

NASA/TM—2000—209891, Vol. 211



**Technical Report Series on the
Boreal Ecosystem-Atmosphere Study (BOREAS)**

Forrest G. Hall and Sara Conrad, Editors

Volume 211

**BOREAS TF-11 CO₂ and CH₄
Concentration Data from the SSA-Fen**

*David W. Valentine
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National Aeronautics and
Space Administration

Goddard Space Flight Center
Greenbelt, Maryland 20771

November 2000

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BOREAS TF-11 CO₂ and CH₄ Concentration Data from the SSA-Fen

David Valentine

Summary

The BOREAS TF-11 team collected several data sets in its efforts to fully describe the flux and site characteristics at the SSA-Fen site. This data set contains temperature, pH, and concentration profiles of methane and carbon dioxide within the surface 50 cm of peat. The measurements were conducted as part of a 2 x 2 factorial experiment in which we added carbon (300 g/m² as wheat straw) and nitrogen (6 g/m² as urea) to four replicate locations in the vicinity of the TF-11 tower. The data set covers the period from the first week of June 1994 through the second week of September 1994. The data are stored in tabular ASCII files.

Table of Contents

- 1) Data Set Overview
- 2) Investigator(s)
- 3) Theory of Measurements
- 4) Equipment
- 5) Data Acquisition Methods
- 6) Observations
- 7) Data Description
- 8) Data Organization
- 9) Data Manipulations
- 10) Errors
- 11) Notes
- 12) Application of the Data Set
- 13) Future Modifications and Plans
- 14) Software
- 15) Data Access
- 16) Output Products and Availability
- 17) References
- 18) Glossary of Terms
- 19) List of Acronyms
- 20) Document Information

1. Data Set Overview

1.1 Data Set Identification

BOREAS TF-11 CO₂ and CH₄ Concentration Data from the SSA-Fen

1.2 Data Set Introduction

This data set contains temperature, pH, and concentration profiles of methane and carbon dioxide within the surface 50 cm of peat. The measurements were conducted as part of a 2 x 2 factorial experiment in which we added carbon (300 g/m² as wheat straw) and nitrogen (6 g/m² as urea) to four replicate locations in the vicinity of the Tower Flux (TF)-11 tower. In addition to siting and treatment variables, it reports air temperature and water table height relative to the average peat surface during each measurement. The data set covers the period from the first week of June 1994 through the second week of September 1994.

1.3 Objective/Purpose

Much of the area within the boreal forest biome consists of wetlands, in which large carbon stores and high water tables drive fundamentally different atmospheric interactions than occur under the other forest types studied by the BOREal Ecosystem-Atmosphere Study (BOREAS). One key difference is in the form carbon is emitted following soil microbial respiration; in wetlands, much of it is emitted as methane. Wetlands are the dominant influence of boreal forests on atmospheric methane.

This study was undertaken in order to assess responses of methane emissions in northern wetlands to potential changes in plant productivity, nitrogen availability or both. Whiting and Chanton (1993) recently observed that methane emissions from wetlands across the globe are well related to net primary productivity (NPP). This may be for a variety of reasons, including enhanced plant transport, increased methanogenic substrates from root exudates, increased litter input cascading to enhanced substrate availability for methanogenesis, or enhanced C and N mineralization of decomposing residues. Previous work by us (Valentine et al., 1994) and others has shown that substrate availability is a key constraint on methane production in wetlands. The present study was an effort to test whether substrate manipulation results from laboratory studies could be mirrored under field conditions.

1.4 Summary of Parameters

We measured concentrations of methane and carbon dioxide dissolved in the porewater of the surface 50 cm of peat using probes integrating a series of "swampsuckers" similar to the device described by Burton and Beauchamp (1994). Measurements are reported for 0 (surface), 5, 10, 20, 30, 40, and 50 cm depths. One profile probe was established per platform and treatment combination, yielding 16 probes. Measurements were made weekly coincident with the flux measurements.

Temperature profiles of the surface 50 cm of peat were obtained using thermocouples integrated with the swampsucker probe. Measurements are reported for 0 (surface), 5, 10, 20, 30, 40, and 50 cm depths. Measurements were made weekly to coincide with the flux measurements in data set one.

1.5 Discussion

These data were collected from a set of small locations within the fen, and therefore no one location represented the entire study site. In fact, the fen in which this work was conducted was characterized by a large-scale gradient of vegetation, microtopography, and hydrology such that the study site itself is representative only of the portion of the fen in which it was located (i.e., the lower 1/3).

These data were collected at the same site and over the same time period as Shashi Verma and his team measured methane and carbon dioxide fluxes using eddy correlation. Measurements made using micrometeorological and chamber techniques comprised the two major components of the TF-II effort.

1.6 Related Data Sets

BOREAS TE-05 CO₂ Concentration and stable isotope composition
BOREAS TF-11 Biomass Data over the SSA-Fen
BOREAS TF-11 CO₂ and CH₄ Flux data from the SSA-Fen
BOREAS TGB-01/TGB-03 CH₄ Chamber flux data over the NSA-Fen
BOREAS TGB-01/TGB-03 Water Table and Peat Temperature Data over the NSA
BOREAS TGB-03 Plant Species Composition Data over the NSA-Fen
BOREAS TGB-01/TGB-03 NEE Data over the NSA-Fen
BOREAS TGB-03 CH₄ and CO₂ Chamber Flux Data over NSA Upland Sites

2. Investigator(s)

2.1 Investigator(s) Name and Title

David Valentine
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Fairbanks, AK 99775-7200

2.2 Title of Investigation

Influence of Substrate Characteristics and Other Environmental Factors on Methane Emissions from the BOREAS Southern Study Area Fen Site

2.3 Contact Information

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3. Theory of Measurements

Concentration profiles of methane (CH₄) and carbon dioxide (CO₂) were measured by collecting porewater samples at 0, 5, 10, 20, 30, 40, and 50 cm using a probe integrating a series of swampsuckers, similar in design to the one described by Burton and Beauchamp (1994). Porewater CH₄ concentration was determined by first stripping the porewater by adding an equal amount of air and shaking the syringe for 5 minutes, then measuring the CH₄ concentration in the equilibrated air using a gas chromatograph (GC) equipped with a flame ionization detector (FID), and finally back-calculating the porewater CH₄ concentration based on Henry's law partitioning. Carbon dioxide concentrations in the headspace were measured by passing the chromatographically separated headspace sample through a Shimadzu methanizer just upstream of the FID. Porewater concentrations were calculated using Henry's law partitioning and accounting for pH of the porewater. The porewater pH was measured after headspace equilibration with a Beckman pH meter and a combination pH probe with an Ag/AgCl reference electrode. Temperatures at each porewater sampling depth were measured using twisted wire thermocouples in the gas profile sampling probe.

4. Equipment

4.1 Sensor/Instrument Description

4.1.1 Collection Environment

Data were collected near mid-day (+/- 2 h) at weekly intervals for each platform. Flooding at the site at the end of July 1994 prevented data collection for 1 week.

4.1.2 Source/Platform

The porewater profile samplers were placed, four per platform, at the beginning of the season and remained in place throughout the summer.

4.1.3 Source/Platform Mission Objectives

Not applicable.

4.1.4 Key Variables

Name	Unit	Description
-----	-----	-----
PWCH4	μmol/L	Dissolved methane
PWCO2	μmol/L	Dissolved carbon dioxide
SOILT	°C	Peat temperature
pH	pH unit	Porewater pH after equilibration with headspace

4.1.5 Principles of Operation

A probe combining sampling tubes and twisted wire thermocouples at several depths is used to measure temperatures and to obtain porewater samples for analysis of pH and concentrations of CH₄ and CO₂.

4.1.6 Sensor/Instrument Measurement Geometry

None given.

4.1.7 Manufacturer of Sensor/Instrument

The profile samplers were built by investigator. They were made of series of Teflon (PTFE) or stainless steel tubing (1/8 OD) embedded in epoxy resin within 3/4" PVC pipe. All tubing elements pass through the upper end of the pipe and terminate in a stopcock used for sampling. Each tubing element terminates at a specific point along the side of the pipe corresponding to the depth to be sampled. Opposite each side opening was a twisted wire thermocouple (copper-constantan), with lead wires also exiting through the upper end of the pipe and terminating with plugs.

GC: GC-8A equipped with 1-mL sample injection loop, FID, and methanizer, manufactured by:

Shimadzu Scientific Instruments, Inc.
7102 Riverwood Drive
Columbia, MD 21046 USA
(410) 381-1227
(410) 381-1222 (fax)
(800) 477-1227

The electronic thermometer: Omega microprocessor thermometer model HH21, was manufactured by:

OMEGA Engineering, Inc.
One Omega Drive
Stamford, CT 06907-0047
P.O. Box 4047
(800) 826-6342
(203) 359-1660
(203) 359-7700 (fax)

The pH meter: Beckman 11 portable pH meter with automatic temperature compensation with Beckman combination pH probe (Ag/AgCl reference electrode), manufactured by:

Beckman Instruments Inc.
2500 Harbor Boulevard
Box 3100
Fullerton, CA 92634-3100
(800) 742-2345
(800) 643-4366 (fax)

4.2 Calibration

4.2.1 Specifications

Profile samplers: Air bubbles drawn into the syringe from the sampler reflect gas bubbles in the profile, and are typically very concentrated in CH₄ (often 40% by volume). Our interest in the concentration profile centered on how dissolved gases were distributed, and therefore we decided to expel any sampled bubbles immediately.

Based on comparisons of the electronic thermometer with a laboratory mercury thermometer at typical soil temperature ranges, all temperatures obtained from the thermocouples were adjusted upward by 0.5 °C.

The pH meter was calibrated using commercially available pH buffers at pH 4 and 7.

The GC column oven was operated at 70 °C, FID temperature was 180 °C, and N₂ carrier gas flowed at 35 mL per minute.

4.2.1.1 Tolerance

None given.

4.2.2 Frequency of Calibration

The GC was calibrated at the start of each day using one of two calibration standards, depending on anticipated concentration ranges. Headspace samples from equilibrating porewater gas profile samples had high concentrations of both methane and carbon dioxide, so we used a standard containing 10,000 ppmv (analysis $\pm 2\%$) of both these gases (Scotty IV Can mix 216, obtained from Scott Specialty Gases, Longmont, CO 80501, (303) 442-4700). For flux measurements entailing much lower concentrations of methane and carbon dioxide, we diluted the above-referenced Scotty standard 10:1 with ambient air. This was done by loading a stopcock-fitted, 60-mL polypropylene syringe with excess standard, expelling all but 6 mL, then immediately pulling in outside ambient air to make up 60 mL. Calculation of the diluted standard accounted for average concentrations of methane and carbon dioxide in ambient air. Analyses of diluted standards yielded reproducibility across dilutions of better than 3%. All standards were run on the GC until reproducibility was better than 1% over the course of three standard injections from a single syringe. Calibration was rechecked initially every 10 samples, but detector stability was so high that calibration was checked only at the end of the day for most of the season. End-of-day calibration checks were always within 5% of the starting calibration.

The pH probe was subject to drift, so a 2-point calibration was performed after every 10 samples.

4.2.3 Other Calibration Information

None given.

5. Data Acquisition Methods

Profiles of peat temperature and concentrations of methane and carbon dioxide were measured concurrently with the flux measurements. We sampled 20 mL of the surface water directly with a 60-mL polypropylene syringe, and for subsurface concentrations combined the syringe with a sampler similar to the one described by Burton and Beauchamp (1994) with ports at 5, 10, 20, 30, 40, and 50 cm. We added a twisted wire copper-constantan thermocouple at each depth, and measured temperature using an electronic thermometer. Any bubbles drawn into the syringe were expelled immediately after sampling, and the water samples were stored in a refrigerator at 4 °C until analysis within 4 days. Analysis consisted first making a 1:1 headspace:water mixture in the syringe using ambient air, then shaking vigorously for 5 minutes. The equilibrated headspace was then injected into the GC and analyzed as above for CH₄ and CO₂. Original porewater concentrations were calculated based on Henry's Law partitioning corrected for temperature and, for CO₂, the water pH after equilibration (after Shurpali et al., 1993).

Temperature at each depth was measured simply by plugging each thermocouple lead into the electronic thermometer.

The pH of each porewater sample was measured after headspace equilibration. The sample was stirred gently during the measurement, and the pH was recorded after the reading stabilized (generally after 2 min.).

6. Observations

6.1 Data Notes

Profile and other data are sparse or missing for the end of July 1994 because heavy rains raised the water table above the tops of all chamber collars and of most platforms.

6.2 Field Notes

None given.

7. Data Description

7.1 Spatial Characteristics

7.1.1 Spatial Coverage

All measurements were made along two transects identified by their location relative to the TF-11 micrometeorology tower: a north transect (NA and NB platforms) and a south transect (SA and SB platforms). All measurements were made within 70 m of the TF-11 tower, whose North American Datum of 1983 (NAD83) coordinates are 53.80206°N, 104.61798°W.

7.1.2 Spatial Coverage Map

Not available.

7.1.3 Spatial Resolution

These data are point measurements at the given locations.

7.1.4 Projection

Not applicable.

DATE_OBS	instrument.
TIME_OBS	The date on which the data were collected.
C_ADDED	The Greenwich Mean Time (GMT) when the data were collected.
N_ADDED	Estimated amount of carbon contained in the wheat straw that was added to the plot.
PEAT_DEPTH	Estimated amount of nitrogen contained in the urea that was added to the plot.
PEAT_TEMP	The depth below the peat surface.
DISS_CH4_CONC	The peat temperature measured at the given depth.
DISS_CO2_CONC	Dissolved methane concentration at the given depth.
POREWATER_PH	Dissolved carbon dioxide concentration at the given depth.
SITE_COMMENTS	Porewater pH after equilibrium with headspace.
CRTFCN_CODE	Descriptive information to clarify or enhance the site information.
REVISION_DATE	The BOREAS certification level of the data. Examples are CPI (Checked by PI), CGR (Certified by Group), PRE (Preliminary), and CPI-??? (CPI but questionable).
	The most recent date when the information in the referenced data base table record was revised.

7.3.3 Unit of Measurement

The measurement units for the parameters contained in the data files on the CD-ROM are:

Column Name	Units
SITE_NAME	[none]
SUB_SITE	[none]
DATE_OBS	[DD-MON-YY]
TIME_OBS	[HHMM GMT]
C_ADDED	[grams C][meter ⁻²]
N_ADDED	[grams C][meter ⁻²]
PEAT_DEPTH	[millimeters]
PEAT_TEMP	[Celsius]
DISS_CH4_CONC	[micromoles][liter ⁻¹]
DISS_CO2_CONC	[micromoles][liter ⁻¹]
POREWATER_PH	[pH]
SITE_COMMENTS	[none]
CRTFCN_CODE	[none]
REVISION_DATE	[DD-MON-YY]

7.3.4 Data Source

The sources of the parameter values contained in the data files on the CD-ROM are:

Column Name	Data Source
SITE_NAME	[Assigned by BORIS Staff]
SUB_SITE	[Assigned by BORIS Staff]
DATE_OBS	[Investigator]
TIME_OBS	[Investigator]
C_ADDED	[Investigator]

N_ADDED	[Investigator]
PEAT_DEPTH	[Investigator]
PEAT_TEMP	[Thermometer]
DISS_CH4_CONC	[Gas chromatograph]
DISS_CO2_CONC	[Gas chromatograph]
POREWATER_PH	[pH meter]
SITE_COMMENTS	[Investigator]
CRTFCN_CODE	[Assigned by BORIS Staff]
REVISION_DATE	[Assigned by BORIS Staff]

7.3.5 Data Range

The following table gives information about the parameter values found in the data files on the CD-ROM.

Column Name	Minimum Data Value	Maximum Data Value	Missng Data Value	Unrel Data Value	Below Detect Limit	Data Not Cllctd
SITE_NAME	SSA-FEN-FLXTR	SSA-FEN-FLXTR	None	None	None	None
SUB_SITE	9TF11-CON01	9TF11-CON08	None	None	None	None
DATE_OBS	08-JUN-94	17-SEP-94	None	None	None	None
TIME_OBS	1	2202	None	None	None	None
C_ADDED	0	300	None	None	None	None
N_ADDED	0	6	None	None	None	None
PEAT_DEPTH	0	500	None	None	None	None
PEAT_TEMP	-1.1	20.6	None	None	None	Blank
DISS_CH4_CONC	.01	1750.45	None	None	None	Blank
DISS_CO2_CONC	66	14544	None	None	None	Blank
POREWATER_PH	5.59	7.1	None	None	None	Blank
SITE_COMMENTS	N/A	N/A	None	None	None	None
CRTFCN_CODE	CPI	CPI	None	None	None	None
REVISION_DATE	09-SEP-98	09-SEP-98	None	None	None	None

- Minimum Data Value -- The minimum value found in the column.
- Maximum Data Value -- The maximum value found in the column.
- Missng Data Value -- The value that indicates missing data. This is used to indicate that an attempt was made to determine the parameter value, but the attempt was unsuccessful.
- Unrel Data Value -- The value that indicates unreliable data. This is used to indicate an attempt was made to determine the parameter value, but the value was deemed to be unreliable by the analysis personnel.
- Below Detect Limit -- The value that indicates parameter values below the instruments detection limits. This is used to indicate that an attempt was made to determine the parameter value, but the analysis personnel determined that the parameter value was below the detection limit of the instrumentation.
- Data Not Cllctd -- This value indicates that no attempt was made to determine the parameter value. This usually indicates that BORIS combined several similar but not identical data sets into the same data base table but this particular science team did not measure that parameter.

Blank -- Indicates that blank spaces are used to denote that type of value.
N/A -- Indicates that the value is not applicable to the respective column.
None -- Indicates that no values of that sort were found in the column.

7.4 Sample Data Record

The following are wrapped versions of data record from a sample data file on the CD-ROM.

```
SITE_NAME, SUB_SITE, DATE_OBS, TIME_OBS, C_ADDED, N_ADDED, PEAT_DEPTH, PEAT_TEMP,  
DISS_CH4_CONC, DISS_CO2_CONC, POREWATER_PH, SITE_COMMENTS, CRTFCN_CODE, REVISION_DATE  
'SSA-FEN-FLXTR', '9TF11-CON02', 08-JUN-94, 1011, 300, 6, 0, , .01, 66.0, 6.92,  
'Hollow: North of Tower, along transect A', 'CPI', 09-SEP-98  
'SSA-FEN-FLXTR', '9TF11-CON02', 08-JUN-94, 1011, 300, 6, 50, 14.5, 1.53, 1458.0, 6.92,  
'Hollow: North of Tower, along transect A', 'CPI', 09-SEP-98
```

8. Data Organization

8.1 Data Granularity

The smallest unit of data tracked by the BOREAS Information System (BORIS) is the measurement(s) made for a given site on a given day.

8.2 Data Format(s)

The Compact Disk-Read-Only Memory (CD-ROM) files contain American Standard Code for Information Interchange (ASCII) numerical and character fields of varying length separated by commas. The character fields are enclosed with single apostrophe marks. There are no spaces between the fields.

Each data file on the CD-ROM has four header lines of Hyper-Text Markup Language (HTML) code at the top. When viewed with a Web browser, this code displays header information (data set title, location, date, acknowledgments, etc.) and a series of HTML links to associated data files and related data sets. Line 5 of each data file is a list of the column names, and line 6 and following lines contain the actual data.

9. Data Manipulations

9.1 Formulae

9.1.1 Derivation Techniques and Algorithms

None given.

9.2 Data Processing Sequence

9.2.1 Processing Steps

None given.

9.2.2 Processing Changes

None given.

9.3 Calculations

9.3.1 Special Corrections/Adjustments

None given.

9.3.2 Calculated Variables

Porewater CH₄ concentration (PWCH₄) was calculated as:

$$CH_4hs * P/Po * Kh + Vhs/Vw * (CH_4hs - CH_4amb) * P/RT$$

where: CH₄hs is the equilibrated headspace concentration of CH₄ (ppm)
P and Po are ambient and sea-level atmospheric pressures (101325 Pa)
Kh is the Henry's law constant for CH₄ at ambient temperature (298 K)
Vhs and Vw are the volumes of headspace and water in the syringe
CH₄amb is the ambient CH₄ concentration (1.8 ppmv)
R is the universal gas constant (8314.4 [L Pa]/[K mol])
T is the ambient temperature (298 K)

Porewater CO₂ concentration (PWCO₂) was calculated as:

$$CO_2hs * P/Po * Kh * (1 + Ka1/10^{pH} * (1 + Ka2/10^{pH})) + Vhs/Vw * (CO_2hs - CO_2amb) * P/RT$$

where: CO₂hs is the equilibrated headspace concentration of CO₂ (ppm)
P and Po are ambient and sea-level atmospheric pressures (101325 Pa)
Kh is the Henry's law constant for CO₂ at ambient temperature (298 K)
Ka1 and Ka2 are the 1st and 2nd dissociation constants for aqueous CO₂
pH is the pH of the porewater after stripping
Vhs and Vw are the volumes of headspace and water in the syringe
CO₂amb is the ambient CO₂ concentration (350 ppmv)
R is the universal gas constant (8314.4 [L Pa]/[K mol])
T is the ambient temperature (298 K)

9.4 Graphs and Plots

None given.

10. Errors

10.1 Sources of Error

During sampling, a leak in the syringe or elsewhere in the profile probe system could generate bubbles that would strip gases from the sample. As bubbles were expelled after sampling, this may have reduced the calculated porewater contents of CH₄ and CO₂.

Samples could not be run immediately because the time involved in retuning the GC normal flux measurement mode to lower sensitivity in order to run porewater samples was prohibitive. We stored the porewater samples in a refrigerator at ~4 °C for up to 4 days in order to run all weekly porewater samples once per week. We do not know what error may have accrued from the storage, but microbial respiration may have boosted CO₂ levels and decreased CH₄ levels.

10.2 Quality Assessment

10.2.1 Data Validation by Source

None given.

10.2.2 Confidence Level/Accuracy Judgment

The degree to which CH₄ and CO₂ concentrations may have changed during sample storage is unknown and likely variable. Nevertheless, the existence of good correlations between CH₄ flux and concentration data suggest that these errors are likely small and do not dominate real patterns of gas concentrations in time or space.

10.2.3 Measurement Error for Parameters

None given.

10.2.4 Additional Quality Assessments

None given.

10.2.5 Data Verification by Data Center

Data were examined for general consistency and clarity.

11. Notes

11.1 Limitations of the Data

See Section 10.1.

11.2 Known Problems with the Data

See Section 10.1.

11.3 Usage Guidance

None given.

11.4 Other Relevant Information

None given.

12. Application of the Data Set

Several avenues are being pursued in publications now being produced to answer the following questions. For this data set, we are most interested in how and why CH₄ concentration profiles and flux measurements vary through time and across the landscape.

13. Future Modifications and Plans

None.

14. Software

14.1 Software Description

We used only commercially available software, mostly Quattro Pro spreadsheet and the Statistical Analysis System (SAS).

14.2 Software Access

None given.

15. Data Access

The CO₂ and CH₄ concentration data are available from the Earth Observing System Data and Information System (EOSDIS) Oak Ridge National Laboratory (ORNL) Distributed Active Archive Center (DAAC).

15.1 Contact Information

For BOREAS data and documentation please contact:

ORNL DAAC User Services
Oak Ridge National Laboratory
P.O. Box 2008 MS-6407
Oak Ridge, TN 37831-6407
Phone: (423) 241-3952
Fax: (423) 574-4665
E-mail: ornl daac@ornl.gov or ornl@eos.nasa.gov

15.2 Data Center Identification

Earth Observing System Data and Information System (EOSDIS) Oak Ridge National Laboratory (ORNL) Distributed Active Archive Center (DAAC) for Biogeochemical Dynamics
<http://www-eosdis.ornl.gov/>.

15.3 Procedures for Obtaining Data

Users may obtain data directly through the ORNL DAAC online search and order system [<http://www-eosdis.ornl.gov/>] and the anonymous FTP site [<ftp://www-eosdis.ornl.gov/data/>] or by contacting User Services by electronic mail, telephone, fax, letter, or personal visit using the contact information in Section 15.1.

15.4 Data Center Status/Plans

The ORNL DAAC is the primary source for BOREAS field measurement, image, GIS, and hardcopy data products. The BOREAS CD-ROM and data referenced or listed in inventories on the CD-ROM are available from the ORNL DAAC.

16. Output Products and Availability

16.1 Tape Products

None.

16.2 Film Products

None.

16.3 Other Products

These data are available from the BOREAS CD-ROM series.

17. References

17.1 Platform/Sensor/Instrument/Data Processing Documentation

None given.

17.2 Journal Articles and Study Reports

Burton, D.L. and E.G. Beauchamp. 1994. Profile nitrous oxide and carbon dioxide concentrations in a soil subject to freezing. *Soil Science Society of America Journal* 58(1):115-22.

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17.3 Archive/DBMS Usage Documentation

None.

18. Glossary of Terms

None.

19. List of Acronyms

ASCII	- American Standard Code for Information Interchange
BOREAS	- BOReal Ecosystem-Atmosphere Study
BORIS	- BOREAS Information System
CD-ROM	- Compact Disk-Read-Only Memory
DAAC	- Distributed Active Archive Center
EOS	- Earth Observing System
EOSDIS	- EOS Data and Information System
GIS	- Geographic Information System
GSFC	- Goddard Space Flight Center
HTML	- HyperText Markup Language
NAD83	- North American Datum of 1983
NASA	- National Aeronautics and Space Administration
NEE	- Net Ecosystem Exchange
NPP	- Net Primary Productivity
NSA	- Northern Study Area
ORNL	- Oak Ridge National Laboratory
PANP	- Prince Albert National Park
PBR	- Productivity/Biomass Ratio
SAS	- Statistical Analysis System
SSA	- Southern Study Area
TE	- Terrestrial Ecology
TF	- Tower Flux
TGB	- Trace Gas Biogeochemistry
URL	- Uniform Resource Locator

20. Document Information

20.1 Document Revision Date

Written: 29-Jan-1997

Last Updated: 06-Aug-1999

20.2 Document Review Date(s)

BORIS Review: 06-Oct-1998

Science Review:

20.3 Document ID

20.4 Citation

When using these data, please include the following acknowledgment as well as citations of relevant papers in Section 17.2:

Valentine, D.W. 1996. Influence of substrate characteristics and other environmental factors on methane emissions from the BOREAS Southern Study Area fen site. II. Profiles of gas concentration, temperature, and pH in the surface peat.

If using data from the BOREAS CD-ROM series, also reference the data as:

Valentine, D., "Influence of Substrate Characteristics and Other Environmental Factors on Methane Emissions from the BOREAS Southern Study Area Fen Site." In *Collected Data of The Boreal Ecosystem-Atmosphere Study*. Eds. J. Newcomer, D. Landis, S. Conrad, S. Curd, K. Huemmrich, D. Knapp, A. Morrell, J. Nickeson, A. Papagno, D. Rinker, R. Strub, T. Twine, F. Hall, and P. Sellers. CD-ROM. NASA, 2000.

Also, cite the BOREAS CD-ROM set as:

Newcomer, J., D. Landis, S. Conrad, S. Curd, K. Huemmrich, D. Knapp, A. Morrell, J. Nickeson, A. Papagno, D. Rinker, R. Strub, T. Twine, F. Hall, and P. Sellers, eds. *Collected Data of The Boreal Ecosystem-Atmosphere Study*. NASA. CD-ROM. NASA, 2000.

20.5 Document Curator

20.6 Document URL

REPORT DOCUMENTATION PAGEForm Approved
OMB No. 0704-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

1. AGENCY USE ONLY (Leave blank)		2. REPORT DATE November 2000	3. REPORT TYPE AND DATES COVERED Technical Memorandum	
4. TITLE AND SUBTITLE Technical Report Series on the Boreal Ecosystem-Atmosphere Study (BOREAS) BOREAS TF-11 CO ₂ and CH ₄ Concentration Data from the SSA-Fen			5. FUNDING NUMBERS 923 RTOP: 923-462-33-01	
6. AUTHOR(S) David W. Valentine Forrest G. Hall and Sara Conrad, Editors				
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS (ES) Goddard Space Flight Center Greenbelt, Maryland 20771			8. PERFORMING ORGANIZATION REPORT NUMBER 2000-03136-0	
9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS (ES) National Aeronautics and Space Administration Washington, DC 20546-0001			10. SPONSORING / MONITORING AGENCY REPORT NUMBER TM—2000—209891 Vol. 211	
11. SUPPLEMENTARY NOTES D.W. Valentine: University of Alaska, Fairbanks; Sara Conrad: DynCorp				
12a. DISTRIBUTION / AVAILABILITY STATEMENT Unclassified—Unlimited Subject Category: 43 Report available from the NASA Center for AeroSpace Information, 7121 Standard Drive, Hanover, MD 21076-1320. (301) 621-0390.			12b. DISTRIBUTION CODE	
13. ABSTRACT (Maximum 200 words) <p>The BOREAS TF-11 team collected several data sets in its efforts to fully describe the flux and site characteristics at the SSA-Fen site. This data set contains temperature, pH, and concentration profiles of methane and carbon dioxide within the surface 50 cm of peat. The measurements were conducted as part of a 2 x 2 factorial experiment in which we added carbon (300 g/m² as wheat straw) and nitrogen (6 g/m² as urea) to four replicate locations in the vicinity of the TF-11 tower. The data set covers the period from the first week of June 1994 through the second week of September 1994. The data are stored in tabular ASCII files.</p>				
14. SUBJECT TERMS BOREAS, tower flux.			15. NUMBER OF PAGES 16	
			16. PRICE CODE	
17. SECURITY CLASSIFICATION OF REPORT Unclassified	18. SECURITY CLASSIFICATION OF THIS PAGE Unclassified	19. SECURITY CLASSIFICATION OF ABSTRACT Unclassified	20. LIMITATION OF ABSTRACT UL	

