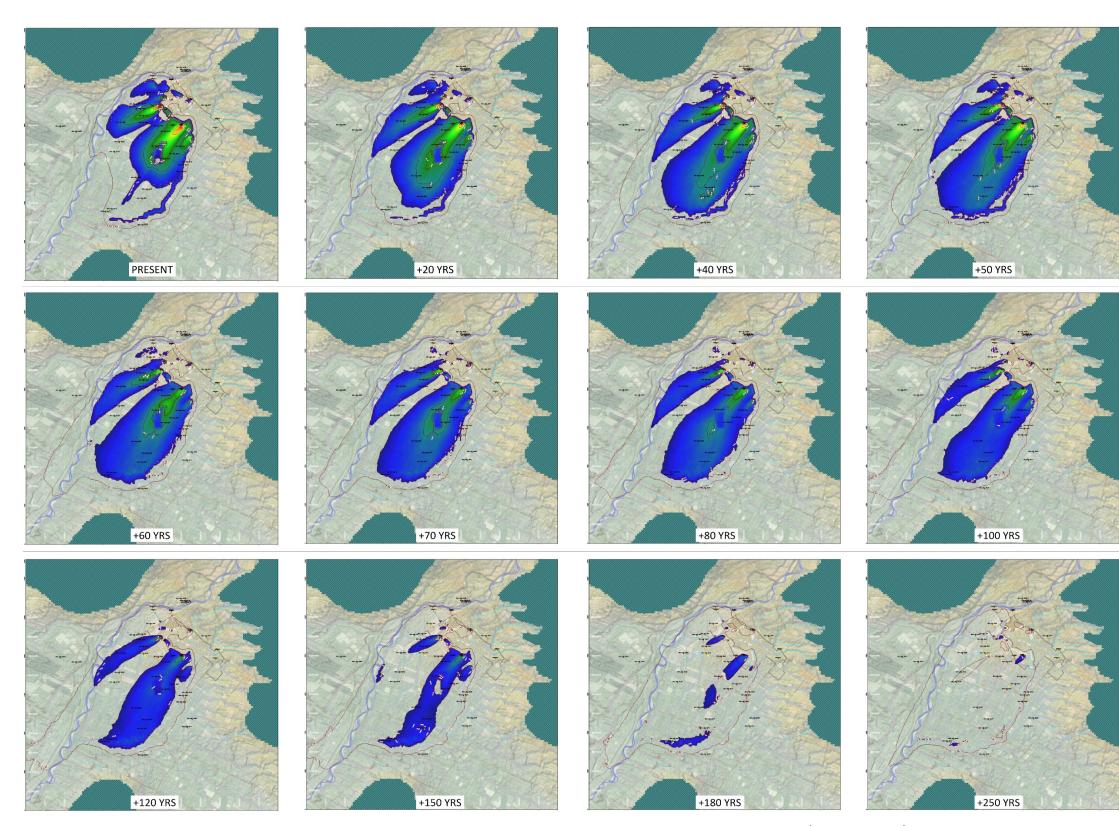
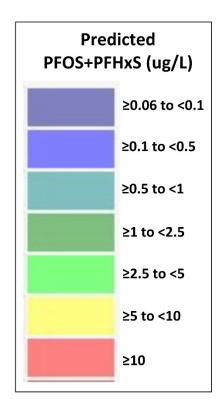
pop



**Diagram 9: Longer Source Depletion Future Plume Prediction (no retardation)** 





pop

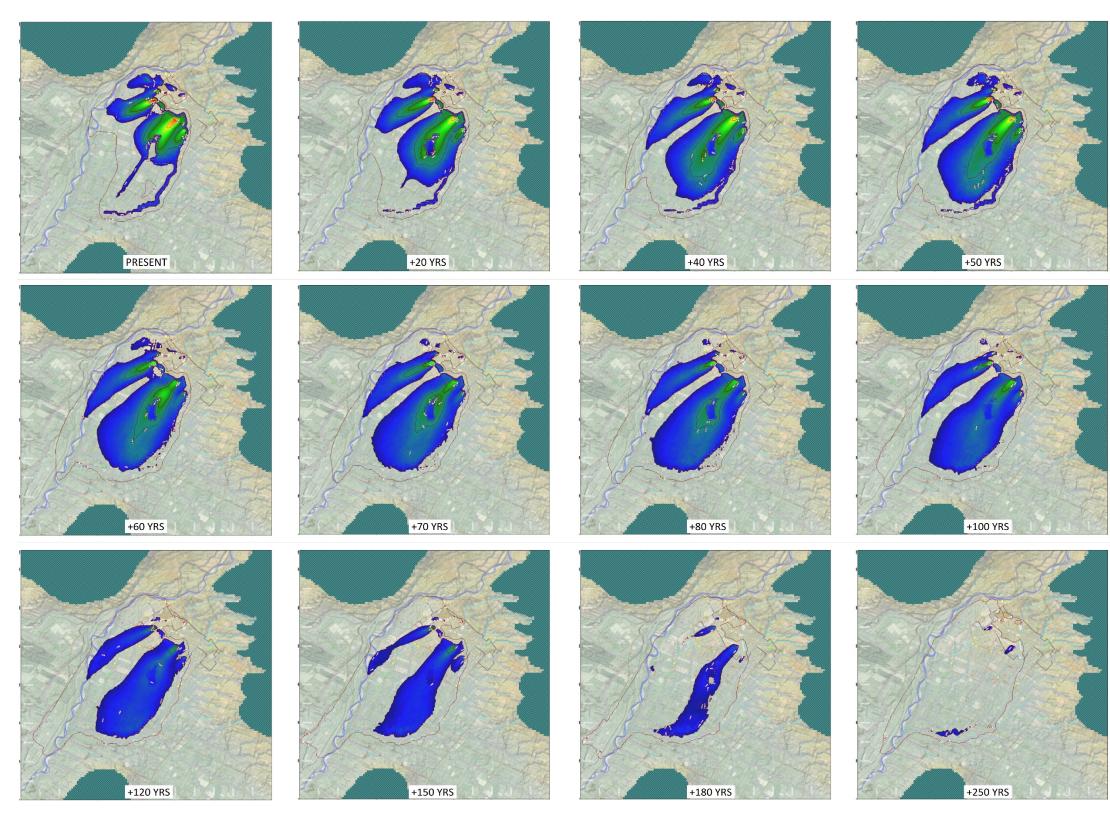
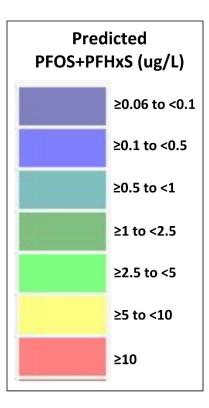


Diagram 10: Longer Source Depletion Future Plume Prediction (with retardation)



64



### 7.7 Plume Predictions - Discussion and Summary

A groundwater plume of PFOS + PFHxS has been modelled and interpreted based on all available observation data obtained during NZDF PFAS investigations. Mapping and interpretation of a PFOS + PFHxS plume has been utilised for the purposes of this project, rather than other PFAS species, due to the current relevance of these two species for drinking water toxicity guidelines, and because they are both terminal products. Other PFAS, many of which are not well understood, may behave differently.

The plume is sourced from at least 13 individual onsite Soil Source Zones. In theory, each Soil Source Zone is likely to be/have been producing an individual plume, but due to the geographical spread and nature of the groundwater flow system beneath the wider Ohakea site, these individual plumes have coalesced into essentially a single plume. Surface Water Source Zones have also been identified as key sources because they can transport contaminant mass, which can enter the groundwater system, long distances and quickly.

The existing plume (PFOS + PFHxS  $\geq 0.06 \text{ ug/L}$ ) has an estimated area of 1100 ha to 1600 ha, and has an estimated total PFOS + PFHxS mass (in solution) between 50 kg to 70 kg. An estimated 'above detection' extent has also been developed for the existing plume, with an estimated area of approximately 3600 ha. Whilst these are considered best estimates, there are gaps in data and knowledge on the plume extent, concentration distribution, and geochemical processes. Consequently, there is significant uncertainty associated with the aforementioned estimates. However, despite the uncertainties, the general plume extent is reasonably well covered spatially by physical observation data, and the present-day plume is interpreted to be well constrained in the northern and eastern direction (and in the western direction to a lesser extent). This has enabled development of a predictive assessment which is considered 'fit *for purpose'* with respect to the project objectives.

Into the future, the plume is expected to continue migration and expansion before beginning a slow process of depletion. This is primarily because while the source is not being added to (i.e. AFFF containing PFOS is no longer used) ongoing leaching from soil is occurring. The individual 'arms' of the plume are generally expected to continue advancing in their current direction of travel – generally west through south-southwest from Base Ohakea - until they encounter a major groundwater discharge boundary (i.e. Rangitikei River or Makowhai Stream). Surface water, particularly the Rangitikei River and Makowhai Stream, are the primary receptors of the plume. The plume discharges to these receptors (and their tributaries) as baseflow.



The hydrogeological setting in which the plume resides provides control on the fate and form of the plume into the future. In general, higher topography and groundwater pressures exist north, east and south of the existing plume. This effectively bounds the plume from migrating much further afield in these directions. Rather, the plume is expected to migrate north-west through south-southwest towards into the aforementioned Rangitikei River and Makowhai Stream (the regional groundwater sinks). It must be noted that plume migration/transport under and beyond these surface water bodies is possible, but as these are the regional groundwater sinks, they are the ultimate receivers, and migration back into these surface water bodies would ultimately occur, albeit slightly further downgradient.

Shallow wells (i.e. <50 m depth) which abstract groundwater from within the extent of the plume and the plumes predicted future migration path are also likely to be receptors. Deep wells e.g. >100 m depth, are less likely to be receptors of the plume. This is because the plume is generally predicted to be present and remain in the top portion of the groundwater system e.g. top 40 m to 60 m of saturation. Significant groundwater abstraction and/or poorly sealed boreholes do however have the potential to locally 'drag' the plume to greater depths.

The 'best estimate' of the likely time period for the existing plume (PFOS + PFHxS >0.06 ug/L) to decrease below its current area is estimated at approximately 75 years (no retardation) to 100 years (with retardation). The time to halve the existing plume area (PFOS + PFHxS >0.06 ug/L) is estimated at approximately 95 years (no retardation) to 125 years (with retardation). Even in a theoretical scenario where all source zones are instantaneously removed, it is expected that the plume (PFOS + PFHxS  $\geq$ 0.06 ug/L) would remain approximately the same area (as the existing plume) for at least the next 25 years (approximately). Consequently, all predictions and interpretations point towards the existing plume having a significant presence for time periods on the multidecade scale.

A maximum future extent of 'above detection' or ≥0.001ug/L (PFOS + PFHxS) is estimated at approximately 4300 ha. This extent should be considered as a probability extent e.g. PFOS + PFHxS detection outside of this extent is considered unlikely, but not impossible. The timing of when this maximum extent could be reached is likely to be in the long-term future i.e. >50 years.



# 8.0 Summary and Conclusions

Investigations at Ohakea have identified PFAS in soil and water on base, as well as in the surrounding environment and neighbouring properties.

The sampling programme completed between 2015 and 2018 included several rounds of groundwater and surface water monitoring, and sampling soil, sediment, animal tissue and plant tissue in various locations within and in the vicinity of the site.

In summary:

- PFAS was detected in all media sampled with the exception of goat's milk.
- Exceedances of applicable guidelines and trigger values were observed for groundwater (drinking water), surface water, eggs, fish tissue and watercress.
- The maximum PFAS concentrations observed were for PFOS for all media on-site. Similarly, off-site the maximum PFAS concentration was for PFOS for all media, except groundwater and surface water. Maximum PFHxS concentrations were higher than PFOS in off-site groundwater and surface water.
- Comparison of the sample results on-site and off-side shows that a significant proportion of PFAS mass in groundwater remains on-site. Median sum of PFOS + PFHxS is an order of magnitude higher than the median of off-site samples. PFAS concentrations off-site generally decreased with distance from the base with the exception of PFHxS.
- There is potentially significantly greater mass of PFAS (particularly PFOS) in the unsaturated soil than in the groundwater on-site.
- In general, PFAS concentrations in surface water decreased with increasing distance from the Base. The exception to this is the Makowhai Stream, where the lowest concentrations of PFOS were observed closest to the site, the highest concentrations approximately 1.5 km downstream from the base, before decreasing again downstream.
- Higher concentrations of PFOS were observed in the Makowhai Stream in summer months and lower concentrations in the wetter winter months.
- Evidence of transformation of PFAS was examined by comparing the molar concentration of PFAS from several groundwater wells extending south-west from the base. Some limited evidence of transformation of PFAS compounds in the plume was found.



Predictions have been made for the existing and future groundwater plume using 3D groundwater and flow and solute transport modelling:

- The existing plume (PFOS + PFHxS ≥0.06 ug/L) has an estimated area of 1100 ha to 1600 ha, and has an estimated total PFOS + PFHxS mass (in solution) in the order of 50 kg to 70 kg. An estimated 'above detection' extent has also been developed for the existing plume, with an estimated area of approximately 3600 ha. This estimate excludes other PFAS compounds. There is significant uncertainty associated with these estimates.
- Into the future, the plume is expected to continue migration and expansion before beginning a slow process of depletion. The plume is generally expected to continue advancing in the current direction of travel – generally west through south-southwest from Base Ohakea until encountering a major groundwater discharge boundary (i.e. Rangitikei River or Makowhai Stream). Surface water is the primary receptor of the plume.
- The 'best estimate' of the likely time period for the existing plume (PFOS + PFHxS >0.06 ug/L) to decrease below its current area is estimated to be in the order of 75 years (no retardation) to 100 years (with retardation).
- The time to halve the existing plume area (PFOS + PFHxS >0.06 ug/L) is estimated to be greater than 100 years; best estimate 95 years (no retardation) to 125 years (with retardation).
- A maximum future extent of 'above detection' or ≥0.001ug/L (PFOS + PFHxS) is estimated at approximately 4300 ha, and predicted to occur >50 years into the future. This extent should be considered as a probability extent e.g. PFOS + PFHxS detection outside of this extent is considered unlikely, but not impossible.
- A prediction was also completed whereby the existing sources were assumed to have already completely depleted. This scenario is considered analogous to a 'Best Possible Case' estimate and its purpose is to provide a prediction which tends towards the fastest perceivable (but unlikely) plume depletion.
- Under this scenario plume depletion is likely to be significantly more rapid than for the 'best estimate' scenario, however plume depletion is still on the multiple decade scale.
- A longer source depletion prediction scenario where the existing sources were assumed to take longer to deplete than for the 'best estimate' scenario was also undertaken. This produced a plume with an overall similar shape and aerial extent (as per the 'best estimate'); however, plume depletion took significantly longer e.g. approximately twice the duration.



The following conclusions have been drawn following interpretation of the sample results and modelling of the groundwater plume:

- The results and the literature indicate that there is potentially significantly greater mass of PFAS in the unsaturated soil than in the groundwater. Leaching of PFAS from the unsaturated soil could potentially provide an ongoing and long term source of PFAS to groundwater.
- Surface water has been identified as an important pathway for the migration of PFAS into groundwater and vice versa. Surface water flow can move contaminants much faster than groundwater flow, and due to the strong connection between groundwater-surface water within the region, contaminant transport via surface water is a key influencing factor for the groundwater plume.
- Interconnectedness of groundwater and surface water is further illustrated in the Makowhai Stream where PFOS concentrations are lower near Ohakea, reaching their maximum approximately 1.5 km from the site, before decreasing with increasing distance from the site.

One potential mechanism for this pattern is the influence of groundwater discharge (to the Makowhai Stream) where PFOS concentrations in the stream are highest.

- The plume of PFAS-containing groundwater emanating from historic use of AFFF at RNZAF Base Ohakea is expected to be constrained in the longer term by topography and higher groundwater pressures to the north, east and south and by the Rangitikei River to the west. It is expected that shallow groundwater in the investigation area is prevented from moving further south than approximately the Makowhai Stream and is instead directed towards the Rangitikei River. Plume migration/transport under and beyond these surface water bodies is possible, but as these are the regional groundwater sinks, they are the ultimate receivers, and migration back into these surface water bodies would ultimately occur, albeit slightly further downgradient.
- The plume is expected to persist in concentrations > 0.06 ug/L for many decades.



## 9.0 References

- AECOM, 2017. Environmental Site Assessment December 2017, RAAF Base Williamtown Stage 2B Monitoring: July 2017 to May 2018. Report prepared by AECOM Australia Pty Ltd.
- AECOM, 2018. Stage 2C Environmental Investigation– groundwater and Surface Water Monitoring: July 2017 to May 2018. Army Aviation Centre Oakey, Oakey QLD. Report prepared by AECOM Australia Pty Ltd.
- Alexander, D. (2012). The Rangitikei River its Tributary Waterways, and other Taihape Waterways - Scoping Report. Report prepared for Crown Forestry Rental Trust.
- Anderson, R. H., Long, G. C., Porter, R. C., & Anderson, J. K. (2016). Occurrence of select perfluoroalkyl substances at US Air Force aqueous film-forming foam release sites other than fire-training areas: Field-validation of critical fate and transport properties. *Chemosphere*, 150, 678-685.
- Awad, E., Zhang, X., Bhavsar, S. P., Petro, S., Crozier, P. W., Reiner, E. J., & Braekevelt, E. (2011). Long-term environmental fate of perfluorinated compounds after accidental release at Toronto airport. *Environmental* science & technology, 45(19), 8081-8089.
- Baduel, C., Paxman, C. J., & Mueller, J. F. (2015). Perfluoroalkyl substances in a firefighting training ground (FTG), distribution and potential future release. *Journal of hazardous materials*, 296, 46-53.
- Baduel, C., Mueller, J. F., Rotander, A., Corfield, J., & Gomez-Ramos, M. J. (2017). Discovery of novel per-and polyfluoroalkyl substances (PFASs) at a fire fighting training ground and preliminary investigation of their fate and mobility. *Chemosphere*, 185, 1030-1038.
- Bakke, T., Kailquist, T., Ruus, A., Breedveld, G. and Huylland, K. (2010). Norway Sediment Quality Guidelines. Journal of Soils and Sediment, 10, pp 172-178.
- Bartholomaeus, A. (2016). *Procedural review of health reference values established by enHealth for PFAS.* University of Canberra and University of Queensland, Australia.
- Batley, G., van Dam, R., Broadgate, K., Warne, M., Cooper, Naomi., and Brumley, C. (2018). Application of revised methodologies for the default guideline value derivation: PFOS in freshwater. SETAC North America 39<sup>th</sup> Annual Meeting. California, USA.
- Buck, R. C., Franklin, J., Berger, U., Conder, J. M., Cousins, I. T., de Voogt, P., Jensen, A. A., Kannan, K., Mabury, S. A., ... van Leeuwen, S. P. (2011).Perfluoroalkyl and polyfluoroalkyl substances in the environment:

terminology, classification, and origins. *Integrated environmental assessment and management*, 7(4), 513-41.

- Casson, R., & Chiang, S. Y. (2018). Integrating total oxidizable precursor assay data to evaluate fate and transport of PFASs. *Remediation Journal*, *28*(2), 71-87.
- CONCAWE, 2016. Environmental Fate and Effects of Poly- and Perfluoroalkyl Substances (PFAS). Report No. 8/16. 23 June 2016.
- Dauchy, X., Boiteux, V., Bach, C., Rosin, C., & Munoz, J. F. (2017). Per-and polyfluoroalkyl substances in firefighting foam concentrates and water samples collected near sites impacted by the use of these foams. *Chemosphere*, *183*, 53-61.
- Dauchy, X., Boiteux, V., Colin, A., Hémard, J., Bach, C., Rosin, C., & Munoz, J. F. (2019). Deep seepage of per-and polyfluoroalkyl substances through the soil of a firefighter training site and subsequent groundwater contamination. *Chemosphere*, 214, 729-737.
- EnRiskS, 2017. *Livestock Uptake Modelling and Screening Criteria Development for PFAS, draft*. Revision C. Environmental Risk Sciences Pty Ltd. 1 November 2017.
- FSANZ, 2017. Assessment of potential dietary exposure to perfluorooctane sulfonate (PFOS), perfluorooctanoic acid (PFOA) and perfluorohexane sulfonate (PFHxS) occurring in foods sampled from contaminated sites Table 8, Supporting Document 2. Food Standards Australia New Zealand, April 2017.
- Gellrich, V., Stahl, T., & Knepper, T. P. (2012). Behavior of perfluorinated compounds in soils during leaching experiments. *Chemosphere*, *87*(9), 1052-1056.
- HEPA, 2018. *PFAS National Environmental Management Plan.* Heads of EPAs Australia and New Zealand, January 2018.
- HEPA, 2019. *PFAS National Environmental Management Plan 2.0 Draft*. Heads of EPAs Australia and New Zealand, April, 2019.
- HC, 2019. Health Canada's Drinking Water Screening Values for Other PFAS.
  Health Canada. April 2019.
  https://www.canada.ca/en/services/health/publications/healthy-living/water-talk-drinking-water-screening-values-perfluoroalkylated-substances.html Accessed. 4/06/19.
- Houtz, E. F., Sedlak, D. L. (2102). Oxidative Conversion as a Means of Detecting Precursors to Perfluoroalkyl Acids in Urban Runoff. *Environmental science* & technology, 46, 9342–9349.

- Houtz, E. F., Higgins, C. P., Field, J. A., & Sedlak, D. L. (2013). Persistence of perfluoroalkyl acid precursors in AFFF-impacted groundwater and soil. *Environmental science & technology*, 47(15), 8187-8195.
- Houtz, E. F., Sutton, R., Park, J. S., & Sedlak, M. (2016). Poly-and perfluoroalkyl substances in wastewater: Significance of unknown precursors, manufacturing shifts, and likely AFFF impacts. *Water research*, 95, 142-149.
- ITRC, 2018a. PFAS Fact Sheet: Environmental Fate and Transport for Per- and Polyfluoroalkyl Substances. Interstate Technology Regulatory Council, March 2018.
- ITRC, 2018b. *PFAS Fact Sheet: Aqueous Film Forming Foam (AFFF).* Interstate Technology Regulatory Council, October 2018.
- ITRC, 2017. PFAS Fact Sheet: History and Use of Per- and Polyfluoroalkyl Substances (PFAS). Interstate Technology Regulatory Council, November 2017.
- Kwadijk, C. J., Kotterman, M., & Koelmans, A. A. (2014). Partitioning of perfluorooctanesulfonate and perfluorohexanesulfonate in the aquatic environment after an accidental release of aqueous film forming foam at Schiphol Amsterdam Airport. *Environmental toxicology and chemistry*, 33(8), 1761-1765.
- McKenzie, E. R., Siegrist, R. L., McCray, J. E., & Higgins, C. P. (2016). The influence of a non-aqueous phase liquid (NAPL) and chemical oxidant application on perfluoroalkyl acid (PFAA) fate and transport. *Water research*, *92*, 199-207.
- MoH, 2017. Poly-fluoroalkyl substances (PFASs), also called perfluoroalkyl substances (PFASs) draft, Ministry of Health November 2017.
- PDP, 2015a. Contaminated Site Investigation: RNZAF Base Ohakea Fire Training Area Preliminary Site Investigation. Report prepared for New Zealand Defence Force, March 2015.
- PDP, 2015b. Contaminated Site Investigation: RNZAF Base Ohakea Fire Training Area Detailed Site Investigation. Report prepared for New Zealand Defence Force, November 2015.
- PDP, 2017a. *PFAS Preliminary Site Investigation: RNZAF Base Ohakea*. Report prepared for New Zealand Defence Force, May 2017.
- PDP, 2017b. *PNZDF Ohakea Groundwater Assessment and PFAS Fate Prediction.* Report prepared for New Zealand Defence Force, November 2017.

- PDP, 2018a. *Stage 1 PFAS Detailed Site Investigation: RNZAF Base Ohakea*. Report prepared for New Zealand Defence Force, September 2018.
- PDP, 2018b. NZDF PFAS Investigation Summary Report: RNZAF Base Ohakea, Stage 1. Revised report prepared for New Zealand Defence Force, February 2018.
- PDP, 2018c. NZDF PFAS Investigation Summary Report: RNZAF Base Ohakea, Stage B. Report prepared for New Zealand Defence Force, April 2018.
- PDP, 2018d. NZDF PFAS Investigation Summary Report: RNZAF Base Ohakea, Stage C. Report prepared for New Zealand Defence Force, June 2018.
- PDP, 2018e. NZDF PFAS Investigation Summary Report: RNZAF Base Ohakea, Stage D. Report prepared for New Zealand Defence Force, November 2018.
- PDP, 2018f. NZDF PFAS Investigation Summary Report: RNZAF Base Ohakea, Stage D. Report prepared for New Zealand Defence Force, August 2018.
- PDP, 2018g. Sampling Protocols for Monitoring Per and Poly-fluorinated Compounds in Groundwater and Surface Water for New Zealand Defence Force. Report prepared for the Ministry for the Environment, May 2018.
- PDP, 2018f. *Impact of Per and Poly Fluoroalkyl Substances on Ecosystems.* Report prepared for the Ministry for the Environment, November 2018.
- PDP, 2019. Supplementary Report Groundwater Modelling. Report prepared for New Zealand Defence Force, May, 2019.
- Place, B. J., & Field, J. A. (2012). Identification of novel fluorochemicals in aqueous film-forming foams used by the US military. *Environmental* science & technology, 46(13), 7120-7127.
- Prevedourous, K., Cousins, I., Kaiser, M., and Buxton, L. (2006). Sources, fate and transport of perfluorocarboxylates. Environmental Science and technology, 40, pp. 32-44.
- Scher, D. P., Kelly, J. E., Huset, C. A., Barry, K. M., Hoffbeck, R. W., Yingling, V. L., & Messing, R. B. (2018). Occurrence of perfluoroalkyl substances (PFAS) in garden produce at homes with a history of PFAS-contaminated drinking water. *Chemosphere*, 196, 548-555.
- Townsend, D., Vonk, A., Kamp, P.J.J. (compilers), 2008: Geology of the Taranaki area: scale 1:250,000. Lower Hutt: GNS Science. Institute of Geological & Nuclear Sciences 1:250,000 geological map 7. 77 p.
- Wang, Z., M. MacLeod, I.T. Cousins, M. Scheringer, and K. Hungerbuhler (2011). Using COSMOtherm to predict Physicochemical Properties of Poly- and Perfluorinated Alkyl Substances (PFASs). *Environmental Chemistry* (8). 389-398



- Weber, A. K., Barber, L. B., LeBlanc, D. R., Sunderland, E. M., & Vecitis, C. D.
  (2017). Geochemical and hydrologic factors controlling subsurface transport of poly-and perfluoroalkyl substances, Cape Cod, Massachusetts. *Environmental science & technology*, *51*(8), 4269-4279.
- Zareitalabad, P., Siemens, J., Hamer, M., Amelung, W. (2013). Perfluorooctanoic acid (PFOA) and perfluorooctanesulfonic acid (PFOS) in surface waters, sediments, soils and wastewater A review on Concentrations and distribution coefficients. *Chemosphere*, (91) 725–732.
- Zhang, S., Lu, X., Wang, N., & Buck, R. (2016) Biotransformation potential of 6:2 fluorotelomer sulfonate (6:2 FTSA) in aerobic and anaerobic sediment. Chemosphere, 154, 224-230.