

Canadian National Committee for SCOR Comité national canadien pour SCOR

Scientific Committee on Oceanic Research

CANADIAN OCEAN SCIENCE NEWSLETTER LE BULLETIN CANADIEN DES SCIENCES DE L'OCÉAN

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Centre for Earth Observation Science (CEOS), Faculty of Environment, Earth, and Resources, The University of Manitoba

The Centre for Earth Observation Science (CEOS), Clayton H. Riddell Faculty of Environment, Earth, and Resources, The University of Manitoba, is seeking qualified candidates for several full time research and student positions, focusing on investigations of oceanic and atmospheric forcing of sea ice in the northern hemisphere. These positions will become part of a collaborative team working with Dr. David G. Barber through his Canada Research Chair (<u>http://www.chairs.gc.ca/web/home_e.asp</u>), the International Polar year (<u>www.ipy-cfl.ca</u>) and ArcticNet (a Network of Centres of Excellence; <u>www.arcticnet.ulaval.ca</u>) Field work is supported by the Canadian Research Icebreaker, Amundsen (<u>http://www.amundsen.quebec-ocean.ulaval.ca</u>). We seek several candidates to fill the following positions:

1) A full time (contingent term) appointment at a 'Research Associate' level to work in climate forcing of sea ice dynamic and thermodynamic processes. Candidates with a background in remote sensing, numerical modelling field observations, and or statistical approaches to understanding sea ice processes are encouraged to apply. A Ph.D. or masters in meteorology, oceanography, physical geography or related field is desirable.

2) A full time (contingent term) appointment at a 'Research Associate' level to work on oceansea ice-atmosphere (OSA) processes relating to how changes in sea ice dynamic and thermodynamic processes affect biogeochemical, ecological, and climate processes operating across the OSA. Candidates with a background in Arctic marine ecology, remote sensing, numerical modelling, field observations and or statistical approaches to understanding sea ice processes are encouraged to apply. A Ph.D. or masters in biology, meteorology, oceanography, physical geography or related field is desirable.

3) A three-year term position (with possibility of extension) as a data manager for the Centre for Earth Observation Science. Duties will include coordination, management, and expediting of data warehousing from various Arctic research programs. The candidate will coordinate management of data through a computerized data warehousing system known as the Canadian Cryospheric Information Network (CCIN). The candidate should have a bachelors or masters degree in physical or computer science and experience with computer database programs (e.g., MySQL and or Unix based systems).

4) Graduate studentships (Ph.D. or masters level) to work on atmospheric forcing of sea ice dynamic processes at the local or hemispheric scale using in situ data from ocean buoys and ship based observations. A background in remote sensing, meteorology, oceanography, Geographic Information Systems, limnology or related field is desirable.

5) Graduate studentships (Ph.D. or masters level) to work on Freshwater-marine coupling in Hudson Bay. These projects will examine the role of freshwater in ocean surface mixed layer processes (both physical and biological) and the role freshwater plumes play in moderating the ocean surface under sea ice in winter. A background in remote sensing, meteorology, oceanography, limnology, Geographic Information Systems or related field is desirable. Candidates should send a CV and letter of intent via email to Prof. David Barber, c/o Ms. Denise Whynot (<u>dwhynot@cc.umanitoba.ca</u>). Pay and benefits are competitive internationally and commensurate with qualifications. The University of Manitoba is an equal opportunity employer. Reviews will begin April 1 and continue until filled. <u>www.umanitoba.ca/ceos</u>

Improved seawater thermodynamics: How should the proposed change in salinity be implemented?

by SCOR/IAPSO Working Group 127, February 2008

Background

The SCOR/IAPSO Working Group 127 on the "Equation of State and Thermodynamics of Seawater" is charged with providing improved algorithms and descriptions of the thermodynamic properties of seawater. The working group has made significant progress on many of its goals, and it is now time to seek the advice of the oceanographic community regarding the best practical ways of adopting these developments into oceanographic practice. The Working Group has met twice to date, once in Warnemünde in 2006, then in Reggio Calabria in 2007. Our next meeting is in Berlin in September 2008.

The working group will soon provide the most accurate algorithms to date for the thermodynamic properties of seawater (such as density, entropy, enthalpy, specific heat capacity, etc). In order to achieve such accuracy it became evident that a salinity variable is required that more accurately represents absolute salinity than does the conductivity-based Practical Salinity. Spatial variations in the composition of seawater upsets the relationship between Practical Salinity *S* (which is a function of conductivity, temperature and pressure) and Absolute Salinity S_A (defined as the mass of dissolved material per mass of seawater solution). If the thermodynamic properties of seawater are to be written in terms of just one type of salinity, then they are much closer to being functions of (S_A, t, p) than being functions of (S,t,p). Moreover, Absolute Salinity is a conservative property (that is, it is conserved when turbulent mixing occurs) whereas Practical Salinity is not conservative.

Absolute salinity for seawater of Reference Composition

In order to progress toward evaluating Absolute Salinity our first task was to define the relative concentrations of the constituents of Standard Seawater. This we have done, and this work is published in Millero et al (2008a). The abstract of this paper is as follows.

Fundamental determinations of the physical properties of seawater have previously been made for Atlantic surface waters, referred to as "Standard Seawater". In this paper a Reference Composition consisting of the major components of Atlantic surface seawater is determined using these earlier analytical measurements. The stoichiometry of sea salt introduced here is thus based on the most accurate prior determination of the composition, adjusted to achieve charge balance and making use of the 2005 atomic weights. Reference Seawater is defined as any seawater that has the Reference Composition and a new Reference-Composition Salinity S_R is defined

to provide the best available estimate of the Absolute Salinity of both Reference Seawater and the Standard Seawater that was used in the measurements of the physical properties. From a practical point of view, the value of S_R can be related to the Practical Salinity S by

 $S_{\rm R} = (35.165\ 04\ /\ 35)\ {\rm g\ kg^{-1}} \times S$

Reference Seawater that has been "normalized" to a Practical Salinity of 35 has a Reference-Composition Salinity of exactly $S_R = 35.16504 \text{ g kg}^{-1}$.

The new independent salinity variable S_{R} is intended to be used as the concentration variable for future thermodynamic functions of seawater, as an SI-based extension of Practical Salinity, as a reference for natural seawater composition anomalies, as the currently best estimate for Absolute Salinity of IAPSO Standard Seawater, and as a theoretical model for the electrolyte mixture "seawater".

As described in this abstract, *for seawater of standard composition* we have been able to relate the Absolute Salinity to the Practical Salinity; for example, at a Practical Salinity of 35, seawater of Reference Composition has an Absolute Salinity of $35.165\ 04\ g\ kg^{-1}$. We expect shortly to be able to recommend an algorithm that accounts for the variation of seawater composition from the standard composition. That is, we soon expect to be able to recommend an algorithm $S_A = S_A(S_R, , ,)$ where the extra arguments will be either measured parameters (such as total alkalinity, silicate and nitrate) or more simply the spatial locations longitude, latitude and pressure. Millero and Kremling (1976), Millero (2000) and Millero et al (2008b) are precursor papers to such an algorithm.

Advantages of Absolute Salinity over Practical Salinity

Absolute Salinity has the following advantages over Practical Salinity for oceanographic use.

- 1. The definition of Practical Salinity S on the PSS-78 scale is separate from the system of SI units. Absolute Salinity can be expressed in the unit (g kg⁻¹). Adopting this SI unit for salinity would terminate the ongoing controversies in the oceanographic literature about the use of "psu" or "pss" and make research papers more readable to the outside scientific community and consistent with SI.
- 2. The freshwater mass fraction of seawater is not (1 0.001 S). Rather, it is $(1 0.001 S_A / (g kg^{-1}))$, where S_A is the Absolute Salinity, defined as the mass fraction of dissolved material in seawater. The values of $S_A / (g kg^{-1})$ and S are known to differ by about 0.5%. There seems to be no good reason for continuing to ignore this known difference, e.g., in ocean models.
- 3. PSS-78 is limited to the salinity range 2 to 42. For a smooth crossover on one side to pure water, and on the other side to concentrated brines up to saturation, as e.g. encountered in sea ice at very low temperatures, salinities beyond these limits need to be defined. While this poses a challenge for S, it is not an issue for S_A .
- 4. The theoretical Debye-Hückel limiting laws of seawater behavior at low salinities, used for example in the determination of the Gibbs function of seawater, can only be computed from a chemical composition model, which is available for S_R but not for *S*.

- 5. For artificial seawater of Reference Composition, S_R has a fixed relation to Chlorinity, independent of conductivity, salinity, temperature, or pressure.
- 6. The next largest improvement in the equation of state of seawater will come from incorporating variations in the composition of seawater, that is, from calling the equation of state with Absolute Salinity rather than with Reference Salinity. The determination of Absolute Salinity is facilitated by the introduction of the Reference Composition and Reference Salinity.
- 7. Absolute Salinity S_A is a conservative variable, whereas, in the presence of compositional variations, Practical Salinity *S* (which is essentially determined by conductivity alone) is not a conservative variable. All of our oceanographic practice assumes that "salinity" is a conservative variable (e.g. ocean model codes, the practice of mixing along straight lines on salinity-potential temperature diagrams, inverse modelling etc).

Expanding on point 7 above, it seems clear that we presently use Practical Salinity *S* as though it is a conservative variable, and yet we now know that it is not; for a given Absolute Salinity, Practical Salinity varies by up to 0.02 between different major ocean basins (Millero, 2000). This non-conservative regional variation in Practical Salinity is at least seven times the error with which salinity can be measured by modern instrumentation at sea. This difference of 0.02 in Practical Salinity causes differences in density that are also several times greater than the remaining uncertainty in the best algorithms for the density of seawater. It seems that in our oceanographic practice we intuitively ascribe the conservative properties of Absolute Salinity to our "salinity" variable, which to date has been Practical Salinity. For example, if we were intent on interpreting the salinity of an ocean model as Practical Salinity, then the salt conservation equation should contain a non-conservative source term to take account of the spatial variations in the composition of seawater.

Here we summarize the reasons why Absolute Salinity is the preferred salinity variable for oceanographic research.

- It will be preferred by journals since it is an SI unit.
- It is the natural salinity variable for ocean models since they assume that their salinity variable is conservative, hence it should be used to initialize ocean models at all depths.
- It is the natural variable to use in inverse models, budget studies and on salinitytemperature diagrams because its conservative nature justifies turbulent mixing occurring along straight lines on such a diagram.
- The freshwater fraction and the meridional freshwater flux follow naturally when using Absolute Salinity but not when using Practical Salinity.
- By using Absolute Salinity in the algorithm for the equation of state, the effects of the spatial variations of seawater composition are accounted for, while if Practical Salinity is used in such a call to the equation of state, a density error is incurred.
- It is the common salinity variable used in engineering, natural and geosciences outside oceanography, where Practical Salinity is often unknown or misconstrued.
- It is applicable to low concentrations in brackish lagoons and river mouths, to high concentrations in freezing or desiccating brines, as well as at higher temperatures in desalination plants, whereas Practical Salinity is defined only in the range 2 < S < 42.
- If necessary for chemical or biological reasons, all partial ion concentrations in a sample are easily available, to which Practical Salinity is unrelated.

The SCOR/IAPSO Working Group 127 regards these as compelling reasons for adopting Absolute Salinity as the new preferred salinity variable in the analysis of oceanographic data. Accordingly we are formulating new algorithms for density, enthalpy, entropy, potential temperature, sound speed, etc in terms of Absolute Salinity, temperature and pressure (Feistel (2008)). The extended validity range of the new formulas in temperature and salinity precludes using Practical Salinity as the independent variable of these thermodynamic quantities. For example, in situ density will have the functional form $\rho(S_A, t, p)$ and potential temperature will have the functional form $\theta(S_A, t, p, p_r)$. Absolute Salinity S_A will be defined as

$$S_{\rm A} = S_{\rm R} + \delta S_{\rm A}$$

where Reference salinity S_R is simply proportional to Practical Salinity S as described in Millero et al (2008), namely by

$$S_{\rm R} = (35.165\ 04\ /\ 35)\ {\rm g\ kg^{-1}} \times S$$
,

and δS_A is the difference between Absolute and Reference Salinities. δS_A will be available as a look up table as a function of latitude, longitude and pressure and also as an alternative linear relationship of nutrient and silicate concentrations, or for example, as a Calcium excess estimate from the river discharge into estuaries. We expect to have algorithms available before the end of 2008.

How to adopt Absolute Salinity?

Having made the case that Absolute Salinity possesses many advantages over Practical Salinity, how should present oceanographic practice adapt to incorporate these advantages?

The obvious thing to do would be to decide on a date on which the whole community ceases to use Practical Salinity and switches to using Absolute Salinity. However the algorithm to convert Reference Salinity to Absolute Salinity is less mature and will probably remain a "work in progress" for several years. Moreover, data that is stored in archives should have a very close connection to a measurement (like temperature or conductivity) rather than being the result of an algorithm that is likely to change with time. Hence one cannot really imagine storing Absolute Salinity in data bases. Rather, the closest thing to do in this vein is to store Reference Salinity

Storing Reference Salinity in data centres would have the advantage that it is an SI unit. However before the equation of state (or other thermodynamic quantities) can be evaluated using the new software, the Reference Salinity data needs to be converted to Absolute Salinity using the most up-to-date version of this software. Moreover, the community cannot completely abandon Practical Salinity since it will remain as the salinity variable in the archives for cruises undertaken before the change-over date. By changing the salinity variable that is reported from cruises to data bases from Practical Salinity to Reference Salinity the possibility of contamination of the data archives arises as salinity of one type is incorrectly labeled and stored as the other type of salinity. In the long run, as with many other historical non-SI units like torr, cal or dyn, it would seem to be an advantage to use only Reference Salinity and abandon the use of Practical Salinity completely. If Reference Salinity were the salinity variable to be used in all of the revised thermodynamic algorithms, the argument for "biting the bullet" and abandoning Practical Salinity as much as possible would seem to be the correct path. But it is Absolute Salinity that we seek, and Reference Salinity is only part way towards the evaluation of Absolute Salinity. Given this, is it worthwhile changing the present archiving practice in favour of a variable (Reference Salinity) that is still not the final salinity that we will use (Absolute Salinity)?

Any choice of action inherently involves compromises, and the best course of action is not obvious to the Working Group. As a way of focusing the discussion we outline two possible routes for adopting the advantages of Absolute Salinity, labeled Option 1 and Option 2.

Option 1

- Change from reporting Practical Salinity to reporting Reference Salinity to national and international data bases. This implies that the data bases store Practical Salinity from the old cruises and store Reference Salinity from new cruises (from say 1st January 2010).
- Provide software (for example, of the form $S_A(S_R, x, y, p)$) to produce the best available estimate of Absolute Salinity from Reference Salinity (using additional information on position or water properties).
- Have all the thermodynamic software in the form $\rho(S_A, t, p)$.

Discussion of Option 1

The main advantage of Option 1 is that the community eventually ceases to use the non-SI unit Practical Salinity, and instead uses the two SI salinity measures, Reference Salinity and Absolute Salinity.

A drawback of Option 1 is that there will be cases of contamination of the data bases where cruise salinity is labeled and stored as Reference salinity whereas in fact it is Practical Salinity data, and vice versa. This kind of error presently contaminates the temperature, oxygen and pressure/depth data bases.

Since both S and S_{R} are simply measures of conductivity, and since they are simply proportional to each other, will it be seen that we are taking a course of action that has potential for confusion for only academic benefit?

Recall that scientific work and papers are mostly done with potential temperature θ rather than in situ temperature *t* so the first thing that one usually does with the *S*, *t*, *p* data from a data centre is to form θ . Similarly, scientific work and papers should be done with Absolute Salinity rather than Reference Salinity so the first thing that one needs to do under Option 1 with the S_R , *t*, *p* data from a cruise or from a data centre is to form not only θ but also S_A . This analogy with what we already do with storing the measured variable *t* but using the derived variable θ is very close.

Under Option 1 we cannot imagine that the community can altogether forget about Practical Salinity however, as the data from older cruises (e.g. all of WOCE) is stored in data centres in terms of Practical Salinity. This data will need converting first to Reference Salinity

and then to Absolute Salinity before the thermodynamic routines such as potential temperature, density, potential enthalpy etc, can be called by oceanographic researchers.

There will be some instances when the new software is called with the salinity data being *S* and in those instances an error will be made. This type of error is an undesirable consequence of both Options 1 and 2.

Option 1 requires manufacturers (such as Seabird) to change what they presently do. The instruments will need to output their salinity in terms of Reference Salinity. Also the ampoules of standard seawater will need to quote their salinity in terms of Reference Salinity. The transition date of say 1st January 2010 has to be handled very carefully in these respects. Further, anyone wanting to make use of older ampoules will have to be aware of the transition and how to deal with it.

Option 2

- Continue to report Practical Salinity *S* from cruises and to have <u>only</u> Practical Salinity *S* stored at national and international data centres.
- Provide software (for example, of the form $S_A(S, x, y, p)$) to produce the best available estimate of Absolute Salinity from Practical Salinity (using additional information on position or water properties).
- Have all the thermodynamic software in the form $\rho(S_A, t, p)$

Discussion of Option 2

By reporting <u>only</u> S in data bases we would expect to greatly reduce the possibility of salinity data being mislabeled in data bases.

Since both S and S_R are simply measures of conductivity, option 2 is consistent with the argument that there is little value in replacing one measure of conductivity (namely Practical Salinity) with another (namely Reference Salinity) in data bases. Rather, under Option 2 data centres store S and S alone.

As mentioned above, scientific work and papers are mostly done with potential temperature θ rather than in situ temperature *t* so the first thing that one usually does with the *S*, *t*, *p* data from a data centre is to form θ . Similarly, scientific work and papers will be mostly done with Absolute Salinity rather than Practical Salinity so the first thing that one needs to do under Option 2 with the *S*, *t*, *p* data from a cruise or from a data centre is to form not only θ but also S_A . This analogy with what we already do with storing the measured variable *t* but using the derived variable θ suggests that storing *S* but using S_A will not cause oceanographers any serious difficulties.

There will be some instances when the new software is called with the salinity data being *S* and in those instances an error will be made. This type of error is an undesirable consequence of both Options 1 and 2. However this error will affect the results and the publications arising out of those who make this error, but this error will not contaminate an archived data set.

Option 2 does not require manufacturers (such a Seabird and the Standard Seawater Service) to change what they presently do. Rather, Option 2 puts the responsibility for the changes in the hands of practicing research oceanographers.

Request for your input

The above two options are just two of many options; please do not feel constrained in your comments to these options. We seek input from the oceanographic community on how to gain the advantages of adopting Absolute Salinity in our oceanographic research work. The key issue seems to revolve around which type of salinity is required to be reported to and archived by oceanographic data centres. We encourage frank responses. Each response will be thoughtfully considered by the Working Group. Please email your comments to trevor.mcdougall@csiro.au with the words "Comment for WG127 on how to adopt Absolute Salinity" as the message title.

References

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Membership of SCOR/IAPSO Working Group 127 on

"Thermodynamics and the Equation of State of Seawater"

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Position of SCOR¹ and GESAMP² on Deliberate Nutrient Additions to the Ocean PRESS RELEASE, 4 March 2008

Deliberate fertilization of the ocean, until recently a subject of mostly scientific interest, has caught the attention of the commercial sector because of its potential to sequester carbon and to increase the production of living marine resources. To be effective for either of these purposes, eventual fertilization would add iron or nitrogen to large areas of the world's ocean. Proposals to realize the potential of ocean fertilization on such scales suffer a major weakness: one does not know how the oceanic ecosystem will respond. Current understanding of how the ocean operates is increasing rapidly, but is still not sufficient to predict the effects of large-scale nutrient manipulations.

Field experiments, carried out in various parts of the world ocean to study the role of iron in ocean ecosystems, have not been able to demonstrate a significant net increase in carbon export to the deep ocean on short or long time scales. These experiments have also raised important and, as yet, unanswered questions about changes in community structure. Ocean fertilization on any significant scale will (by design) impact the species succession and the ecosystem structure and function in the affected areas. Furthermore, the impacts of fertilization are unlikely to be confined to the specific region that receives the fertilizer. Ocean currents mix and move water continuously and so can transport nutrients, the resulting biomass, and decomposition products beyond the target areas, with unknown consequences. Inadvertent anthropogenic additions of nutrients to the coastal ocean are presently causing significant problems such as hypoxia, anoxia and harmful algal blooms. At the present, the long-term consequences of ecosystem alterations from nutrient additions are unforeseeable and may be harmful. The effects of deliberate large-scale nutrient addition may therefore range from the desired and positive to the unintended and negative.

The Scientific Committee on Oceanic Research (SCOR) of the International Council for Science and the Joint Group of Experts on the Scientific Aspects of Marine Environmental Protection (GESAMP) of the United Nations agree that any deliberate large-scale addition of nutrients to the ocean must be conducted in such a way that the outcomes of these experiments are statistically quantified and independently verified with respect to but not limited to:

- Changes in new primary production and total community respiration rates at the fertilization site and "downstream" of the site;
- Assimilative capacity of selected ocean regions;
- Changes in the drawdown of carbon dioxide from the overlying atmosphere, and carbon dioxide and essential macro-nutrients (P, N, and Si) from the surface waters;
- Changes in the production of carbon dioxide and other gases relevant to climate change (e.g., nitrous oxide, methane, and dimethyl sulfide) in surface and mesopelagic waters;
- Changes in denitrification rates within the oxygen minimum zone;
- Changes in the production of toxins that might be detrimental to other organisms, for example, by harmful algal blooms;
- Changes in the export of carbon to a depth where sequestration for at least 100 years is likely;
- Changes in pH and oxygen concentrations in the water column;

- Changes in biomass, composition, and biodiversity of phytoplankton, bacteria, and zooplankton, and recruitment of fish and shellfish; and
- Changes in food web structure.

To be scientifically credible the design and implementation of large-scale nutrient addition experiments must be transparent and the results must be clearly stated and made available to the scientific community and the general public. Transparency is essential, because any appearance of lack of independence from vested interests lowers the credibility of the results among ocean scientists, environmental organizations, policymakers, and potential investors in carbon credits. Carbon credits for fertilization should not be allowed unless and until reliable methods have been developed to estimate and verify the amount of carbon actually sequestered, and side effects have been properly understood and taken into account. We commend efforts by some commercial ventures to create codes of conduct and obtain outside reviews. It is essential that each stage of these experiments is reviewed by well-qualified experts free of vested interests. The goal of any new experiment on the effects of nutrient addition should be to increase our understanding of ocean processes at adequate spatial and temporal resolution; experiments should build on the lessons and the insights of previous experiments.

For further information please contact:

General Questions about the Scientific Committee on Oceanic Research (SCOR) and SCOR's interests in this topic: Prof. Bjorn Sundby, SCOR President (Canada)—Can be reached at +1 514 398- 4883.

General Questions about the Joint Group of Experts on the Scientific Aspects of Marine Environment Protection (GESAMP) and GESAMP's interests in this topic: Dr. Michael E. Huber, Chairman of GESAMP (Australia)— Can be reached at +61 7 3244 7336.

Questions about the effects of iron in ocean ecosystems:

Dr. Ken Buesseler, Senior Scientist, Woods Hole Oceanographic Institution (USA, but on sabbatical in New Zealand) — Can be reached at +64 2 1056 0521 between 9 a.m. and 5 p.m. (New Zealand time).

Questions about iron chemistry in the ocean: Prof. Tim Jickels, School of Environmental Sciences, University of East Anglia (United Kingdom)—Can be reached at +441603 593117.

General questions about GESAMP: Fredrik Haag, GESAMP Officer, International Maritime Organization (United Kingdom), Can be reached at +44 20 7463 4139, or through gesamp@gesamp.org.

This statement contains views expressed or endorsed by members of SCOR and GESAMP who act in their individual capacities; their views may not correspond with those of their sponsoring organizations or Governments.

¹ SCOR is an international nongovernmental organization created in 1957 by the International Council for Science to promote international cooperation in all areas of ocean science (see <u>www.scor-int.org</u>).

 2 GESAMP is an independent group of experts, formed in 1969, that advises the United Nations (UN) system on the scientific aspects of marine environmental protection. It is sponsored by eight UN organizations with responsibilities for the marine environment and provides a mechanism for coordination and collaboration among them (see www.gesamp.org).

Proceedings of IUGG XXIV General Assembly - Perugia 2007 Italy

All abstracts presented at the XXIV IUGG General Assembly, held in Perugia from July 2nd to the 13th, 2007 are now available on line, including those of IAPSO. The abstracts as pdf files can be viewed and/or downloaded from the <u>www.iugg2007perugia.it</u> web site at the following link: <u>http://www.iugg2007perugia.it/webbook/</u>

ICES Annual Science Conference 2008 – Halifax, Nova Scotia

The 2008 ICES Annual Science Conference to be held at the World Trade and Convention Centre in Halifax, Nova Scotia, Canada, Monday September 22 – Friday September 26, 2008. Theme sessions include:

- Incorporating microbial dynamics in studies of shelf ecosystems
- Role of sea ice in polar ecosystems
- Mid-ocean ridges and seamounts: oceanography, ecology and exploitation
- New trends in diseases of marine organisms: causes and effects
- Marine spatial planning in support of integrated management tools, methods, and approaches
- Size is almost everything! Size and trait based processes and models in ecosystems and management
- Sediment biota interactions and mapping marine habitats
- Ecological carrying capacity in shellfish culture
- Fishing capacity, effort and fishing mortality; The understanding of fishery dynamics and their links to management
- Comparative dynamics of populations in the Baltic Sea and Gulf of St. Lawrence ecosystems
- Small-scale and recreational fisheries surveys, assessment, and management
- Coupled physical and biological models: parameterization, validation, and applications
- How much habitat is enough? Evaluating habitats in terms of their ecosystem function, goods and services
- Problems and solutions for the assessment, conservation and restoration of rare, threatened and endangered fish species
- Governmental quality and risk management
- New methodology for tracking fish, mammal and sea bird behavior and migrations

- Evidence of global warming effects on zooplankton populations and communities, including larvae of benthic invertebrates and fish
- Environmental and fisheries data management, access, and integration

Abstracts must be submitted by no later than Monday 21 April 2008. Additional information may be found at: <u>http://www.ices.dk/iceswork/asc/2008/index.asp</u>

Second Symposium on The Ocean in a High-CO₂ World

Registration and abstract submissions is now open for the Second Symposium on The Ocean in a High-CO2 World in Monaco on Oct. 6-9, 2008. The symposium Web site can be found at <u>http://www.highco2world-ii.org</u>.

Deceased Oceanographers

Each year CMOS recognizes in their publications, and with a moment of silence at their annual Congress, the passing of Canadian meteorologists and oceanographers. Should you be aware of the passing of any Canadian oceanographers in 2007, or up to May 2008, it would be very much appreciated if you could pass relevant information to both the Secretary of CNC/SCOR (dick.stoddart@sympatico.ca) and to the Executive Director of CMOS, Ian Rutherford (cmos@cmos.ca)

Call for SCOR Working Group Proposals for 2008

The 29th SCOR General Meeting will take place in Woods Hole, Massachusetts, USA on **22-24 October 2008** and it will evaluate proposals for new SCOR Working Groups at that time. New Working Group proposals will be entertained until **May 31, 2008.** Model proposals and other information about working groups can be found at http://www.scor-int.org/wkgrpinfo.htm.

SCOR examines the disciplinary balance of its working groups annually. The 2007 evaluation of found that the set of SCOR working groups is relatively balanced, given that other organizations are already focusing on climate issues. However, it was recommended that more attention be given to interdisciplinary topics. Each proposal will be evaluated by national SCOR committees in terms of scientific merit and quality, timeliness, and achievability of the proposed terms of reference.

SCOR tentatively plans to start two new working groups in 2009, pending availability of adequate funding. National SCOR committees are an important aspect of SCOR's operation and can play a key role in reviewing working group proposals and in seeking new funds to pay for working group activities. Proponents should consider submitting their proposals through their national SCOR committees, although SCOR will also accept proposals from individuals and other organizations. Details on the call for proposals may be found at: <u>http://www.scor-int.org/2008GM/2008GM.htm</u> (the second entry).

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Previous newsletters may be found on the CNC/SCOR web site. Les bulletins antérieurs se retrouvent sur le site web du CNC/SCOR.

Newsletter #36 will be distributed on May 15, 2008. Please send contributions to <u>dick.stoddart@sympatico.ca</u> Bulletin #36 sera distribué le 15 mai 2008. Veuillez faire parvenir vos contributions à <u>dick.stoddart@sympatico.ca</u>

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