



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
WASHINGTON, D.C. 20460

OCT 8 2010

OFFICE OF
RESEARCH AND DEVELOPMENT

Dr. Gary S. Sayler
Chair, Board of Scientific Counselors
University of Tennessee
Center for Environmental Biotechnology

^{George}
Dear Dr. ~~Sayler~~:

The Office of Research and Development (ORD) would like to take this opportunity to thank you and the members of the Board of Scientific Counselors (BOSC) for the September 2009 progress review of the National Center for Computational Toxicology (NCCT). We especially thank the members of the subcommittee who conducted the review, Drs. George Daston, James Clark, Richard DiGiulio, Ali Faqi, Lawrence Hunter, Muiz Mumtaz, Dennis Paustenbach, John Quackenbush, Santiago Schnell, Cynthia Stokes, and Katrina Waters.

ORD asked the BOSC to provide input on the progress made by ORD's Computational Toxicology Research Program over the past 4.5 years. ORD also requested the BOSC's advice on whether the NCCT should continue as an established organization beyond its 5-year charter. We are pleased that the BOSC was supportive of the NCCT and the direction we are taking in this important research area.

Enclosed with this letter is our response to the comments and recommendations in your April 2010 letter. If further information is needed, please feel free to contact me or Dr. Robert Kavlock (kavlock.robert@epa.gov).

Again, thank you for your advice to ORD.

Sincerely,

A handwritten signature in black ink, appearing to read "Kevin Teichman".

Kevin Teichman, Ph.D.
Deputy Assistant Administrator
for Science

Enclosure

cc: Dr. George Daston
Dr. James Clark
Dr. Richard DiGiulio
Dr. Ali Faqi
Dr. Lawrence Hunter
Dr. Muiz Mumtaz
Dr. Dennis Paustenbach
Dr. John Quackenbush
Dr. Santiago Schnell
Dr. Cynthia Stokes
Dr. Katrina Waters



**Office of Research and Development's Response to the
Board of Scientific Counselors April 2010 Final Report on
ORD's National Center for Computational Toxicology**

September 2010

BOSC Computational Toxicology Subcommittee

George Daston (Chair)

James Clark

Richard DiGiulio

Ali Faqi

Lawrence Hunter

Muiz Mumtaz

Dennis Paustenbach

John Quackenbush

Santiago Schnell

Cynthia Stokes

Katrina Waters

Submitted by:

Robert J. Kavlock

Director, National Center for Computational Toxicology

Office of Research and Development

The following is a narrative response to the comments and recommendations of the BOSC final letter report on the review of ORD's National Center for Computational Toxicology (NCCT), held September 29 and 30, 2009, in Research Triangle Park, NC. The review was conducted by a standing subcommittee of the BOSC. The subcommittee had previously reviewed the NCCT three times--in April 2005, June 2006, and February 2007. ORD responded to those reviews previously. In this fourth review, the BOSC was asked to provide the Computational Toxicology Research Program (CTRP) with advice on (1) the progress the Center has made, in the past 4.5 years, in fulfilling its mission and strategic goals; and (2) whether the NCCT should continue as an established organization beyond its original 5-year charter.

In the BOSC's final letter report of April 15, 2010 to ORD Assistant Administrator, Paul Anastas, they summarized, "During the 4.5 years between its establishment and this review, the CTRP has made substantial progress in establishing and meeting its priorities and goals, collaborating within and outside EPA to leverage the staff's expertise and transforming the field of toxicity testing. Many of the recommendations made by the BOSC during its earlier reviews have been acted on by the CTRP. This includes improved capabilities in bioinformatics through the funding of two external centers and in informatics and systems biology through staff hires; expansion of the CTRP's technical approaches to even more programs within the Agency; and the formation of an extensive collaboration with the National Institute of Environmental Health Sciences (NIEHS) and the National Human Genome Research Institute (NHGRI) for its ToxCast project."

In particular, the BOSC addressed five charge questions that focused on the progress and future of the NCCT. Each charge question is shown below in bold, followed by the BOSC's comments and recommendations in italics (the recommendations have been numbered for ease of reference when addressing them). ORD's response to the comments and recommendations follow. A summary of the BOSC recommendations and ORD's responses is provided in Table 1 at the end of this report.

Charge Question 1: Does the scope and involvement of expertise in the project reflect activities consistent with the function of a Center?

BOSC Comments:

The mission statement of the CTRP is to integrate modern computing and information technologies with molecular biology to provide the Agency with decision support tools for high-throughput risk assessment.

The three initial long-term goals of the CTRP were as follows:

- 1. Risk assessors use improved methods and tools to better understand and describe the linkages of the source-to-outcome paradigm,*
- 2. EPA Program Offices use advanced hazard characterization tools to prioritize and screen chemicals for toxicological evaluation, and*

3. *EPA assessors and regulators use new and improved methods and models based on the latest science for enhanced dose-response assessment and quantitative risk assessment.*

These are ambitious goals for the Program, particularly the last one, the objective of which is to identify non-classical methods for assigning an approximate acute and chronic toxicological hazard rating to hundreds, if not thousands, of chemicals. The vast majority of these ratings are to be conducted without the traditional reliance on animal testing and human epidemiology.

The BOSC members believe that the CTRP has made substantial progress toward meeting the original long-term goals, and that the progress is appropriate given the duration of the Program's existence and the resources involved. Although the goals have not been fully met, the CTRP has made significant advancements to creating and providing the methods, tools, and models that will enable risk assessors, program offices, and regulators to use 21st century science and technology for their work, as indicated in the goals. Notably, a number of new tools already have been released to support decision-making, including ACToR, DSSTox, ToxRefDB, and ExpoCast.

Underpinning all of these, the CTRP has focused on integrating modern computational approaches with molecular and cellular biology and physiology to create new methods for EPA's chemical prioritization and risk assessment efforts. The activities of the CTRP in pursuit of these goals have included the assembly and integration of vast quantities of existing toxicological and toxicogenomics data; creation of new database and data warehousing tools to house the data; development of methods and tools for discerning actionable knowledge from the data; additional data acquisition on chemicals of Agency interest; and the development of various types of computational models to understand biological and toxicological mechanisms and provide predictive tools for hazard evaluation and prioritization and risk assessment. As several of the CTRP's projects have matured, they have started to link together in ways that allow ToxCast and Tox21 data to be effectively captured and managed, and allow the various sources of information to be leveraged against each other in productive ways.

Overall, during its inaugural funding period, the CTRP has built the infrastructure necessary to bring computational tools to risk assessment; assembling the data and building the tools that are needed to collect the high-throughput screening (HTS) data that the Program is now facing. In the process, the Program staff has learned the limitations of the existing data and has begun to explore how it might address the limitations of the existing systems, data, and models that will be necessary to move forward. Issues of bioavailability and bioactivity, correlations between in vitro and in vivo results, and dose response relationship and toxicity are cases in point.

The BOSC concludes that the CTRP has laid a strong foundation and has put forth good ideas to move forward, and that there is clearly a need to continue to build on the success of the Program.

Because of the large number of projects and collaborations involved, it is difficult to assess in detail whether the CTRP has utilized the available resources efficiently. Nonetheless, it is the conclusion of the BOSC that the progress made is substantial and appropriate in light of the time the Program has been in existence, its budget, the number of personnel involved, and the collaborations in place within and outside the Agency.

As expected with such an ambitious program, the CTRP has been faced with a number of challenges as the Program has strived to meet its goals. The following are the BOSC's observations about some of these challenges along with suggestions for meeting them that are relevant to continued progress.

One of the challenges that the CTRP has taken on is the assembly and integration of the vast quantities of existing, available toxicological and toxicogenomics data. Although it might seem that this should be a relatively easy task, much of the data are not electronic, much is poorly annotated, and many ambiguities exist within the available data. The CTRP has initiated a number of data base and data warehousing projects to begin to address these issues, including ACToR, DSSTox, ToxRefDB, and ExpoCast, each of which approaches a different aspect of the problem. Each of these has made significant progress over the past few years, along the way discovering the need to do a significant amount of manual curation, but using existing tools and developing new ones as appropriate, and they are beginning to be linked together in productive ways. Despite all that has been achieved, much of the work is clearly in an early stage, and many of the current resources are more like a structured index to the underlying data rather than a comprehensive linked data resource. These projects need to continue to build on things that are in place, drilling deeper into the data and continuing the problem of structuring, standardizing, and organizing the data so that they can be more easily subjected to comprehensive meta-analyses. In addition, much of what is available is focused on research scientists and the data are presented "as is" without context. The potential problem is that as these data resources are publicly available, the public may access the information and, without context, may misinterpret what is there. For example, the data in ACToR identifies compounds that have been tested for carcinogenicity or genotoxicity, but not those that have been found to be carcinogenic or genotoxic. In this context, it might be worthwhile to obtain some public feedback on how people may interpret the available data.

Database development (sometimes called knowledge warehouse) is the foundation on which much of the CTRP is being built and doing it successfully requires a significant intellectual investment. Fundamentally, a database is a model, and building the model requires two components: understanding the relationship between the data elements and understanding how people will use the data. In many ways, the CTRP is in a critical phase in that, having assembled the data into one place, building the linkages in a systematic fashion now will require understanding the value of data to the community and the manner in which those data will be used.

Much of the success of the CTRP relies on the development of computational models to interpret data and make predictions. A major part of the modeling effort focuses on interrogating the databases. The BOSC noted that a substantial part of these efforts utilizes machine-learning methods. Although the BOSC has no specific objection to such approaches, it should be noted that many biostatisticians are sometimes apprehensive about such methods. Because many of the ultimate customer clients and stakeholders of the CTRP's efforts are expected to have biostatistical backgrounds, the BOSC encourages the Program to consult with biostatisticians early and often to assure they can address any objections; the CTRP also should consider attempting some additional methods.

There also is a need to interact more extensively with the broader scientific user community in the process of developing and rolling out tools and software. One of the driving principles

behind the program is that the resources it produces should be both useful and used. For the resources to be useful, they must address important questions; the CTRP seems to have a strong focus on addressing relevant problems in environmental toxicology and exposure and risk assessment. For the resources to be used, the Program must understand how risk assessors outside the CTRP and within the broader toxicological and toxicogenomics community will use these resources. This process should allow the development of appropriate use cases that can guide how the tools are created and integrated. If this is done in a systematic manner, it has the potential to rapidly advance the evolution of the resources the CTRP is developing. This could be achieved through an annual or biannual conference by bringing together the data generators, the data users, and the risk assessors/managers—the ultimate users of these alternative methods/models.

There were concerns expressed by some BOSC members that associations are not causation and this should be recognized by the EPA management, both at the CTRP level and at the level of the Office of the Administrator. The results of a computer-generated association should be carefully examined through traditional testing and careful scientific study. Some of the BOSC members recalled that the Ames test once was thought to serve in a similar manner as that being proposed for the various tools that are being built by the CTRP. Thus, although the BOSC fully supports providing more resources to CTRP efforts, it offers the precautionary warning that, at best, the results of these efforts will be the temporary placement of a chemical into a bin that could likely initiate a “science forcing event.” In turn, the manufacturer or user of the chemical can be put on notice that this chemical appears to have certain characteristics that give it likelihood for being a hazard and that they will need to conduct further toxicological testing. This will bring the chemical industry on par with the pharmaceutical industry that has been conducting such cost/risk/benefit analyses for several decades.

BOSC Recommendations:

- ✧ 1. Several CTRP projects have undertaken structuring, standardizing, and organizing the data so that they can be more easily subjected to comprehensive meta-analyses. At this point, the CTRP should obtain some public feedback on how people are using and interpreting the available data.
- ✧ 2. Acceptance of products, methods, and databases by the risk assessment community is the key to success. Hence, the NCCT should organize an annual or biannual conference that brings together the data generators, data users, and risk assessors/managers—the ultimate users of these alternative methods/models.
- ✧ 3. As more data from high-throughput assays and computer models become available, the NCCT should provide guidance on how to interpret this information in the context of more traditional testing and scientific examination so that risk assessment practitioners in the EPA program offices can apply these findings.

ORD Response:

- ✓ 1. The CTRP plans to survey the public to determine awareness, interest and usage of CTRP data, tools and research. This is an integral part of the newly drafted Computational Toxicology communications and outreach strategy. A survey has been drafted and is currently being reviewed by the CTRP research team. However, to survey

more than nine non-EPA people, approval from the Office of Management and Budget (OMB) is required. The plan is to work through the OMB approval process and then survey the CTRP database of more than 25,000 people who have expressed an interest in CTRP. According to OMB's approval timeline for surveys, it will take at least a year for approval. CTRP is aiming to release the survey in January 2012 unless the approval process is faster than expected, or the survey can be distributed using a different process.

- ✓ 2. The NCCT continues to host and sponsor the Computational Toxicology Community of Practice meetings and teleconferences that bring together all the groups on a monthly basis. In regard to an annual or biannual conference, we are planning a second ToxCast Data Analysis Summit (TDAS) that will bring together data generators, data users, risk assessors and managers in the fall of 2011. The purpose of the second data analysis summit will be for the groups to discuss ToxCast Phase II screening data and determine next steps. In addition, we have been working with EPA's Risk Assessment Forum (RAF) on training sessions for CompTox tools. Plans are to have that training early in 2011.
- ✓ 3. In the first publications of ToxCast data, explicit statistical and biological linkages to traditional testing results have been made for cancer and non-cancer endpoints. Future publications and other forms of guidance (e.g., Risk Assessment Forum training on CompTox tools scheduled for early 2011) will focus on this transitional application of new NCCT tools and resources within the context of traditional data and approaches. New tools will provide weight of evidence comparative data and help develop confidence and understanding in how these new approaches can supplement and perhaps replace existing methodologies. We are also working with EPA's OPP PPDC on planning a workshop for 2011 on 21st Century chemical evaluation tools.

Charge Question 2: To what extent and how effectively has the CTRP utilized internal and external partnerships to foster its goals?

BOSC Comments:

It is clear that the CTRP staff has been effective at finding professional colleagues in various institutions with whom to collaborate. Certainly, the Program has been able to identify various research activities and data sets through EPA and other organizations in an attempt to assemble sufficient information to achieve some of its goals. For example, the exposure assessment work by Hubal, et al., is an example of a group that has a good idea about where various data sets reside in the government and they are committed to make more of those data sets available. When those data are gathered, they will be accessed by other groups in the CTRP to combine with data the Program has on toxicity or predictors of toxicity.

It appears that the CTRP has successfully engaged those in the NCCT, National Health and Environmental Effects Research Laboratory (NHEERL), National Exposure Research Laboratory (NERL), National Risk Management Research Laboratory (NRMRL), National Center for Environmental Assessment (NCEA), and other groups within the Federal Government. There are two major government agencies, however, which support research work relevant to the Computational Toxicology program—the National Science Foundation (NSF) and

the Department of Energy (DOE). These agencies do not appear to be CTRP partners. NSF has basic science programs that are relevant not only to the biological sciences, but also computer science and the area of qualitative research into the activities (and information needs) of scientists. These are essential components of the CTRP research. DOE programs and data, particularly in radiation safety and environmental remediation, would seem natural sources of valuable information for the CTRP. In addition, the collaboration with the National Institutes of Health (NIH) appears to be mediated by NIEHS and the NIH Chemical Genomics Center (NCGC). Also, a more formal relationship can be established with the National Library of Medicine, particularly with the National Center for Biotechnology Information and its PubChem program. The BOSC members believe that the development of these relationships would be appropriate for the CTRP.

It appears that the CTRP has been less effective in developing equally strong research groups at various universities in the United States and internationally, or partnerships with other scientific and regulatory bodies outside the United States (e.g., the United Kingdom, Australia, New Zealand, Switzerland, Germany, and a few other countries have substantial databases on exposure, toxicology, and predictive toxicology). No doubt, these relationships will follow as the CTRP matures and receives additional funding. At the moment, the Program is establishing collaborations in multi-scale modeling of developmental toxicity and virtual tissues with some U.S. academic partners (Indiana University and University of Texas). Although these collaborations clearly are of value, it also seems appropriate to expand these partnerships to include multi-scale modeling work in Europe, Asia, and elsewhere in the United States. An excellent example is the virtual physiological human initiative that is intended to support the development of patient-specific computer models and their applications in personalized and predictive medicine. This constitutes an integral part of the international Physiome Project, a worldwide public domain effort to develop a computational framework for quantitative description of biological processes in living systems across all relevant levels of structural and functional integration, from molecule to organism, including the human. The Physiome Project has established standards, which are used in the multi-scale modeling community. In parallel, the mathematical and computational oncology community has established similar initiatives for assessing cancer progression and treatments. The CTRP will benefit substantially in establishing partnerships with these multi-scale modeling enterprises in the United States, Europe, and Asia.

As an advisory group, the BOSC would very much like to have seen a table which presented the various relationships with the full-time equivalents (FTEs) from each organization committed to a particular "joint" collaboration. It would have been helpful to include in that table an indication of the level of financial resources from the CTRP that were dedicated to the various projects, and a timeline for various milestones. Like so many government initiatives, it is very difficult to rate the efficiency or productivity of the Program because it is unclear if the project's success is the result of CTRP resources or those of another entity. For example, if the CTRP is devoting 0.25 FTE for 3 years to the initiative and NHEERL is devoting 5 FTEs from other programs for 4 years, the success of the project is predominantly the result of NHEERL resources. Another layer of complexity occurs when some portion of the project is conducted by post-docs, summer or more permanent interns, and graduate students (who may or may not be counted as FTEs). Then, beyond that, there are contractors who often participate in projects. Building such teams is not to be discouraged and, indeed, it should be promoted. It is not

possible, however, to determine which person or persons is “leading the project” and providing the driving force to resolution. Only by interviewing each team, would it be clear who was truly providing the leadership and effort.

BOSC Recommendations:

- ✧ 4. *Continue to interact with other scientific bodies, regulatory agencies, and universities both in the United States and globally so as to insure that work conducted elsewhere can be “built upon.” In addition, it is recommended that the group interact with the toxicology groups within pharmaceutical and major chemical companies. One possible benefit of this interaction is that it may promote harmonization regarding the organization of historical data that currently are being assembled, as well as new data. This would eliminate the time-consuming task of extracting data from original studies and then entering them in the databases.*
- ✧ 5. *Routinely (perhaps biannually) sponsor some sort of exchange of information with risk assessment practitioners both inside and outside EPA (corporations, consultants, and government scientists) to be sure that the end products of the Program’s work are both reliable and of use to the future users.*
- ✧ 6. *for the next BOSC review, develop a table that presents the level of effort dedicated to specific projects, by year. This table would contain the number of CTRP FTEs, as well as the approximate level of “collaborative” effort (from other EPA laboratories and other partners and consultants). In kind support and “hard” dollars also should be presented.*

ORD Response:

- ✓ 4. CTRP and NCCT have continued to expand collaborations and other interactions within the Federal government (Tox21 now includes FDA), with academic partners (funded or not by STAR), with scientific societies, with NGOs of various types, and with international groups such as OECD, ECHA, EU JRC and EU ChemScreen. In addition, active collaborations are expanding with numerous pharmaceutical companies, chemical companies (e.g., Dow Chemical) and researchers funded by industry (The Hamner).
- ✓ 5. Following on the September 2009 BOSC, the research presented was brought to Washington DC in January 2010, in a session co-hosted with OCSPP. A large number of program office scientists were able to engage in discussions with scientists from the CTRP at that all day meeting. Additional workshops, colloquia, and training sessions are being developed within EPA for 2011 (see ORD response to recommendations 2 and 3 above). In the future, we anticipate workshops, conferences and stakeholder meetings to integrate CTRP work within the new Safer Products for a Sustainable World (SPSW) research program.
- ✓ 6. With the advent of the Safer Products for a Sustainable World program in FY2012, research involving the risks of chemicals will become more integrated across ORD and resources will be more accountable. The CTRP will become an integrated part of SPSW and we anticipate a new BOSC committee will be formed to review the much larger SPSW. SPSW will include all or parts of six existing multi-year plans within ORD.

Nevertheless, we should be able to provide information on ORD FTEs and overall funding even as we transition to SPSW during FY2011. However, obtaining this information from external collaborators/partners will be more difficult. As part of our efforts to track and update our partnerships, we will follow-up with our partners and request information on the level of effort and resources they are providing.

Charge Question 3: What evaluation can you provide relative to the contributions of the CTRP to the advancement of transforming the field of toxicity testing?

BOSC Comments:

The CTRP appears to be at the cutting edge of transforming the field of toxicity testing. Its original goals, defined at its creation in 2005, were consistent with recommendations subsequently described in the National Academy of Sciences' 2007 report, entitled Toxicology in the 21st Century: A Vision and a Strategy. Particularly important contributions in this area include major advances in HTS, advances in approaches for data-mining from various data sources including HTS and other ToxCast efforts, advanced model development of virtual tissues, and the incorporation of uncertainty analysis into model development and, ultimately, risk assessments.

The incorporation of modern computing with molecular biology as developed by the CTRP is necessary to move the discipline of toxicology from the current stage, which is primarily descriptive science, to a more predictive one. The concept of utilization cell-based in vitro testing will assist the understanding of the key biological pathways by which chemicals induce adverse effects. The development of knowledge bases of toxicity pathways, toxicological responses, and key information on biological networks will lead to the use of solid science in the risk assessment. The HTS will lead to more cost-effective testing, which will save money and reduce the use of animal testing. In addition, virtual tissues (liver and embryo) that link across levels of biological organization from molecular to cellular to tissue level responses will be good predictive tools for general and developmental toxicity. The NCCT is establishing collaborations with other institutions across the world with similar goals, which may lead to the expansion of the number of predictive virtual tissues. Moreover, it is expected that the attrition rate for pharmaceutical compounds will be reduced as the computational toxicology tools provide a better prediction of human toxicity.

The BOSC applauds the efforts of the NCCT to embark on the challenges of the Tox21 paradigm, elucidating the strengths and limitations of the "toxicity pathway" approach and the challenges of generating truly predictive HTS platforms. The CTRP has demonstrated rapid progress in the development of HTS. To date, during Phase I of ToxCast, 467 assays have been employed with nine platforms. These assays have been applied to an initial set of 309 chemicals for which relatively extensive toxicological information is available (largely active chemicals in pesticides). In Phase II, this number of assays is expanding and will be applied to a new set of 700+ chemicals for which less information is available. This phase will provide for a critical evaluation of the large array of assays under consideration for their relative utility of chemical prioritization. The challenge for this group is to apply the lessons learned from the Phase I ToxCast efforts and iterate with Phase II to add new assays and define a strategy for attrition of those that provide limited or inconsistent information. The BOSC recommends that the Program

keep the statisticians and mathematical modelers involved in assay evaluation so that they can move from qualitative prediction to quantitative prediction of outcomes from exposure data. Although a strategy has been outlined to include this information, it has not been sufficiently communicated to all members of the team. In addition, to achieve their future milestones related to prediction of outcome, the group needs to define metrics by which they can measure their success (such as specificity and sensitivity goals) and declare victory for specific classes of compounds or cell types. The identification of new toxicity pathways, and therefore new assays, will be essential to gain the predictive power necessary to predict outcome from exposure beyond a single class of compounds. Again, a strategy to define new pathways/assays may exist but, when questioned at the poster session, many of the Program researchers were unaware of who was doing it or how it was being done.

The initial evaluation of Endocrine Disruptor profiling with ToxCast data is extremely promising in its ability to identify new modes of action (MOA) that are not traditionally considered in addition to known developmental or reproductive endpoints. Likewise, the ToxCast data display good sensitivity for predicting neoplastic liver lesions, and statistical models have demonstrated a data gap for new assays to be developed for non-neoplastic lesions. Although there is some suspicion of these statistical approaches because “they don’t make sense based on what we know”, we recommend an unbiased evaluation of the usefulness of particular assays to achieve prediction beyond a single class of compounds and to define knowledge gaps for new assay design. This is a huge step forward for the NCCT to demonstrate the potential utility of the ToxCast approach, and it justifies continued funding to maintain the momentum of this research team.

The engagement of collaborators in NCEA to transform risk assessment into a “NexGen” paradigm using tools and databases coming out of the NCCT is certainly impressive, although daunting. Translating the predictions from the HTS assays to human population risk will require strong connections between the MOA data and statistical genetic diversity, such as those being provided by the Carolina Center for Computational Toxicology, led by Dr Ivan Rusyn. The ability to put real uncertainty factors into exposure limits that protect 99 percent of the population will truly revolutionize human health risk assessment.

It is very difficult to determine how the work of the CTRP will be implemented from a regulatory standpoint. The Program’s work will likely help organizations identify the chemicals most deserving of significant study and the “type” of additional testing that needs to be conducted. Ultimately, the CTRP research will establish a methodology that can be relied upon by researchers around the world for quickly identifying those chemicals that “have a red flag.” The BOSC recommends that the NCCT develop case studies that demonstrate a strategy for incorporation of CTRP tools/research into the risk assessment process. This could help significantly in optimizing the resources of the various organizations such that they will not be conducting routine toxicology “screening tests” on chemicals that raise too many red flags after being run through the ultimate program offered by the CTRP.

BOSC Recommendations:

- ❖ 7. *Keep the statisticians and mathematical modelers involved in assay evaluation so that they can move from qualitative prediction to quantitative prediction of outcomes from exposure data.*

- ✧ 8. *Conduct an unbiased evaluation of the usefulness of particular assays to achieve prediction beyond a single class of compounds and to define knowledge gaps for new assay design.*
- ✧ 9. *Develop case studies that demonstrate a strategy for incorporation of CTRP tools/research into the risk assessment process.*

ORD Response:

- ✓ 7. The majority of assay evaluation and data analysis within NCCT is conducted by statisticians and mathematical modelers who are also working on the virtual tissues and other projects. As the quantity, quality and scope of data increases, it will be possible to transition from qualitative to quantitative predictions and models that will be useful for risk assessments and other regulatory decisions.
- ✓ 8. With completion of ToxCast Phase II in 2011, it will be possible to conduct such an evaluation across 1000 chemicals for over 400 assays. Tox21 will allow similar evaluations across 10,000 chemicals for dozens of assays. This type of analysis will carry forward beyond 2013 as part of an integrated evaluation system within the SPSW program. Gaps in data and knowledge, which in some cases have already been identified, are being filled by new NCCT contracts and partners (internal and external to EPA). At the close of this analysis and expansion period, a robust, proven and comprehensive set of predictive assays and models will be available for addressing the Agency's most critical needs.
- ✓ 9. CTRP and NCCT staff is actively engaged with program office colleagues and NCEA NexGen researchers developing case studies for pesticides and industrial chemicals. Peer-reviewed publications are expected on these case studies, as well as presentations at various internal and external meetings and workshops (e.g., the November 2010 NexGen risk assessment workshop coordinated by NCEA). Finally, this translation of CTRP tools and research to risk assessment applications will be a part of the integrated efforts with the SPSW program beginning in FY2012.

Charge Question 4: To what extent do the ORD intramural projects, the extramural Science To Achieve Results (STAR) centers, and the five stated CTRP management priorities described in the FY09-12 implementation plan combine to efficiently support the goal of providing high-throughput decision support tools for screening and assessing chemical exposure, hazard, and risk to human health?

BOSC Comments:

Ultimately, the implementation of the CTRP's goals is going to require an iterative approach to developing models as the data available grows in both quantity and complexity. To do this, the CTRP must develop methods that can be used to make verifiable predictions and to generate appropriate data to test those predictions. This clearly is going to require developing a strong partnership that reaches beyond the boundaries of the CTRP and takes advantage of other existing programs, including ORD intramural projects, other intramural EPA projects, the extramural STAR Centers, and partnerships with other agencies.

The FY09-12 implementation plan lays out these interactions and the role that they will play in helping to direct and develop the CTRP. This document serves as a useful reference and guide as to the roles and activities of all parties contributing to the CTRP. The new start programs outlined for the ORD laboratories and centers are well leveraged with the CTRP. Results from those projects have the potential to be quickly incorporated into ongoing CTRP activities, providing for effective and efficient use of R&D efforts. Continuing cooperation with the STAR Centers and the innovations that are taking place at these university sites will enhance the development and robustness of the decision support tools under development. In total, NCCT interactions with other ORD laboratories and the STAR Centers are essential as they will provide not only the starting data needed to fully develop the CTRP, but also some sampling of the potential users of the systems that will be essential for providing feedback and ideas for the next iteration of tools.

In its implementation plan, the CTRP lays out five priority management areas:

- 1. Toxicity Predictions and Chemical Prioritizations Incorporating Exposure*
- 2. Strengthening Cross-ORD Collaborations*
- 3. Tox21: A Federal Partnership Transforming Toxicology*
- 4. Communicating Computational Toxicology*
 - a. EPA Program Office Training and Implementation of Computational Tools*
 - b. Communities of Practice for Chemical Prioritization and Exposure Science*
- 5. Developing Clients for Virtual Tissues*

Overall, it is evident that addressing the five priority management areas will further support program efficiency and effectiveness, and help sustain the progress the CTRP has achieved to date. The BOSC agrees that the combined programs outlined in the implementation plan are key contributions that are needed to maintain the CTRP on a path to achieve the stated goals.

The following paragraphs present some of the BOSC's observations about some challenges along with suggestions for meeting them that are relevant to the Program's continued progress.

The CTRP needs to be more integrative, both internally and externally, to ensure all parties are working from common assumptions, data development schedules, and deliverable planning. As an example, at the review, there were instances where the theoretical and modeling groups noted that they needed access to quantitative data to inform certain aspects of their models while in the same session the experimental groups were presenting precisely those data. Some of this speaks to the stage of the various projects, but as the ultimate goal is to have computational tools that are useful for informing risk assessment, these groups need to begin to work together more closely.

It will be important to detail specific roles for the STAR Centers as part of the integrated approach to managing the Program's mission. One of the omissions in this management plan is the role that the STAR Centers will play in the future. Although the STAR Centers and their importance in the CTRP's future plans are laid out elsewhere in the implementation plan, it will be important to include these into a more integrated approach to managing the CTRP's mission. The BOSC's understanding is that this omission is deliberate and related to the fact that the STAR Program funding is independent of the CTRP budget; consequently, CTRP management cannot rely on it. The Centers, however, play a key role in the activities of the CTRP; therefore,

it is crucial to have a firm plan for their continued contribution. The CTRP should consider the implications for the research program if a STAR Center is not renewed, and prepare some contingencies on how the research gaps might be filled if these key contributors are lost from the research program.

As the overall CTRP continues to progress, there should be a higher priority on incorporation of ecological receptors and greater focus on assessment of exposure factors. The BOSC noted an absence of ecological health as an endpoint for the high-throughput decision support tools for screening and assessing chemical exposure, hazard, and risk. To become fully integrated and supportive of the Agency's regulatory activities, the CTRP will have to move into the field of ecological risk assessment at some point. Acknowledging this need and developing a forward plan to incorporate it as part of the CTRP should be part of the longer term plan.

The ExpoCast initiative, as described in the Hubal & Egeghy poster, is an important direction for the NCCT to pursue. Expansion of this effort to include real exposure and outcomes data, as well as the additional development of software resources to take advantage of these data for exposure and outcome predictions, should be a priority of the NCCT.

Training:

One area that deserves particular comment in regard to the management plan is its training component, and in particular the training of postdoctoral fellows. The BOSC was universally impressed with the quality of the postdoctoral fellows and their work. The BOSC encourages the CTRP to continue its emphasis on training postdoctoral fellows because these scientists have the potential to be ambassadors to the rest of the community to help extend the understanding and acceptance of the types of computational tools the CTRP is trying to develop, and in doing so, ultimately help to improve those tools and their efficacy.

Quality Assurance for Software and Models:

As the CTRP continues and expands its efforts to develop complex models and software systems, such as the virtual tissues, testing and quality assurance of these tools becomes even more important. Software and model testing is the practice of probing for errors (or “bugs”), typically by using a structured set of manually constructed inputs to generate a list of specific performance errors. Static testing of software and models involves inspection of the source code and formulas, usually in a structured fashion called a walk-through. Dynamic testing involves executing the code or model on a set of test inputs. Structured input sets can test how these tools perform in the face of boundary conditions (e.g., null or very long inputs), and systematically vary combinations of representative inputs. These methods are called “black box” tests because they do not require any knowledge of the implementation. Dynamic methods that make use of knowledge of the implementation are called “white box” approaches; for example, code coverage metrics that test what proportion of the source code is reached while processing an input suite. “White box” methods can provide information about how to improve test suites themselves. Open source approaches to development of software can be exploited as an extension of “white box” methods bringing large communities of software engineers to the evaluation of code.

A frequent cause of software or model failures is a lack of compatibility with another application, an operating system, or Web browser. Compatibility testing is particularly important in a plan for distributed development, where the separately developed components must be compatible with each other. Test suites that exercise all aspects of each component's interface to the others address this issue in distributed software and model development.

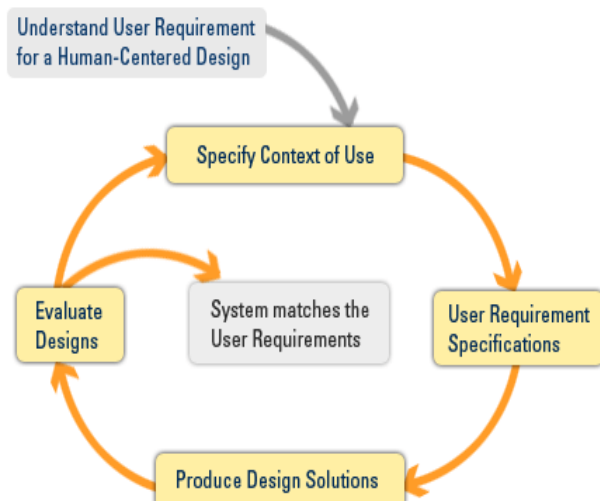
Testing in each of the proposed computational research areas is complicated by the fact that structured input sets and code coverage metrics have blind spots in these advanced systems. The range of possible inputs to a tissue model or natural language processing system cannot be exhausted by a structured test set. For these reasons, a software and model testing approach must be augmented with a sophisticated evaluation approach that probes how the tools and systems produced work in the hands of users.

User-Centered Design:

Developing computer models and software systems to support complex and incompletely specified activities (such as modeling toxicity, assessing risk, and prioritizing exposures) is a difficult task. Users are generally unable to specify a priori what would be most useful for a computer system to do, although generally they do not have trouble describing what they like and do not like about any particular implementation. Furthermore, customer communities are heterogeneous, with needs that vary with analytical goals, methodological approaches, the types of data being analyzed, the amount and quality of relevant background knowledge available, and a wide variety of other factors. Customers and developers both can be frustrated by this seemingly circular need to produce models or software before defining requirements, which can themselves change in different circumstances and as new databases and software are produced.

User-centered design is an approach that grounds the process of design in information about the people who will use the product. There is an international standard (ISO 13407: Human-centered design process) that defines the general approach. Figure 1 shows the iterative nature

Figure 1: The ISO User-Centered Design Process



of this process: starting with an understanding of the context of use (who will use it, under what conditions, to what ends) leads to a set of requirements that must be met, which in turn leads to a design solution, which then is evaluated through usability testing with actual users, which may lead to additional understanding of the context of use, and so on.

Qualitative methodology is a proven approach for effectively characterizing and explaining such issues as how scientists and policy makers make meaning while proceeding through complex analyses and how they inscribe visualized representations of knowledge into these problem-solving practices (Neressian, 2008). Field

observations can reveal the ways in which users generate preferred, new, or augmented

analytical practices as they progressively use a new technology (Mirel, 2009; Vicente, 2002). Additionally, qualitative ethnographic methods can reveal ambiguities that scientists negotiate amid biological uncertainty and incomplete data and show the processes by which they disambiguate them. NSF recently issued a report stressing the importance of the study of scientists' research processes through qualitative methods (Lamont & White, 2005).

In general, trained observers using longitudinal fieldwork are better than domain experts themselves at discerning discrete steps in their own reasoning and the role that artifacts such as software play in facilitating their work (Schon, 1983; Hutchins, 1996). Field observations and semi-structured post hoc interviews can importantly complement focus groups or structured interviews or surveys for elucidating longitudinal views of scientists' and policy makers' thinking and behaving (Schon, 1983; Cresswell and Plano, 2006).

BOSC Recommendations:

- ✧ 10. *Be more integrative, both internally and externally, to ensure all parties are working from common assumptions, data development schedules, and deliverable planning.*
- ✧ 11. *Expand outreach to the broader community, both within EPA and in the extramural community. This is not to say that the CTRP has not been effective in building a strong outreach program, but only that this needs to be a priority, and possibly a higher priority.*
- ✧ 12. *Detail specific roles for the STAR Centers as part of the integrated approach to managing the Program's mission.*
- ✧ 13. *Place a higher priority on incorporation of ecological receptors and greater focus on assessment of exposure factors.*
- ✧ 14. *Develop a forward, longer term plan to incorporate the field of ecological risk assessment as part of the CTRP.*
- ✧ 15. *Expand the ExpoCast program to include real exposure and outcomes data, as well as the additional development of software resources to take advantage of these data for exposure and outcome predictions. This should be a priority of the Center.*
- ✧ 16. *Continue training postdoctoral fellows because these scientists have the potential to be ambassadors to the rest of the community to help extend the understanding and acceptance of the types of computational tools the CTRP is trying to develop, and in doing so, ultimately help to improve those tools and their efficacy.*
- ✧ 17. *Highlight quality assurance for software and models with a specific testing approach augmented with a sophisticated evaluation approach that probes how the systems produced work in the hands of users.*
- ✧ 18. *Promote "user-centered design", an approach that grounds the process of design in information about the people who will use the product.*

ORD Response:

- ✓ 10. ORD is moving rapidly to a more integrated, transdisciplinary research planning process that involves the EPA program offices in problem formulation and research implementation. A vanguard effort, entitled Safer Products for a Sustainable World (SPSW), is being developed that incorporates the chemical-centric research in six of ORDs current multi-year plans (Nano, comptox, EDCs, SP2, HHRP and HHRA). The

NCCT Director has been the executive lead developing this vanguard effort for ORD. It is scheduled to become operational in FY2012 budget year and the BOSC Executive Council will be briefed on progress in the fall of 2010.

- ✓ 11. NCCT hired a public affairs specialist in January 2010. Working with the NCCT team, a Computational Toxicology Research Program (CTRP) communications and outreach strategy has been drafted and is currently being implemented. The communications strategy lists CTRP communication goals, measurable objectives, target audiences, strategies and timeline. Some examples of the outreach efforts implemented to date include an updated CTRP webpage, positive news coverage (local and national) resulting from distribution of news releases and fact sheets that summarize CTRP research for a non-technical audience. The NCCT public affairs specialist is currently on the ORD Communication Strategy working group, ensuring close coordination with ORD and EPA wide efforts. The strategy is being implemented now. The main goal of the communications strategy is to implement communications and outreach tactics that will increase target audiences awareness, interest and usage of CTRP tools and research. The success of implementing the strategy will be based on meeting measurable objectives in the strategy. The CTRP communications strategy will be updated annually based on what strategies are the most effective at reaching target audiences.
- ✓ 12. The first 3 CompTox STAR centers complemented the intramural aspects of the program and helped provide fundamental tools and scope of the overall program. The TVIS center is partnering closely with the intramural Virtual Embryo project. The 2010 STAR RFP focused on developing experimental and computational models for the Virtual Liver project, although no award has been issued as of August 2010. Future STAR RFPs are being designed to support the CompTox programs integration with SPSW and bring in new elements and principles of sustainability.
- ✓ 13 & 14. Working with the SPSW process (see response to recommendation #10), the OECD Molecular Screening Workgroup that is chaired by the NCCT Director, and the intergovernmental Tox21 program that is developing a screening library of 10,000 chemicals that will be assayed in up to 100 assays over the next two years, efforts are underway to bring elements of ecologically relevant toxicity pathways into the screening process. The major challenge is to determine what are the unique toxicity pathways for ecotoxicity, and how HTS assays can be developed to evaluate them. As we mature the research directed at making human health toxicity more efficient and effective through Phase II of ToxCast, resources should be available to help bring this transformation to ecotoxicity studies. Future STAR RFAs funded by CompTox will also target development of ecologically critical assays and approaches to HTS.
- ✓ 15. We are currently in the process of incorporating exposure data sources into ACToR. In addition, we are developing ExpoCast database, an exposure database that will have a separate interface with inner workings of ACToR to take advantage of linkages to toxicity information. ExpoCast database will provide capability to download customized datasets, and obtain summary statistics and distributional parameters. The data model has been developed and a prototype is under construction. Together with our partners, we are

developing an exposure ontology. Based on this ontology, we will pilot curation of exposure information from the literature first into the Comparative Toxicogenomic Database (high level exposure information) and then into ExpoCast database (more detailed data).

- ✓ 16. NCCT recognizes the importance of postdoctoral fellows to the success of our program and continues to hire and backfill those who move on to permanent positions in industry, academia and government. In fact, one third of the EPA staff are postdoctoral fellows, and we have a commitment to continue this level of staffing. We continue to strengthen our postdoctoral program and launched a Mentoring Program for pre- and post-docs during FY 2010.
- ✓ 17. NCCT recently reassigned a chemist from existing NCCT staff to serve as our Quality Assurance Manager. This individual previously served in that capacity as a contractor to EPA, and has extensive experience in computational solutions ongoing in the CTRP and is working with NCCT staff and management to develop and implement appropriate quality metrics and systems.
- ✓ 18. NCCT is actively setting up numerous opportunities for CTRP tools to be presented and utilized by end-users both within and outside the Agency (see also the action items for recommendations #1-3, 5, 9). The first formal iteration of training to an external group on CTRP tools took place in July 2010, and this type of training will be carried out repeatedly. The result of this outreach and training will be a large intramural and extramural cadre of users that will inform continuing tool development. The integration of the CTRP with SPSW in FY2012, and the problem formulations feeding into the development of the SPSW research program, will formalize the "user-centered design" approach around defined ORD and Agency teams working on specific applications and projects.

Charge Question 5: The NCCT was established as an organization with a 5-year charter ending in February 2010, which would continue dependent on: 1) meeting established goals; and 2) having continuing mission-critical goals and objectives. What recommendation(s) can you provide the Agency regarding continuation of the NCCT as an established organization, and the criticality of its goals and objectives to EPA?

BOSC Comments:

The BOSC strongly supports action by EPA to make the NCCT permanent. It is clear from EPA's Strategic Plan for Evaluating the Toxicity of Chemicals that computational toxicology will be integral to the future of toxicology, risk assessment, and regulatory decision-making by the Agency. EPA will not be able to fulfill its strategy, or indeed its mission, without significant expertise and an active research program in computational toxicology. The NCCT has made significant contributions during the short time it has been in existence. The Center's work products have had an impact on Agency activities. Center products such as DSSTox, Actor, and ToxRefDB have been of great assistance to EPA program offices and to the toxicology community at large. The longer-term projects underway at the Center have been productive and have demonstrated their potential value to EPA. The staff of the NCCT has proved that the

structure of the Center—a core of strong expertise with additional expertise leveraged through collaborations outside the NCCT—is an ideal organizational structure in a resource-scarce environment. The BOSC recommends in the strongest terms that the NCCT be made permanent.

The CTRP has been enormously successful in developing the tools and resources necessary to bring computational predictions to the science of toxicology and risk assessment. In the process, the staff members have learned many of the lessons needed to move forward, including many of the reasons why simply building the framework they have in place has been so challenging. In moving forward, the Program cannot rest on its laurels, but must continue to address these fundamental infrastructure problems and questions, building on what it already has done. The challenge will be to do this, which can become all-absorbing, while continuing to achieve its goals and leverage resources effectively.

The CTRP also has assembled a tremendous intellectual resource, ranging from the more senior personnel involved in the Program to the staff and postdoctoral trainees. It is this intellectual infrastructure as much as the data, databases, and software that represent the real value of the Program. The various scientists who are the backbone of the Program are extraordinarily focused on the end goals and while there is always room for improvement in any program, it is these scientists and their commitment that will assure that the CTRP retains its focus.

One recommendation would be the establishment of performance metrics. Although this group has been publishing at a reasonable rate, the primary goal is not academic publication but rather the development of tools and resources for informing risk assessment, and there are potential objectives that can be used to assess the relative impact of these tools on the field, including Web hits, software downloads, and the citation rate of these tools in publications and grants. Although some of this was presented at the review, the Program should establish a subset of these as benchmarks and provide some measure of the historical change in these metrics during future funding periods.

The NCCT has done a very good job of outreach to the risk assessors at EPA, and to other stakeholders within and outside the Agency. The BOSC saw ample evidence that the Program is focused on developing models and tools that can be applied to the mission of the Agency. Given that computational toxicology is on the leading edge of science, and that risk assessment decisions are generally made using established science, it will be important for the NCCT to continue to broadcast its message and to engage in constructive dialog with the prospective recipients of its work products.

Center management mentioned hiring a communications person to help disseminate NCCT's plans and products. The BOSC endorses this intention.

The NCCT has hired several highly accomplished scientists under Title 42 authorization. These individuals are providing critical leadership in computational toxicology and have contributed materially to both short-term and long-term initiatives. They also have provided instant credibility and stature to the CTRP, which has been critical in recruiting post-docs and in establishing EPA as a leader in computational toxicology. The BOSC strongly believes that the Title 42 positions have been crucial in establishing direction for the CTRP and in providing continuing scientific leadership in this complex and cutting-edge area of research.

BOSC Recommendations:

- ✧ 19. *Establish performance metrics that track the development of tools and resources for informing chemical prioritization, toxicity testing, and risk assessment.*
- ✧ 20. *Before CTRP presents its views on various chemicals in the coming years, it was recommended that the CTRP randomly identify 10 percent of the predictions of the models for comparison to the results of traditional toxicological testing so as to insure that there is no bias by the researchers prior to claiming that the the models are accurate predictors of adverse effects.*
- ✧ 21. *Continue to meet with customers, clients, and stakeholders on a regular basis to ensure that the Program is meeting the needs of the risk assessors and risk managers in the Agency.*

ORD Response:

- ✓ 19. Working with our communication strategy, we are developing processes for translating the CTRP tools into resources that are utilized by EPA Program Offices. The ToxPi concept is an early effort in this as we are scheduled to utilize the concept to assay and prioritize up to 2000 chemicals for EDC potential in FY11. Our website has been completely revamped to better display our research activities. We are also using a number of information sheets to distribute basic information about our tools and to assess their utilization via metrics such as requests for assistance, URL hits, and inclusion in the decision process of program offices.
- ✓ 20. See response to recommendation #8. In addition to the data from ToxCast Phase II and Tox21, data from a subset of assays relevant to endocrine activity will be generated on a total of 2000 chemicals, supporting comparisons to endocrine and reproductive models as well.
- ✓ 21. Two-way communication with our customers and partners in the program offices remains a priority for us. We are working with EPAs Risk Assessment Forum to develop and host a multi-tiered training workshop on computational toxicology. This would include modules targeted at senior managers, office directors and operations staff of the program offices. We have also undertaken a major overhaul of the NCCT website to better present and describe the tools we are developing and to track the activity of them. The SPSW planning process (see #10) includes a close and interactive participatory process in problem formulation and research implementation that will help ensure that ORD is meeting the highest priority needs for chemical safety assessment.

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TABLE 1

ORD Response to BOSC Report on CTRP April 15, 2010

Summary of Recommendations and Proposed Actions

Recommendation	Action Item	Timeline
1. At this point, the CTRP should obtain some public feedback on how people are using and interpreting the available data.	CTRP plans to survey the public to determine awareness, interest and usage of CTRP data, tools and research. This is an integral part of the newly drafted Computational Toxicology communications and outreach strategy. A survey has been drafted and is currently being edited by the CTRP research team. However, to survey more than nine non-EPA people, approval from the Office of Management and Budget (OMB) is required. The plan is to work through the OMB approval process and then survey the CTRP database of more than 25,000 people who have expressed an interest in CTRP.	According to OMB's approval timeline for surveys, it will take at least a year for approval. CTRP is aiming to release the survey in January 2012 unless the approval process is faster than expected or the survey can be distributed using a different process.
2. NCCT should organize an annual or biannual conference that brings together the data generators, data users, and risk assessors/managers—the ultimate users of these alternative methods/models	NCCT will plan a second data analysis summit (TDAS) that will bring together data generators, data users, and risk assessors and managers. The purpose of the second data analysis summit will be for these groups to discuss ToxCast Phase II screening data and determine next steps. Working with EPA's Risk Assessment Forum (RAF), training sessions on CompTox tools are being developed.	Early 2011 for RAF training; Fall 2011 for the 2 nd TDAS.

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Recommendation	Action Item	Timeline
<p>3. NCCT should provide guidance on how to interpret this information in the context of more traditional testing and scientific examination so that risk assessment practitioners in the EPA program offices can apply these findings.</p>	<p>In the first publications of ToxCast data, explicit statistical and biological linkages to traditional testing results have been made for cancer and non-cancer endpoints. Future publications and other forms of guidance (e.g., Risk Assessment Forum training on CompTox tools scheduled for early 2011) will focus on this transitional application of new NCCT tools and resources within the context of traditional data and approaches. New tools will provide weight of evidence comparative data and help develop confidence and understanding in how these new approaches can supplement and perhaps replace existing methodologies. Also working with EPA's OPP PPDC on planning a workshop for 2011 on 21st century chemical evaluation tools.</p>	<p>2009-2013</p>
<p>4. Continue to interact with other scientific bodies, regulatory agencies, and universities both in the United States and globally so as to insure that work conducted elsewhere can be "built upon." In addition, it is recommended that the group interact with the toxicology groups within pharmaceutical and major chemical companies.</p>	<p>CTRP and NCCT have continued to expand collaborations and other interactions within the Federal government (Tox21 now includes FDA), with academic partners (funded or not by STAR), with scientific societies, with NGOs of various types, and with international groups such as OECD, ECHA, EU JRC and EU ChemScreen. In addition, active collaborations are expanding with numerous pharmaceutical companies, chemical companies (e.g., Dow Chemical) and researchers funded by industry (The Hamner).</p>	<p>Ongoing</p>

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Recommendation	Action Item	Timeline
<p>5. Routinely (perhaps biannually) sponsor some sort of exchange of information with risk assessment practitioners both inside and outside EPA (corporations, consultants, and government scientists) to be sure that the end products of the Program’s work are both reliable and of use to the future users.</p>	<p>Following on the data summit hosted by NCCT in 2009, a session was co-hosted with OCSPP in early 2010 showcasing CTRP research for program office staff. Additional workshops, colloquia, and training sessions are being developed within EPA for 2011, and will be developed in the future within the new Safer Products for a Sustainable World (SPSW) research program. Informational sessions and external stakeholder meetings are also planned for 2011, and will carry forward with SPSW as well.</p>	<p>Ongoing</p>
<p>6. For the next BOSC review, develop a table that presents the level of effort dedicated to specific projects, by year. This table would contain the number of CTRP FTEs, as well as the approximate level of “collaborative” effort (from other EPA laboratories and other partners and consultants). In kind support and “hard” dollars also should be presented.</p>	<p>With the advent of the Safer Products for a Sustainable World program in FY2012, work across ORD will become more integrated and resources will be more accountable. We should be able to provide information on ORD FTEs and overall funding even as we transition to SPSW during FY2011. However, obtaining this information from external collaborators/partners will be more difficult. As part of our efforts to track and update our partnerships, we will follow-up with our partners and request information on the level of effort and resources they are providing.</p>	<p>FY 2011</p>

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Recommendation	Action Item	Timeline
<p>7. Keep the statisticians and mathematical modelers involved in assay evaluation so that they can move from qualitative prediction to quantitative prediction of outcomes from exposure data.</p>	<p>The majority of assay evaluation and data analysis within NCCT is conducted by statisticians and mathematical modelers who are also working on the virtual tissues and other projects. As the quantity, quality and scope of data increases, it will be possible to transition from qualitative to quantitative predictions and models that will be useful for risk assessments and other regulatory decisions.</p>	<p>Ongoing</p>
<p>8. Conduct an unbiased evaluation of the usefulness of particular assays to achieve prediction beyond a single class of compounds and to define knowledge gaps for new assay design</p>	<p>With completion of ToxCast Phase II in 2011, it will be possible to conduct such an evaluation across 1000 chemicals for over 400 assays. Tox21 will allow similar evaluations across 10,000 chemicals for dozens of assays. This type of analysis will carry forward beyond 2013 as part of an integrated evaluation system within the SPSW program. Gaps in data and knowledge, which in some cases have already been identified, are being filled by new NCCT contracts and partners (internal and external to EPA). At the close of this analysis and expansion period, a robust, proven and comprehensive set of predictive assays and models will be available for addressing the Agency's most critical needs.</p>	<p>2011-2013</p>

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Recommendation	Action Item	Timeline
<p>9. Develop case studies that demonstrate a strategy for incorporation of CTRP tools/research into the risk assessment process.</p>	<p>CompTox and NCCT staff is actively engaged with program office colleagues and NCEA NexGen researchers developing case studies for pesticides and industrial chemicals. Peer-reviewed publications are expected on these case studies, as well as presentations at various internal and external meetings and workshops (e.g., the November 2010 NexGen risk assessment workshop coordinated by NCEA). Finally, this translation of CTRP tools and research to risk assessment applications will be a part of the integrated efforts with the SPSW program beginning in FY2012.</p>	<p>2010-2013</p>
<p>10. Be more integrative, both internally and externally, to ensure all parties are working from common assumptions, data development schedules, and deliverable planning.</p>	<p>ORD is moving rapidly to a more integrated, transdisciplinary research planning process that involves the EPA program offices in problem formulation and research implementation. A vanguard effort, entitled Safer Products for a Sustainable World (SPSW), is being developed that incorporates the chemical-centric research in six of ORDs current multi-year plans (Nano, comptox, EDCs, SP2, HHRP and HHRA). The NCCT director has been the executive lead developing this vanguard effort for ORD. It is scheduled to become operational in FY2012 budget year and the BOSC Executive Council will be briefed on progress in the Fall of 2010.</p>	<p>Implementation of the new SPSW is anticipated in FY12, with the ground work being conducted in FY10 and FY11.</p>

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Recommendation	Action Item	Timeline
<p>11. Expand outreach to the broader community, both within EPA and in the extramural community. This is not to say that the CTRP has not been effective in building a strong outreach program, but only that this needs to be a priority, and possibly a higher priority.</p>	<p>NCCT hired a public affairs specialist in January 2010. Working with the NCCT team, a Computational Toxicology Research Program (CTRP) communications and outreach strategy has been drafted and is currently being implemented. The communications strategy lists CTRP communication goals, measurable objectives, target audiences, strategies and timeline. Some examples of the outreach efforts implemented to date include an updated CTRP webpage, positive news coverage (local and national) resulting from distribution of news releases and fact sheets that summarize CTRP research for a non-technical audience. The NCCT public affairs specialist is currently on the ORD Communication Strategy working group, ensuring close coordination with ORD and EPA wide efforts.</p>	<p>The strategy is being implemented now. The main goal of the communications strategy is to implement communications and outreach tactics that will increase target audiences awareness, interest and usage of CTRP tools and research. The success of implementing the strategy will be based on meeting measurable objectives in the strategy. The CTRP communications strategy will be updated annually based on what strategies are the most effective at reaching target audiences.</p>
<p>12. Detail specific roles for the STAR Centers as part of the integrated approach to managing the Program's mission.</p>	<p>The first 3 CompTox STAR centers complemented the intramural aspects of the program and helped provide fundamental tools and scope of the overall program. The TVIS center is partnering closely with the intramural Virtual Embryo project. The 2010 STAR RFP focused on developing experimental and computational models for the Virtual Liver project. Future STAR RFPs are being designed to support the CompTox programs integration</p>	<p>Ongoing</p>

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Recommendation	Action Item	Timeline
	with SPSW and bring in new elements and principles of sustainability.	
13. Place a higher priority on incorporation of ecological receptors and greater focus on assessment of exposure factors.	Working with the SPSW process (see recommendation #10), the OECD Molecular Screening Workgroup that is chaired by the NCCT Director, and the intergovernmental Tox21 program that is developing a screening library of 10,000 chemicals that will be assayed in up to 100 assays over the next two years, efforts are underway to bring elements of ecologically relevant toxicity pathways into the screening process. The major challenge is to determine what are the unique toxicity pathways for ecotoxicity, and how can HTS assays be developed to evaluate them. As we mature the research directed at making human health toxicity more efficient and effective through Phase II of ToxCast, resources should be available to help bring this transformation to ecotoxicity studies. Future STAR RFAs funded by CompTox will also target development of ecologically critical assays and approaches to HTS.	Ongoing

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Recommendation	Action Item	Timeline
14. Develop a forward, longer term plan to incorporate the field of ecological risk assessment as part of the CTRP.	See response to #13	
15. Expand the ExpoCast program to include real exposure and outcomes data, as well as the additional development of software resources to take advantage of these data for exposure and outcome predictions. This should be a priority of the Center.	We are currently in the process of incorporating exposure data sources into ACToR. In addition, we are developing ExpoCast database, an exposure database that will have a separate interface with inner workings of ACToR to take advantage of linkages to toxicity information. ExpoCast database will provide capability to download customized datasets, and obtain summary statistics and distributional parameters. The data model has been developed and a prototype is under construction. Together with our partners, we are developing an exposure ontology. Based on this ontology, we will pilot curation of exposure information from the literature first into the Comparative Toxicogenomic Database (high level exposure information) and then into ExpoCast database (more detailed data).	2010-2013

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Recommendation	Action Item	Timeline
<p>16. Continue training postdoctoral fellows because these scientists have the potential to be ambassadors to the rest of the community to help extend the understanding and acceptance of the types of computational tools the CTRP is trying to develop, and in doing so, ultimately help to improve those tools and their efficacy.</p>	<p>NCCT recognizes the importance of postdoctoral fellows to the success of our program and continues to hire and backfill those who move on to permanent positions in industry, academia and government. In fact, a third of the EPA staff are postdoctoral fellows, and we have a commitment to continue this level of staffing. We continue to strengthen our postdoctoral program and launched a Mentoring Program for pre- and post-docs during FY 2010.</p>	<p>ongoing</p>
<p>17. Highlight quality assurance for software and models with a specific testing approach augmented with a sophisticated evaluation approach that probes how the systems produced work in the hands of users.</p>	<p>NCCT recently reassigned a chemist from existing NCCT staff to serve as our Quality Assurance Manager. This individual had previously served in that capacity as a contractor to EPA, and has extensive experience in computational solutions ongoing in the CTRP. He is working with NCCT staff and management to develop and implement appropriate quality metrics and systems.</p>	<p>2010-</p>
<p>18. Promote "user-centered design," an approach that grounds the process of design in information about the people who will use the product.</p>	<p>NCCT is actively setting up numerous opportunities for CTRP tools to be presented and utilized by end-users both within and outside the Agency (see also the action items for recommendations #1-3, 5, 9). The first formal iteration of training to an external group on CTRP tools took place in July 2010, and this type of training will be carried out repeatedly. The result of this outreach and training will be a</p>	<p>2010-2012</p>

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Recommendation	Action Item	Timeline
	<p>large intramural and extramural cadre of users that will inform continuing tool development. The integration of the CTRP with SPSW in FY2012, and the problem formulations feeding into the development of the SPSW research program, will formalize the "user-centered design" approach around defined ORD and Agency teams working on specific applications and projects.</p>	
<p>19. Establish performance metrics that track the development of tools and resources for informing chemical prioritization, toxicity testing, and risk assessment.</p>	<p>Working with our communication strategy, we are developing processes for translating the CTRP tools into resources that are utilized by EPA Program Offices. The ToxPi concept is an early effort in this as we are scheduled to utilize the concept to assay and prioritize up to 2000 chemicals for EDC potential in FY11. We are also using a number of information sheets to distribute basic information about our tools and to assess their utilization via metrics such as requests for assistance, URL hits, and inclusion in the decision process of program offices</p>	<p>Ongoing</p>

ORD Response to April 2010 BOSC Computational Toxicology Letter Report
September 2010

Recommendation	Action Item	Timeline
<p>20. Before CTRP presents its views on various chemicals in the coming years, it was recommended that the CTRP randomly identify 10 percent of the predictions of the models for comparison to the results of traditional toxicological testing so as to insure that there is no bias by the researchers prior to claiming that the models are accurate predictors of adverse effects.</p>	<p>See answer to #8. In addition to the data from ToxCast Phase II and Tox21, data from a subset of assays relevant to endocrine activity will be generated on a total of 2000 chemicals, supporting comparisons to endocrine and reproductive models as well.</p>	<p>2010-2013</p>
<p>21. Continue to meet with customers, clients, and stakeholders on a regular basis to ensure that the Program is meeting the needs of the risk assessors and risk managers in the Agency.</p>	<p>Two-way communication with our customers and partners in the program offices remains a priority for us. We are working with EPAs Risk Assessment Forum to develop and host a multi-tiered training workshop on computational toxicology. This would include modules targeted at senior managers, office directors and operations staff of the program offices. We have also undertaken a major overhaul of the NCCT website to better present and describe the tools we are developing and to track the activity of them. The SPSW planning process (see #10) includes a close and interactive participatory process in problem formulation and research implementation that will help ensure that ORD is meeting the highest priority needs for chemical safety assessment.</p>	<p>Fall 2010 and beyond</p>