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**REPORT NO. DEC14231.2**

**AIR QUALITY IMPACT ASSESSMENT  
OF THE ABC BIRKENHEAD CEMENT  
FACILITY – FY 2014**

***INCREASED PLASTICS IN WOOD TRIAL***

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## 1.0 INTRODUCTION

AirLabs Environmental Pty Ltd (AirLabs) was commissioned by Adelaide Brighton Cement Limited (ABC) to conduct an air quality assessment of the ABC Birkenhead Cement Plant (the facility) located in Birkenhead, South Australia.

This assessment is based on data obtained during source sampling undertaken by AirLabs during October 2014 (AirLabs Report OCT14181.2, issued on 16<sup>th</sup> January, 2015). The objective of this assessment is to numerically determine the air quality impacts beyond the facility's boundary. This is achieved by undertaking dispersion modelling.

This report presents a brief overview of the assessment methodology, an outline of the regulatory framework and a summary of the findings of the assessment.

## 2.0 SOURCE LOCATION AND DESCRIPTION

The Birkenhead facility manufactures cement used in the construction and mining industry. Based on previous modelling studies undertaken and AirLabs' knowledge of the facility, the main sources include the licenced Dry Process Kiln Stack (Stack 4A) and the Pre-Calcliner Plant Stack (Stack 4B). Source parameters are discussed in the subsequent sections of this report.

**Figure 1:** Stack 4A and Stack 4B – ABC Birkenhead Facility



Dry Process Kiln Stack (Stack 4A)



Pre-Calcliner Plant Stack (Stack 4B)

### 3.0 REGULATORY GUIDELINES

#### 3.1 National Legislation

In June 1988 (revised in 2003), the National Environment Protection Council (NEPC) developed the Ambient Air Quality National Environmental Protection Measure (NEPM) which set uniform standards for air quality at the national level and included ambient air quality standards for carbon monoxide, nitrogen dioxide, photochemical oxidants (as ozone), sulphur dioxide, lead and particulate matter with a nominal aerodynamic diameter of less than or equal to 10 microns (PM<sub>10</sub>). The National Environmental Protection (Air Toxics) Measure (Air Toxics NEPM) includes standards for toxics such as Polycyclic Aromatic Hydrocarbons (PAHs).

#### 3.2 South Australia Legislation

The South Australia EPA (EPA, 2006) specifies Design Ground Level Concentration (DGLC) criteria for the assessment of ground level concentrations of air pollutants for new developments and established facilities. The EPA also recognises the air quality standards set by National Environmental Protection Council under the National Environmental Protection (Ambient Air Quality) Measure (Air NEPM) and National Environmental Protection (Air Toxics) Measure (Air Toxics NEPM).

#### 3.3 Summary of Air Criteria for the Project Goals

The air quality criteria adopted for the Project are summarised in **Table 1**.

**Table 1:** Air Quality Criteria adopted for the Project

Pollutant	Averaging Period	Criteria (µg/m <sup>3</sup> )	Source	Environmental Value
<b>Criteria Pollutants</b>				
Nitrogen dioxide (NO <sub>2</sub> )	1 hour	113	EPA	Toxicity
Sulfur dioxide (SO <sub>2</sub> )	1 hour	450	EPA	Toxicity
Carbon monoxide (CO)	1 hour	29,000	EPA	Toxicity
Particulate matter (PM <sub>10</sub> )	24 hours	50	Air NEPM	Health
<b>Metals, Metalloids and Metal Fumes</b>				
Arsenic and compounds	3 minutes	0.17	EPA	International Agency for Research on Cancer (IARC) Group 1
Antimony and compounds	3 minutes	17	EPA	Toxicity
Barium	3 minutes	17	EPA	Toxicity
Benzene	3 minutes	53	EPA	IARC Group 1 Carcinogen
Beryllium and compounds	3 minutes	0.007	EPA	IARC Group 1 Carcinogen
Cadmium and compounds	3 minutes	0.033	EPA	IARC Group 1 Carcinogen

Pollutant	Averaging Period	Criteria (µg/m <sup>3</sup> )	Source	Environmental Value
Chromium (III) compounds	3 minutes	17	EPA	Toxicity
Chromium (VI) compounds	3 minutes	0.17	EPA	IARC Group 1 Carcinogen
Copper	3 minutes	6.7	EPA	Toxicity
Iron oxide	3 minutes	170	EPA	Toxicity
Lead	Annual	0.5	Air NEPM	Health
Magnesium oxide fumes	3 minutes	330	EPA	Toxicity
Manganese and compounds	3 minutes	33	EPA	Toxicity
Mercury - inorganic	3 minutes	3.3	EPA	Bioaccumulation
Mercury - organic	3 minutes	0.33	EPA	Bioaccumulation
Nickel and nickel compounds	3 minutes	0.33	EPA	IARC Group 1 Carcinogen
Zinc oxide	3 minutes	170	EPA	Toxicity
<b>Polycyclic Aromatic Hydrocarbons (PAH)</b>				
PAH (as Benzo(a)pyrene)	Annual	0.0003	Air Toxics NEPM	Monitoring investigation level
<b>Acid Gases</b>				
Hydrogen chloride	3 minutes	250	EPA	Toxicity
<b>Halogens</b>				
Chlorine	3 minutes	100	EPA	Toxicity
Fluoride	24 hours	2.9	EPA	Bioaccumulation
	7 days	1.7	EPA	Bioaccumulation
	90 days	0.5	EPA	Bioaccumulation
<b>Polychlorinated Dibenzo-p-Dioxins and Polychlorinated Dibenzofurans (PCDD and PCDF)</b>				
Dioxins and Furans (as TCDD I-TEQ) – Toxic Equivalency based on North Atlantic Treaty Organisation (NATO)	3 minutes	0.0000037	VIC – EPA Gazette S Schedule 240A	Health

## 4.0 ASSESSMENT METHODOLOGY

In order to predict the impacts from 4A and 4B stacks, dispersion modelling has been used, details of which are presented in this section of the report.

### 4.1 Meteorological Modelling

Meteorological mechanisms govern the generation, dispersion, transformation and eventual removal of pollutants from the atmosphere. The local meteorology at the site plays a significant role in understanding the pollutant transport and dispersion mechanisms. For the project, the meteorological model – The Air Pollution Model (TAPM) (Version 4) was used to predict local meteorological data.

TAPM, developed by the Commonwealth Scientific and Industrial Research Organisation (CSIRO) is a prognostic model which is used to predict three-dimensional meteorological data and air pollution concentrations. The model predicts a time-series dataset comprising of parameters like wind speed, wind direction, temperature, pressure, water vapour, cloud, rain, mixing heights, atmospheric stability classes which are essential for dispersion modelling.

The software allows users to generate synthetic observations by referencing in-built databases (e.g. terrain information, synoptic scale meteorological observations, vegetation and soil type etc.) which are subsequently used in generating site-specific hourly meteorological observations.

Details of the TAPM configuration for this Project are outlined in **Table 2**.

**Table 2:** TAPM Model Configuration for the Project

Parameter	Value
Year of Analysis	2013 (01/01/2013 to 31/12/2013)
Centre of Analysis (Easting, Northing ) (m)	271384, 6142589 (UTM Zone 54 S)
Number of grids (spacing)	5 (30km, 10km, 3km, 1km, 0.3km)
Grid dimensions (nx, ny, nz)	25, 25, 25

Meteorological information obtained from TAPM modelling is presented in the form of wind roses. Wind roses are a way of presenting a summary of wind speed and directional data for a particular time and location. Seasonal and diurnal wind roses along with Annual wind roses obtained from TAPM modelling are presented in **Figure 2** and **Figure 3**, respectively.

Figure 2: Seasonal Wind Roses Modelled for 2013

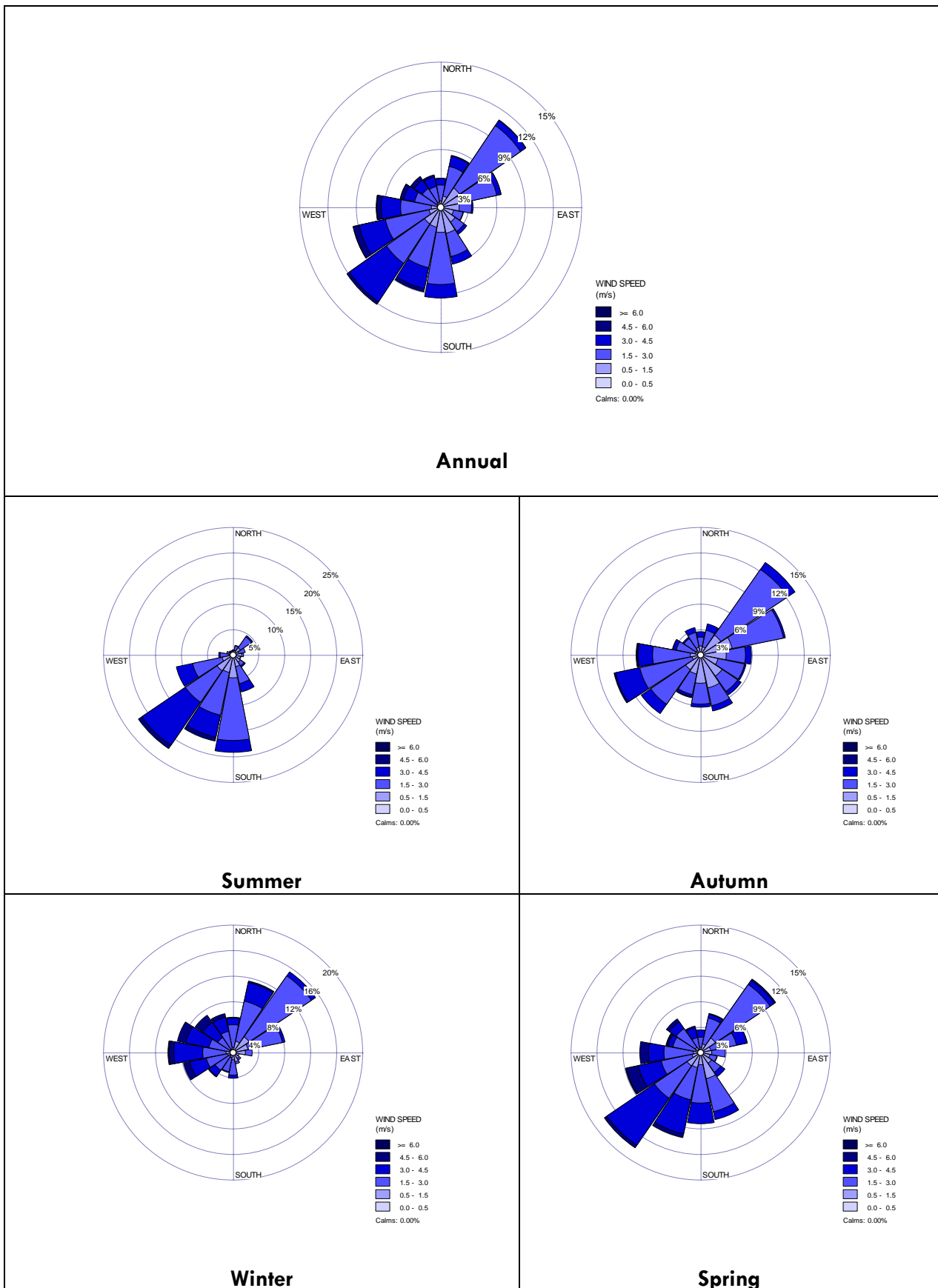
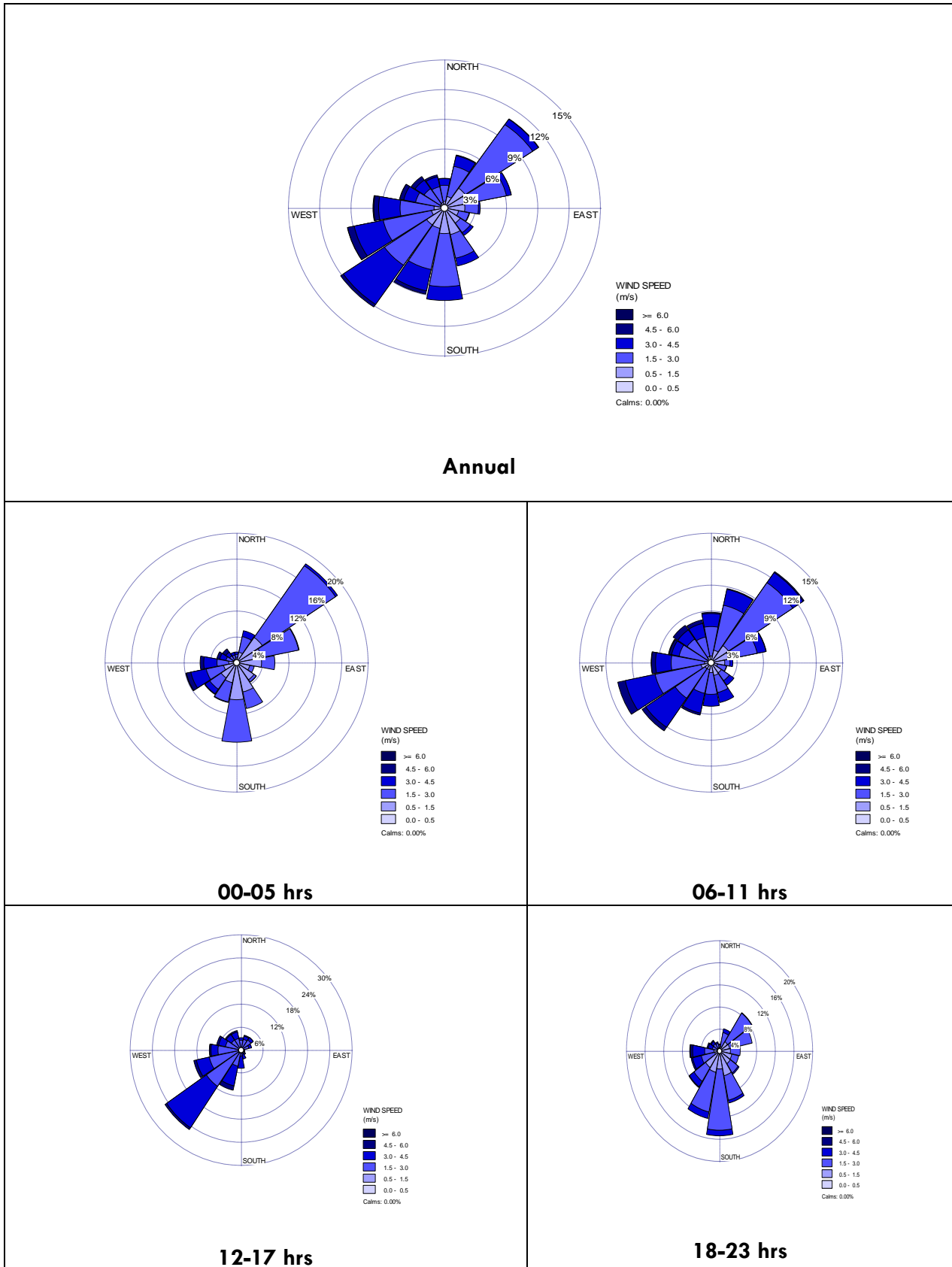


Figure 3: Diurnal Wind Roses Modelled for 2013





From the wind roses shown in **Figure 2** and **Figure 3**, the following observations are made:

- Wind flow is predominantly from the south, south-westerly sectors;
- Light to moderate wind speeds (1.5 – 4.5 m/sec) are observed for most of the year;
- Distinct seasonal variation is observed, with the spring and summer seasons characterised by south, south-westerly flow whereas the wind flow is more scattered during the winter and autumn seasons;
- On a daily cycle, highest wind speeds are typically observed during the afternoon period (12 – 17 hrs) mainly from the south-westerly direction (Refer to **Appendix A** for seasonal and diurnal wind speed frequency distribution analyses); and
- Lowest wind speeds are typically observed during the evening periods (18 – 23 hrs).

#### 4.2 Emission Rates and Source characteristics

Stack testing was conducted in October 2014 (Airlabs Report OCT14181.2, issued on 16<sup>th</sup> January, 2015) to sample the emission rates and source characteristics from Stacks 4A and 4B. Dispersion modelling parameters obtained from the stack testing undertaken in October 2014 are presented in **Table 3**.

**Table 3:** Stack Parameters (Stack 4A and Stack 4B)

Parameter	Stack 4A	Stack 4B
Sampling Date	22 <sup>nd</sup> – 23 <sup>rd</sup> October 2014	23 <sup>rd</sup> – 24 <sup>th</sup> October 2014
Stack height (m)	75.5	96
Stack diameter (m)	3.23	3.0
Stack temperature (°C)	102.6	105
Exit velocity (m/s)	14.7	18.5

Emission rates obtained from the sampling undertaken in October, 2014 and used in the dispersion modelling assessment are presented in **Table 4**.

**Table 4:** Modelled Emission Rates (Stack 4A and Stack 4B)

Pollutant	Emission Rates (g/s)	
	Stack 4A (22 -23 October 2014)	Stack 4B (23- 24 October 2014)
<b>Criteria Pollutants</b>		
Oxides of nitrogen (as NO <sub>2</sub> ) - Minimum	11.8 <sup>(1)</sup>	12.1 <sup>(1)</sup>
Oxides of nitrogen (as NO <sub>2</sub> ) - Average	13.8 <sup>(1)</sup>	13.4 <sup>(1)</sup>
Oxides of nitrogen (as NO <sub>2</sub> ) - Maximum	16.1 <sup>(1)</sup>	15.3 <sup>(1)</sup>
Sulfur dioxide (SO <sub>2</sub> )	7.5 <sup>(2)</sup>	0.23
Carbon monoxide (CO)	3.17	31
Particulate matter (PM <sub>10</sub> )	2.15	0.57
<b>Metals, Metalloids and Metal Fumes</b>		
Arsenic and compounds	0.00002	0.00001 <sup>(2)</sup>
Antimony and compounds	0.00002	0.000014
Barium	0.00023	0.00028
Benzene	0.007 <sup>(2)</sup>	0.009
Beryllium and compounds	0.00001 <sup>(2)</sup>	0.00001 <sup>(2)</sup>
Cadmium and compounds	0.000055 <sup>(2)</sup>	0.000006 <sup>(2)</sup>
Chromium (III) compounds	0.0003	0.0002
Chromium (VI) compounds	0.00011	0.00002
Copper oxide	0.0005	0.00014
Iron oxide	0.05167	0.0233
Lead	0.00085	0.00014
Magnesium oxide fumes	0.01833	0.01
Manganese and compounds	0.00057	0.0002
Mercury - inorganic	0.000003	0.000001
Mercury - organic	0.000163 <sup>(2)</sup>	0.0004
Nickel and nickel compounds	0.00016	0.00011
Zinc oxide	0.00583	0.00048
<b>Polycyclic Aromatic Hydrocarbons (PAH)</b>		
PAH (as Benzo (a)pyrene)	0.000001	0.000001
<b>Acid Gases</b>		
Hydrogen chloride	0.1 <sup>(2)</sup>	0.07
<b>Halogens</b>		
Chlorine	0.003 <sup>(2)</sup>	0.0083 <sup>(2)</sup>
Fluoride	0.003 <sup>(2)</sup>	0.003 <sup>(2)</sup>
	0.003 <sup>(2)</sup>	0.003 <sup>(2)</sup>
	0.003 <sup>(2)</sup>	0.003 <sup>(2)</sup>
<b>Polychlorinated Dibenzo-p-Dioxins and Polychlorinated Dibenzofurans (PCDD and PCDF)</b>		
Dioxins and Furans (as TCDD I-TEQ)	1.1E-10	9.3E-11

**Notes:**

(1): It is to be noted that the modelled emission rates for nitrogen dioxide (NO<sub>2</sub>) are 30% of the measured emissions rates for oxides of nitrogen (NO<sub>x</sub>) based on assumed photochemical NO<sub>x</sub> to NO<sub>2</sub> conversion rate of 30% (Katesone, 2012)

(2): Where the measured results were less than the detection limit, the detection limit was used for modelling purposes.

### 4.3 Dispersion Modelling

AUSPLUME (V6) dispersion model was used in conjunction with prognostic meteorological data obtained from TAPM for predicting impacts from Stack 4A and Stack 4B. AUSPLUME is a steady state Gaussian dispersion model, developed by the Victorian Environmental Protection Agency (Vic-EPA). Ground level concentrations at sensitive receivers in the surrounds of the facility were predicted based on stack parameters (**Table 3**) and stack emission rates (**Table 4**) and assessed for compliance by comparing the predicted ground level concentrations with the air quality criteria (**Table 1**).

Buildings in the vicinity of the sources play a major role in pollutant dispersion as they may create zones of strong turbulence and enhance downward mixing resulting in higher ground-level concentrations. For the Project, the building downwash effect from both the stacks were determined using the US Building Input Program – Plume Rise Model Enhancements (BPIP -PRIME) which is a utility program built into the AUSPLUME (V6) dispersion model. Results obtained from the dispersion modelling are presented in the following sections.

## 5.0 RESULTS AND DISCUSSION

Predicted ground level concentrations of all air pollutants from Stack 4A and Stack 4B are outlined in **Table 5**.

**Table 5:** Dispersion Modelling Results from Stack 4A and Stack 4B

Pollutant	Averaging Period	Criteria ( $\mu\text{g}/\text{m}^3$ )	Max Predicted Impacts Outside Facility Boundary	
			Conc. <sup>(1)</sup> ( $\mu\text{g}/\text{m}^3$ )	% of Criteria
<b>Criteria Pollutants</b>				
Oxides of nitrogen (as NO <sub>2</sub> ) – Minimum	1 hour	113	25	22%
Oxides of nitrogen (as NO <sub>2</sub> ) - Average	1 hour	113	29	26%
Oxides of nitrogen (as NO <sub>2</sub> ) - Maximum	1 hour	113	33	29%
Sulfur dioxide (SO <sub>2</sub> )	1 hour	450	8	1.8%
Carbon monoxide (CO)	1 hour	29000	36	0.1%
Particulate matter (PM <sub>10</sub> )	24 hours	50	0.7	1.5%
<b>Metals, Metalloids and Metal Fumes</b>				
Arsenic and compounds	3 minutes <sup>(2)</sup>	0.17	0.0001	0.06%
Antimony and compounds	3 minutes <sup>(2)</sup>	17	0.0001	0.0006%
Barium	3 minutes <sup>(2)</sup>	17	0.002	0.0093%
Benzene	3 minutes <sup>(2)</sup>	53	0.05	0.09%
Beryllium and compounds	3 minutes <sup>(2)</sup>	0.007	0.0001	1%
Cadmium and compounds	3 minutes <sup>(2)</sup>	0.033	0.0002	0.6%
Chromium (III) compounds	3 minutes <sup>(2)</sup>	17	0.0015	0.009%

Pollutant	Averaging Period	Criteria ( $\mu\text{g}/\text{m}^3$ )	Max Predicted Impacts Outside Facility Boundary	
			Conc. <sup>(1)</sup> ( $\mu\text{g}/\text{m}^3$ )	% of Criteria
Chromium (VI) compounds	3 minutes <sup>(2)</sup>	0.17	0.0004	0.2%
Copper oxide	3 minutes <sup>(2)</sup>	6.7	0.002	0.029%
Iron oxide	3 minutes <sup>(2)</sup>	170	0.2	0.135%
Lead	Annual	0.5	0.00008	0.015%
Magnesium oxide fumes	3 minutes <sup>(2)</sup>	330	0.1	0.026%
Manganese and compounds	3 minutes <sup>(2)</sup>	33	0.002	0.007%
Mercury - inorganic	3 minutes <sup>(2)</sup>	3.3	0.000012	0.0004%
Mercury - organic	3 minutes <sup>(2)</sup>	0.33	0.0017	0.5%
Nickel and nickel compounds	3 minutes <sup>(2)</sup>	0.33	0.001	0.25%
Zinc oxide	3 minutes <sup>(2)</sup>	170	0.02	0.011%
<b>Polycyclic Aromatic Hydrocarbons (PAH)</b>				
PAH (as Benzo(a)pyrene)	Annual	0.0003	9.5E-08	0.03%
<b>Acid Gases</b>				
Hydrogen chloride	3 minutes <sup>(2)</sup>	250	0.5	0.2%
<b>Halogens</b>				
Chlorine	3 minutes <sup>(2)</sup>	100	0.036	0.036%
Fluoride	24 hours	2.9	0.002	0.1%
	7 days	1.7	0.001	0.1%
	90 days	0.5	0.00079	0.2%
<b>Polychlorinated Dibenzo-p-Dioxins and Polychlorinated Dibenzofurans (PCDD and PCDF)</b>				
Dioxins and Furans (as TCDD I-TEQ)	3 minutes <sup>(2)</sup>	0.0000037	6.2E-10	0.017%

**Notes:**

(1): The 99.9th percentile concentrations are reported as the maximum concentrations

(2): 3-minute average concentrations were estimated from predicted hourly averages using the CSIRO Peak to Mean Scaling approach

From the results outlined in **Table 5**, it is observed that ground level concentrations of all pollutants are well below their respective criteria. From all the modelled pollutants, highest concentrations were observed for NO<sub>2</sub>, with their maximum 1-hour average concentrations predicted outside the facility boundary ranging from 22% to 29% of the air quality criteria for lower bound and upper bound emission rates, respectively.

As mentioned earlier, the NO<sub>2</sub> emissions were conservatively estimated by assuming that 30% of total NO<sub>x</sub> emissions released would be converted to NO<sub>2</sub> (Katestone, 2012). For visual illustration, the highest 1-hour average NO<sub>2</sub> concentration isopleths for lower, average and upper bound emission rates are presented in **Figure 4**, **Figure 5** and **Figure 6**, respectively.

With respect to PM<sub>10</sub>, It is to be noted that the 24-hour average concentrations have been conservatively determined by assuming that all the particulates released from Stack 4A and Stack 4B would be in the PM<sub>10</sub> size range.



**Figure 4:** Predicted Maximum Ground Level Concentrations of NO<sub>2</sub> (µg/m<sup>3</sup>) (1 hour averaging period) based on lower bound of emission rates



**Figure 5:** Predicted Maximum Ground Level Concentrations of NO<sub>2</sub> (µg/m<sup>3</sup>) (1 hour averaging period) based on average emission rates



**Figure 6:** Predicted Maximum Ground Level Concentrations of NO<sub>2</sub> (µg/m<sup>3</sup>) (1 hour averaging period) based on upper bound of emission rates



Please note that the coloured area in **Figure 4**, **Figure 5** and **Figure 6**, represent the Birkenhead Facility Precincts

## 6.0 CONCLUSIONS

The AUSPLUME (V6) Gaussian plume dispersion model, in conjunction with TAPM generated meteorological data, was used to predict the ground level concentrations of various air contaminants outside the boundaries of the Adelaide Brighton Cement facility in Birkenhead. Emission rates considered in this assessment were based on results from stack testing conducted by Airlabs Environmental on 22<sup>nd</sup> - 24<sup>th</sup> October, 2014 (Airlabs Report No. OCT14181.2, issued on 16<sup>th</sup> January, 2015).

Results from the dispersion modelling (**Table 5**) show that all the modelled pollutants are below their respective regulatory criteria, indicating that there were no air quality issues observed from the operation of Stacks 4A and 4B.



## 7.0 REFERENCES

AirLabs Environmental (2014) Test report No: OCT14181.2 *Air Emissions Monitoring of Release Points 4A & 4B At Adelaide Brighton Cement Ltd In Birkenhead, 22<sup>nd</sup> – 24<sup>th</sup> October, 2014*. Prepared for Adelaide Brighton Cement. 17<sup>th</sup> December, 2014.

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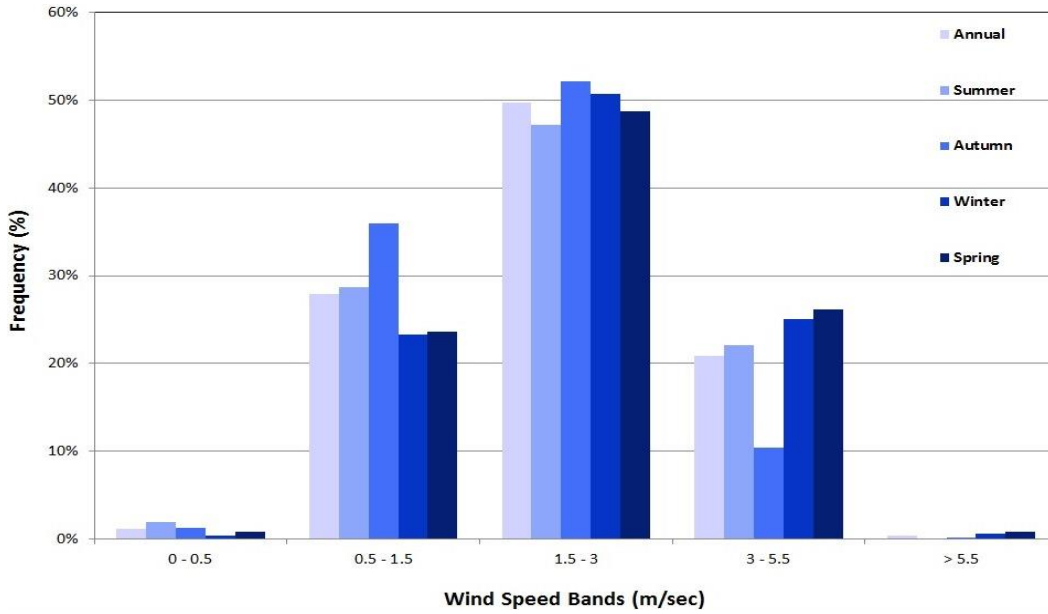
Victoria Environmental Protection Agency (VIC-EPA) Air Quality Guidelines – VIC EPA Gazette S 240 Schedule A

## APPENDIX - A

### 2013 TAPM MODELLED METEOROLOGICAL DATASET – WIND SPEED FREQUENCY DISTRIBUTION ANALYSIS (SEASONAL AND DIURNAL CYCLE)



**Figure A-1:** Seasonal Wind Speed Frequency Distribution for the 2013 TAPM Modelled Meteorological Dataset for the Birkenhead Facility



**Figure A-2:** Diurnal Wind Speed Frequency Distribution for the 2013 TAPM Modelled Meteorological Dataset for the Birkenhead Facility

