# Uniform format surface fCO<sub>2</sub> database Benjamin Pfeil and Are Olsen Bjerknes Centre for Climate Research, Bergen, Norway

### Introduction

Over the last few decades several million measurements of the surface ocean  $CO_2$  concentration have been made, in particular following the advent of infrared based systems which determines the  $CO_2$  concentration in an air headspace in equilibrium with a continuous stream of sea water. The concentration can be expressed as the mole fraction of  $CO_2$  in the headspace ( $xCO_2$ ), the  $CO_2$  partial pressure ( $pCO_2$ ), and the fugacity of  $CO_2$  ( $fCO_2$ ) in the headspace, which takes into account the non-ideal behavior of  $CO_2$  gas. It is this latter which should be used for gas exchange calculations. Conversion between these can be carried out using a set of standard procedures (DOE, 1984, Dickson et al., 2007).

Unfortunately investigators have reported data differently. Some have reported  $xCO_2$ , some  $pCO_2$ , and some  $fCO_2$  of the surface seawater. Given this, and given that the format of the files have varied it has always been a time consuming task to handle and use data from publicly available data repositories like CDIAC (Carbon Dioxide Information Analysis Center) for instance.

To alleviate this situation a uniform format global surface ocean  $fCO_2$  data set has been developed, as encouraged by both SOLAS (at the International CO<sub>2</sub> Conference, Boulder 2005) and IOCCP (workshop on "Ocean Surface pCO<sub>2</sub> Data Integration and Database Development", Tsukuba 2004). This document briefly describes the work that has been done.

#### Equations

To ensure consistency we decided to recompute  $fCO_2$  data whenever possible using a set of standard equations.

Calculations were carried out done according to *Recommendation for autonomous* underway  $pCO_2$  measuring systems and data reduction routines by Pierrot et al, 2008, which follows the DOE handbook (DOE, 1994). Unless otherwise specified, reported  $xCO_2$  data were assumed to be dry mole fractions standardized by each investigator with respect to calibration gas runs. Calculation of  $CO_2$  partial pressures from these data follows:

$$(pCO_2)_{equT}^{wet} = (xCO_2)_{equT}^{dry}(P_{equ} - pH_2O)$$
(1)

where  $(xCO_2)_{equT}^{dry}$  is the CO<sub>2</sub> mole fraction and  $pH_2$ Ois the water vapor pressure at equilibrator temperature.

Water vapor pressure is calculated according to Weiss and Price (1980):

$$pH_2O = \exp(24.4543 - 67.4509(100/T) - 4.8489\ln(T/100) - 0.000544S)$$
(2)

The correction for difference in intake and equilibrator temperatures was carried out using the emperical relationship derived by Takahahsi et al (1993)

$$(pCO_2)_{SST}^{wet} = (CO_2)_{equT}^{wet} \exp\left\{0.0423(SST - equT)\right\}$$
(3)

where SST is the sea surface temperature in the same units as equT. If only fCO<sub>2</sub> at equilibrator temperature was provided, the conversion to in situ temperature was carried out on this.

Although several approaches are available (Copin- Montegut, 1988; Goyet et al., 1993; Takahashi et al., 1993; Weiss et al., 1982), the one of Takahashi et al. (1993) was preferred as it does not require knowledge of the alkalinity and TCO<sub>2</sub> of the waters and was determined for isochemical conditions, while the others were not.

The conversion of pCO<sub>2</sub> to fCO<sub>2</sub> values is carried out according to:

$$(fCO_2)_{SST}^{wet} = (pCO_2)_{SST}^{wet} \exp\left\{\frac{\left|B(CO_2, SST) + 2\left(1 - \left(x_{CO_2}\right)_{equT}^{wet}\right)^2 \delta(CO_2, SST)\right| P_{equ}}{R \times SST}\right\}$$
(4)

where and  $P_{equ}$  is the and pressure (atm) of the equilibrator, and SST is the sea surface temperature (K). R = 82.0578 cm<sup>3</sup> atm mol<sup>-1</sup> K<sup>-1</sup>, and B(CO<sub>2</sub>,T) and  $\delta$ (CO<sub>2</sub>,T) are the virial coefficients for CO<sub>2</sub> (Weiss, 1974).

 $B(CO_2,T)$  in cm<sup>3</sup> mol<sup>-1</sup> is given by:

$$B(CO_2, T) = 1636.75 + 12.0408 T - 3.27957 10^{-2} T^2 + 3.16528 10^{-5} T^3$$
(5)

and  $\delta(CO_2,T)$  in cm<sup>3</sup> mol<sup>-1</sup> by:

$$\delta(CO_2, T) = 57.7 - 0188 T \tag{6}$$

#### Implementation

The sea surface  $CO_2$  concentration data in the files were reported in 11 different ways, and the large majority of the files contained data expressed in at least two different manners (e.g.  $xCO_2$  and  $fCO_2$ ).

Ideally we would like to have always computed or recomputed fCO<sub>2</sub> values from dry mole fractions along with reported equilibrator and intake temperatures, equilibrator pressure, and surface salinity using the set of equations given above. However, on many occasions not all of the required data were reported in the data files, and this necessitated the use of different starting points for our calculations and/or the use of data from external sources. In particular, atmospheric pressure and/or salinity data were sometimes missing. When pressure was missing we used 6 hourly sea level pressure data from the NCEP/NCAR reanalysis project (Kalnay et al., 1996). When salinity was not reported we used climatological monthly mean salinity data from the World Ocean Atlas 2005 (Antonov et al., 2005).

All in all this means that  $fCO_2$  has been recomputed from different starting points and with different ancillary parameters. The different recomputed  $fCO_2$  parameters are given in Table 1. Most times it was possible to recompute  $fCO_2$  from different starting points in the same file, and we therefore provide the  $fCO_2$  recommended value in each file ( $fCO2\_rec$ ). This is the recomputed  $fCO_2$  value we recommend is used. If two or more recomputed fCO<sub>2</sub> parameters is present, we recommend that the one calculated from closest to dry  $xCO_2$  values is used, and this is fCO<sub>2</sub>\_rec in the data file. The order of preference is given in Table 1 as well. That is, if (1) was possible this was used as fCO2\_rec, if (1) was not possible but (2) was, then (2) was used. If neither (1) nor (2) was possible but (3) was then this was used and so on. The philosophy behind this scheme was to (a) start out as close to dry  $xCO_2$  values as possible and (b) to limit use of external data to those cases were absolutely required (i.e. when no in situ fCO<sub>2</sub> data could be obtained without resorting to WOA salinities or NCEP/NCAR pressures.). This also means for instance, that if fCO<sub>2</sub> data were provided, but no  $xCO_2$  or pCO<sub>2</sub>, like from for instance a CARIOCA bouy, the fCO<sub>2</sub> values were retained and are the fCO2\_rec data in the file.

Finally, if either atmospheric pressure or NCEP/NCAR were used, 3 hPa were added to account for the overpressure normally maintained in ships (Takahashi and Sutherland, 2007)

Table 1: reported used for the recalculations in order of preference			
CO <sub>2</sub> parameter	Number	required	
	of cases	extra var.	
fCO2_insitu_from_xCO2_water_eq	2250925		
ui_temp_dry_ppm			
fCO2_insitu_from_xCO2_water_sst	204951		
_dry_ppm			
fCO2_from_pCO2_water_water_eq	724892		
ui_temp			
fCO2_from_pCO2_water_sst_100hu	334085		
midity_uatm			
fCO2_insitu_from_fCO2_water_equ	155969		
i_uatm			
fCO2_insitu_from_fCO2_water_sst	1978648		
_100humidty_uatm			
fCO2_from_pCO2_water_water_eq	26606	Pressure	
ui_temp_ncep			
fCO2_from_pCO2_water_sst_100hu	1263959	Pressure	
midity_uatm_ncep			
fCO2_insitu_from_xCO2_water_eq	2281	Salinity	
ui_temp_dry_ppm_woa			
fCO2_insitu_from_xCO2_water_sst	2876	Salinity	
_dry_ppm_woa			
fCO2_insitu_from_xCO2_water_eq	164	Pressure	
ui_temp_dry_ppm_ncep			
	CO <sub>2</sub> parameter fCO2_insitu_from_xCO2_water_eq ui_temp_dry_ppm fCO2_insitu_from_xCO2_water_sst _dry_ppm fCO2_from_pCO2_water_water_eq ui_temp fCO2_from_pCO2_water_sst_100hu midity_uatm fCO2_insitu_from_fCO2_water_equ i_uatm fCO2_insitu_from_fCO2_water_sst _100humidty_uatm fCO2_from_pCO2_water_water_eq ui_temp_ncep fCO2_from_pCO2_water_sst_100hu midity_uatm_ncep fCO2_insitu_from_xCO2_water_eq ui_temp_dry_ppm_woa fCO2_insitu_from_xCO2_water_sst _dry_ppm_woa fCO2_insitu_from_xCO2_water_eq	CO2 parameterNumber of casesfCO2_insitu_from_xCO2_water_eq ui_temp_dry_ppm2250925fCO2_insitu_from_xCO2_water_sst _dry_ppm204951fCO2_from_pCO2_water_water_eq ui_temp724892fCO2_from_pCO2_water_sst_100hu midity_uatm334085fCO2_insitu_from_fCO2_water_equ i_uatm155969fCO2_insitu_from_fCO2_water_sst 100humidty_uatm1978648fCO2_from_pCO2_water_water_eq ui_temp_ncep26606fCO2_from_pCO2_water_sst_100hu fCO2_from_pCO2_water_sst_100hu i_temp_ncep1263959fCO2_insitu_from_xCO2_water_eq ui_temp_dry_ppm_woa2281fCO2_insitu_from_xCO2_water_sst fCO2_insitu_from_xCO2_water_sst_28762876dry_ppm_woa fCO2_insitu_from_xCO2_water_eq164	

Table 1: reported used for the recalculations in order of preference

12	fCO2_insitu_from_xCO2_water_sst	5860	Pressure
	_dry_ppm_ncep		
13	fCO2_insitu_from_xCO2_water_eq	776	Pressure,
	ui_temp_dry_ppm_ncep_woa		Salinity
14	fCO2_insitu_from_xCO2_water_sst		Pressure,
	_dry_ppm_ncep_woa		Salinity

<sup>b</sup>Number is also used within the data file for identifying which reported CO<sub>2</sub> variable is used as the recommended one (fCO2\_rec).

## **Reported data**

All scripts and in- and output data have been made available along with this report. Transparency is essential for assuring the best quality data product and we encourage all to evaluate our calculations to identify errors, which may occur.

The data file contains all of the reported data, the NCEP/NCAR pressures and the WOA salinities. In addition bottom depth from ETOPO2 (<u>http://www.ngdc.noaa.gov/mgg/global/global.html</u>) has been included for identification of shelf and coastal data.

The file also contains an identifier which shows what input parameter was used. It should also be evident whether the original data were provided at equilibrator or intake temperature.

Abbreviation	Description	unit
cruise	Cruise name (the file name)	
sta	internal station number	
mon	month	
day	day	
yr	year	
hh	hour	
mm	minute	
long	longitude (0-360)	deg
lat	latitude (-90-90)	deg

The following parameters are reported in the output file:

bottomD	bottom depth as reported in file (can be deleted)	m
depth	intake depth used if no intake depth was reported	m
depthW	water intake depth as reported	m
temp	sea surface temperature	deg C
sal	sea surface salinity	PSU
XCO2_water_sst_	xCO <sub>2</sub> water at sea surface temperature in wet air	ppm
wet_ppm		
XCO2_water_equi	xCO <sub>2</sub> water at equilibrator temperature in wet air	ppm
_temp_wet_ppm		
XCO2_water_sst_	xCO <sub>2</sub> water at sea surface temperature in dry air	ppm
dry_ppm		
XCO2_water_equi	xCO <sub>2</sub> water at equilibrator temperature in dry air	ppm
_temp_dry_ppm		
fCO2_water_sst_1	fCO <sub>2</sub> water at sea surface temperature in wet air	µatm
00humidity_uatm	(100 % humidity)	
fCO2_water_corr2	fCO <sub>2</sub> water corrected to 25 deg C sea surface	µatm
5_uatm	temperature in wet air	
fCO2_water_corr_	fCO <sub>2</sub> water corrected to 20 deg C sea surface	µatm
to_20	temperature in wet air	
fCO2_water_equi_	fCO <sub>2</sub> water at equilibrator temperature in wet air	µatm
atm		
pCO2_water_sst_1	pCO <sub>2</sub> water at sea surface temperature in wet air	µatm
00humidity_atm	(100 % humidity)	
pCO2_water_equi	pCO <sub>2</sub> water at equilibrator temperature in wet air	µatm
_temp		
pCO2_theta_SW_c	pCO <sub>2</sub> water at sea surface temperature in wet air	µatm
orrected_to_sst		
Temperature_equi	temperature at equilibration	deg C
Pressure_atm	atmospheric pressure as reported	hPa
Pressure_equi	pressure in the equilibrator as reported	hPa
wind_direc_deg	wind direction as reported	deg
wind_speed	wind speed as reported	m/s
ship_speed	ship speed	knot

ship_direc	ship heading direction	deg
Humidity	humidity	%
woa_sss	salinity extracted from WOA 2005	PSU
woa_land_marker	0 sea	
	1 land	
ncep_slp	atmospheric pressure extracted from NCEP/NCAR	hPa
	6 hourly data	
speed_calc_knots	Ship speed calculated	knot
ETOPO2	Bottom depth extracted from ETOPO2	m
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at equilibrator	µatm
_xCO2_water_equ i_temp_dry_ppm	temperature in dry air; salinity and either	
	atmospheric pressure or pressure at equilibration	
	provided in the file	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at sea surface	µatm
_xCO2_water_sst_ dry ppm	temperature in dry air; salinity and either	
	atmospheric pressure or pressure at equilibration	
	provided in the file	
fCO2_from_pCO2	fCO <sub>2</sub> recomputed from pCO <sub>2</sub> water at equilibrator	µatm
_water_water_equi temp	temperature in wet air; salinity and either	
t	atmospheric pressure or pressure at equilibration	
	provided in the file	
fCO2_from_pCO2	fCO <sub>2</sub> recomputed from pCO <sub>2</sub> water at sea surface	µatm
_water_sst_100hu midity uatm	temperature in wet air (100 % humidity); salinity	
	and either atmospheric pressure or pressure at	
	equilibration provided in the file	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from fCO <sub>2</sub> water at equilibrator	µatm
_fCO2_water_equi uatm	temperature in wet air; salinity and either	
	atmospheric pressure or pressure at equilibration	
	provided in the file	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from fCO <sub>2</sub> water at sea surface	µatm
_fCO2_water_sst_ 100humidty uatm	temperature in wet air (100 % humidity); salinity	
uuuii	and either atmospheric pressure or pressure at	
	equilibration provided in the file	

fCO2_from_pCO2	fCO <sub>2</sub> recomputed from pCO <sub>2</sub> water at equilibrator	µatm
_water_water_equi _temp_ncep	temperature in wet air, NCEP pressure used	
fCO2_from_pCO2	fCO <sub>2</sub> recomputed from pCO <sub>2</sub> water at sea surface	µatm
_water_sst_100hu midity_uatm_ncep	temperature in wet air (100 % humidity), NCEP	
	pressure used	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at equilibrator	µatm
_xCO2_water_equ i_temp_dry_ppm_ woa	temperature in dry air, salinity from WOA used	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at sea surface	µatm
_xCO2_water_sst_ dry_ppm_woa	temperature in dry air, salinity from WOA used	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at equilibrator	μatm
_xCO2_water_equ i_temp_dry_ppm_	temperature in dry air, NCEP pressure used	
ncep		
fCO2_insitu_from	$fCO_2$ recomputed from $xCO_2$ water at sea surface	µatm
_xCO2_water_sst_ dry_ppm_ncep	temperature in dry air, NCEP pressure used	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at equilibrator	µatm
_xCO2_water_equ i temp dry ppm	temperature in dry air, NCEP pressure and salinity	
ncep_woa	from WOA used	
fCO2_insitu_from	fCO <sub>2</sub> recomputed from xCO <sub>2</sub> water at sea surface	µatm
_xCO2_water_sst_ dry_ppm_ncep_wo	temperature in dry air, NCEP pressure and salinity	
a	from WOA used	
fCO2_rec	recommended fCO <sub>2</sub>	µatm
fCO2 source	Identifies which reported CO <sub>2</sub> value was used for	
_	calculations (see Table 1 for details)	

#### References

- Antonov, J.I., Locarnini, R.A., Boyer, T.P., Mishonov, A.V., Garcia, H.E., 2006.
  World Ocean Atlas 2005, Volume 2: Salinity. Levitus, S., ed, NOAA Atlas
  NESDIS 62, U.S. Government Printing Office, Washington, D.C., 182 pp.
- Copin-Montegut, C., 1988. A new formula for the effect of temperature on the partial pressure of CO<sub>2</sub> in seawater. Marine Chemistry 25 (1), 29-37.
- DOE, 1994. Handbook of methods for the analysis of the various parameters of the carbon dioxide system in sea water. Version 2, Dickson, A.G., Goyet, C., eds, ORNL/CDIAC-74.
- Dickson, A.G., Sabine, C.L., Christian, J.R, 2007. Guide to best practices for ocean CO<sub>2</sub> measurements. PICES Special Publication 3,191pp.
- Goyet, C., Millero, F.J., Poisson, A., Shafer, D.K., 1993. Temperature dependence of CO<sub>2</sub> fugacity in seawater. Marine Chemistry 44 (2-4), 205-219.
- Kalnay, E. et al, 1996. The NCEP/NCAR 40-year reanalysis project. Bulletin of American Meteorological Society 77, 437-470.
- Pierrot, D., Neill, C., Sullivan, K., Castle, R., Wanninkhof, R., Lüger, H., Johannessen, T., Olsen, A., Feely R.A., Cosca, C.E., 2008. Recommendations for Autonomous Underway pCO2 Measuring Systems and Data Reduction Routines. Deep-Sea Research Part II. To be published.
- Takahashi, T., Olafsson, J., Goddard, J.G., Chipman, D.W., Sutherland, S.C., 1993.
  Seasonal-Variation of CO<sub>2</sub> and Nutrients in the High-Latitude Surface Oceans
   a Comparative-Study. Global Biogeochemical Cycles 7 (4), 843-878.
- Weiss, R.F., 1974. Carbon dioxide in water and seawater: the solubility of a non-ideal gas. Marine Chemistry 2, 203-215.
- Weiss, R.F., Jahnke, R.A., Keeling, C.D., 1982. Seasonal effects of temperature and salinity on the partial pressure of CO<sub>2</sub> in seawater (Pacific Ocean). Nature 300 (5892), 511-513.
- Weiss, R.F., Price, B.A., 1980. Nitrous oxide solubility in water and seawater. Marine Chemistry 8, 347–359.