

# **Long-Term Trends in Atmospheric Pollutants at Valentia Observatory, Ireland**

## **Environmental Research Centre Report**

### **Authors:**

**Wasim Bashir, Frank McGovern, Margaret Ryan, Liam Burke and Brett Paull**

### **ENVIRONMENTAL PROTECTION AGENCY**

An Ghníomhaireacht um Chaomhnú Comhshaoil  
PO Box 3000, Johnstown Castle, Co. Wexford, Ireland

Telephone: +353 53 916 0600 Fax: +353 53 916 0699

E-mail: [info@epa.ie](mailto:info@epa.ie) Website: [www.epa.ie](http://www.epa.ie)

Lo Call 1890 33 55 99

## **ACKNOWLEDGEMENTS**

This report has been prepared through the Environmental Research Centre, a measure of the ERTDI Programme which is financed by the Irish Government under the National Development Plan 2000–2006. It is administered on behalf of the Department of the Environment, Heritage and Local Government by the Environmental Protection Agency which has the statutory function of co-ordinating and promoting environmental research.

The authors wish to acknowledge the excellent work on sample collection and analysis carried out by Met Éireann staff over many years which has enabled the compilation of this report.

## **DISCLAIMER**

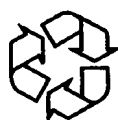
Although every effort has been made to ensure the accuracy of the material contained in this publication, complete accuracy cannot be guaranteed. Neither the Environmental Protection Agency nor the author(s) accept any responsibility whatsoever for loss or damage occasioned or claimed to have been occasioned, in part or in full, as a consequence of any person acting, or refraining from acting, as a result of a matter contained in this publication. All or part of this publication may be reproduced without further permission, provided the source is acknowledged.

Reports produced through the Environmental Research Centre are intended as contributions to inform policy makers and other stakeholders to the necessary debate on the environment.

## **ENVIRONMENTAL RESEARCH CENTRE PROGRAMME 2000–2006**

Published by the Environmental Protection Agency, Ireland

PRINTED ON RECYCLED PAPER



ISBN: 1-84095-193-1

Price: Free

09/06/300

## Details of Project Partners

**Wasim Bashir**

Environmental Protection Agency  
Environmental Research Centre  
McCumiskey House  
Richview, Clonskeagh Road  
Dublin 14  
Ireland

*and*

NCSR  
Dublin City University  
Glasnevin  
Dublin 9  
Ireland

**Margaret Ryan**

Met Éireann, Headquarters  
Glasnevin Hill  
Dublin 9  
Ireland

**Brett Paull**

National Centre for Sensor Research (NCSR)  
Dublin City University  
Glasnevin  
Dublin 9  
Ireland

**Frank McGovern**

Environmental Protection Agency  
Environmental Research Centre  
McCumiskey House  
Richview, Clonskeagh Road  
Dublin 14  
Ireland

Tel.: +353 1 2680160

E-mail: [f.mcgovern@epa.ie](mailto:f.mcgovern@epa.ie)

**Liam Burke**

Met Éireann, Headquarters  
Glasnevin Hill  
Dublin 9  
Ireland



# Table of Contents

|  |            |
|--|------------|
| <b>Acknowledgements</b>  | <b>ii</b>  |
| <b>Disclaimer</b>  | <b>ii</b>  |
| <b>Details of Project Partners</b>                                     | <b>iii</b> |
| <b>Executive Summary</b>   | <b>vii</b> |
| <b>1 Introduction</b>  | <b>1</b>   |
| <b>2 Background</b>  | <b>2</b>   |
| <b>3 National Monitoring of Transboundary Pollutants</b>               | <b>3</b>   |
| <b>4 Results and Analysis of Data from the Valentia Site</b>           | <b>4</b>   |
| 4.1 Sulphur Dioxide (SO <sub>2</sub> )                                 | 4          |
| 4.2 Sulphate (SO <sub>4</sub> <sup>2-</sup> )                          | 5          |
| 4.3 Nitrogen Dioxide (NO <sub>2</sub> )                                | 5          |
| <b>5 Wind Direction-Based Analysis</b>                                 | <b>6</b>   |
| 5.1 Trends for Sectoral SO <sub>2</sub> -S and SO <sub>4</sub> -S Data | 7          |
| <b>6 Discussion</b>  | <b>8</b>   |
| <b>7 Conclusions</b>   | <b>9</b>   |
| <b>References</b>  | <b>10</b>  |
| <b>Appendix 1 Linear Regression Analysis for Sectoral Data</b>         | <b>11</b>  |



# Executive Summary

Concerns in the United Nations Economic Commission for Europe (UNECE) area with respect to the environmental impacts of acidifying species have resulted in a series of international agreements to reduce emissions of such species. Most recently, these have culminated in the targets established under the Gothenburg Protocol. The Gothenburg targets have also been introduced into EU legislation through the National Emission Ceilings (NEC) Directive.

The European Monitoring and Evaluation Programme (EMEP) network, established in the 1970s under the UNECE, provides the structure for development of scientific understanding of emissions and their impacts. EMEP also acts as the co-ordinating body for measurements of ambient levels of key pollutants across Europe.

Since the 1980s, considerable progress has been reported in Europe and nationally on reductions of sulphur species. These emissions reductions have resulted in reduced ambient sulphur dioxide (SO<sub>2</sub>) and sulphate (SO<sub>4</sub><sup>2-</sup>) levels over Ireland. The greatest reductions are seen in SO<sub>2</sub>-S levels for which an average decrease of 0.036 µg m<sup>-3</sup> per annum has been observed. This amounts to a decrease of about 60% in ambient SO<sub>2</sub> levels during the measurement period.

The reduction in ambient SO<sub>4</sub>-S levels is less pronounced at 0.019 µg m<sup>-3</sup> per annum. Nevertheless, this also amounts to a decrease of approximately 40% over the

measurement period. This analysis may be influenced by the presence of natural SO<sub>4</sub><sup>2-</sup> associated with sea spray at the measurement location. The NO<sub>2</sub>-N data series is not as extensive as that for the sulphur species. No trend is evident for the annual average data.

Wind-sector analysis of the SO<sub>2</sub>-S and SO<sub>4</sub>-S data shows that the highest concentrations of these species are correlated with winds coming from an easterly direction. These levels also displayed the greatest rate of decrease over the measurement period. The SO<sub>2</sub>-S levels observed in air from westerly directions are also decreasing. This observation requires further analysis but is considered to be largely associated with reduced local emissions of SO<sub>2</sub>.

The overall reductions in ambient SO<sub>2</sub> and SO<sub>4</sub><sup>2-</sup> levels are considered to show the success of international actions in addressing the challenges of acidification. These reductions have positive consequences both in terms of impacts on ecosystems and human health, and on commercial crops and buildings.

As SO<sub>4</sub><sup>2-</sup> is a significant component of ambient particulate matter (PM), the reduction of SO<sub>4</sub><sup>2-</sup> levels will also contribute towards the achievement of air quality targets for PM<sub>10</sub> and PM<sub>2.5</sub>. However, further significant reductions in acidifying species, particularly nitrogenous species, are required in order to meet current emissions targets by the 2010 deadline.





# 1 Introduction

This EPA National Environmental Research Centre (ERC) report, produced in co-operation with Met Éireann, is the first in a series on air pollution and acidification. This series of reports will provide background information and national analyses of measures in relation to the international Convention on Long-Range Transboundary Air Pollution (CLRTAP) and inform national responses to the EU Clean Air For Europe (CAFE) initiative. Synergies with other areas such as climate change will also be explored.

This report provides an analysis of changes in ambient levels of pollutant species, covered under international agreements on transboundary air pollution and, more

recently, regulated under the EU National Emissions Ceiling Directive, that are measured as part of national monitoring programmes. The data were collected over a period of more than 20 years at the Met Éireann Observatory at Valentia. This site is also a meteorological observatory and houses a phenological garden.

The analysis of these data provides an opportunity to evaluate if and how reported reductions in emissions of species covered under international transboundary pollutant agreements have been observed in Ireland. The analysis presented here is confined to air pollutant data. Analyses of wet deposition/rainfall data will be published in subsequent reports.

## 2 Background

Scientific and public concerns in relation to the impacts of acidifying species on ecosystems and human health, which emerged during the 1970s, have resulted in a series of international agreements to reduce or eliminate such emissions. The European Monitoring and Evaluation Programme (EMEP) was initiated in 1977 as a special programme under the United Nations Economic Commission for Europe (UNECE) under the CLRTAP. EMEP provides scientific support for the Convention and acts to co-ordinate monitoring activities at an extensive range of sites throughout the UNECE area. Data from EMEP sites, which are operated by the Member States, contribute to efforts to understand transport and impacts of the monitored pollutants.

Considerable reductions in European emissions of sulphur dioxide (SO<sub>2</sub>) and other acidifying species have been reported. These reductions have been brought about through a series of factors including change of fuel

use, the introduction of new clean technologies and economic restructuring. These achievements have been coupled with greater understanding of the complexities and interlinkages of pollution emission, transport and transformation, as well as health and ecological impacts, and other damages. This has resulted in a broadening of the scope of species described as transboundary pollutants and the concomitant evolution of monitoring requirements. The increased understanding of the role of transboundary pollutants such as sulphate and ozone in climate change is of particular note (IPCC, 2001).

Ireland's location on the western boundary of Europe, combined with the dominance of mid-latitude North Atlantic westerly winds, favours good air quality conditions. However, long-range pollution transport from North America and particularly short- and medium-range transport from Europe increases ambient levels of pollutant species.

### 3 National Monitoring of Transboundary Pollutants

The Met Éireann Valentia Observatory positioned at 51° 56' N, 10° 15' W, altitude 9 m above sea level (a.s.l.), was established as an EMEP monitoring site in 1980. The Electricity Supply Board (ESB) subsequently established additional sites at Turlough Hill (1991), the Burren (1997) and the Ridge of Capard (1997). The location of these sites is shown in Fig. 3.1 and other details are given in Table 3.1. An extensive suite of atmospheric measurements is also carried out at the National University of Ireland (NUI) Galway atmospheric research station at Mace Head, some of which are reported to the EMEP database.

The three ESB sites were closed in 2003. However, three replacement sites have been established at Carnsore, Co. Wexford, Oakpark, Co. Carlow and Malin Head, Co. Donegal in 2004. These have been established by the EPA in co-operation with Met Éireann, as part of an EPA COE project on transboundary air pollution and acidification. The locations of the new sites are also shown in Fig. 3.1. The sites provide enhanced geographical coverage and operate under the new EMEP operational protocol, which has been developed to address current transboundary pollution issues, such as PM<sub>10</sub> and PM<sub>2.5</sub>.

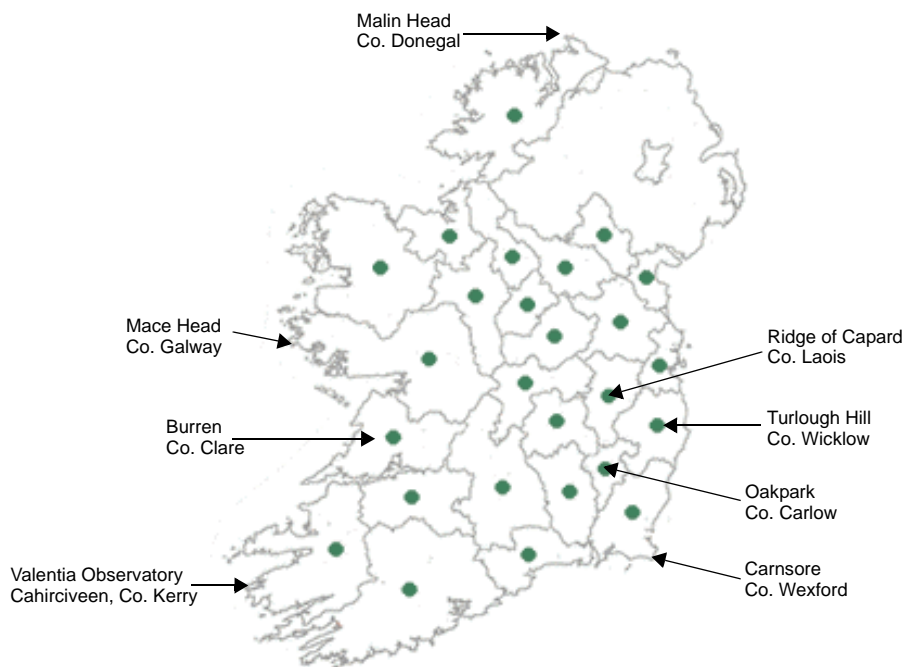


Figure 3.1. Map of Ireland showing the locations of Met Éireann Valentia Observatory, Burren, the NUI Galway atmospheric research station at Mace Head, Malin Head, Ridge of Capard, Turlough Hill, Oakpark, and Carnsore.

Table 3.1. Details of EMEP sites in Ireland. The Valentia and Mace Head sites are regional and Global Atmospheric Watch sites. The ESB sites were closed down in 2002. Three new sites are being established at Oakpark, Malin Head and Carnsore.

| Organisation | Measurement site | Database code | Geographical co-ordinates | EMEP co-ordinates (50 km) | Altitude a.s.l. (m) | In operation since |
|--------------|------------------|---------------|---------------------------|---------------------------|---------------------|--------------------|
| Met Éireann  | Valentia         | IE0001R       | 51° 56' N, 10° 15' W      | 73.39, 44.83              | 9                   | 1980               |
| ESB          | Turlough Hill    | IE0002R       | 53° 02' N, 06° 24' W      | 77.35, 49.33              | 420                 | 1991               |
| ESB          | Burren           | IE0003R       | 53° 00' N, 96° 00' W      | 73.96, 47.74              | 90                  | 1997               |
| ESB          | Ridge of Capard  | IE0004R       | 05° 37' N, 07° 27' W      | 75.94, 48.88              | 340                 | 1997               |
| NUI Galway   | Mace Head        | IE0031R       | 53° 20' N, 09° 54' W      | 73.30, 47.86              | 340                 | 1988               |

## 4 Results and Analysis of Data from the Valentia Site

This report concentrates on results for the Valentia Observatory. This site has the most extensive data record for SO<sub>2</sub>-S, SO<sub>4</sub>-S and NO<sub>2</sub>-N available for Ireland. Measurements of SO<sub>2</sub> and sulphate (SO<sub>4</sub><sup>2-</sup>) commenced in 1980. The measurements of nitrogen dioxide (NO<sub>2</sub>) commenced in 1988. The data record was interrupted during 1998/99 due to instrumentation problems and site reorganisation.

### 4.1 Sulphur Dioxide (SO<sub>2</sub>)

The time series of annual average SO<sub>2</sub>-S concentration data along with standard deviation values are shown in Fig. 4.1. A clear downward trend towards lower concentration levels is evident from these data. The linear regression trend line shown in Fig. 4.2 indicates an average concentration reduction rate of 0.036 µg m<sup>-3</sup> per

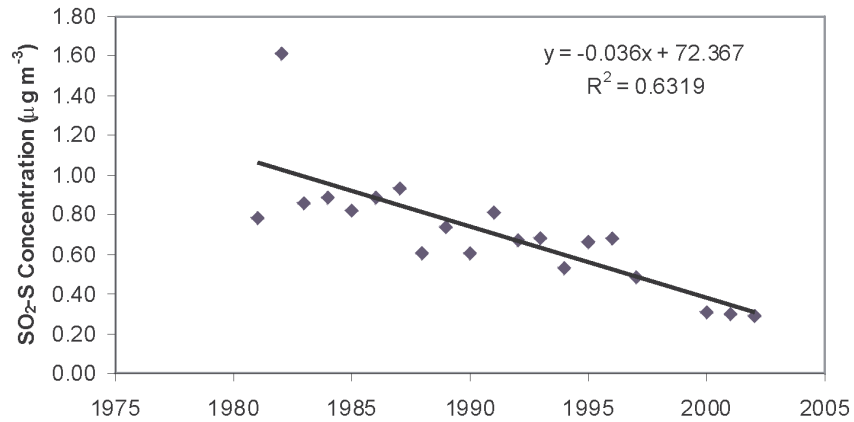


Figure 4.1. Annual average SO<sub>2</sub>-S levels (µg m<sup>-3</sup>) for the period 1981 to 2002. The linear regression trend line shows a significant reduction in ambient levels over the measurement period.

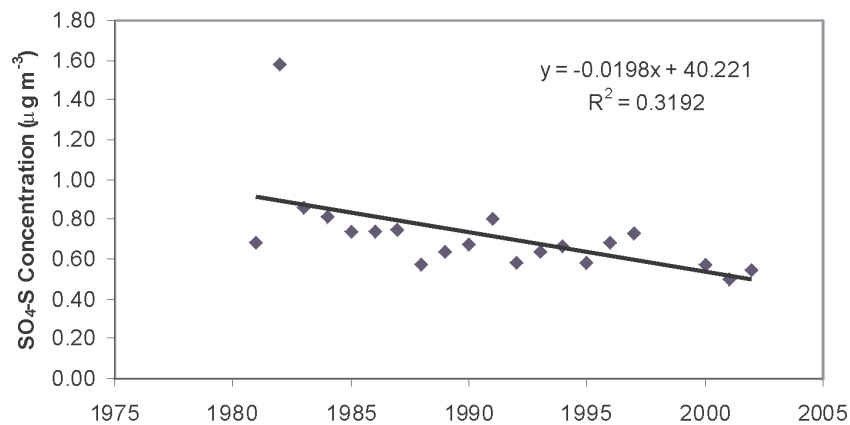


Figure 4.2. Annual average SO<sub>4</sub>-S levels (µg m<sup>-3</sup>) for the period 1981 to 2002. The linear regression trend line indicates a significant reduction in ambient levels over the measurement period.



**Figure 4.3. Annual average NO<sub>2</sub>-N levels (µg m<sup>-3</sup>) for the period 1989 to 2002. The linear regression trend line shows that ambient levels remained relatively stable over the measurement period with no distinct trend.**

annum over the measurement period<sup>1</sup>. This analysis suggests that annual average levels have decreased from approximately 1.0 µg m<sup>-3</sup> in 1981 to less than 0.4 µg m<sup>-3</sup> in 2002. This represents a 60% reduction in annual average levels. Further analysis of the data shows that this was primarily due to a reduction in peak levels, which is supported by the observation that the standard deviation values also decrease over this period. This is considered to reflect a reduction in the magnitude of variation in peak concentration levels experienced at the site.

## 4.2 Sulphate (SO<sub>4</sub><sup>2-</sup>)

Annual average SO<sub>4</sub>-S data and standard deviation values are shown in Fig. 4.2. As was the case for SO<sub>2</sub>-S, a clear trend towards lower levels is evident for the annual average data. The linear regression trend line indicates a concentration reduction rate of 0.012 µg m<sup>-3</sup> per annum. This rate of decrease is not as large as that observed for

SO<sub>2</sub>-S. A number of factors including the contribution of sea-salt sulphate<sup>2</sup> at this site may have contributed to the observed levels. However, SO<sub>4</sub><sup>2-</sup>, which is largely formed in the atmosphere through oxidation of SO<sub>2</sub>, may be transported over long distances. This means that a range of anthropogenic sources, including North American sources, may have contributed to these observations.

## 4.3 Nitrogen Dioxide (NO<sub>2</sub>)

The time series of annual average NO<sub>2</sub>-N data for Valentia is shown in Fig. 4.3. The levels have remained relatively stable throughout the measurement period, which commenced in 1989. The data set is therefore less extensive than that for SO<sub>2</sub>-S and SO<sub>4</sub>-S. An annual wintertime peak is apparent in the daily data. Further analysis of this feature is required.

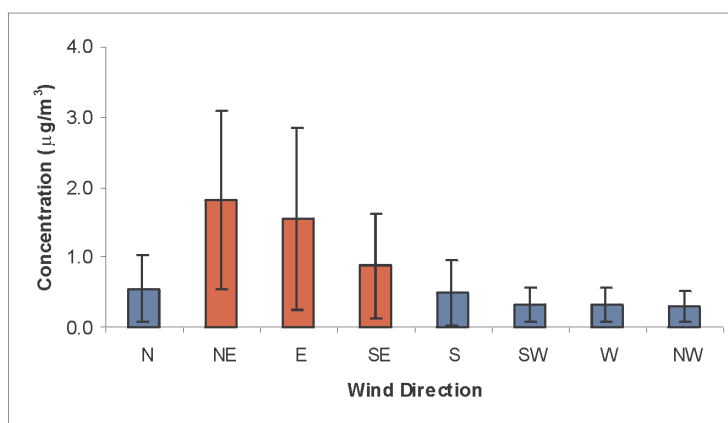
1. Linear regression analysis is not fully applicable to these data but is used as a first approximation.

2. Salt arising from sea spray is a source for sulphate. Sea-salt ions such as sodium (Na<sup>+</sup>) and chlorine (Cl<sup>-</sup>) were not determined. Therefore it was not possible to determine the sea-salt contribution, i.e. to determine non-sea-salt/anthropogenic sulphate levels. Current sample analysis allows for this.

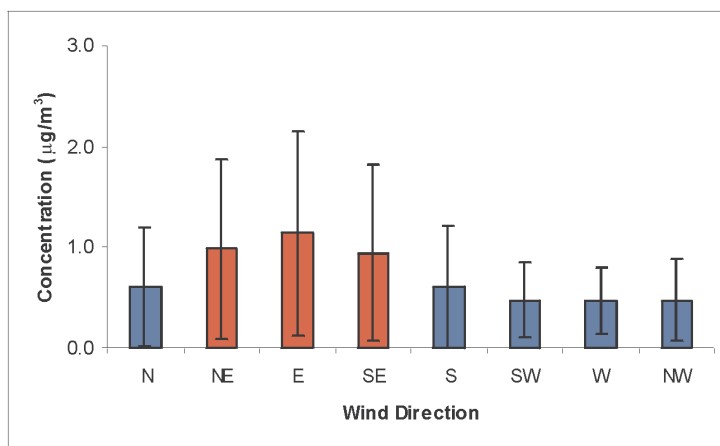
## 5 Wind Direction-Based Analysis

The daily concentration data were sectorized according to the daily average wind direction. Average concentrations of  $\text{SO}_2\text{-S}^3$ ,  $\text{SO}_4\text{-S}^4$  and  $\text{NO}_2\text{-N}^5$  data for six wind local sectors are shown in Figs 5.1–5.3, respectively. The data

and figures clearly show that the highest pollutant concentrations were recorded for east and north-easterly directions. The lowest concentrations were found for westerly directions.



**Figure 5.1. Average  $\text{SO}_2\text{-S}$  levels ( $\mu\text{g m}^{-3}$ ) over the period 1981 to 2002 for different wind sectors at the Valentia Observatory. Standard deviation values are also shown.**

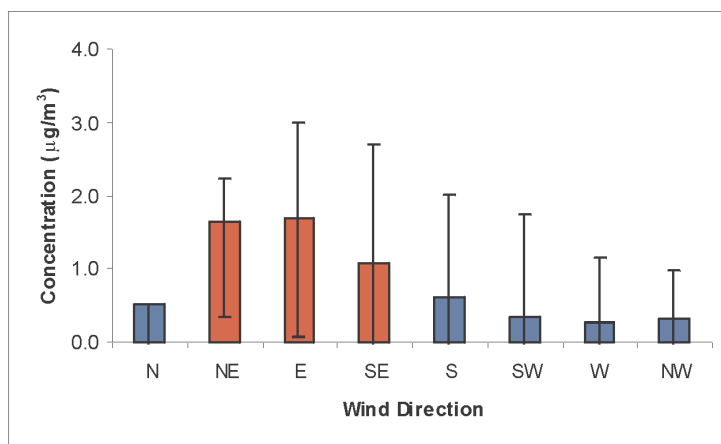


**Figure 5.2. Average  $\text{SO}_4\text{-S}$  levels ( $\mu\text{g m}^{-3}$ ) over the period 1981 to 2002 for different wind sectors at the Valentia Observatory. Standard deviation values are also shown.**

3. To convert  $\text{SO}_2\text{-S}$  ( $\mu\text{g m}^{-3}$ ) to  $\text{SO}_2$  ( $\mu\text{g m}^{-3}$ ), multiply  $\text{SO}_2\text{-S}$  ( $\mu\text{g m}^{-3}$ ) by 2.

4. To convert  $\text{SO}_4\text{-S}$  ( $\mu\text{g m}^{-3}$ ) to  $\text{SO}_4^{2-}$  ( $\mu\text{g m}^{-3}$ ), multiply  $\text{SO}_4\text{-S}$  ( $\mu\text{g m}^{-3}$ ) by 3.

5. To convert  $\text{NO}_2\text{-N}$  ( $\mu\text{g m}^{-3}$ ) to  $\text{NO}_2$  ( $\mu\text{g m}^{-3}$ ), divide  $\text{NO}_2\text{-N}$  ( $\mu\text{g m}^{-3}$ ) by 0.3.



**Figure 5.3. Average NO<sub>2</sub>-N levels (µg m<sup>-3</sup>) over the period 1989 to 2002 for different wind sectors at the Valentia Observatory. Standard deviation values are also shown.**

The difference in concentrations between easterly and westerly data is clearest for the SO<sub>2</sub>-S and NO<sub>2</sub>-N data. The difference in concentration is not as large for SO<sub>4</sub>-S. As was outlined in the previous section, this can in part be attributed to the contribution from sea salt in air coming from the west over the open ocean. This contribution is reduced during easterly conditions. It is also notable that the SO<sub>2</sub>-S and NO<sub>2</sub>-N values are non-zero for westerly directions.

### 5.1 Trends for Sectoral SO<sub>2</sub>-S and SO<sub>4</sub>-S Data

It has been shown above that the annual average SO<sub>2</sub>-S and SO<sub>4</sub>-S data display a significant decrease in concentration levels since measurements commenced in 1980. It is apparent that reductions in ambient levels occur for all wind directions. Linear regression analysis shows that the greatest rate of decrease, i.e. 0.06 µg m<sup>-3</sup> per annum, was observed in SO<sub>2</sub>-S levels measured during

easterly winds. This is considered to reflect largely local emissions reductions, i.e. in Ireland and Britain. These analyses are given in [Appendix 1](#).

The NE data are less well defined and may be influenced by industrial emissions in the Shannon basin region. More surprisingly, the concentrations measured in samples from westerly air display a small but significant reduction over the measurement period, i.e. 0.012 µg m<sup>-3</sup> per annum for the south-westerly sector. Long-range transport of SO<sub>2</sub> from North America is not considered to be a significant influence on the measurements. Natural oceanic sources such as seasonal emissions from phytoplankton are also not expected to display such a decline. Therefore, this observation is considered to reflect decreased SO<sub>2</sub> levels in recirculated air from largely local sources. However, further examination of this feature is required.

Trends in the NO<sub>2</sub>-N data were less defined.

## 6 Discussion

The Valentia data provide a relatively long-term perspective on variation of ambient  $\text{SO}_2$ ,  $\text{SO}_4^{2-}$  and  $\text{NO}_2$  levels at the western boundary of Europe. The data show the success of international emissions reduction actions in relation to ambient levels of sulphur species. The decrease in observed  $\text{SO}_2$ -S levels is most pronounced for easterly air masses but is also, and somewhat surprisingly, evident in westerly air masses. Similar reductions are observed in ambient  $\text{SO}_4^{2-}$  levels, although these are less pronounced. The  $\text{NO}_2$ -N data remain relatively stable and probably reflect the difficulties that exist in addressing emissions of oxides of nitrogen from combustion sources.

The  $\text{SO}_2$  levels would be expected to decrease as local emissions are reduced. This observation therefore confirms the success of local emissions reduction actions.

Sulphate is produced by oxidation of  $\text{SO}_2$  in the atmosphere and is typically associated with  $\text{NH}_4$ . It is well known that  $\text{SO}_4^{2-}$  is an important and sometimes dominant component of ambient particulate matter (PM) levels. As particulate sulphate is typically found in size range less than  $1\ \mu\text{m}$  it contributes to both  $\text{PM}_{10}$  and  $\text{PM}_{2.5}$  levels. McGovern and McGettigan (2003) show that

the concurrence of peaks in ambient  $\text{SO}_4^{2-}$  levels at Valentia and peaks in urban  $\text{PM}_{10}$  levels, e.g. Dublin and Cork cities. The reduction in the observed  $\text{SO}_4^{2-}$  peak is therefore also welcome, as an indication that background PM levels have also been reduced.

The analysis of the  $\text{SO}_4$ -S data is somewhat hampered by the absence of analysis of the contribution from sea salt to these levels. Sea-salt ions, such as  $\text{Na}^+$  and  $\text{Cl}^-$  are now part of the regular EMEP chemical analysis. The newer data may provide a mechanism to estimate the contribution of sea salt for historical concentration levels. Integrated analysis of these data with similar data for Mace Head, which have been corrected for sea salt, may also help to estimate this contribution.

There is no clear trend apparent in the  $\text{NO}_2$ -N data although the daily data suggest that an annual winter peak exists. Further work on these data is required, including analysis of levels of linked species, such as ozone. This work will be enhanced through the current work being developed in the COE transboundary air pollution project. In addition, the new EMEP analysis protocol requires a wider range of particulate/aerosol ion species analysis.



## 7 Conclusions

The Valentia data constitute a valuable record of atmospheric change since 1981. These data show the success of national and international actions to reduce emissions of sulphur species. This has resulted in a decrease in ambient SO<sub>2</sub> and SO<sub>4</sub><sup>2-</sup> levels observed in Ireland. These reductions have positive consequences in term of impacts on ecosystems and human health, as well as for commercial crops and buildings. The main findings from the observational records are:

- SO<sub>2</sub>-S levels have decreased at a rate of approximately 0.036 µg m<sup>-3</sup> per year since 1981. The rate of decrease is greater for air masses originating from an easterly direction. Levels in westerly air have also decreased.
- SO<sub>4</sub>-S levels have decreased at a rate of 0.02 µg m<sup>-3</sup> per year; however, the real rate of decrease may be masked by local influences.

- The results for NO<sub>2</sub>-N are not yet as clear. No trend has been observed in the ambient levels, which have remained relatively stable since measurements commenced.

Further and more detailed analysis of these data is required. This includes integration with new data for the Valentia site and inter-comparison with similar data for other national and international sites. This will contribute to the development of a broader understanding of these results.

It is envisaged that further reduction of emissions of acidifying species will occur in the coming years. This will lead to further reductions in ambient levels of these species and contribute to improvement of our environment. The ongoing monitoring and analysis of the data from the Valentia site, and other sites in Ireland, will provide evidence of the success of these reductions. The full data set can be obtained from <http://coe.epa.ie>.

## References

Intergovernmental Panel on Climate Change (IPCC), 2001. *Working Group I Third Assessment Report*. Cambridge University Press. Cambridge, UK. 881 pp.

Intergovernmental Panel on Climate Change (IPCC), 2001. *Synthesis Report, Third Assessment Report*. Cambridge University Press. Cambridge, UK. 397 pp.

Intergovernmental Panel on Climate Change (IPCC), 2001. *Working Group II Third Assessment Report*. Cambridge University Press. Cambridge, UK. 1032 pp.

McGovern, F. & McGettigan, M., 2003. *Air Quality Monitoring Annual Report 2001*. Environmental Protection Agency, Johnstown Castle Estate, Co. Wexford, Ireland.

## Appendix 1 Linear Regression Analysis for Sectorised Data

The SO<sub>2</sub>-S, SO<sub>4</sub>-S and NO<sub>2</sub>-N concentration data were segregated according to wind sectors based on wind direction data for the Valentia site. Table A1 shows linear regression analysis for annual average values for these data. It is notable that all the slope values are negative for

the SO<sub>2</sub>-S and SO<sub>4</sub>-S data, with the highest gradient/rate of decrease being observed for samples obtained from easterly air masses. There is no clear trend in the NO<sub>2</sub>-N data.

**Table A1. Linear regression analysis for annual average values for SO<sub>2</sub>-S, SO<sub>4</sub>-S and NO<sub>2</sub>-N concentration data.**

|                         | Slope   | Error  | Constant | Error | R <sup>2</sup> |
|-------------------------|---------|--------|----------|-------|----------------|
| <b>SO<sub>2</sub>-S</b> |         |        |          |       |                |
| W                       | -0.018  | 0.0020 | 36.3     | 4.02  | 0.82           |
| NW                      | -0.015  | 0.0023 | 29.2     | 4.65  | 0.68           |
| SW                      | -0.012  | 0.0019 | 24.8     | 3.97  | 0.68           |
| E                       | -0.065  | 0.0113 | 130.0    | 22.7  | 0.64           |
| SE                      | -0.035  | 0.0062 | 69.6     | 12.5  | 0.63           |
| S                       | -0.017  | 0.0034 | 34.4     | 6.86  | 0.58           |
| NE                      | -0.045  | 0.0266 | 91.0     | 53.1  | 0.14           |
| N                       | -0.016  | 0.0183 | 32.4     | 36.   | 0.04           |
| <b>SO<sub>4</sub>-S</b> |         |        |          |       |                |
| SE                      | -0.022  | 0.005  | 44.8     | 10.0  | 0.52           |
| W                       | -0.006  | 0.002  | 12.0     | 4.2   | 0.3            |
| S                       | -0.005  | 0.002  | 10.5     | 4.0   | 0.25           |
| E                       | -0.023  | 0.0097 | 47.4     | 19.4  | 0.24           |
| NW                      | -0.0073 | 0.004  | 15.0     | 8.0   | 0.15           |
| N                       | -0.022  | 0.0132 | 45.3     | 26.2  | 0.14           |
| SW                      | -0.0029 | 0.0018 | 6.3      | 3.6   | 0.12           |
| NE                      | -0.017  | 0.0144 | 34.7     | 28.6  | 0.07           |
| <b>NO<sub>2</sub> N</b> |         |        |          |       |                |
| N                       | 0.0334  | 0.0124 | -66.3    | 24.73 | 0.42           |
| NE                      | 0.0981  | 0.0387 | -194.0   | 77.16 | 0.39           |
| E                       | 0.0629  | 0.0249 | -125.1   | 49.59 | 0.39           |
| SE                      | 0.0348  | 0.0149 | -69.0    | 29.79 | 0.35           |
| S                       | 0.0270  | 0.0205 | -53.3    | 40.81 | 0.15           |
| SW                      | 0.0152  | 0.0217 | -29.2    | 43.24 | 0.05           |
| W                       | 0.0138  | 0.0335 | -26.9    | 66.78 | 0.02           |
| NW                      | -0.0058 | 0.0298 | 13.3     | 59.47 | 0.00           |