

GEMS/AMIS Air Quality Monitoring Programme Annual Report 2005

Prepared by
Watercare Services Limited
Laboratory Services Air Quality Department
New Zealand Accredited Laboratory

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1 Summary

This report contains the 2005 annual ambient air quality dataset measured from two sites in Auckland and one site in Christchurch. These sites form New Zealand's contribution to the global environmental monitoring system (GEMS).

The GEMS monitoring sites were established to measure key air pollutants associated with adverse effects on people's health and well-being. The GEMS sites include some of the longest-running air quality monitoring sites in New Zealand. For example, various air quality monitoring has been undertaken at the Gavin Street, Penrose site in Auckland since 1964. These sites were established to determine the effects of policies for air quality management and are important for recording trends in pollution levels in New Zealand. The GEMS sites have provided long-term continuity in monitoring data for Auckland and Christchurch and provide an important snapshot of air quality for two of New Zealand's largest cities.

The two Auckland monitoring sites include a site dominated by residential and vehicle emissions (Kowhai Intermediate School, Kingsland) and a site representative of vehicle and industrial emissions (Gavin Street, Penrose). The Greers Road, Burnside site in Christchurch is located within a residential area and represents emissions from domestic properties.

Air pollutants arise from a number of different sources. Particulate matter (PM₁₀) arises from stationary and mobile combustion sources, principally domestic heating, industrial processes and vehicle emissions (as well as natural processes such as soil erosion and sea salt). Sulphur dioxide is produced from the burning of fossil fuels, particularly coal and oil. Carbon monoxide is a product of incomplete combustion of carbon containing fuels, especially from motor vehicles. Volatile organic compounds (VOCs) are organic chemicals, such as hydrocarbons, that are closely tied to vehicle emissions and many industrial processes. Historically, lead was a petrol additive but since the significant reduction of lead in petrol from 1996, levels have declined in New Zealand.

Five ambient air quality standards for carbon monoxide, nitrogen dioxide, ozone, PM₁₀ and sulphur dioxide were promulgated in October 2004. These standards are the minimum requirements that outdoor air quality must meet to guarantee a set level of protection for human health and the environment. The ambient standards are based on existing ambient air quality guidelines. Guideline levels for pollutants (and averaging periods) not covered by the standards still apply.

For most of the time, air pollutants at all sites during 2005 were below the standards. However, there were times when these standards were exceeded. One exceedance of the nitrogen dioxide 1-hour standard at Gavin Street, Penrose was found to be caused by local construction works where machinery was operating several metres from the monitoring site. The ambient air standard for NO₂ makes allowance for nine 1-hour exceedances per year before the standard is breached. At Greers Road, Burnside, there were 18 exceedances of the 24-hour standard for PM₁₀, mainly during the winter months from June to August 2005. These exceedances were most likely caused by home heating emissions. The standard allows for one exceedance of the PM₁₀ 24-hour threshold per year.

2 Introduction

This report presents the 2005 ambient air quality data set for Auckland and Christchurch, New Zealand. The monitoring is conducted by Watercare Services Ltd, on behalf of the Ministry for the Environment (the Ministry, MfE).

The Ministry has a Memorandum of Understanding with the New Zealand Ministry of Health (MoH) to collect and supply air quality monitoring data to the World Health Organization (WHO) from three sites – two in Auckland and one in Christchurch.

This data has historically formed New Zealand's contribution to the WHO's global environmental monitoring system/air pollution programme (GEMS/AIR) which began in 1973.¹

In 1996, the WHO developed the air management information system (AMIS), the successor to GEMS/AIR. The objective of AMIS is to transfer information on air pollutant concentrations and air quality management between countries. It also aims to support and assist in the maintenance of air quality in parts of New Zealand that enjoy clean air and to improve air quality in places where it has deteriorated.

As a result, monitoring from the AMIS programme is used by the Ministry to support and enhance ambient air quality monitoring and management in Auckland and Christchurch. In fact, the GEMS/AMIS ambient air quality sites are the longest-running sites in New Zealand and, as such, are very important in identifying local long-term trends in air pollution.

The Auckland sites are located in the industrial area of Penrose, which is to the southeast of the city centre, and in Mt Eden and Kingsland, both of which are older residential areas just south of the city centre. Air quality monitoring has been performed in Penrose since 1964 and at Mt Eden between 1982 and 2004. In October 2004, the Mt Eden site was decommissioned pending redevelopment of the site and replaced by the Kingsland site at Kowhai Intermediate School.

Between 1989 and 2002, monitoring was undertaken in Christchurch at a site located in the older residential area of St Albans which is just north of the city centre. Due to impending redevelopment of this site, the monitoring station was relocated in November 2002 to a site in Burnside/Bishopdale which is a newer residential area to the northwest of the city centre.

Environment Canterbury provided sampling services for gravimetric and passive monitoring methods at the two Christchurch sites (Coles Place, St Albans and Greers Road, Burnside).

This report includes graphical and statistical presentations of the data as well as any data collection issues that may have arisen during the monitoring period.

All data in this report has been completely validated. Quality assurance checks have been carried out to ensure that invalid and calibration data is not reported.

¹ Schwela DH. 1999. Public health and the air management information system (AMIS). *Epidemiology* 10(5): 647-55.

3 Air Pollutants Monitored

3.1 Particulate matter

Particulate matter can be a significant air pollutant that is associated with a variety of health and environmental effects.

Sources of particulates vary widely from location to location reflecting the diverse range of emission sources that contribute to particulate concentrations in New Zealand. Typical sources can include:

- fine particulates emitted as a result of fuel combustion such as road vehicles, power generation, industrial processes, domestic heating appliances etc
- particulates formed by chemical reactions in the atmosphere, comprising mainly sulphates and nitrates
- coarse particulates arising from a wide range of sources, including re-suspended dusts from road vehicles, construction works and mineral extraction processes, wind-blown dusts and soils, sea salt and biological particles such as pollen.

There are a variety of measurements that can be used to determine the different health and environmental effects of particulate matter. As part of the GEMS/AMIS programme two particle size fractions are monitored:

- fine particulates (PM₁₀)
- total suspended particulates (TSP).

3.1.1 Fine particulates (PM₁₀)

As described above, particles with a diameter of 10 µm or less can be inhaled into the respiratory system and affect human health. The coarser fraction of airborne particles (2.5 to 10 µm) are deposited in the trachea bronchial region where asthma attacks are triggered.

Particulate matter refers to numerous substances that exist in the atmosphere. It is a somewhat complex pollutant, encompassing a wide range of chemically and physically diverse substances. Particulate matter includes all solids and aerosols that exist in ambient conditions.

3.1.2 Total suspended particulates (TSP)

TSP consists of all particles that range in size up to 50 µm in diameter. TSP is sufficiently small to be inhaled, however the larger particles (10-50 µm) are readily filtered out in the nasal cavity. Particles 10 µm and less can be drawn into the respiratory system. TSP affects both the aesthetic and health quality of the ambient air.

3.2 Lead

Lead is a toxic metal emitted into the air both from motor vehicles that use leaded fuel and some industries. Since lead was removed from fuel in 1996, concentrations of lead in the air have dropped markedly. In October 2000, monitoring of lead was reduced from monthly samples to samples taken over a three-month period during winter only (June – August).

3.3 Sulphur dioxide

Sulphur dioxide is an acidic gas with a pungent odour which is mainly produced by the burning of fossil fuels. The gas is quite corrosive and can cause damage to buildings and other materials.

It can also have significant effects on the human respiratory system. Inhalation of high ambient concentrations of sulphur dioxide can cause stimulation of the nerves in the air passages, resulting in a reflex cough, irritation and chest tightness.

In addition, sulphur dioxide can also cause narrowing of the air passages, particularly in people suffering from asthma and chronic lung disease. These people frequently have narrowed airways, and any further restriction will have a disproportionately large effect compared to people with uncompromised respiratory systems.

3.4 Carbon monoxide

This colourless, odourless, toxic gas is formed as a product of incomplete combustion in the burning of fossil fuels. The main sources in most parts of New Zealand are motor vehicle exhaust emissions, and, as such, elevated levels are mainly found in areas of significant traffic congestion, particularly at busy intersections on inner-city streets.

Carbon monoxide acts on humans by displacing oxygen from the blood. Prolonged exposure at moderate levels can lead to symptoms such as headaches and dizziness, while at high levels it can lead to loss of consciousness and even death. At the lower levels typically encountered in urban areas, carbon monoxide measurements can serve as a useful indicator for objectionable levels of vehicle exhaust fumes.

3.5 Nitrogen oxides

Nitrogen oxides incorporates several species that exist in the atmosphere which are collectively referred to as NO_x . The two main oxides are nitrogen dioxide (NO_2), which is of concern due to its potential to cause health effects, and the monoxide form nitric oxide (NO), which is less toxic but may oxidise to NO_2 in the atmosphere.

Nitrogen oxides are formed in most combustion processes by oxidation of the nitrogen present in the atmosphere. Nitric oxide is the predominant primary product but, as indicated, this can then be oxidised to nitrogen dioxide in ambient air. Emissions from motor vehicles are the major source of NO_x in most parts of the country, although power stations and other large combustion units may be significant localised sources as well.

The main health effects of the oxides of nitrogen are due to NO_2 which is a respiratory irritant. Nitric oxide is believed to be quite harmless at the levels normally encountered in urban air.

NO_x is also an important air pollutant because of its role in photochemical smog. NO_2 is a reddish brown gas and has synergistic effects with other pollutants such as SO_2 and particulate.

3.6 Volatile organic compounds

Volatile organic compounds are chemicals that easily evaporate at room temperature. The term 'organic' indicates that the compounds contain carbon.

To rationalise air quality guidelines, the Ministry for the Environment has compiled a list of priority contaminants based on a review of international literature. The priority list includes the volatile organic compounds (VOC) benzene and 1,3-butadiene and provides ambient air quality guidelines for these contaminants (MfE 2002).

4 Ambient Air Quality Guidelines and Standards

In October 2004, the Ministry for the Environment introduced the National Environmental Standards (NES) for Air Quality. The NES includes five standards for ambient (outdoor) air quality. These and other New Zealand guidelines are described in Table 1 below.

The criteria applied to TSP, $60 \mu\text{g}/\text{m}^3$ (seven-day average), was previously applied by the Ministry of Health. This has been superseded by the Ministry for the Environment's Ambient Air Quality Guidelines but is useful for analysing the results of the monitoring data.

Table 1: National Environmental Standards, guidelines and regional targets²

Air pollutant	National Environmental Standards 2004	Ministry for the Environment Ambient Air Quality Guidelines 2002 and other	Averaging period	National Environmental Standards permissible excess
Carbon monoxide	$10 \text{ mg}/\text{m}^3$	$30 \text{ mg}/\text{m}^3$	8-hour average 1-hour average	One 8-hour period in a 12-month period
Nitrogen dioxide	$200 \mu\text{g}/\text{m}^3$	$100 \mu\text{g}/\text{m}^3$	24-hour average 1-hour average	9 hours in a 12-month period
Sulphur dioxide	$350 \mu\text{g}/\text{m}^3$ $570 \mu\text{g}/\text{m}^3$	$120 \mu\text{g}/\text{m}^3$	24-hour average 1-hour average 1-hour average	9 hours in a 12-month period Not to be exceeded at any time
Benzene – Year 2000 – Year 2010		$10 \mu\text{g}/\text{m}^3$ $3.6 \mu\text{g}/\text{m}^3$	Annual average Annual average	
1,3-Butadiene		$2.4 \mu\text{g}/\text{m}^3$	Annual average	
Fine particulate (PM_{10})	$50 \mu\text{g}/\text{m}^3$	$20 \mu\text{g}/\text{m}^3$ $50 \mu\text{g}/\text{m}^3$ (ARC and ECAN)	Annual average 24-hour average	One 24-hour period in a 12-month period
Total suspended particulates (TSP)		$60 \mu\text{g}/\text{m}^3$ (DoH) ²	7-day average	
Lead		$0.2 \mu\text{g}/\text{m}^3$	3-month average	

² See <http://www.mfe.govt.nz/laws/standards/air-quality-standards.html>

5 Monitoring Sites


5.1 Site descriptions

A brief description of all the monitoring sites in the GEMS/AMIS air quality monitoring programme is given below. This includes the two Auckland sites – at Gavin Street, Penrose and Kowhai Intermediate School, Kingsland – and two sites in Christchurch – at Greers Road, Burnside and Coles Place, St Albans. (Note: at Coles Place only TSP and lead are monitored by the Ministry for the Environment.)


5.1.1 MfE Kowhai, Auckland – Site AKL073

Site name	MfE Kowhai, Kingsland		Site ID	AKL073
Address	Kowhai Intermediate School, Sandringham Road, Auckland		Site class	Residential – peak
Description				
This site is located within the grounds of Kowhai Intermediate School. It is surrounded by residential properties on three sides as well as the school buildings which lie about 100 metres to the east. The busy New North Road is approximately 100 metres to the north of the site while Sandringham Road runs northwest to southeast past the site and Eden Park rugby ground is within 300 metres to the southeast of the site. The site is representative of emissions arising from road vehicles as well as domestic properties in the older inner-city area of Kingsland which lies to the south of Auckland city centre. This is a new site, commissioned in 2004, and designed to replace the neighbouring Kelly Street site in Mt Eden which is due to be redeveloped. The new Kowhai site lies about 500 metres to the west of the Kelly Street and both sites have the same 'residential – peak' classification. During 2004, a period of parallel monitoring between the two sites was undertaken before all monitoring was relocated to the new Kowhai site in October 2004.				
Pollutants monitored	CO N	NO ₂ Y	SO ₂ N	VOCs Y
	PM ₁₀ Y	TSP Y	Lead Y	
Meteorological parameters monitored	Wind speed Y	Wind direction Y	RH Y	
	Temperature (6 m) Y	Temperature (10 m) N	Temperature (2 m) N	

Location map



Photograph



5.1.2 MfE Gavin Street, Penrose, Auckland – Site AKL009

Site name	MfE Gavin Street, Penrose		Site ID	AKL009
Address	Transpower, Gavin Street, Penrose, Auckland		Site class	Industrial – dense / traffic – peak
Description				
This site is located within the grounds of the Transpower NZ Ltd electrical substation on Gavin Street. It is representative of road vehicle and industrial emissions in the Penrose area which lies to the southeast of Auckland city centre and is also approximately 50 metres northeast of the Southern Motorway. There are residential properties immediately to the northeast of the site. During 2003, parallel monitoring was undertaken between this site and the neighbouring ACI site on the Great South Road in Penrose with a view to consolidating all monitoring at the Gavin Street site early in 2004.				
Pollutants monitored	CO N	NO ₂ Y	SO ₂ Y	VOCs Y
	PM ₁₀ Y	TSP Y	Lead Y	
Meteorological parameters monitored	Wind speed Y	Wind direction Y	RH Y	
	Temperature (6 m) Y	Temperature (10 m) N	Temperature (2 m) N	

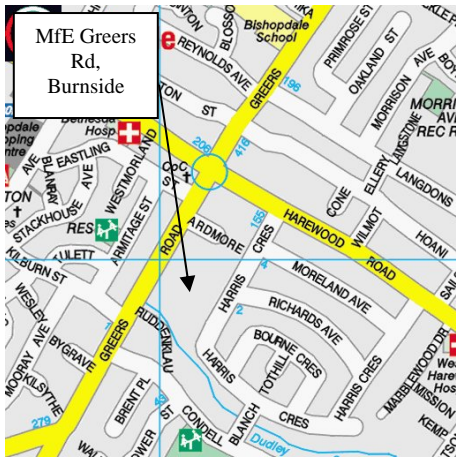
Location map

Photograph


5.1.3 MfE Greers Road, Burnside, Christchurch – Site CAN002

Site name	MfE Greers Road, Burnside		Site ID	CAN002
Address	Transpower, Greers Road, Burnside, Christchurch		Site class	Residential – neighbourhood
Description				
This site is located in a paddock to the rear of the Transpower New Zealand Ltd electrical substation on Greers Road and is surrounded by residential properties on four sides. It is representative of emissions arising from domestic properties in the newer suburban areas of Burnside and Bishopdale which lie to the northwest of Christchurch city centre.				
Pollutants monitored	CO Y	NO ₂ Y	SO ₂ Y	VOCs Y
	PM ₁₀ Y	TSP N	Lead N	
Meteorological parameters monitored	Wind speed Y	Wind direction Y	RH Y	
	Temperature (6 m) N	Temperature (10 m) Y	Temperature (2 m) Y	

Location map



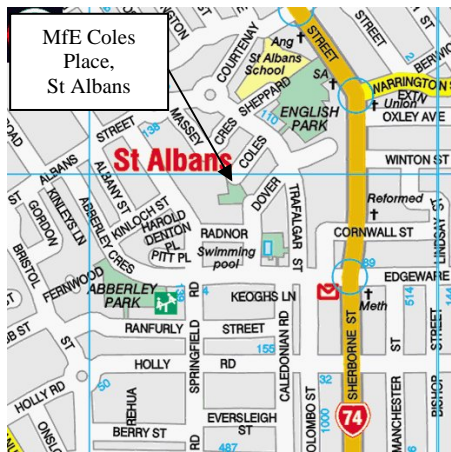
Photograph




5.1.4 MfE Coles Place, St Albans, Christchurch – Site CAN003

Site name	MfE Coles Place, St Albans		Site ID	CAN003
Address	Coles Place, St Albans, Christchurch		Site class	Residential – neighbourhood
Description				
This site is operated by Environment Canterbury and is located on an area of public open space at the end of Coles Place. It is surrounded by residential properties on four sides. It is representative of emissions arising from domestic properties in the older suburban area of St Albans which lies to the north of Christchurch city centre.				
Pollutants monitored	CO	NO ₂	SO ₂	VOCs
	N	N	N	Y
Meteorological parameters monitored	PM ₁₀	TSP	Lead	
	N	Y	Y	
	Wind speed	Wind direction	RH	
	N	N	N	
	Temperature (5 m)	Temperature (10 m)	Temperature (2 m)	
	N	N	N	

Location map



Photograph



5.2 Air pollutants monitored at GEMS/AMIS sites

An overview of the air pollutants monitored at each site as part of the GEMS/AMIS programme is given in Table 2 below.

Table 2: Air pollutants monitored at GEMS/AMIS sites

Site	CO	NO ₂	SO ₂	VOC	PM ₁₀	TSP	Lead*
Kowhai Intermediate, Kingsland AKL073N		✓		✓	✓	✓	✓
Transpower, Gavin Street, Penrose, Auckland AKL009		✓	✓	✓	✓	✓	✓
Greers Road, Burnside, Christchurch CAN002	✓	✓	✓	✓	✓		
Coles Place, St Albans, Christchurch CAN003				✓		✓	✓

Note:

* Lead is monitored in the months of June, July and August only.

6 Methods

6.1 Quality assurance

All monitoring services are undertaken by Watercare Services Ltd in accordance with the Ministry for the Environment's *Good Practice Guide for Air Quality Monitoring and Data Management* and, wherever applicable, the appropriate Australian/New Zealand and US EPA monitoring methods.

Watercare Laboratory Services is accredited by IANZ (International Accreditation New Zealand) and, since October 2003, has held accreditation for the following methods:

- Australian Standard AS 3580.7.1 – 1992 'Method 7.1: Determination of carbon monoxide – direct reading instrumental method'.
- Australian Standard AS 3580.5.1 – 1993 'Method 5.1: Determination of oxides of nitrogen – chemiluminescence method'.
- Australian Standard AS 3580.4.1 – 1990 'Method 4.1: Determination of sulphur dioxide – direct reading instrumental method'.
- US EPA Method 40, Part 50, Appendix J 'Reference Method for the Determination of Particulate Matter as PM₁₀ in the Atmosphere'.

As part of the GEMS/AMIS programme, Watercare Laboratory Services provides the following monitoring services:

- instrument operation, calibration and maintenance: this includes the use of automatic daily calibration systems for all continuous ambient gas monitors and ensures that the requirements of the relevant Australian Standards for weekly calibration of continuous analysers
- site maintenance as well as, when necessary, the commissioning of new sites and decommissioning of old sites
- data logging, polling, checking, rescaling, validation, ratification and reporting: this encompasses the entire data quality assurance process and ensures that the final dataset reported is fit for the purpose of the GEMS/AMIS programme.

6.2 Analytical methods

6.2.1 Carbon monoxide

Measurements are made in accordance with AS 3580.7.1 – 1992 ‘Determination of carbon monoxide – direct reading instrumental method’. The performance of the instruments is checked using an automatic calibration system. This ensures compliance with the method which requires instrumentation to be calibrated on a weekly basis.

The instruments themselves are IR-absorption gas analysers which continuously measure carbon monoxide. This allows data to be analysed and reported over a variety of average periods, including 10 minutes, 24 hours and one year.

6.2.2 Nitrogen oxides

Measurements are made in accordance with AS 3580.5.1 – 1993 ‘Determination of oxides of nitrogen –chemiluminescence method’. The performance of the instruments is checked using an automatic calibration system. This ensures compliance with the method which requires instrumentation to be calibrated on a weekly basis.

The instruments themselves are chemiluminescence gas analysers which continuously measure nitrogen oxides. This allows data to be analysed and reported over a variety of average periods, including 10 minutes, 24 hours and one year.

6.2.3 Sulphur dioxide

Measurements are made in accordance with AS 3580.4.1 – 1990 ‘Determination of sulphur dioxide – direct reading instrumental method’. The performance of the instruments is checked using an automatic calibration system. This ensures compliance with the method which requires instrumentation to be calibrated on a weekly basis.

The instruments themselves are UV-fluorescence gas analysers which continuously measure sulphur dioxide. This allows data to be analysed and reported over a variety of average periods, including 10 minutes, 24 hours and one year.

6.2.4 Volatile organic compounds

VOCs are measured each quarter (January – March, April – June, July – September and October – December) in accordance with Watercare’s Air Quality Group Test Method T114 (ref – NIOSH Methods 1500 and 1501).

VOC samples are taken using passive (3M) sampling badges which are exposed for a three-month period. The VOCs diffuse on to the badges which are coated with activated carbon. Following exposure, the samples are forwarded to AgriQuality who extract the VOCs using carbon disulphide and analyse them using GC-MS.

Note: Determination of 1,3-butadiene over a three-month exposure period is not reliable. Investigations have determined that samples are unstable when held above -4°C (OSHA Method 54, NIOSH Method 1024), with significant desorption occurring. Technical information supplied with 3M badges reports a 10 percent loss of 1,3-butadiene over three weeks' storage at room temperature. Due to the potential for error over a three-month exposure period, 1,3-butadiene has not been analysed and reported.

6.2.5 Particulate matter (PM₁₀)

Measurements are made in accordance with the US EPA equivalent method for measuring PM₁₀ EQPM-1102-150 'Thermo Andersen Series FH62-C14 Continuous PM₁₀ Ambient Particulate Monitor'. This method was designated as an equivalent method by the US EPA in accordance with 40 CFR Part 53 on 11 December 2002.

The Thermo Andersen FH62-C14 is fitted with a size-selective PM₁₀ head and measures particle mass as it accumulates during sampling. As a result, the instrument is able to record and output real-time measurements of PM₁₀ data which allows measurements to be reported over a variety of average periods, including 10 minutes, 24 hours and one year. The inlet temperature of all beta-gauges operated by the Ministry for the Environment is 40°C.

6.2.6 Particulate matter (TSP)

Measurements of TSP are made in accordance with Watercare's Air Quality Group Test Method T101. It is a gravimetric method of measuring particulates and is modelled upon the high-volume sampler method. The technique has been used to provide TSP data at existing GEM/AMIS sites since 1982.

The equipment used to measure TSP is quite basic and involves ambient air being pulled through a glass fibre filter by a pump with a gas metre being used to measure the air volume drawn through the filter. The filter is weighed before and after sampling. The TSP concentration is determined from the weight of particulates collected and the air volume sampled.

6.2.7 Lead

Lead is sampled during the winter months (June – August) with the same instrumentation used to measure TSP according to Watercare's Air Quality Group Test Method T101. This is a gravimetric technique used to measure particulates and is modelled upon the high volume sampler method. The technique has been used to provide TSP data at existing GEMS sites since 1987.

Analysis of lead is performed by Watercare Laboratory Services according to US EPA methods 3051 and 2007. This involves analysing each individual TSP filter exposed during the winter period using mixed acid digestion. This sample is then analysed for lead using the technique of ICP-OES. The concentration of lead is then determined from the amount of lead detected and the total volume of air sampled during that sample period. Concentrations are then averaged for the three-month monitoring period.

7 Results and Discussion

7.1 Site performance and quality assurance

Continuously monitored pollutant's instrument performance during 2005 was generally very good with all sites having data capture rates well over 90 percent.

Valid NO₂, SO₂, CO, and PM₁₀ data from the Greers Road, Burnside site were below 90 percent for the month of February due to data-logger problems early in the month (8 to 11 February 2005) and at the end of the month (25 to 28 February 2005). Valid data capture rates for these pollutants were greater than 90 percent for 2005.

Overall site performance is shown in Table 3 below based on 10-minute averages for continuously monitored data. Percent data capture is the percent of total instrument availability and includes downtime for calibration and routine maintenance. Percent valid data is defined as the percent valid data following quality assurance eg, invalidation of data resulting from calibrations, routine maintenance, spurious data and excessively negative data.

Table 3: Percentage valid and capture data 2005

Analyte	Site	Percentage valid data (V) and percentage data capture (C)																											
		January		February		March		April		May		June		July		August		September		October		November		December		Annual			
		V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C	V	C		
CO	Burnside	94.9	99.5	69.3	77.7	95.1	100	95.3	100	94.7	100	95.2	100	95.6	100	95.6	100	91.7	100	94.3	100	97.6	100	97.66	100	95.7	98.1		
NO ₂	Penrose	94.3	99.3	87.5	99.4	95.9	100	95.6	100	95.5	100	98.4	100	98.0	100	98.5	100	98.4	100	98.5	100	86.1	100	97.9	100	93.5	100		
NO ₂	Burnside	96.1	99.5	69.1	77.5	94.9	100	95.3	100	94.8	100	95.3	100	95.0	100	95.3	100	91.3	100	94.2	100	98.4	100	98.0	100	93.5	98.1		
NO ₂	Kowhai	96.7	99.6	93.4	99.8	96.3	100	96.9	100	96.7	100	95.3	100	96.1	100	95.9	100	96.3	100	96.7	100	97.1	100	98.1	100	96.6	100		
SO ₂	Penrose	95.3	99.3	92.4	99.4	96.8	100	96.1	100	96.9	100	98.4	100	98.3	100	98.3	100	97.8	100	98.1	100	97.2	100	95.1	100	97.0	100		
SO ₂	Burnside	96.0	99.5	69.2	77.5	94.8	100	95.3	100	94.9	100	95.3	100	95.7	100	95.5	100	91.5	100	94.3	100	96.3	100	92.4	100	93.0	98.0		
PM ₁₀	Burnside	98.9	100	65.6	77.5	98.3	100	95.0	100	97.6	100	98.4	100	88.5	100	99.1	100	96.2	100	98.8	100	98.7	100	98.8	100	94.5	98.1		
PM ₁₀	Kowhai	98.5	100	94.8	99.8	98.2	100	98.9	100	98.4	100	98.6	100	97.8	100	98.9	100	99.1	100	99.0	100	98.5	100	97.7	100	98.5	100		
PM ₁₀	Penrose	98.7	99.3	95.7	99.4	98.8	100	98.4	100	87.8	100	99.2	100	95.7	100	99.2	100	99.3	100	99.1	100	99.2	100	98.7	100	99.7	100		
VOC	Penrose	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100		
VOC	Burnside	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100		
VOC	St Albans	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100		
TSP	St Albans	100	100	75	75	100	100	100	100	100	100	100	100	75	75	100	100	100	100	100	100	100	100	100	100	95.8	95.8		
TSP	Penrose	100	100	75	75	100	100	100	100	100	100	100	100	50	50	100	100	100	100	100	100	100	100	100	100	93.8	93.8		
TSP	Kowhai	75	75	25	25	100	100	100	100	100	100	100	100	100	100	80	80	75	75	100	100	100	100	100	100	87.9	87.9		
Lead	St Albans	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	80	80	75	75	100	100	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	85.0	85.0		
Lead	Penrose	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	75	75	50	50	100	100	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	75.0	75.0		
Lead	Kowhai	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	100	100	100	100	80	80	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A	93.3	93.3		

Notes: V = valid data; C = percent total captured data (includes calibration and invalid data).

7.2 Carbon monoxide (CO) 2005

CO was monitored at Greers Road, Burnside. One-hour and eight-hour averages have been calculated from 10-minute averages recorded by the instruments.

The maximum results and their dates are described below.

Site	1-hour max (mg/m ³)	99.9 percentile 1-hour (mg/m ³)	8-hour max (mg/m ³)	99.9 percentile 8-hour (mg/m ³)
Greers Road, Burnside	10.5 (29 June – 01:00)	8.3	8.1 (29 June – 03:20)	5.4

Results are given in Figures 6 to 9.

Concentrations of CO at Greers Road Burnside were below the ambient air quality 1-hour guideline (30 mg/m³) and 8-hour national environmental standard (10 mg/m³) during 2005.

7.3 Nitrogen oxides (NO₂ and NO) 2005

Oxides of nitrogen were monitored at Gavin Street, Penrose; Kowhai Intermediate, Kingsland; and Greers Road, Burnside. One hour and 24-hour averages have been calculated from 10-minute averages recorded by the instruments.

The maximums for NO₂ and their dates for each site are described below.

Site	1-hour max (µg/m ³)	99.9 percentile 1-hour (µg/m ³)	24-hour max (µg/m ³)	99.5 percentile 24-hour (µg/m ³)
Gavin Street, Penrose	236.0 (19 Dec 12:00)	88.7	52.5 (19 Dec)	50.5
Kowhai Intermediate School, Kingsland	103.3 (21 Jun 18:00)	72.7	47.2 (5 Aug)	45.5
Greers Road, Burnside	83.3 (28 Jun 11:00)	73.2	49.9 (28 Jun)	38.0

Nitrogen dioxide results are in Figures 10 to 15 (Kowhai), 16 to 21 (Gavin Street), and 22 to 27 (Burnside).

There was one exceedance of the NO₂ ambient air quality 1-hour standard (200 µg/m³) at Gavin Street, Penrose on 19 December 2005. The NES allow for nine 1-hour NO₂ exceedances in a 12-month period. There were no exceedances of the 24-hour guideline (100 µg/m³) during 2005 at any site in Auckland or Christchurch.

7.4 Sulphur dioxide (SO₂) 2005

Sulphur dioxide was monitored at Gavin Street, Penrose and Greers Road, Burnside. One-hour and 24-hour averages have been calculated from 10-minute averages recorded by the instruments.

The maximums for SO₂ and their dates for each site are described below.

Site	1-hour max (µg/m ³)	99.9 percentile 1-hour (µg/m ³)	24-hour max (µg/m ³)	99.5 percentile 8-hour (µg/m ³)
Gavin Street, Penrose	54.2 (21 Oct 10:00)	42.0	20.7 (21 Oct)	17.7
Greers Road, Burnside	42.2 (26 May 12:00)	32.9	14.0 (28 Jun)	12.7

Results for Gavin Street, Penrose, are shown in Figures 28 and 29 and Greers Road, Burnside, are shown in Figures 30 and 31. There were no exceedances of the SO₂ ambient air quality 1-hour standard (350 µg/m³) or the 24-hour guideline (125 µg/m³) during 2005 at any site.

7.5 Volatile organic compounds (VOC) January – December 2005

Monitoring of VOCs was conducted at four sites: Kowhai Intermediate School, Kingsland; Gavin Street, Penrose; Greers Road, Burnside; and Coles Place, Christchurch. VOC monitoring utilises passive sampling badges exposed over a three-month period. A set of results for each 2005 quarter are shown in Tables 4 to 7. See monitoring method in section 4.

The benzene guideline is 10 µg/m³ as an annual average with an average value of 3.6 µg/m³ to be achieved by 2010. The 2005 six-month and 12-month averages are described below.

Site	Six-month average (January – June 2005) benzene (µg/m ³)	2005 annual average benzene (µg/m ³)
Coles Place, St Albans	3.1	2.1
Greers Road, Burnside	2.4	1.9
Gavin Street, Penrose	2.2	1.9
Kowhai Intermediate School, Kingsland	2.5	2.2

Table 4: VOC results (January – March 2005)

January February March 2005 Analyte	Limit of detection ($\mu\text{g}/\text{m}^3$)	Results ($\mu\text{g}/\text{m}^3$)			
		Coles Place	Burnside	Gavin Street	Kowhai
Target VOCs					
ethanol	ND				
isopropyl alcohol	ND				
acetone	ND				
pentane	ND				
dichloromethane	ND				
butan-2-one	ND				
hexane	0.5	0.6	1.2	1.1	1.2
ethyl acetate	ND				
trichloromethane	ND				
1,1,1-trichloroethane	ND				
n-butanol	ND				
benzene	0.3	1.0	0.8	1.6	1.8
2-methylhexane	ND				
2,3-dimethylpentane	ND				
3-methylhexane	ND				
heptane	0.5		0.6		
trichloroethene	0.5				0.6
propyl acetate	0.6			0.8	
methylcyclohexane	ND				
4-methylpentan-2-one	ND				
toluene	0.3	3.8	3.7	7.2	7.5
octane	ND				
tetrachloroethene	ND				
butyl acetate	ND				
ethylbenzene	0.3	0.5	0.4	1.1	0.9
m+p-xylene	0.3	1.4	1.2	2.8	2.7
styrene	ND				
o-xylene	0.3	0.6	0.5	1.1	1.1
nonane	ND				
alpha pinene	ND				
propylbenzene	ND				
1,3,5-trimethylbenzene	ND				
beta pinene	ND				
decane	0.6			0.9	
1,2,4-trimethylbenzene	0.6	0.7		1.2	1.1
limonene	ND				
undecane	0.8			1.5	
dodecane	ND				

Table 5 VOC results (April – June 2005)

April May June 2005 Analyte	Limit of detection ($\mu\text{g}/\text{m}^3$)	Results ($\mu\text{g}/\text{m}^3$)			
		Coles Place	Burnside	Gavin Street	Kowhai
Target VOCs					
ethanol	ND				
isopropyl alcohol	ND				
acetone	ND				
pentane	1.9	2.5	1.9	2.3	2.2
dichloromethane	ND				
butan-2-one	ND				
hexane	0.4	1.8	1.7	1.5	1.6
ethyl acetate	ND				
trichloromethane	ND				
1,1,1-trichloroethane	ND				
n-butanol	ND				
benzene	0.2	5.1	4.0	2.8	3.2
2-methylhexane	0.5	0.8	0.7	0.6	0.6
2,3-dimethylpentane	ND				
3-methylhexane	0.5	1.0	0.8	0.7	0.6
heptane	0.4	0.8	0.9	0.7	0.5
trichloroethene	0.4				0.8
propyl acetate	0.5		0.5	0.9	
methylcyclohexane	ND				
4-methylpentan-2-one	ND				
toluene	0.3	13.3	10.2	12.5	11.1
octane	ND				
tetrachloroethene	ND				
butyl acetate	ND				
ethylbenzene	0.2	1.9	1.4	1.6	1.2
m+p-xylene	0.2	6.7	4.9	5.0	4.9
styrene	ND				
o-xylene	0.3	2.6	2.0	2.0	1.8
nonane	0.5	0.5	0.4	0.6	0.6
alpha pinene	0.5	0.6	0.9	0.6	
propylbenzene	ND				
1,3,5-trimethylbenzene	0.6	0.8	0.6	0.7	0.7
beta pinene	0.6	0.7	1.1	0.8	
decane	0.6	0.6	0.5	0.9	0.7
1,2,4-trimethylbenzene	0.6	2.3	1.8	2.0	2.0
limonene	ND				
undecane	0.8			1.0	
dodecane	ND				

Table 6: VOC results (July – September 2005)

July August September 2005 Analyte	Limit of detection ($\mu\text{g}/\text{m}^3$)	Results ($\mu\text{g}/\text{m}^3$)			
		Coles Place	Burnside	Gavin Street	Kowhai
Target VOCs					
ethanol	ND				
isopropyl alcohol	ND				
acetone	ND				
pentane	2.2			2.1	
dichloromethane	ND				
butan-2-one	ND				
hexane	0.5	1.4	1.0	1.4	1.4
ethyl acetate	0.5			0.5	
trichloromethane	ND				
1,1,1-trichloroethane	ND				
n-butanol	ND				
benzene	0.3	1.6	2.1	2.3	2.7
2-methylhexane	0.5			0.5	0.5
2,3-dimethylpentane	ND				
3-methylhexane	0.5	0.7	0.6	0.6	0.6
heptane	0.5	0.8		0.6	0.5
trichloroethene	0.5				0.7
propyl acetate	0.5			0.5	
methylcyclohexane	ND				
4-methylpentan-2-one	ND				
toluene	0.3	5.6	7.3	9.2	9.2
octane	ND				
tetrachloroethene	ND				
butyl acetate	0.6			0.6	
ethylbenzene	0.3	0.7	0.9	1.4	1.1
m+p-xylene	0.3	2.6	3.5	4.6	4.5
styrene	ND				
o-xylene	0.3	1.0	1.4	1.7	1.6
nonane	ND				
alpha pinene	ND				
propylbenzene	ND				
1,3,5-trimethylbenzene	0.6			0.6	0.6
beta pinene	ND				
decane	0.6			0.7	0.5
1,2,4-trimethylbenzene	0.6	0.9	1.2	1.6	1.7
limonene	ND				
undecane	0.8			1.0	
dodecane	ND				

Table 7: VOC results (October – December 2005)

October November December 2005 Analyte	Limit of detection ($\mu\text{g}/\text{m}^3$)	Results ($\mu\text{g}/\text{m}^3$)			
		Coles Place	Burnside	Gavin Street	Kowhai
Target VOCs					
ethanol	ND				
isopropyl alcohol	ND				
acetone	ND				
pentane	ND				
dichloromethane	ND				
butan-2-one	ND				
hexane	0.5	0.5	0.9	0.8	0.8
ethyl acetate	0.4			0.4	
trichloromethane	ND				
1,1,1-trichloroethane	ND				
n-butanol	ND				
benzene	0.3	0.6	0.5	0.9	1.0
2-methylhexane	ND				
2,3-dimethylpentane	ND				
3-methylhexane	0.5				
heptane	0.5				
trichloroethene	ND				
propyl acetate	0.5			0.5	
methylcyclohexane	ND				
4-methylpentan-2-one	ND				
toluene	0.3	3.2	2.8	5.7	6.5
octane	ND				
tetrachloroethene	ND				
butyl acetate	ND				
ethylbenzene	0.3	0.4	0.3	0.7	0.7
m+p-xylene	0.3	1.6	1.2	2.4	2.6
styrene	ND				
o-xylene	0.3	0.6	0.5	0.9	0.9
nonane	ND				
alpha pinene	ND				
propylbenzene	ND				
1,3,5-trimethylbenzene	ND				
beta pinene	ND				
decane	0.6			0.6	
1,2,4-trimethylbenzene	0.6	0.6	0.0	1.0	0.9
limonene	ND				
undecane	0.8			0.8	
dodecane	ND				

7.6 PM₁₀ 2005

PM₁₀ is monitored at Greers Road, Burnside, using a thermo FH62-C14 beta gauge. PM₁₀ results measured by Auckland Regional Council at Gavin Street, Penrose and Kowhai Intermediate, Kingsland sites, also using thermo FH62-C14 beta gauges, are also included in this report. Twenty four-hour averages have been calculated from 10-minute averages recorded by the instruments. All PM₁₀ data is reported at standard temperature and pressure (0°C and 101.3 kPa).

The maximums for PM₁₀ and their dates for each site are described below.

Site	24-hour maximum (µg/m ³)	99.5 percentile 24-hour (µg/m ³)
Gavin Street, Penrose	44.7 (10 June)	33.4
Kowhai Intermediate, Kingsland	49.0 (11 June)	35.7
Greers Road, Burnside	98.4 (28 June)	77.0

There were no exceedances of the ambient air quality standard (50 µg/m³) at Gavin Street, Penrose or Kowhai Intermediate, Kingsland during the 12-month period.

At Greers Road, Burnside there were 18 exceedances of the 24-hour standard. Each exceedance and the date is listed in Table 8 below. Charts describing 24-hour averaged data for 2005 for each site are shown in Figures 32 to 33 (Kowhai), 34 to 35 (Gavin Street), and 36 and 37 (Burnside). As there were exceedances at the Christchurch site, more data analysis was carried out in Figures 38 to 41. All exceedances occurred over the winter period, a time when wood burning is widely used to heat homes. As reported by Environment Canterbury, cold winter conditions strongly influence air pollution in the region. A total of 14 exceedances occurred during the period June to July, with only four exceedances occurring during the months of April, May and August 2005.

Table 8: Greers Road, Burnside PM₁₀ exceedances of the daily NES 2005

Date	Burnside PM ₁₀ (µg/m ³)
28/04/05	60.5
09/05/05	51.8
10/05/05	50.0
07/06/05	77.0
08/06/05	61.8
09/06/05	56.4
12/06/05	59.8
18/06/05	53.4
19/06/05	50.7
27/06/05	61.1
28/06/05	98.4
29/06/05	91.6
01/07/05	61.7
10/07/05	60.8
11/07/05	64.4
22/07/05	57.0
27/07/05	56.9
14/08/05	59.1

Note : National environmental standard for PM10 = 50 µg/m³.

7.7 Total suspended particulates (TSP) 2005

TSP is measured as a seven-day average from Gavin Street, Penrose; Kowhai Intermediate, Kingsland; and Coles Place, St Albans. Maximum results are shown in the table below.

Site	Maximum seven-day average (µg/m ³)
Gavin Street, Penrose	38 (15 June)
Kowhai Intermediate, Kingsland	40 (19 January)
Coles Place, St Albans	52 (30 June)

There were no exceedances of the Ministry of Health's guideline of 60 µg/m³ at any site. The data from each site is described in Figures 1 and 2 below.

Figure 1: Auckland TSP seven-day average 2005

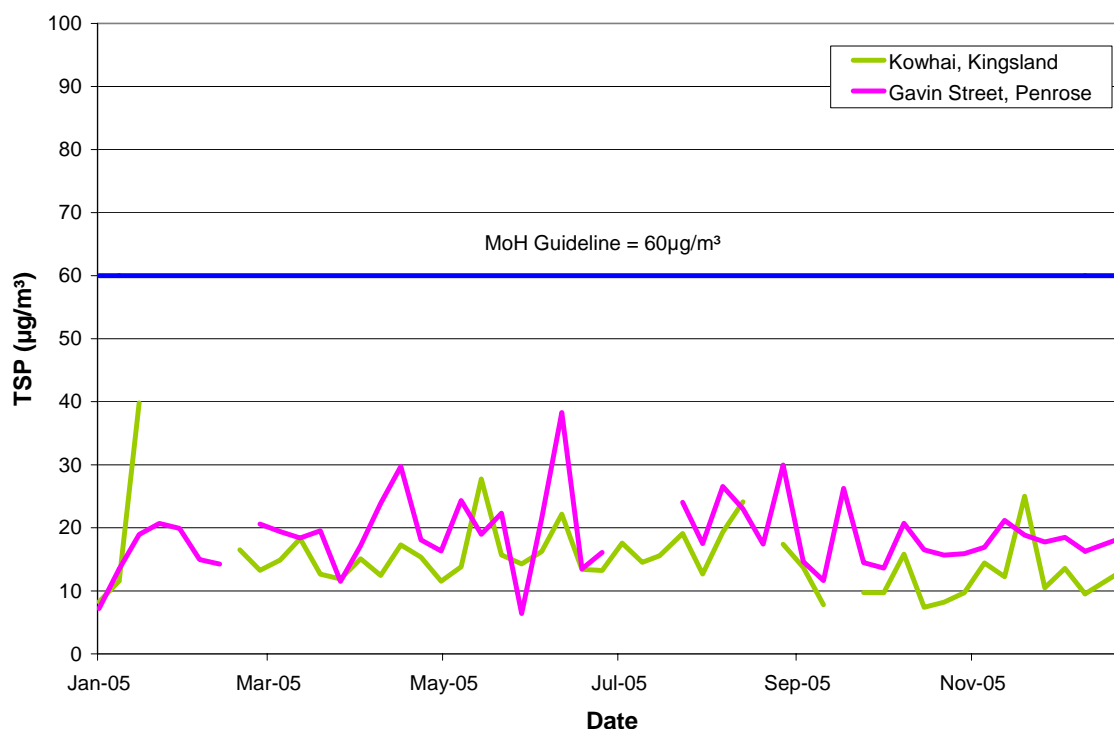


Figure 2: Auckland TSP seven-day average 1994 – 2005

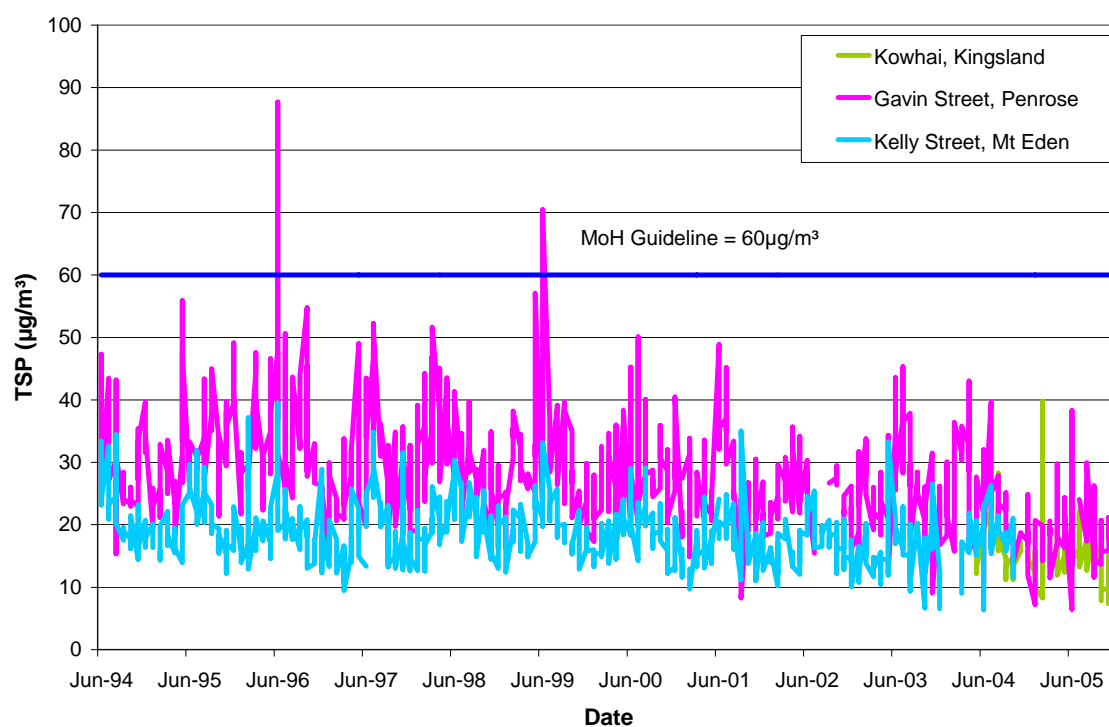


Figure 3: Christchurch TSP seven-day average 2005

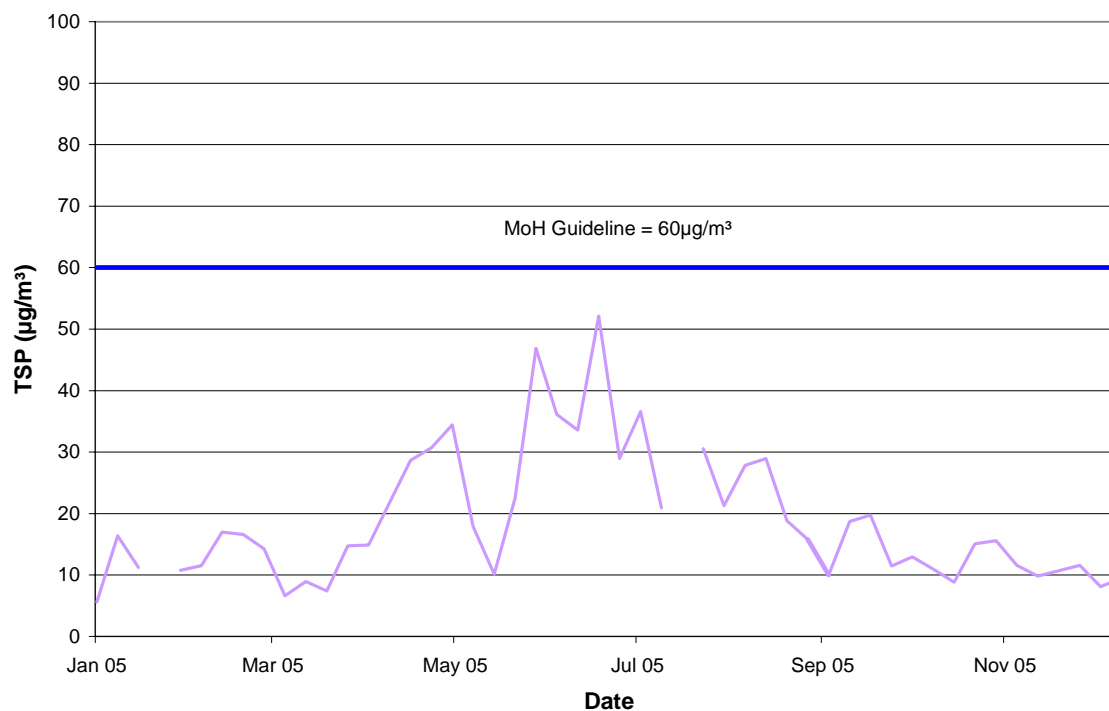
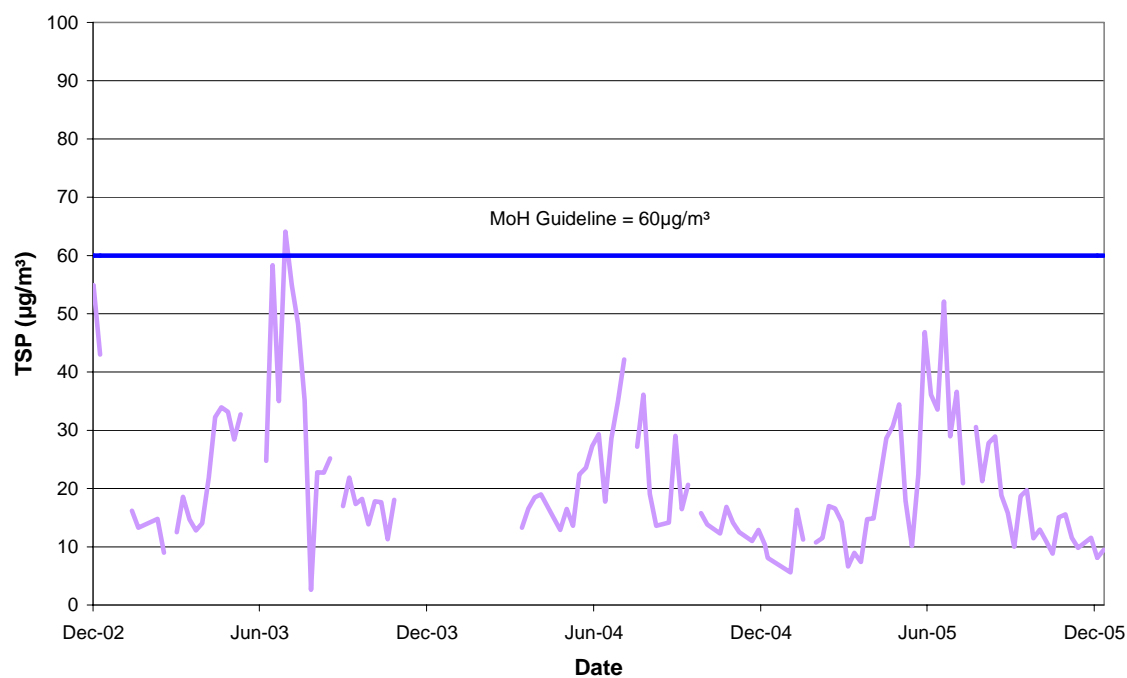


Figure 4: Christchurch TSP seven-day average 2002 – 2005



7.8 Lead (Pb) June – August 2005

Lead is measured from seven-day averaged TSP samples to derive a three-month average. The results are described in the table below. Figure 5 provides moving three-month averaged lead data between January 1996 and September 2000, when lead monitoring was performed on a monthly basis. From this point, lead continued to be monitored over a three-month period (June to August) annually.

Site	June 2005 average ($\mu\text{g}/\text{m}^3$)	July 2005 average ($\mu\text{g}/\text{m}^3$)	August 2005 average ($\mu\text{g}/\text{m}^3$)	Winter 2005 average ($\mu\text{g}/\text{m}^3$)
Kowhai Intermediate, Kingsland	0.022	0.013	0.018	0.017
Gavin Street, Penrose	0.012	0.013	0.016	0.014
Coles Place, St Albans	0.031	0.023	0.018	0.024

No site exceeded the three-month average guideline for lead ($0.2 \mu\text{g}/\text{m}^3$).

Figure 5: MfE lead three-month average results 1996 – 2005

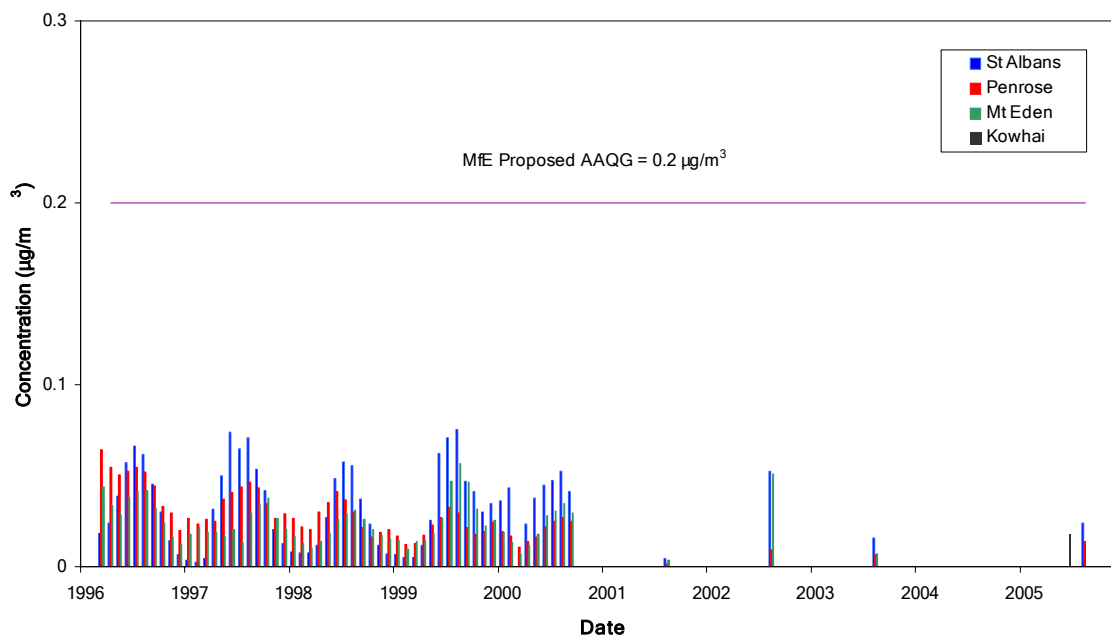


Figure 6: MfE Burnside CO 1-hour fixed average January – December 2005

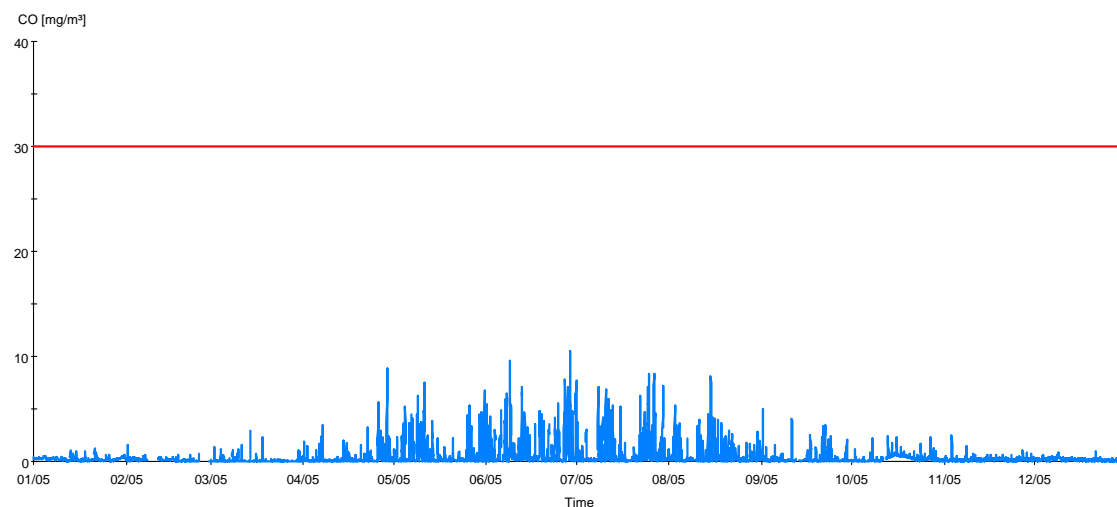


Figure 7: MfE Burnside CO 1-hour fixed average 1 January 2003 – 31 December 2005

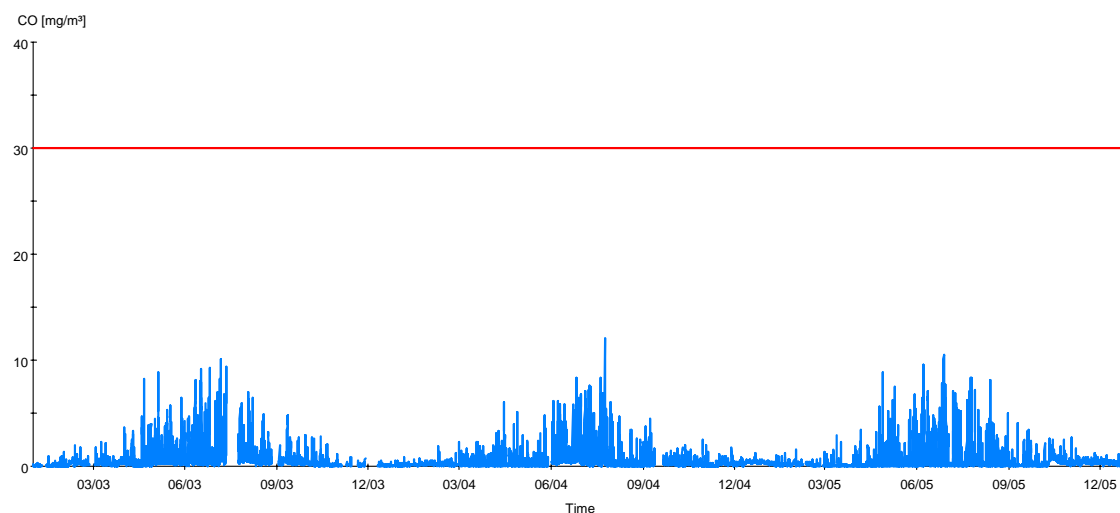


Figure 8: MfE Burnside CO 8-hour rolling average January – December 2005

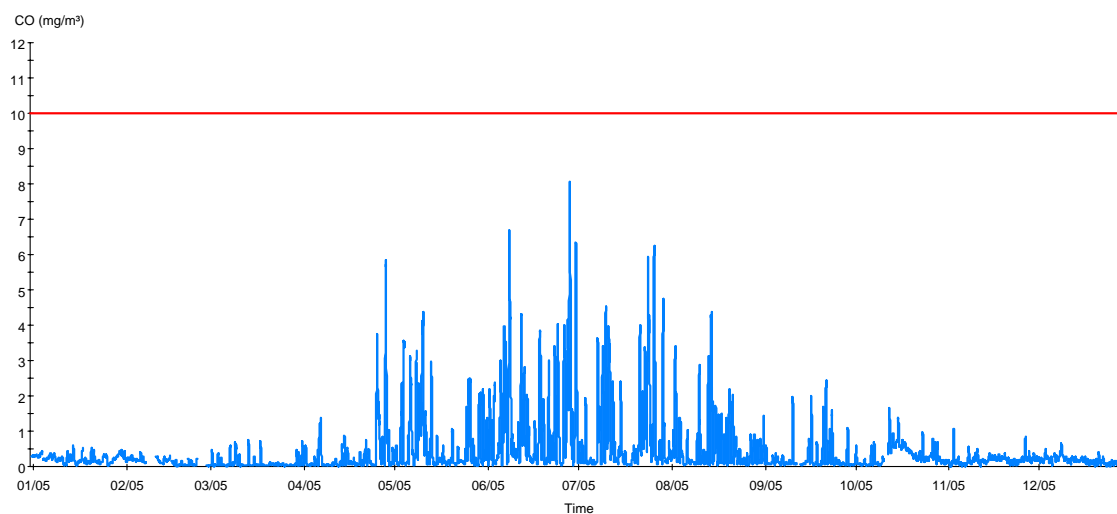


Figure 9: MfE Burnside CO 8-hour rolling average 1 January 2003 – 31 December 2005

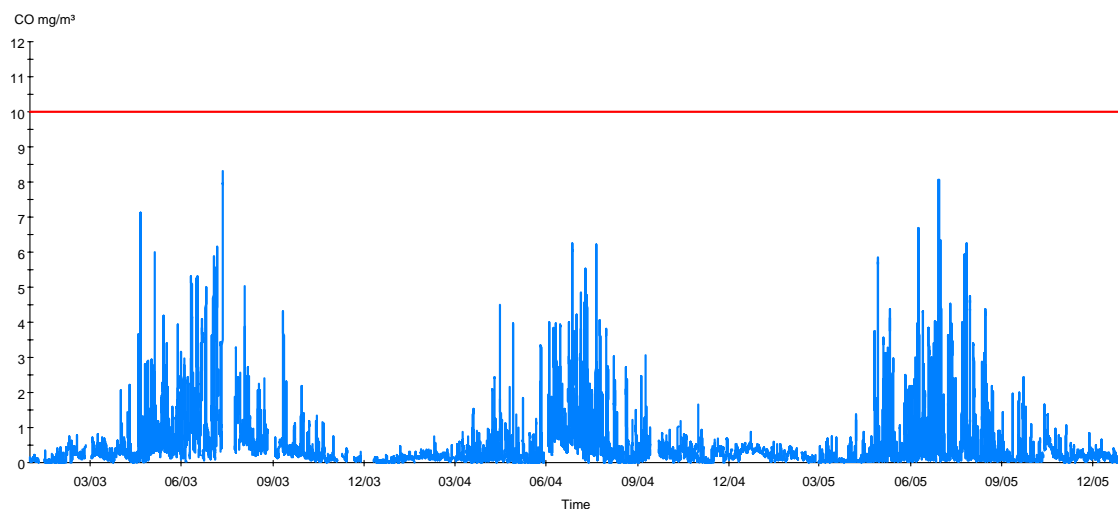


Figure 10: MfE Kowhai NO₂ 1-hour fixed average January – December 2005

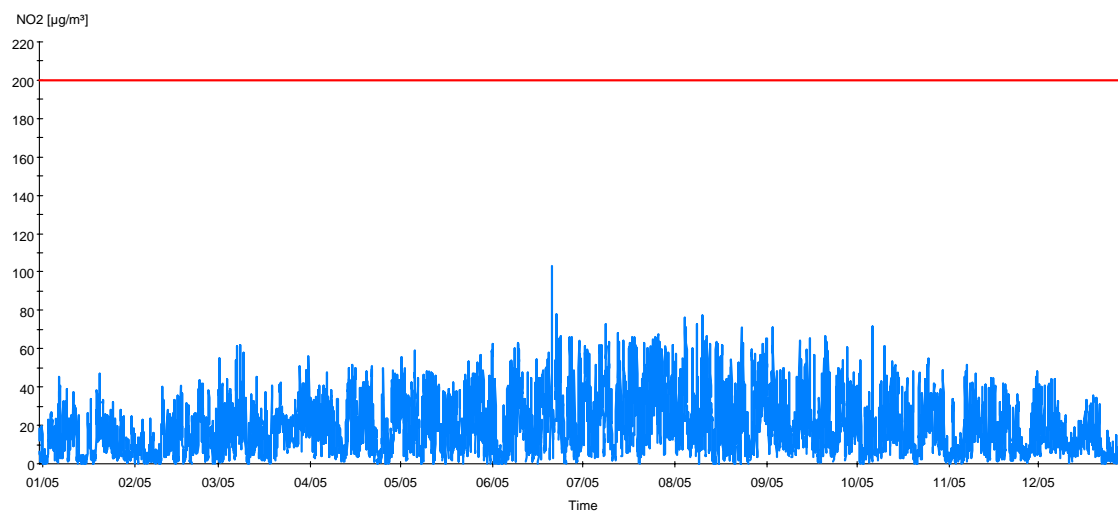


Figure 11: MfE Kowhai NO₂ 1-hour fixed average 1 January 2004 – 31 December 2005

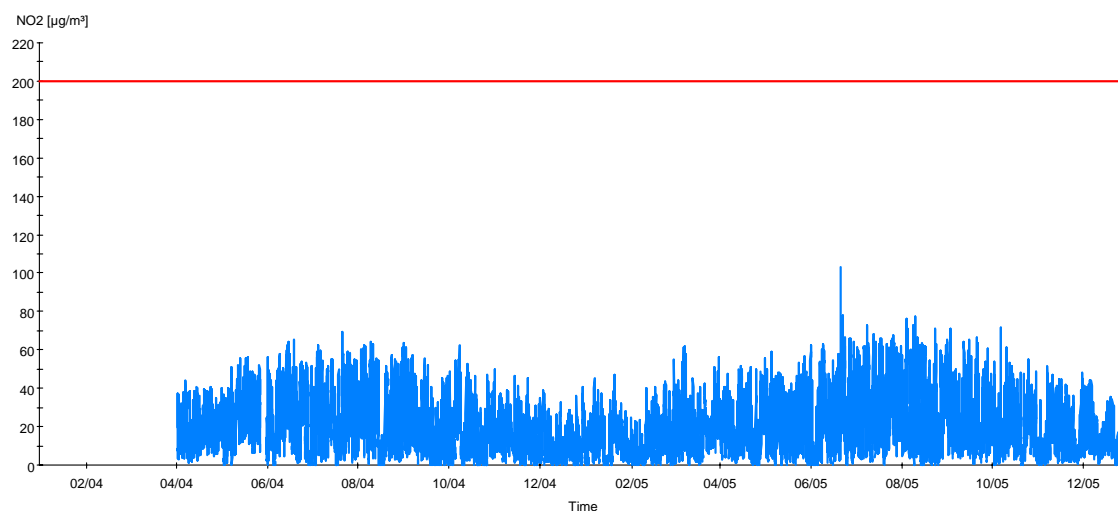


Figure 12: MfE Kowhai NO₂ 24-hour fixed average January – December 2005

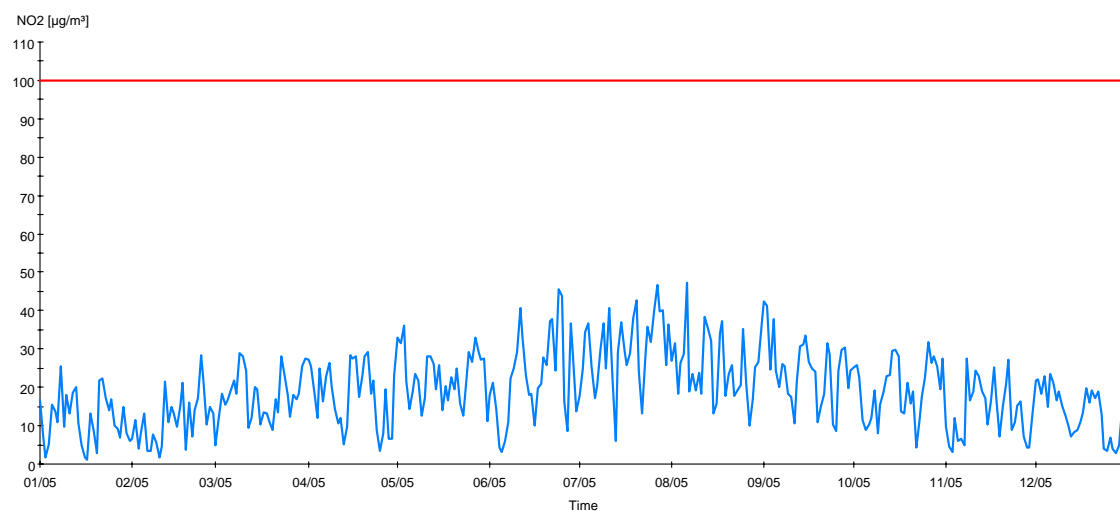


Figure 13: MfE Kowhai NO₂ 24-hour fixed average 1 January 2004 – 31 December 2005

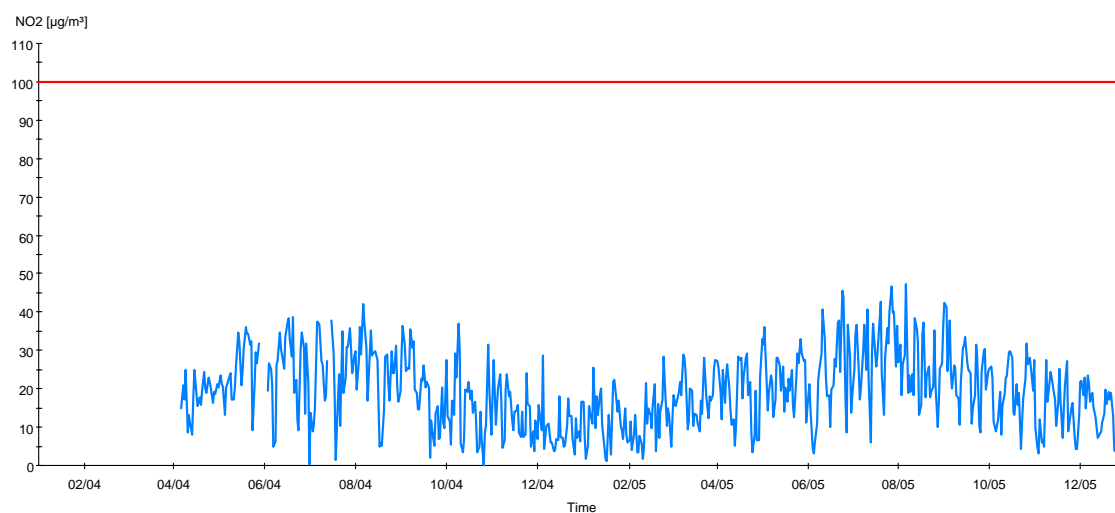


Figure 14: MfE Kowhai NO₂ and NO 1-hour fixed average January – December 2005

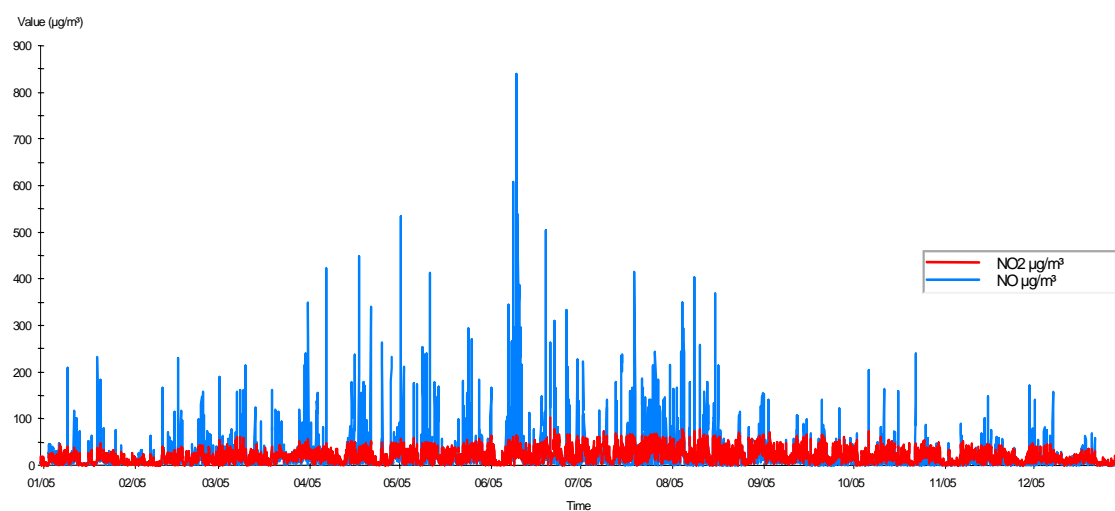


Figure 15: MfE Kowhai NO₂ and NO 24-hour fixed average January – December 2005

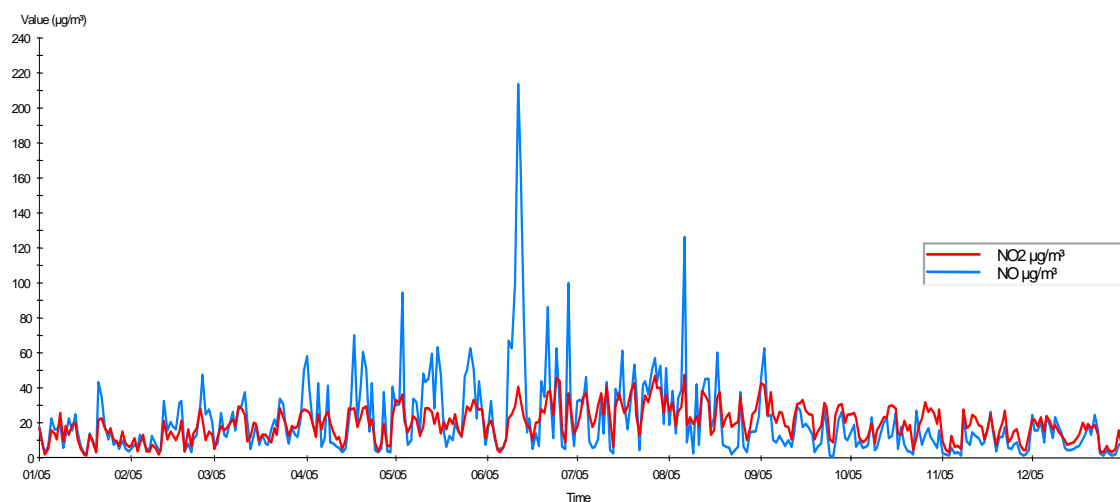


Figure 16: MfE Gavin Street NO₂ 1-hour fixed average January – December 2005

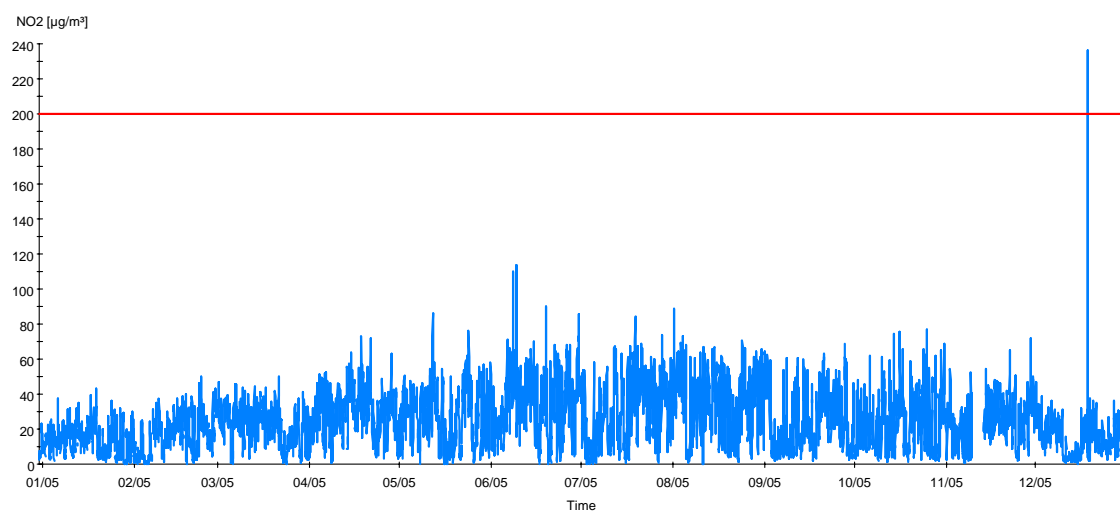


Figure 17: MfE Gavin Street NO₂ 1-hour fixed average 1 January 1997 – 31 December 2005

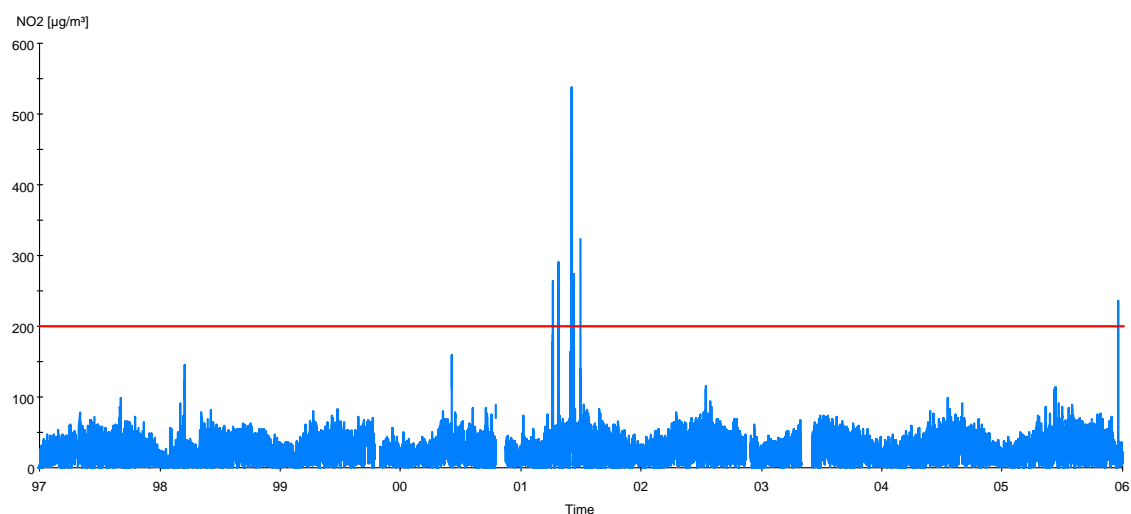


Figure 18: MfE Gavin Street NO₂ 24-hour fixed average January – December 2005

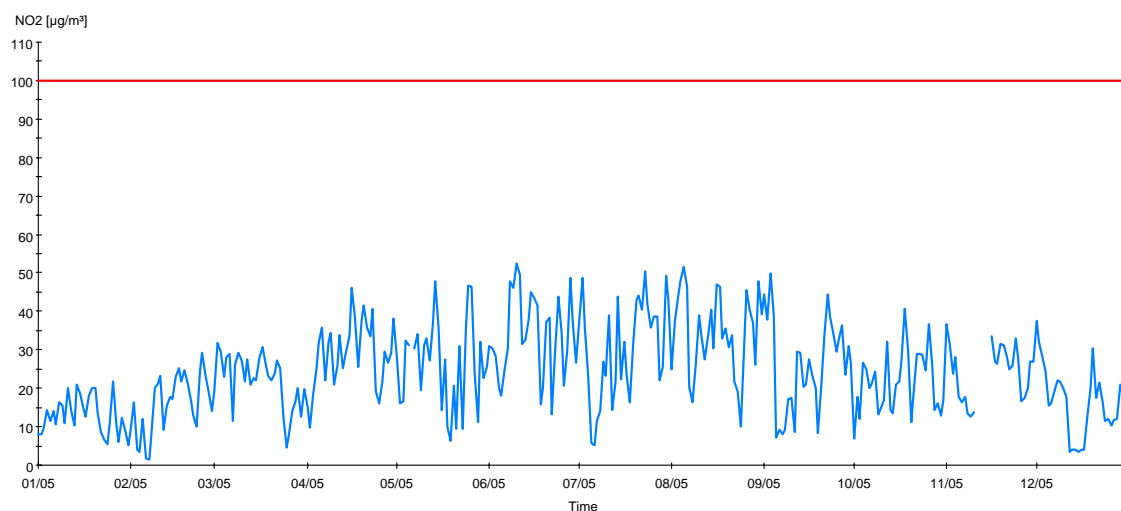


Figure 19: MfE Gavin Street NO₂ 24-hour fixed average 1 January 1997 – 31 December 2005

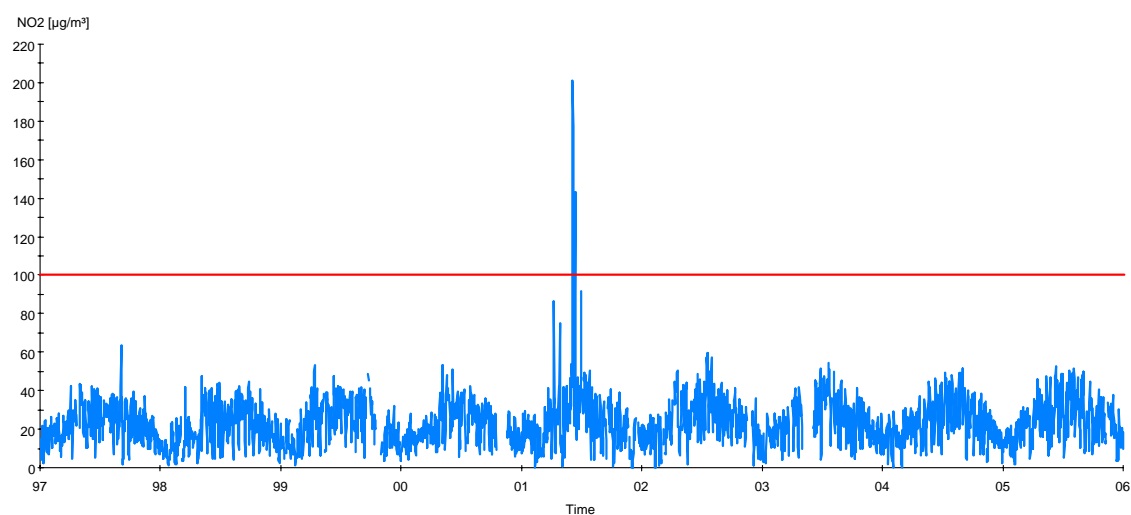


Figure 20: MfE Gavin Street NO₂ and NO 1-hour fixed average January – December 2005

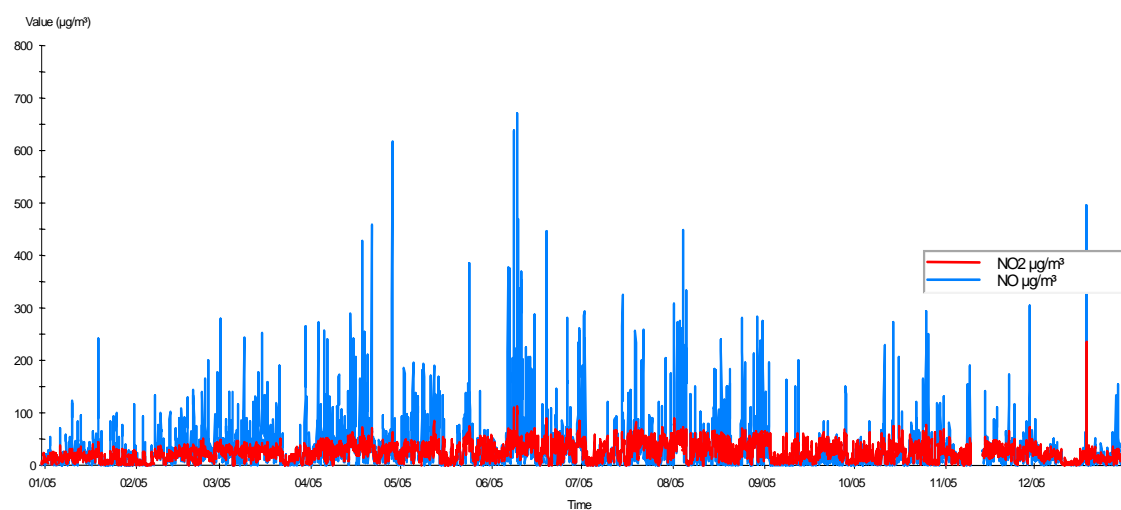


Figure 21: MfE Gavin Street NO₂ and NO 24-hour fixed average January – December 2005

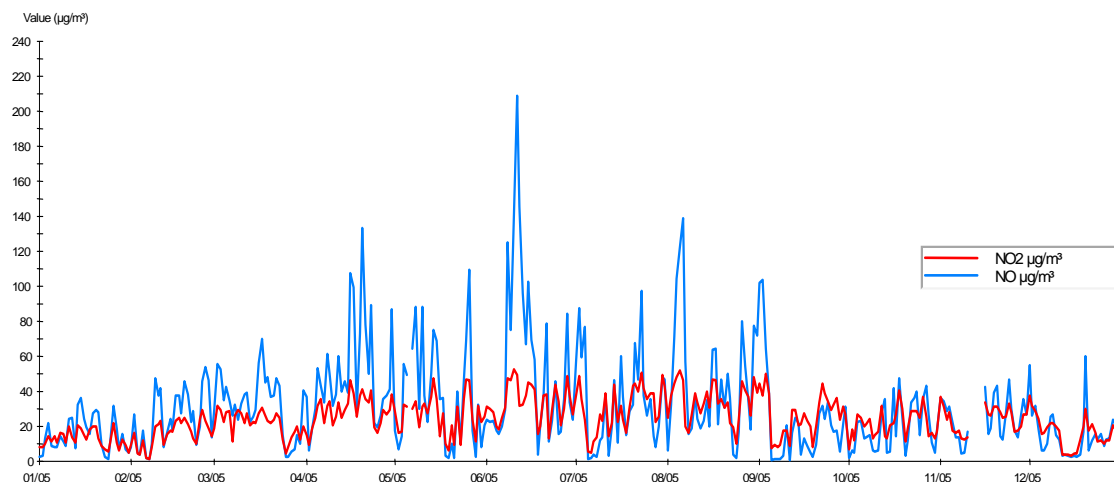


Figure 22: MfE Burnside NO₂ 1-hour fixed average January – December 2005

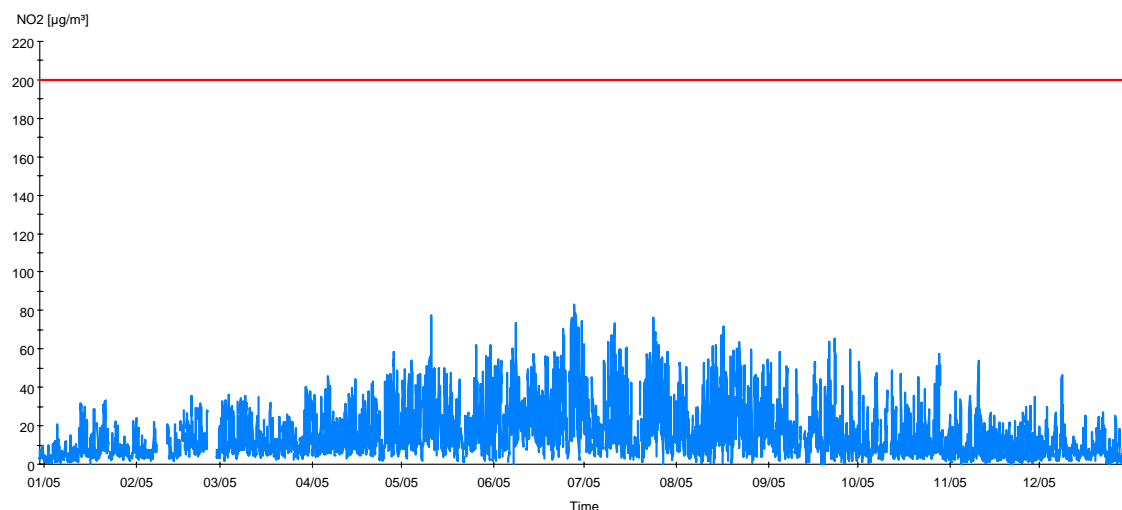


Figure 23: MfE Burnside NO₂ 1-hour fixed average 1 January 2003 – 31 December 2005

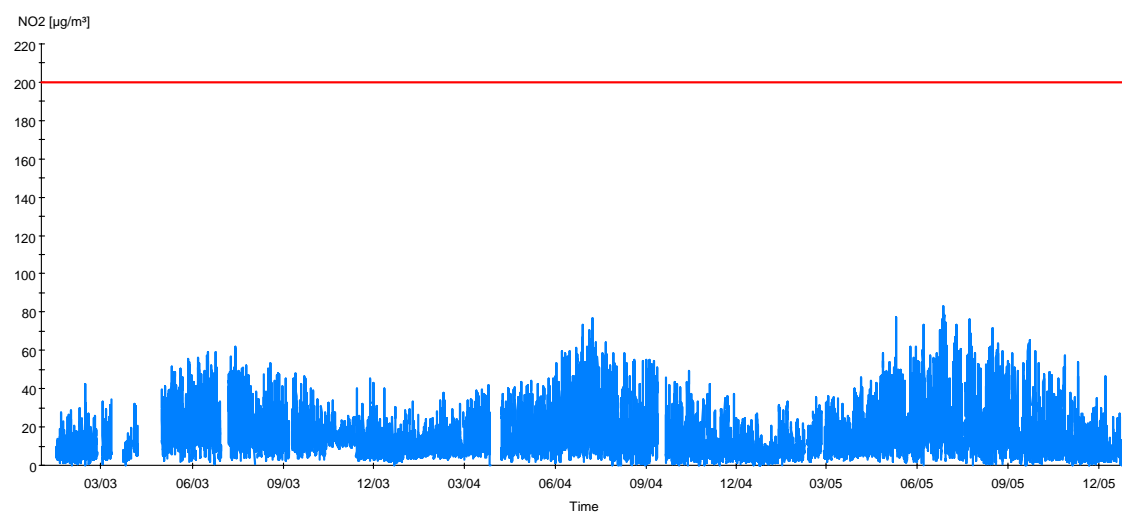


Figure 24: MfE Burnside NO₂ 24-hour fixed average January – December 2005

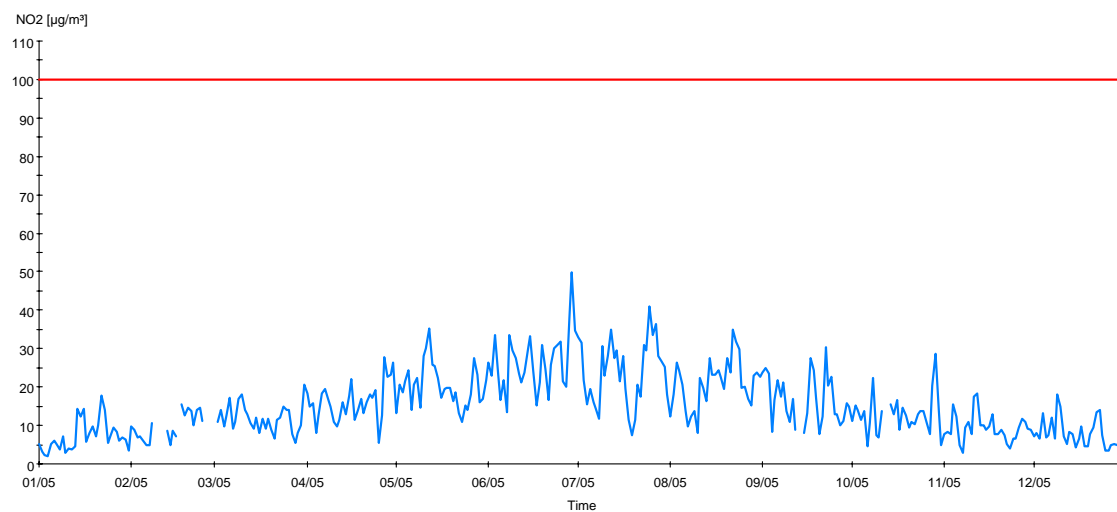


Figure 25: MfE Burnside NO₂ 24-hour fixed average 1 January 2003 – 31 December 2005

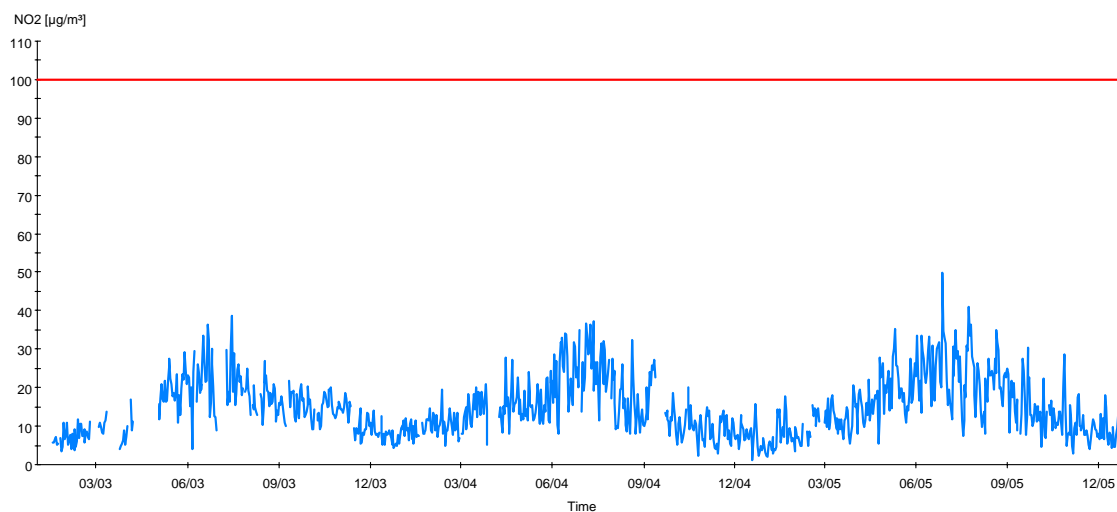


Figure 26: MfE Burnside NO₂ and NO 1-hour fixed average January – December 2005

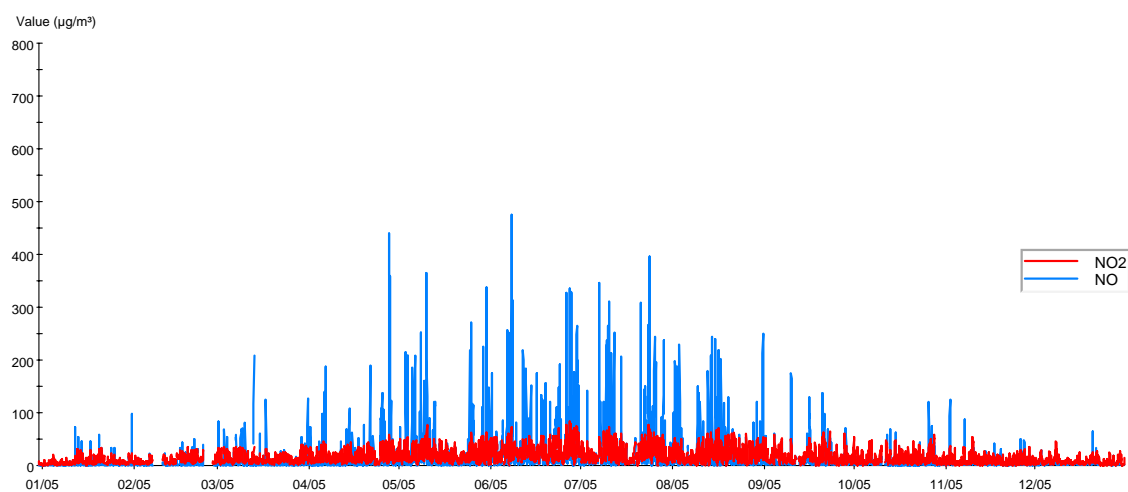


Figure 27: MfE Burnside NO₂ and NO 24-hour fixed average January – December 2005

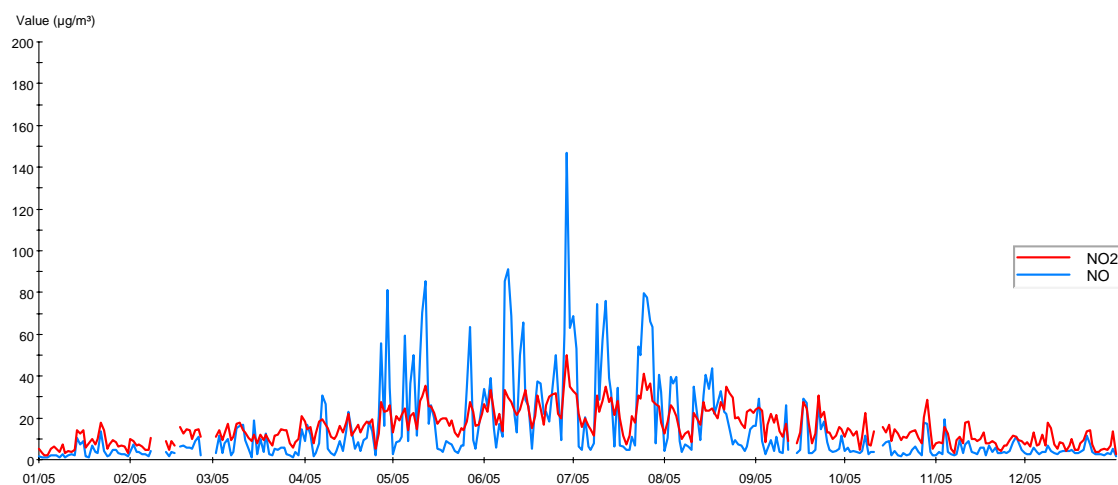


Figure 28: MfE Gavin Street SO₂ 1-hour fixed average January – December 2005

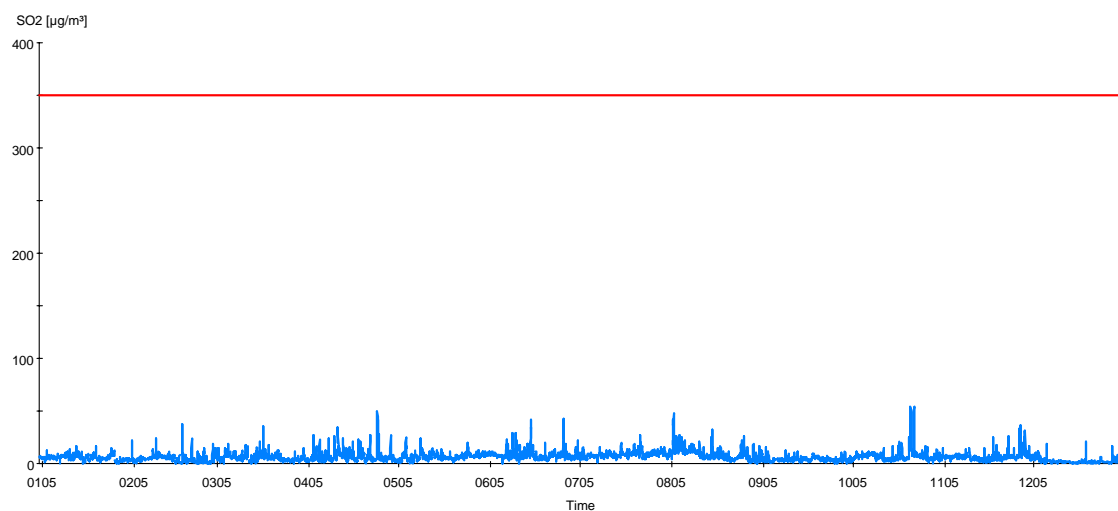


Figure 29: MfE Gavin Street SO₂ 24-hour fixed average January – December 2005

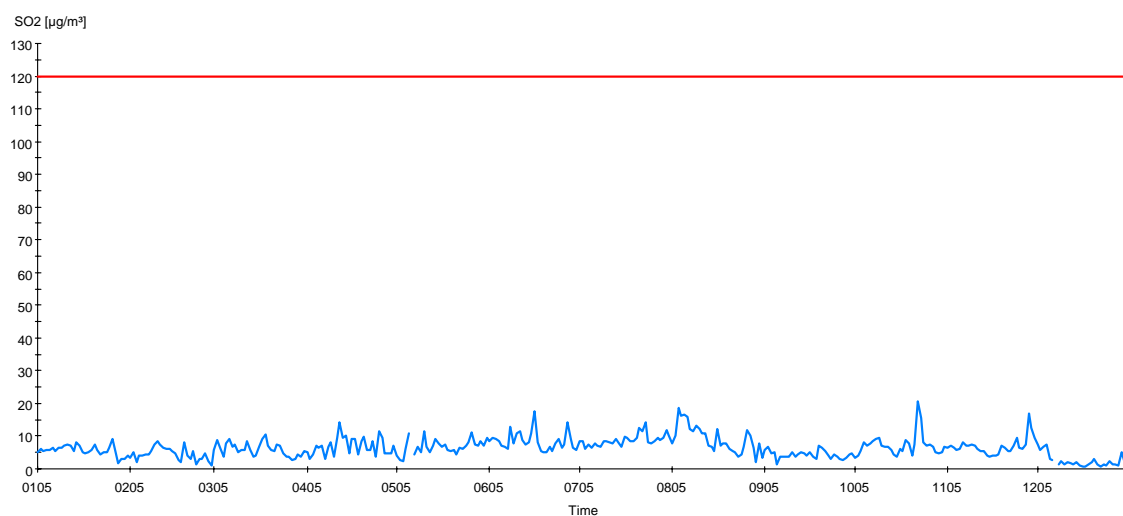


Figure 30: MfE Burnside SO₂ 1-hour fixed average January – December 2005

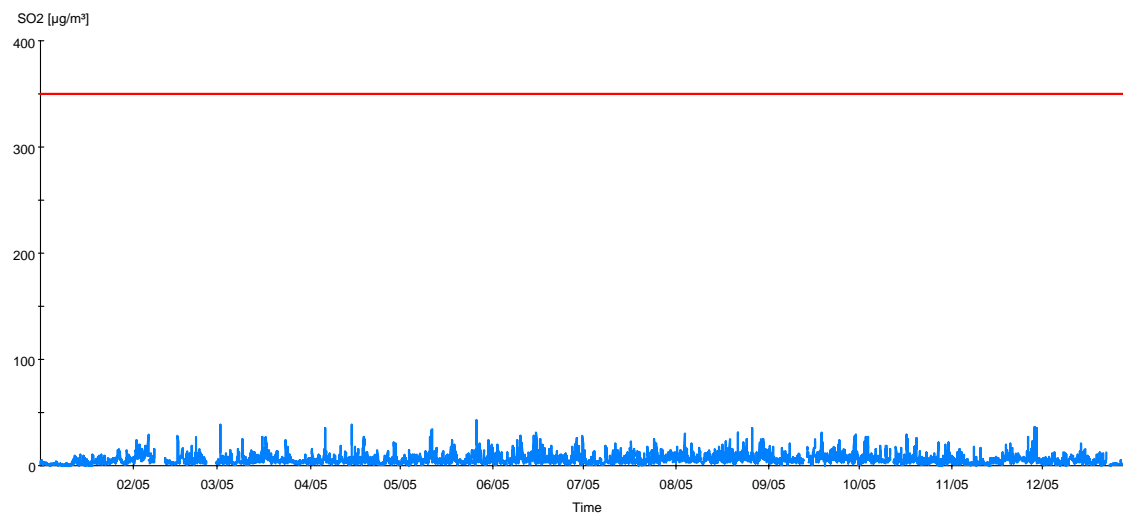


Figure 31: MfE Burnside SO₂ 24-hour fixed average January – December 2005

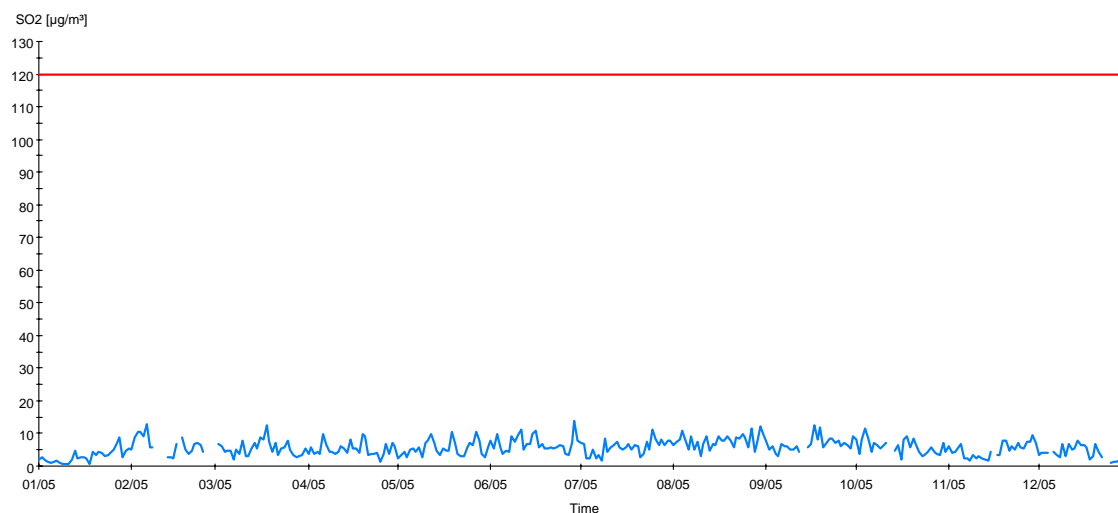


Figure 32: MfE Kowhai PM₁₀ 24-hour fixed average January – December 2005

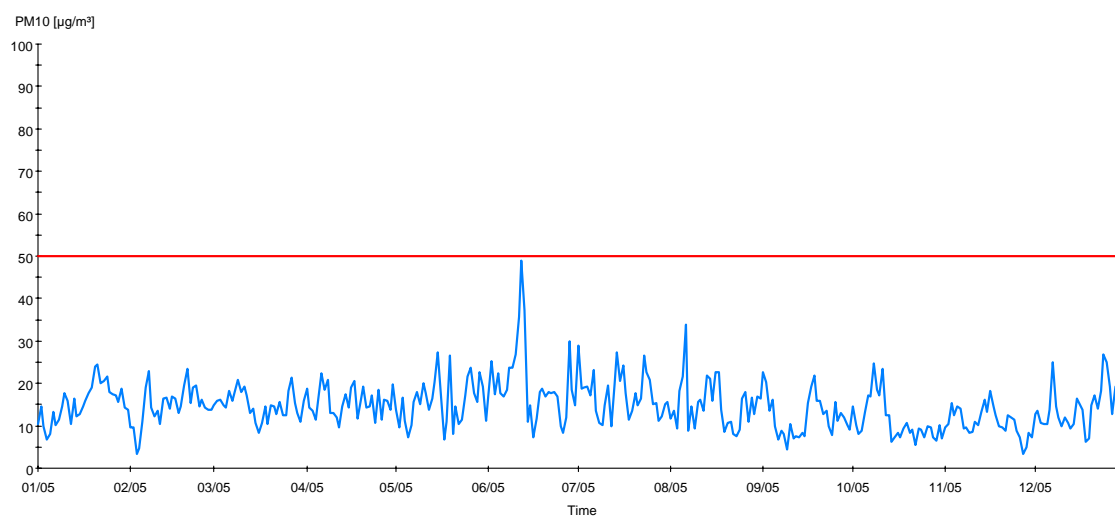


Figure 33: MfE Kowhai PM₁₀ 24-hour fixed average 1 January 2004 – 31 December 2005

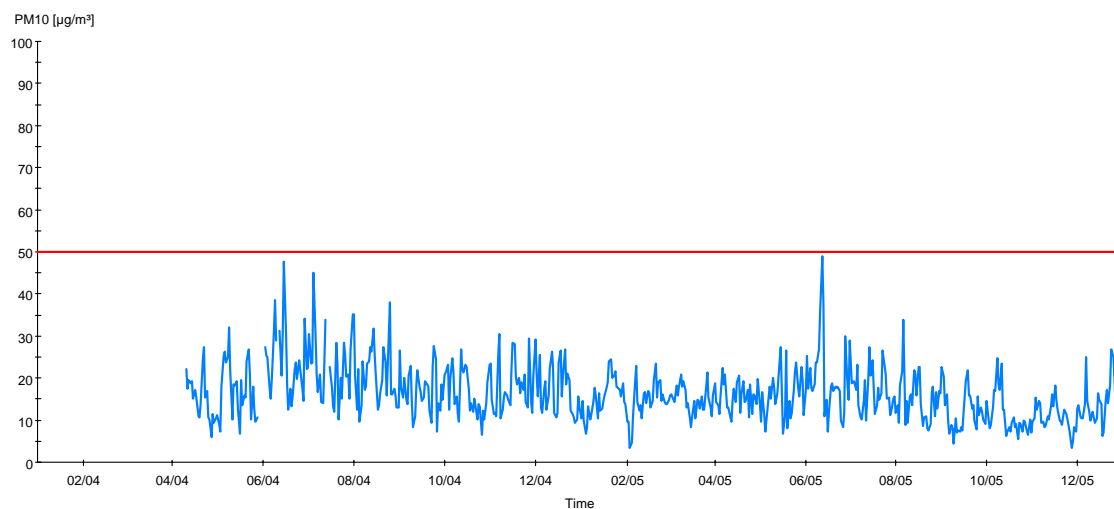


Figure 34: MfE Gavin Street PM₁₀ 24-hour fixed average January – December 2005

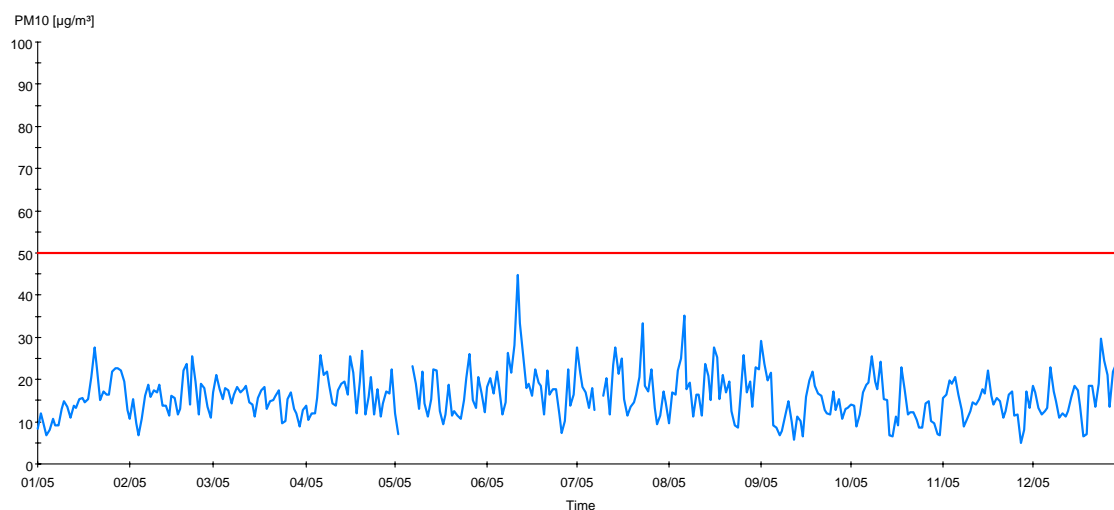


Figure 35: MfE Gavin Street PM₁₀ 24-hour fixed average 1 January 2003 – 31 December 2005

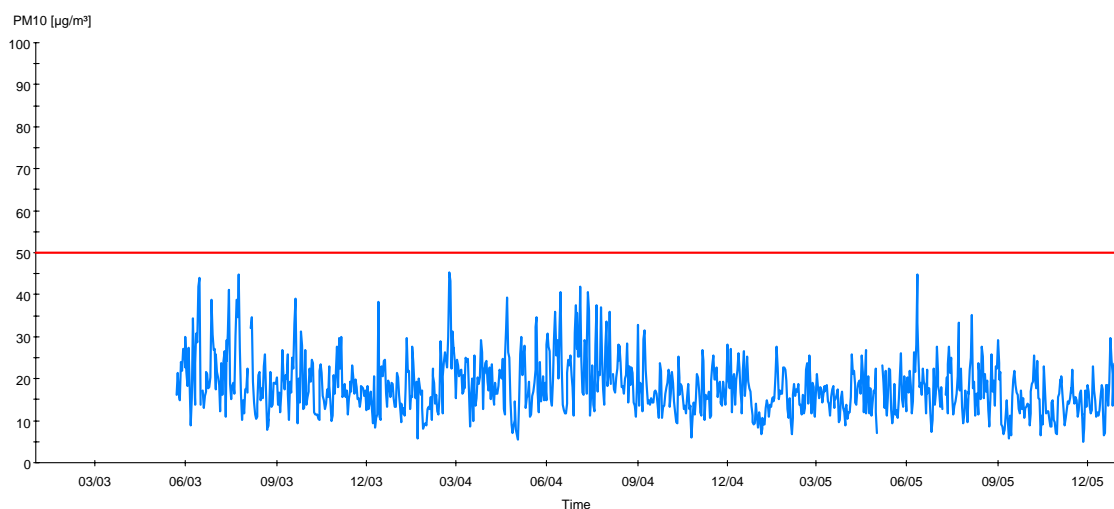


Figure 36: MfE Burnside PM₁₀ 24-hour fixed average January – December 2005

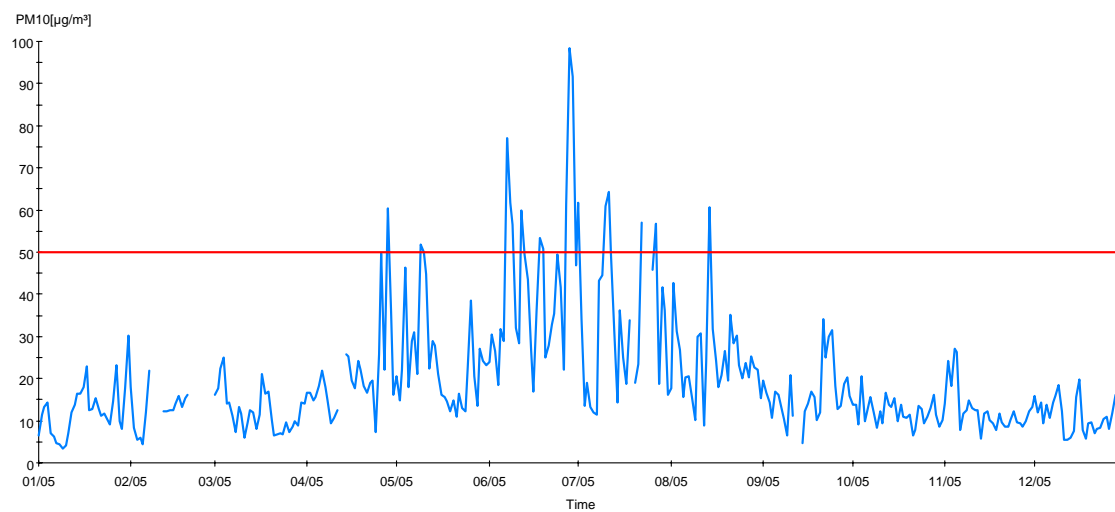
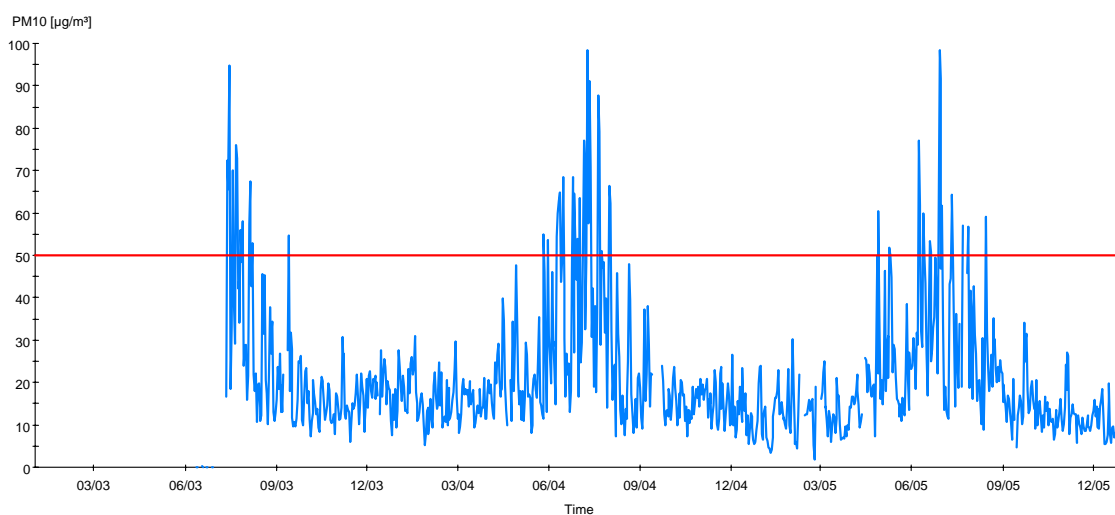


Figure 37: MfE Burnside PM₁₀ 24-hour fixed average 1 January 2003 – 31 December 2005



7.9 Analysis of exceedances

7.9.1 PM₁₀ exceedances at Greers Road, Burnside

Environment Canterbury noted one of the mildest winters on record for Christchurch in 2005. Although light winds and clear sunny days were prevalent during the winter period, evening cloud brought about by a series of anticyclones created warmer temperatures. These favourable conditions may have contributed to a reduction in PM₁₀ exceedances (25 exceedances in 2004 compared with 18 exceedances in 2005).³

A 72-hour period from 7 to 9 June 2005 was chosen as a typical example of winter diurnal trends coinciding with PM₁₀ exceedances to describe the relationship between pollution levels and meteorological conditions. Figure 38 provides an example of the typical diurnal trend in air pollution during this period. Figures 39 to 41 present the meteorological conditions that influence the diurnal trend. It is apparent when comparing wind speed (Figure 39) and PM₁₀ concentrations (Figure 38) that low wind speeds coincide with PM₁₀ peaks and conversely higher wind speeds coincide with low PM₁₀ concentrations. Reduced air mixing caused by low wind speeds results in poor dispersion of pollution while an unstable atmosphere caused by high wind speeds is conducive to pollution dispersion.

Low temperatures, often coinciding with still atmospheric conditions, can cause temperature inversions. This can contribute to higher PM₁₀ concentrations being measured as pollution is trapped at ground level. A comparison of temperatures measured at 1.5 and 10 m (Figure 40) does not indicate the presence or absence of a temperature inversion as the inversion height may be greater than 10m, however, a diurnal trend is apparent. As the temperature drops during the evening, Christchurch residents light their heating appliances causing particle emissions to increase and PM₁₀ levels to peak just before midnight. As the fires die down and the atmosphere becomes more unstable toward morning, concentrations of PM₁₀ drop off.

A pollution rose for the same 72-hour period (Figure 41) that describes the relationship between wind direction and PM₁₀ concentrations shows no obvious pattern. This would suggest that the incidents are not directly related to very localised sources of PM₁₀ but are perhaps more related to prevailing meteorological conditions such as temperature inversions.

Temperature inversions occur when the ground temperature falls below the surrounding air temperature. Air in contact with the ground is cooled to a lower temperature than the air layers above it. As an inversion continues air becomes stagnant and pollution becomes trapped in the mixing layer close to the ground.

³ Environment Canterbury. 2005. Winter Air Report. <http://www.ecan.govt.nz/Our+Environment/Air/Winter+Reports>.

Figure 38: MfE Burnside PM₁₀, typical winter diurnal trend, 10-minute fixed average 7–9 June 2005

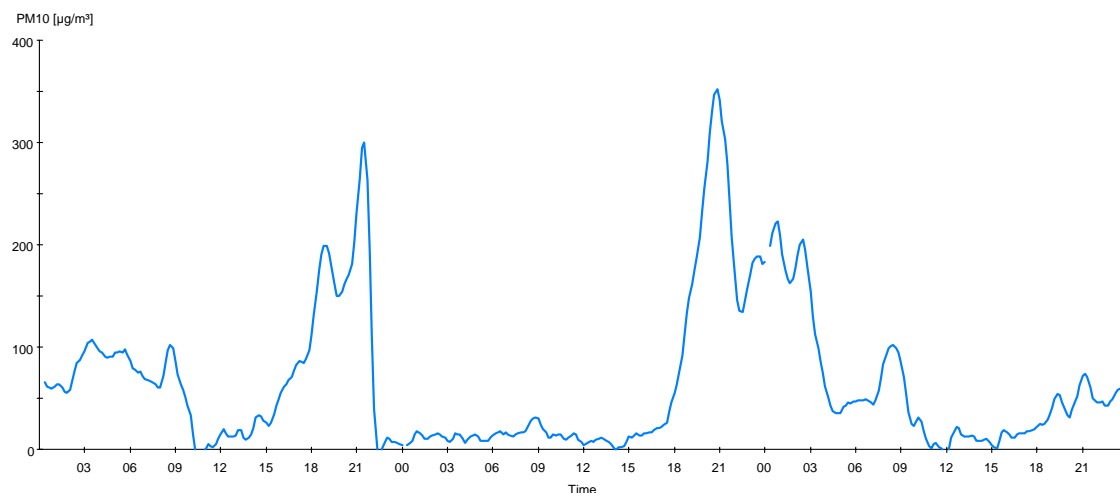


Figure 39: MfE Burnside wind speed, typical winter diurnal trend, 10-minute fixed average 7–9 June 2005

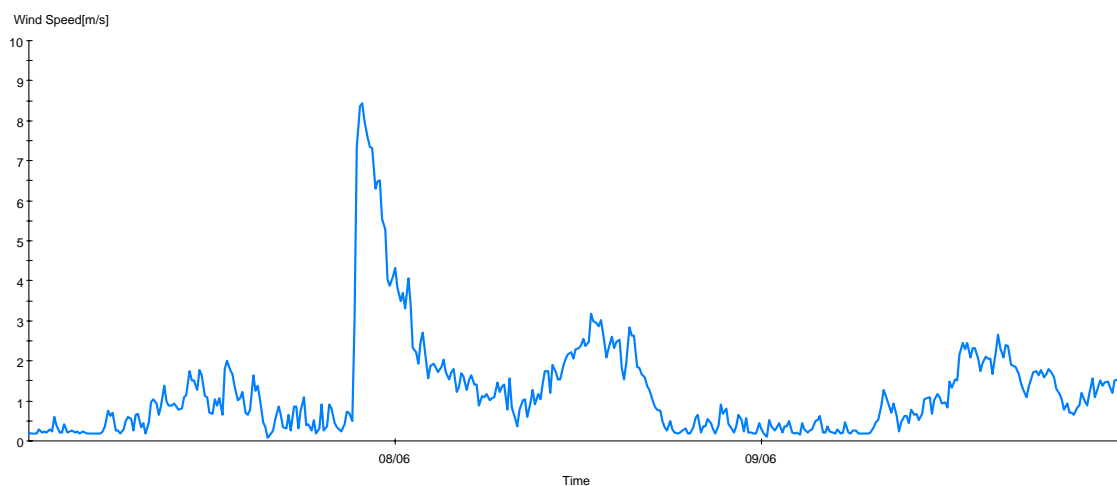


Figure 40: MfE Burnside ambient temperature, typical winter diurnal trend, 10-minute fixed average 7–9 June 2005

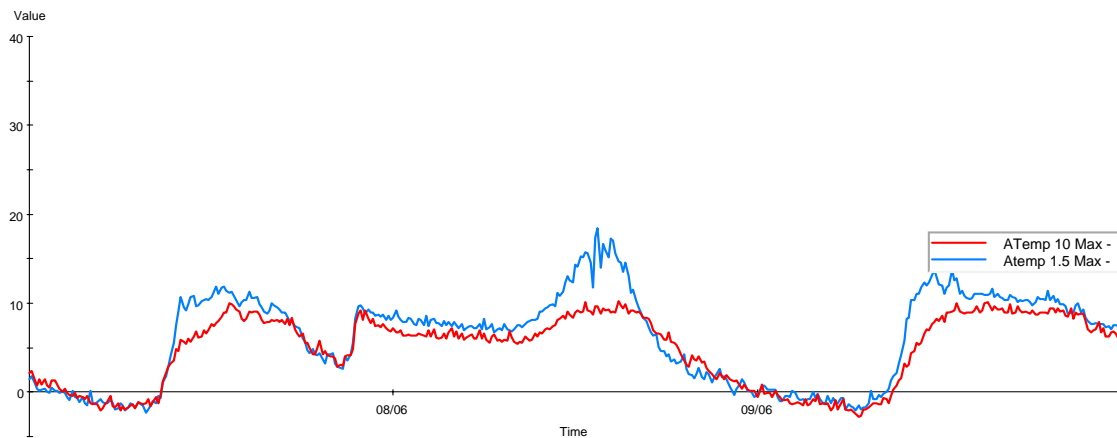
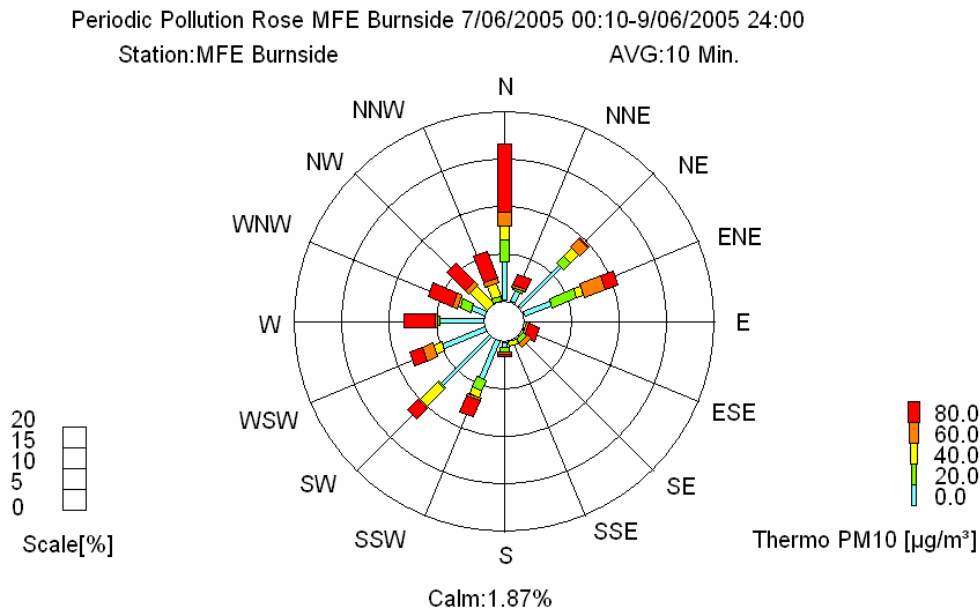


Figure 41: MfE Burnside pollution rose, 10-minute average 7–9 June 2005



7.9.2 NO₂ exceedance at Gavin Street, Penrose

The NO₂ exceedance occurring on 19 December 2005 was investigated and found to be caused by machinery and trucks operating during construction works at the Gavin Street substation, several metres to the east and northeast of the air quality monitoring shed. Figures 42 and 43 provide NO₂ exceedance data for 19 December 2006. Other pollutants were also elevated at Gavin Street during this time (PM₁₀ and SO₂), as described in Figures 44 and 45 below. Figure 46 shows the relationship between wind direction and pollution levels monitored on 19 December 2005.

Figure 42: MfE Gavin Street NO₂ exceedance 10-minute average 19 December 2005

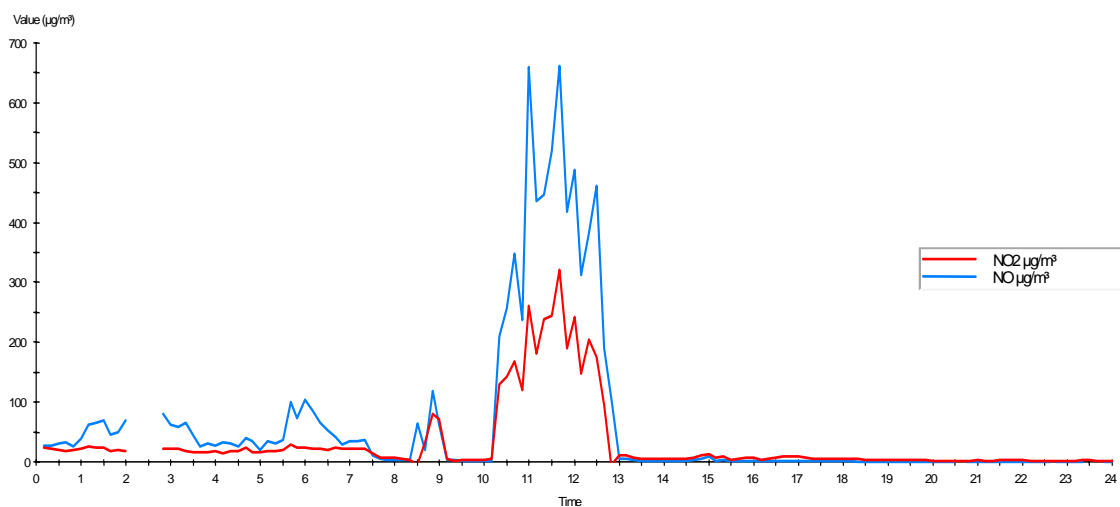


Figure 43: MfE Gavin Street NO₂ exceedance 1-hour average 19 December 2005

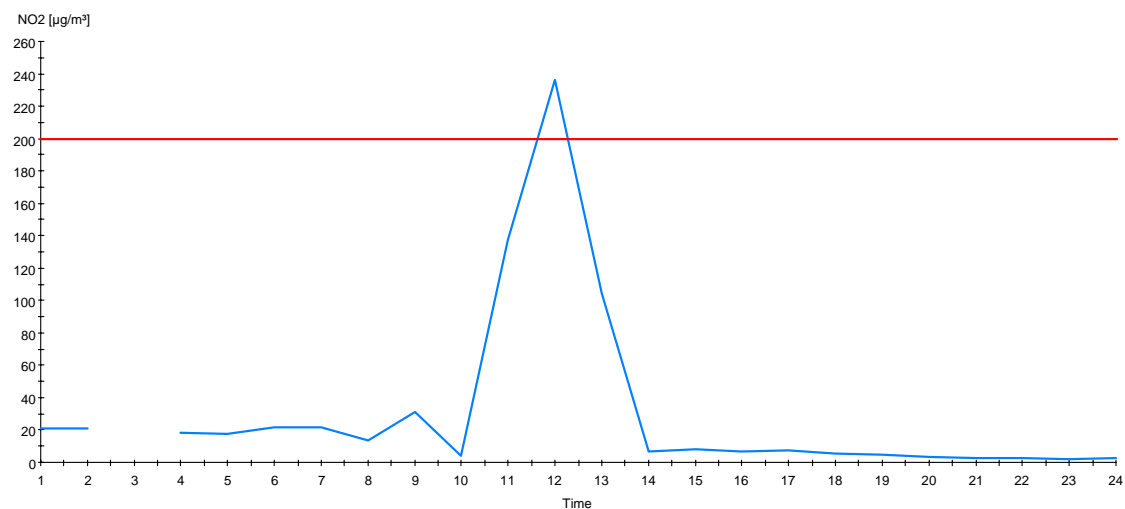


Figure 44: MfE Gavin Street SO₂ 1-hour average 19 December 2005

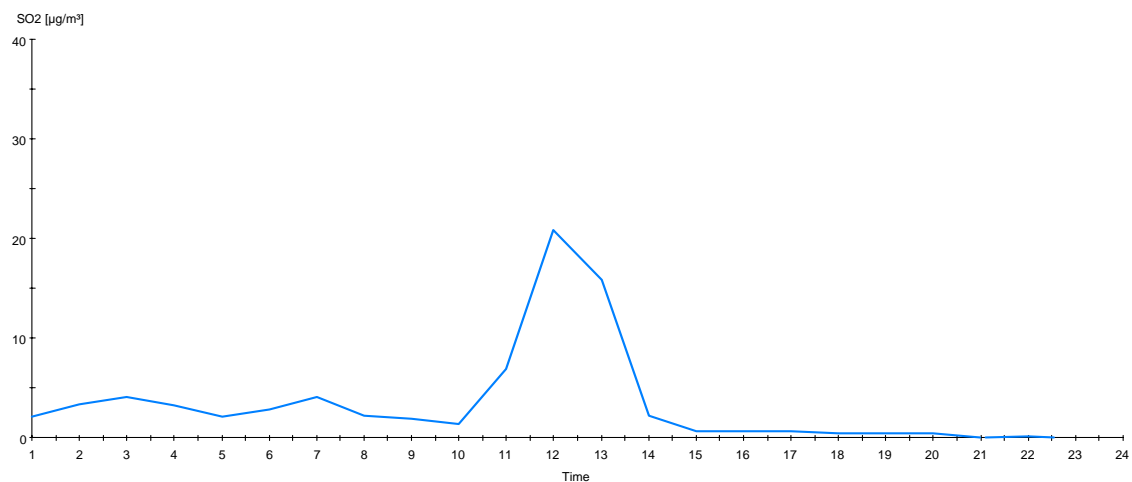


Figure 45: MfE Gavin St PM₁₀ 1-hour average 19 December 2005

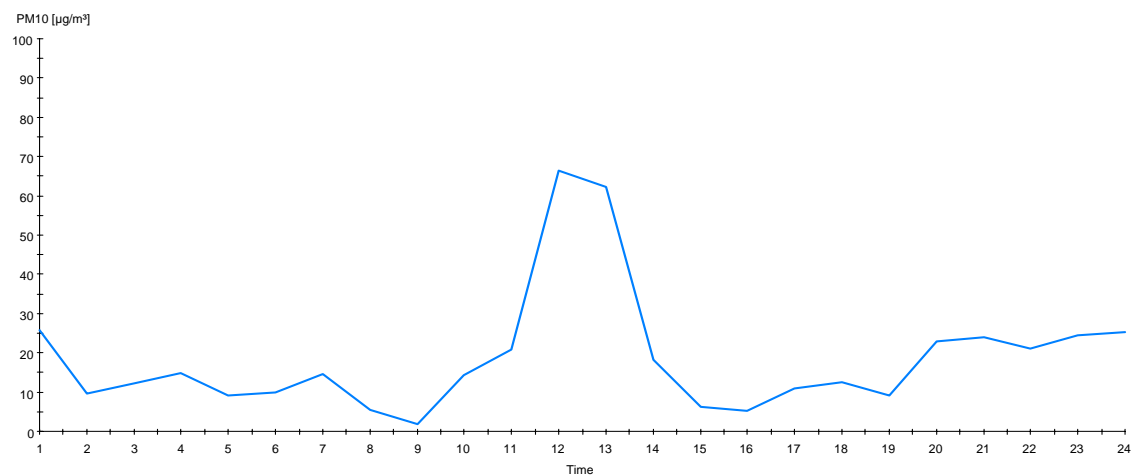


Figure 46: MfE Gavin Street pollution rose 10-minute average 19 December 2005

