Dichlorodifluoromethane	3.2	—	—	—	
1,1-Dichloroethane	0.186	—	—	—	
1,2-Dichloroethane	1.58	0.038	2400	—	
Ethyl Acetate	20.8	—	—	—	
Ethylbenzene	11.8	—	260	22000	1000
4-Ethyltoluene	1.59	—	—	—	
Heptane, N-	8.59		—	-	I
Hexane, N-	28	—	2100	—	700
2-Hexanone	0.49	—	—	—	30
Isopropanol	1250	—	—	—	
Methyl Isobutyl Ketone	1.59	—	—	—	3000
Methyl tert-Butyl Ether (MTBE)	0.22	—	2500	7200	3000
Methylene Chloride	61	63	1000	2100	600
Naphthalene	1.02	—	3.7	—	3
Propene	146	—	—	—	-
Styrene	1.55	—	850	21000	1000
Tetrachloroethylene	6.81	3.8	41	41	40
Tetrahydrofuran	245	—	—	_	2000
Toluene	13	—	3800	7500	5000
Trichloro-1,2,2-trifluoroethane,	33.7	—	—	—	_
Trichloroethane, 1,1,1-	32.6	—	—	11000	5000
Trichloroethylene	0.627	0.22	2.1	—	2
Trichlorofluoromethane	11	_	_	_	_
Trimethylbenzene, 1,2,4-	8.66	-	_	—	60
Trimethylbenzene, 1,3,5-	3.38	-	—	_	60
Vinyl Acetate	8.45	-	—	—	200
Xylene, m&p-	14.3	-	220	8700	100
Xylene, o-	6.73	l – [220	8700	100

Table 1: Maximum Contaminant Concentrations and Health–Based Comparison Values

Cells with "-" = no available comparison value Bold text = contaminant exceeded a CV

Tables 2a-2g: Sub-slab, Basement Indoor Air, and First-Floor Indoor Air Detections for Contaminants that Exceeded CREG Values; µg/m³

Table	2a:	Benzene	(µg/	′m³)
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				,						
Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-Floor Indoor Air			
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	
Unit 34	0.004	0.116	0.004	0.670	0.606	1.970	0.760	1.140	1.670	
Unit 50	0.004	0.426	1.870	0.840	1.370	1.400	2.300	0.941	1.260	
Unit 84	0.340	0.084	0.004	0.700	0.567	1.820	0.940	0.547	1.630	
Unit 70	0.004	0.104	0.004	0.750	1.090	1.370	0.720	1.040	0.939	
Unit 116	1.200	0.266	2.190	0.780	0.591	2.050	1.100	0.542	1.740	
Unit 175	0.004	0.004	0.004	1.300	2.240	1.320	1.300	14.20 *	1.040	
Unit 175	0.004	0.266	0.004	1.300	2.240	1.320	1.100	0.542 14.20*	1.740	

CREG = 0.13, MDL = 0.008

*Cancer risk estimate for maximum exposure concentration: 4.69 x 10⁻⁵

Table 2b: Chloroform $(\mu g/m^3)$ CREG = 0.043, MDI = 0.011

			0=	e e.e.e).		-				
Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-Floor Indoor Air			
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	
Unit 34	3.600	0.705	0.005	0.410	0.266	1.750	0.530	0.616	1.240	
Unit 50	2.400	0.511	0.005	0.220	0.214	0.767	0.190	0.167	0.620	
Unit 84	0.005	1.970	0.793	0.170	0.578	0.138	0.220	0.235	0.143	
Unit 70	0.005	0.123	0.005	0.190	0.758	1.080	0.190	0.788	0.690	
Unit 116	0.770	17.100	0.005	0.210	0.588	0.167	0.220	0.005	0.201	
Unit 175	0.780	5.190	67.00	2.600	0.841	0.780	0.950	9.26 *	0.748	

*Cancer risk estimate for maximum exposure concentration: 9.01 x 10⁻⁵

Table 2c: 1,2-Dichloroethane (µg/m³)

CREG = 0.038, MDL = 0.019

Residential	Base	ement Sub	Slab	Basement Indoor Air			First-Floor Indoor Air			
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	
Unit 34	0.009	0.009	0.009	0.180	0.009	0.187	0.210	1.290	0.260	
Unit 50	0.009	0.009	0.009	0.009	0.167	0.508	0.009	0.114	0.937	
Unit 84	0.009	0.009	0.009	0.009	0.009	0.181	0.082	0.009	0.343	
Unit 70	0.009	0.009	0.009	0.150	1.58*	0.226	0.150	1.570	0.144	
Unit 116	0.009	0.009	0.009	0.009	0.107	0.255	0.130	0.009	0.433	
Unit 175	0.009	0.009	0.009	0.009	0.092	0.755	0.009	0.241	0.217	

*Cancer risk estimate for maximum exposure concentration: 1.74 x 10⁻⁵

Blue cells = Undetected result. Here, MDL/2 is used to estimate level of non-detects.

Green cells = Lab-estimated result

Red text = Maximum exposure concentration

Italicized text = Maximum detected concentration (if different from the maximum exposure concentration)

Tables 2a–2g (continued): Sub-slab, Basement Indoor Air, and First-Floor Indoor Air Detections for Contaminants that Exceeded CREG Values; µg/m³

Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-Floor Indoor Air			
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	
Unit 34	0.008	0.008	0.008	0.082	0.008	0.008	0.140	0.008	0.008	
Unit 50	0.008	0.008	0.008	0.190	0.008	0.008	0.230*	0.008	0.008	
Unit 84	0.008	0.008	0.008	0.081	0.008	0.008	0.170	0.008	0.008	
Unit 70	0.008	0.008	0.008	0.110	0.008	0.008	0.093	0.008	0.008	
Unit 116	0.500	0.008	0.008	0.150	0.008	0.008	0.190	0.008	0.008	
Unit 175	0.008	0.008	0.008	0.130	0.008	0.008	0.210	0.008	0.008	

Table 2d: 1,3-Butadiene (µg/m³)

CREG = 0.033, MDL = 0.016

*Cancer risk estimate for maximum exposure concentration: 2.92 x 10⁻⁶

Table 2e: Carbon Tetrachloride (µg/m³)

Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-	Floor Indo	or Air
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16
Unit 34	0.008	0.636	0.008	0.370	0.417	0.464	0.720*	0.415	0.495
Unit 50	0.008	0.442	0.008	0.410	0.428	0.417	0.008	0.466	0.453
Unit 84	0.008	0.446	0.008	0.360	0.429	0.456	0.008	0.084	0.446
Unit 70	0.008	0.439	0.008	0.350	0.442	0.487	0.410	0.078	0.331
Unit 116	0.008	0.197	0.008	0.410	0.442	0.471	0.008	0.008	0.447
Unit 175	0.008	135.0	0.315	0.550	0.467	0.501	0.390	0.445	0.460

CREG = 0.017, MDL = 0.015

*Cancer risk estimate for maximum exposure concentration: 1.83 x 10⁻⁶

Table 2f: Tetrachloroethylene (µg/m³)

CREG = 3.8, MDL = 0.022

Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-	Floor Indo	or Air
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16
Unit 34	3.800	14.300	0.011	0.550	0.234	1.620	0.580	0.178	6.81 *
Unit 50	0.011	2.520	1.180	0.450	0.237	0.324	0.011	0.179	0.011
Unit 84	1.000	4.420	0.011	0.011	0.011	0.194	0.011	0.011	0.157
Unit 70	1.500	1.260	0.721	0.011	0.197	0.011	0.011	0.011	0.154
Unit 116	0.810	27.100	0.011	0.130	0.526	0.311	0.011	0.011	0.200
Unit 175	0.920	8.560	1.280	0.310	0.324	0.219	0.011	1.130	0.178

*Cancer risk estimate for maximum exposure concentration: 7.49 x 10⁻⁷

Blue cells = Undetected result. Here, MDL/2 is used to estimate level of non-detects.

Green cells = Lab-estimated result

Red text = Maximum exposure concentration

Italicized text = Maximum detected concentration (if different from maximum exposure concentration)

Tables 2a–2g (continued): Sub-slab, Basement Indoor Air, and First-Floor Indoor Air Detections for Contaminants that Exceeded CREG Values; µg/m³

Residential	Base	ement Sub	Slab	Base	ment Indoo	or Air	First-	Floor Indo	or Air
Units	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16	Jan-14	Oct-14	Mar-16
Unit 34	0.009	0.009	0.009	0.009	0.292	0.009	0.009	0.226	0.108
Unit 50	0.550	1.550	0.009	0.009	0.553	0.118	0.009	0.622	0.009
Unit 84	0.009	0.009	0.009	0.009	0.537	0.009	0.009	0.009	0.009
Unit 70	0.009	0.009	0.009	0.009	0.247	0.009	0.009	0.009	0.009
Unit 116	0.009	0.009	0.009	0.009	0.573	0.009	0.009	0.009	0.009
Unit 175	8.400	0.009	0.009	0.009	0.009	0.009	0.009	0.627*	0.009

Table 2g: Trichloroethylene (µg/m³)

CREG = 0.22, MDL = 0.019

*Cancer risk estimate for maximum exposure concentration: 1.09 x 10⁻⁶

Blue cells = Undetected result. Here, MDL/2 is used to estimate level of non-detects.

Green cells = Lab-estimated result

Red text = Maximum exposure concentration

Italicized text = Maximum detected concentration (if different from maximum exposure concentration)

Location	Contaminants	Maximu	ım Indoor Ai	r μg/m³	CREG	EPA IUR	Cance contamin	r Risk Estima ants exceed	ites for ing CREG*
		Jan-14	Oct-14	Ma r-16	µg/m*	(µg/m ⁻)	Jan-14	Oct-14	Ma r-16
	1,2-Dichloroethane	0.18	NA	0.187	0.038	2.60E-05	1.98E-06	NA	2.06E-06
	1,3-Butadiene	0.082	NA	NA	0.033	3.00E-05	1.041E-06	NA	NA
UT34	Benzene	0.67	0.606	1.97	0.13	7.80E-06	2.211E-06	2E-06	6.50E-06
Basement	Carbon Tetrachloride	0.37	0.417	0.464	0.17	6.00E-06	9.392E-07	1.06E-06	1.18E-06
Indoor Air	Chloroform	0.41	0.266	1.75	0.043	2.30E-05	3.99E-06	2.59E-06	1.70E-05
	Tetrachloroethylene	0.55	0.234	1.62	3.8	2.60E-07	6.05E-08	2.57E-08	1.78E-07
	Trichloroethylene	NA	0.292	NA	0.22	4.10E-06	NA	5.065E-07	NA
CUMULATIV	E CANCER RISK - UT34 Ba	sement Indoo	or Air				1.02E-05	6.18E-06	2.69E-05
	1,2-Dichloroethane	0.21	1.29	0.26	0.038	2.60E-05	2.31E-06	1.42E-05	2.86E-06
	1,3-Butadiene	0.14	NA	NA	0.033	3.00E-05	1.78E-06	NA	NA
UT34 First	Benzene	0.76	1.14	1.67	0.13	7.80E-06	2.51E-06	3.76E-06	5.51E-06
Floor	Carbon Tetrachloride	0.72	0.415	0.495	0.17	6.00E-06	1.83E-06	1.05E-06	1.26E-06
Indoor Air	Chloroform	0.53	0.616	1.24	0.043	2.30E-05	5.16E-06	5.99E-06	1.21E-05
	Tetrachloroethylene	0.58	0.178	6.81	3.8	2.60E-07	6.38E-08	1.96E-08	7.49E-07
	Trichloroethylene	NA	0.226	0.108	0.22	4.10E-06	NA	3.92E-07	1.87E-07
CUMULATIV	E CANCER RISK - UT34 Fir	st Floor Indoo	or Air				1.36E-05	2.54E-05	2.26E-05
	1,2-Dichloroethane	NA	0.167	0.508	0.038	2.60E-05	NA	1.84E-06	5.59E-06
	1,3-Butadiene	0.19	NA	NA	0.033	3.00E-05	2.41E-06	NA	NA
UT50	Benzene	0.84	1.37	1.4	0.13	7.80E-06	2.77E-06	4.52E-06	4.62E-06
Basement	Carbon Tetrachloride	0.41	0.428	0.417	0.17	6.00E-06	1.04E-06	1.09E-06	1.06E-06
Indoor Air	Chloroform	0.22	0.214	0.767	0.043	2.30E-05	2.14E-06	2.08E-06	7.46E-06
	Tetrachloroethylene	0.45	0.237	0.324	3.8	2.60E-07	4.95E-08	2.61E-08	3.56E-08
	Trichloroethylene	NA	0.553	0.118	0.22	4.10E-06	NA	9.59E-07	2.05E-07
CUMULATIV	E CANCER RISK - UT50 Ba	sement Indoo	or Air				8.41E-06	1.05E-05	1.90E-05
	1,2-Dichloroethane	NA	0.114	0.937	0.038	2.60E-05	NA	1.25E-06	1.03E-05
	1,3-Butadiene	0.23	NA	NA	0.033	3.00E-05	2.92E-06	NA	NA
UT50 First	Benzene	2.3	0.941	1.26	0.13	7.80E-06	7.59E-06	3.11E-06	4.16E-06
Floor	Carbon Tetrachloride	NA	0.466	0.453	0.17	6.00E-06	NA	1.18E-06	1.15E-06
Indoor Air	Chloroform	0.19	0.167	0.62	0.043	2.30E-05	1.85E-06	1.63E-06	6.03E-06
	Tetrachloroethylene	NA	0.179	NA	3.8	2.60E-07	NA	1.97E-08	NA
	Trichloroethylene	NA	0.622	NA	0.22	4.10E-06	NA	1.08E-06	NA
CUMULATIV	E CANCER RISK - UT50 Fir	st Floor Indoo	or Air				1.24E-05	8.27E-06	2.16E-05
	1,2-Dichloroethane	0.15	1.58	0.226	0.038	2.60E-05	1.65E-06	1.74E-05	2.49E-06
	1,3-Butadiene	0.11	NA	NA	0.033	3.00E-05	1.40E-06	NA	NA
UT70	Benzene	0.75	1.09	1.37	0.13	7.80E-06	2.48E-06	3.60E-06	4.52E-06
Basement	Carbon Tetrachloride	0.35	0.442	0.487	0.17	6.00E-06	8.88E-07	1.12E-06	1.24E-06
Indoor Air	Chloroform	0.19	0.758	1.08	0.043	2.30E-05	1.85E-06	7.38E-06	1.05E-05
	Tetrachloroethylene	NA	0.197	NA	3.8	2.60E-07	NA	2.17E-08	NA
	Trichloroethylene	1.2	1.34	0.909	0.22	4.10E-06	2.08E-06	2.32E-06	1.58E-06
CUMULATIV	E CANCER RISK - UT70 Ba	sement Indoo	or Air			•	1.03E-05	3.18E-05	2.03E-05
	1,2-Dichloroethane	0.15	1.57	0.144	0.038	2.60E-05	1.65E-06	1.73E-05	1.58E-06
	1,3-Butadiene	0.093	NA	NA	0.033	3.00E-05	1.18E-06	NA	NA
UT70 First	Benzene	0.72	1.04	0.939	0.13	7.80E-06	2.38E-06	3.43E-06	3.10E-06
Floor	Carbon Tetrachloride	0.41	NA	0.331	0.17	6.00E-06	1.04E-06	NA	8.40E-07
muoor Alf	Chloroform	0.19	0.788	0.69	0.043	2.30E-05	1.85E-06	7.67E-06	6.71E-06
	Tetrachloroethylene	NA	NA	0.154	3.8	2.60E-07	NA	NA	1.69E-08
	Trichloroethylene	NA	NA	NA	0.22	4.10E-06	NA	NA	NA
	F CANCER RISK - LIT70 Fir	st Floor Indor	or Air				8 10F-06	2 8/E-05	1 225-05

Table 3: Cumulative Cancer Risk per Residential Unit per Sampling Event

NA = Contaminants were undetected or results were estimated

Red text = maximum exposure concentration

Green text = detections did not exceed CREG but were included in cumulative cancer risk evaluation

Location	Contaminants	Maximu	um Indoor Ai	r μg/m³	CREG	EPA IUR	Cance contamin	r Risk Estima ants exceed	ates for ling CREG*
		Jan-14	Oct-14	Ma r-16	µg/m³	(µg/m ³) ⁻	Jan-14	Oct-14	Ma r-16
	1,2-Dichloroethane	NA	NA	0.181	0.038	2.60E-05	NA	NA	1.99E-06
	1,3-Butadiene	0.081	NA	NA	0.033	3.00E-05	1.03E-06	NA	NA
UT84	Benzene	0.7	0.567	1.82	0.13	7.80E-06	2.31E-06	1.87E-06	6.01E-06
Basement	Carbon Tetrachloride	0.36	NA	0.456	0.17	6.00E-06	9.14E-07	NA	1.16E-06
Indoor Air	Chloroform	0.17	0.578	0.138	0.043	2.30E-05	1.65E-06	5.62E-06	1.34E-06
	Tetrachloroethylene	NA	NA	0.194	3.8	2.60E-07	NA	NA	2.13E-08
	Trichloroethylene	NA	NA	NA	0.22	4.10E-06	NA	NA	NA
CUMULATIV	E CANCER RISK - UT84 Ba	sement Indoo	or Air				5.91E-06	7.50E-06	1.05E-05
	1,2-Dichloroethane	0.082	NA	0.343	0.038	2.60E-05	9.02E-07	NA	3.77E-06
	1,3-Butadiene	0.17	NA	NA	0.033	3.00E-05	2.16E-06	NA	NA
UT84 First	Benzene	0.94	0.547	1.63	0.13	7.80E-06	3.10E-06	1.81E-06	5.38E-06
Floor	Carbon Tetrachloride	NA	NA	0.446	0.17	6.00E-06	NA	NA	1.13E-06
Indoor Air	Chloroform	0.22	NA	0.143	0.043	2.30E-05	2.14E-06	NA	1.39E-06
	Tetrachloroethylene	NA	NA	0.157	3.8	2.60E-07	NA	NA	1.73E-08
	Trichloroethylene	NA	NA	NA	0.22	4.10E-06	NA	NA	NA
CUMULATIV	E CANCER RISK - UT84 Fir	st Floor Indoo	or Air				8.30E-06	1.81E-06	1.17E-05
	1,2-Dichloroethane	NA	0.107	0.255	0.038	2.60E-05	NA	1.18E-06	2.81E-06
	1,3-Butadiene	0.15	NA	NA	0.033	3.00E-05	1.90E-06	NA	NA
UT116	Benzene	0.78	0.591	2.05	0.13	7.80E-06	2.57E-06	1.95E-06	6.77E-06
Basement	Carbon Tetrachloride	0.41	0.442	0.471	0.17	6.00E-06	1.04E-06	1.12E-06	1.20E-06
Indoor Air	Chloroform	0.21	0.588	0.167	0.043	2.30E-05	2.04E-06	5.72E-06	1.63E-06
	Tetrachloroethylene	0.13	0.526	0.311	3.8	2.60E-07	1.43E-08	5.79E-08	3.42E-08
	Trichloroethylene	NA	0.573	NA	0.22	4.10E-06	NA	9.94E-07	NA
CUMULATIV	E CANCER RISK - UT116 B	asement Indo	oor Air				7.58E-06	1.10E-05	1.24E-05
	1,2-Dichloroethane	0.13	NA	0.433	0.038	2.60E-05	1.43E-06	NA	4.76E-06
	1,3-Butadiene	0.19	NA	NA	0.033	3.00E-05	2.41E-06	NA	NA
UT116 First	Benzene	1.1	0.542	1.74	0.13	7.80E-06	3.63E-06	1.79E-06	5.74E-06
Floor	Carbon Tetrachloride	NA	NA	0.447	0.17	6.00E-06	NA	NA	1.13E-06
Indoor Air	Chloroform	0.22	NA	0.201	0.043	2.30E-05	2.14E-06	NA	1.96E-06
	Tetrachloroethylene	NA	NA	0.2	3.8	2.60E-07	NA	NA	2.20E-08
	Trichloroethylene	NA	NA	NA	0.22	4.10E-06	NA	NA	NA
CUMULATIV	E CANCER RISK - UT116 F	irst Floor Indo	oor Air				9.61E-06	1.79E-06	1.36E-05
	1,2-Dichloroethane	NA	0.0919	0.755	0.038	2.60E-05	NA	1.01E-06	NA
	1,3-Butadiene	0.13	NA	NA	0.033	3.00E-05	1.65E-06	NA	NA
UT175	Benzene	1.3	2.24	1.32	0.13	7.80E-06	4.29E-06	7.39E-06	4.36E-06
Basement	Carbon Tetrachloride	0.55	0.467	0.501	0.17	6.00E-06	1.40E-06	1.19E-06	1.27E-06
maoor An	Chloroform	2.6	0.841	0.78	0.043	2.30E-05	2.53E-05	8.18E-06	7.59E-06
	Tetrachloroethylene	0.31	0.324	0.219	3.8	2.60E-07	3.41E-08	3.56E-08	2.41E-08
0110 4111 A TD 4	Trichloroethylene	NA	NA	NA	0.22	4.10E-06	NA	NA	NA
CUMULATIV	E CANCER RISK - UT175 B	asement indo	bor Air	0.047	0.000	0.005.05	3.27E-05	1.78E-05	1.32E-05
	1,2-Dichloroethane	NA 0.24	0.241	0.217	0.038	2.60E-05	NA	2.65E-06	2.39E-06
	1,3-Butadiene	0.21	NA	NA 1.01	0.033	3.00E-05	2.67E-06		NA
UT175 First	Carbon Totrachlarida	1.3	14.2	1.04	0.13	7.80E-06	4.29E-06	4.09E-05	3.43E-Ub
FIOOR Indoor Air		0.39	0.445	0.46	0.17	6.00E-06	9.90E-07	1.13E-06	1.1/E-06
	Chiorotorm	0.95	9.26	0.748	0.043	2.30E-05	9.24E-06	9.01E-05	7.28E-06
1 1	Totrachlore athulog -	N1 A	1.10	0.170	2.0	2 605 07	NIA	1 245 07	1 065 00
	Tetrachloroethylene	NA	1.13	0.178	3.8	2.60E-07	NA	1.24E-07	1.96E-08

Table 3 (continued): Cumulative Cancer Risk per Residential Unit per Sampling Event

NA = Contaminants were undetected or results were estimated

Red text = maximum exposure concentration

Green text = detections did not exceed CREG but were included in cumulative cancer risk evaluation

APPENDIX A – Site Maps



The Passyunk Soil Gas Site is outlined in red. Philadelphia Gas Works (PGW) is to the west, Philadelphia Energy Solutions Refinery (PESR) is to the south, outlined in blue.



A close-up of the Passyunk Soil Gas Site perimeter.

APPENDIX B – ATSDR Vapor Intrusion Fact Sheet

Overview of Vapor Intrusion

What is vapor intrusion?

Vapor intrusion is a way that volatile chemicals in soil and groundwater can enter and build-up inside buildings. Volatile chemicals are a class of chemicals that are volatile (evaporate easily) and form a vapor in the air.

- Common sources of volatile chemicals include gas stations, dry cleaners, and industrial operations.
- When a chemical is spilled or leaks into the ground, it can contaminate the soil and the groundwater.
- The chemical can move with the groundwater and travel under a building (migration route).
- If that chemical is volatile, it can become a gas and seep into nearby buildings and contaminate indoor air.

This fact sheet explains how vapor intrusion occurs and what factors can cause vapors (sometimes called gases) to move into indoor air.



Why is vapor intrusion important to me?

When chemicals move indoors, you can be exposed to them by breathing indoor air. This exposure can cause health effects, depending on the type and amount of chemical and the length of exposure.

You can learn more about the possible health effects of individual volatile chemicals in ATSDR's Toxic Substances Portal: <u>http://www.atsdr.cdc.gov/substances/index.asp</u>.

How does vapor intrusion occur?

Vapor intrusion does not occur every time there is contaminated soil or groundwater. It occurs only when volatile chemicals move from a source (like a chemical spill) along an underground migration route and into a building.

The type and amount of chemicals coming from a source will determine whether vapor intrusion occurs at levels of possible health concern.

Agency for Toxic Substances and Disease Registry Division of Community Health Investigations



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What factors affect vapor intrusion?

The following factors affect vapor intrusion:

- The type of soil beneath your building
- The type and condition of your building (foundation, leaks, air exchange)
- The weather conditions in your area

The amount of vapors entering a building can be different over time—changing hourly, daily, weekly, and seasonally.

The amount of vapor intrusion can also be different on different floors and in different rooms of the same building or in buildings right next to each other.

If scientists suspect vapor intrusion in buildings in a specific location, they may decide to conduct an investigation. See ATSDR's fact sheet "Investigating Vapor Intrusion" for information on what to expect if a vapor intrusion investigation is planned for buildings in your area.

Where can I learn more about vapor intrusion?

U.S. Environmental Protection Agency

Vapor intrusion website, visit: <u>http://www2.epa.gov/vaporintrusion</u>

Interstate Technology & Regulatory Council

Vapor intrusion website, visit: <u>http://www.itrcweb.org/Team/Public?teamID=22</u>

Agency for Toxic Substances and Disease Registry

Investigating vapor intrusion, visit: <u>https://www.atsdr.cdc.gov/docs/atsdr_vapor_investigation.pdf</u>