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RECENT RESEARCH RESULTS AND ONGOING
AND PLANNED RESEARCH PROGRAMMES

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A summary of Australian research activities relevant to the ozone layer for the 8th Session of the Co-ordinating Committee on the Ozone Layer

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In Australia atmospheric research programs relevant to studies on the ozone layer are, by in large, carried out by the CSIRO Division of Atmospheric Research and by the Bureau of Meteorology. These organisations jointly operate the Baseline Air Pollution Monitoring Station at Cape Grim, Tasmania, at which several of the following research programs are based.

CSIRO has conducted research on stratospheric ozone in the Southern Hemisphere since the late 1950's. Over a twenty year record variability on several time scales has been identified, the most important are being annual cycles (winter maximum, summer minimum), biennial cycles and a decreasing trend (approximately 4% over 20 years), the latter being barely statistically significant and possibly due to long term changes in the strength of stratospheric circulation. The Bureau of Meteorology operates the five Dobson ozone stations, between 27°S and 54°S, at Brisbane, Perth, Melbourne, Hobart and Macquarie Island. All stations make daily total ozone measurements and weekly Umkehr observations. The Melbourne program is augmented by fortnightly Mast-Brewer ozone sonde measurements at nearby Laverton. The instrument used at Perth is fully automated and operated in conjunction with LIDAR measurements on stratospheric aerosols, the latter through the Western Australian Institute of Technology in a program supported by the Bureau. Future stations are planned for Darwin (12°S) and Alice Springs (23°S) using automated instrumentation. The Hobart station may be closed down, due to its proximity to Melbourne. The ozone data collected at these stations are periodically logged with the World Ozone Data Center. The Bureau of Meteorology have recently appointed a scientist who will supervise the ozone/radiation program and do data analysis and interpretation.

CSIRO has been researching background tropospheric ozone since the 1960's. Surface ozone measurements have been made at Cape Grim since 1976 in a program designed to detect long term changes in ozone concentrations in the lower atmosphere. An ozone annual cycle at Cape Grim has been identified, again with a winter maximum and a summer minimum, caused by seasonal variations in the transport of ozone rich
stratospheric air into the troposphere at mid-latitudes of the southern hemisphere, and by summertime photochemical destruction in this clean southern hemispheric air. Analysis is under way to determine the meteorological influences on ozone levels at Cape Grim. Long term trends in ozone concentration at Cape Grim have not been detected, in agreement with results from other Baseline Stations (Samoa, South Pole). Tropospheric ozone removal processes have also been studied and the loss of ozone from the atmosphere over a variety of natural surfaces has been investigated and ozone destruction rates deduced.

Halocarbons (CCl₃F (Figure 1a), CCl₂F₂, CH₃CCl₃ (Figure 1b), CCl₄) and nitrous oxide (N₂O (Figure 2b)) observational programs have also been conducted at Cape Grim since 1976. Long term trends have been detected and modelled using global release data to deduce the atmospheric lifetimes of these chemicals, which are precursors of catalysts capable of ozone destruction. Halocarbon growth rates at Cape Grim have been used to calibrate the transport component of a 2-dimensional tracer transport model or, conversely, given the known transport fields and atmospheric residence times, to deduce global releases. Since 1978 this program has formed an integral part of the Atmospheric Lifetime Experiment (ALE), involving scientists from CSIRO, the USA (Oregon Graduate Center, the Massachusetts and Georgia Institutes of Technology) and from the UK (University of Bristol). Since early 1985 this program has been expanded to include methane and CCl₂CFCl₂, and renamed GAGE (Global Atmospheric Gases Experiment).

A number of other atmospheric chemical constituents are significant in ozone chemistry. Methane, carbon dioxide and nitrous oxide are important 'greenhouse' gases which help determine average stratospheric temperatures and thus ozone levels. Methane also reduces chlorine catalysed ozone destruction by partial removal of stratospheric chlorine, some of which originates from the above anthropogenically released organochlorine compounds. Carbon monoxide levels control the abundance of hydroxyl (OH) radicals, which in turn regulate the fluxes of certain trace gases from the troposphere to the stratosphere.

Methane (CH₄ (Figure 2a)) and carbon monoxide (CO) have been studied at Cape Grim and over SE Australia since 1978. A long term trend has been measured for CH₄ (ca. 1% per year) and significant seasonal cycles in CH₄ and CO quantified. Modelling experiments suggest
that the trend in CH$_4$ is due to expanding anthropogenic industrial and agricultural activities, and the seasonal cycles appear to be responding to calculated seasonal changes in OH radical levels. CSIRO participants in a global CH$_4$ and CO$_2$ observational program organised by NOAA's Geophysical Monitoring for Climate Change Laboratory.

Carbon dioxide (CO$_2$ (Figure 3)) observations commenced at Cape Grim in 1976 and complement a CSIRO program of CO$_2$ measurement over SE Australia that commenced in 1972. Trends, seasonal cycles and vertical gradients have been measured. The data have been used in conjunction with global data from the Scripps Institute of Oceanography and the National Oceanographic and Atmospheric Administration (USA), in a 2-dimensional model, to refine the magnitudes of the known sources and sinks of CO$_2$, especially the biosphere, that are part of the global carbon cycle.

Oxides of nitrogen are important precursors in tropospheric ozone chemistry, as are non-methane hydrocarbons. A sensitive chemiluminescent nitrogen oxides (NO, NO$_2$) detector has been operating at Cape Grim intermittently since 1978, to study the reactive gas chemistry of the background maritime airmass. Studies are also underway at CSIRO to determine the influence of biological sources of oxides of nitrogen (denitrification and nitrification) on the NO/NO$_2$ budget of the remote continental atmosphere. To this end measurements of NO/NO$_2$ fluxes have been made over various fertilized and unfertilized agricultural areas. A research program has just commenced at Cape Grim investigating the level of and temporal variations in non-methane hydrocarbons (largely ethane) in background maritime air.

CSIRO have also been involved in the measurement of stratospheric constituents over Australia. NO, NO$_2$, HNO$_3$, O$_3$, N$_2$O, CCl$_3$F, H$_2$O, CH$_4$ and aerosols have been analysed at 23°S and 34°S during the Austral autumn and spring, corresponding with the times of O$_3$ minimum and maximum in the stratosphere. Such measurements provide important tests of the chemical and dynamical theories of the stratosphere.

In conjunction with US scientists (Temple University), erythermally effective ultraviolet radiation (UV-B) has been measured at Aspendale and Brisbane since 1975. To date the data show no significant change in UV-B, the radiation responsible for skin damage, consistent with the observation that very little change has occurred in total ozone.
CSIRO is conducting research into atmospheric dynamics relevant to stratospheric ozone. In conjunction with Monash University the climatology of the southern hemisphere stratosphere is being established, and the dynamics of wintertime stratospheric warmings and meridional circulations are being studied. CSIRO is also researching the theory of 2-dimensional transport parameterizations and, in conjunction with Geophysical Fluid Dynamics Laboratory (NOAA) and Goddard Space Center (NASA), is deriving 2-dimensional transport coefficients from the output of general circulation models of the atmosphere as well as from direct observations.

Adelaide University is conducting research into the study of mesospheric winds, planetary waves, and gravity waves, and with CSIRO, large scale atmospheric transport in the mesosphere.

In early 1987 the Bureau of Meteorology is co-ordinating a series of international experiments in northern Australia involving up to 100 scientists and technicians. One experiment, STEP (Stratosphere - Troposphere Exchange Project), sponsored by NASA, will research the vertical transport of $O_3$ and $H_2O$ across the tropical tropopause. The other experiments, AMEX (Australian Monsoon Experiment) conducted by the Bureau, EMEX (Equatorial Mesoscale Experiment) conducted by NOAA and a consortium of US universities, will investigate the synoptic environment and heat exchange processes in the Australian monsoon. The CSIRO F-27 aircraft will be involved in AMEX and will provide a platform for a series of trace gas measurements.
Figure Captions

**Figure 1** Monthly mean CC\textsubscript{3}F (a) and CH\textsubscript{3}CCl\textsubscript{3} (b) concentrations observed at Cape Grim Tasmania (pptv = parts per 10\textsuperscript{12} by volume).

**Figure 2** Monthly means CH\textsubscript{4} (a) and N\textsubscript{2}O concentrations observed at Cape Grim Tasmania (ppbv = parts per 10\textsuperscript{9} by volume).

**Figure 3** Monthly mean CO\textsubscript{2} concentrations observed over SE Australia at 3.5 to 5.5 km (ppmv = parts per 10\textsuperscript{6} by volume).
Figure 1a

$\text{CCl}_3\text{F} \ pptv$

Figure 1b

$\text{CH}_3\text{CCl}_3 \ pptv$