(1) To add a more formal probabilistic model than is used in the INTERNIST system which has greatly influenced our work. This will allow assumptions made by NESTOR to be explicit.

(2) To represent temporal knowledge explicitly and to use it in reasoning.

(3) To represent and use causal information to express the pathophysiology of diseases.

(4) To incorporate abstract diseases into the representation (e.g., it will be possible to include CANCER as a component of a user's hypothesis if this seems more appropriate at the time than specifying BRONCHOGENIC-CARCINOMA or MYELOMA or RENAL-CELL-CARCINOMA).

We have already developed several techniques for utilizing time and causality in determining the probability of a disease. We have also developed search techniques that allow NESTOR to explore efficiently a very large search space in order to find the most probable (multiple disease) hypothesis. This technique is general and can be applied to many nonmedical problems where the goal is to find the most probable hypothesis among many possibilities.

INTEGRATING MATHEMATICAL MODELS WITH AI METHODS

The work to date has concentrated on analyzing physiological function using knowledge of anatomy, physiology and causality. During the coming months, the causal analysis will be generalized. The issue is how to make valid qualitative conclusions based upon the quantitative relations of a physiological model. Rules will be developed and tested for inferring qualitative relations such as the following:

--inferring the direction of change in a dependent parameter following change in an independent parameter;

--inferring whether qualitatively significant change in a dependent parameter may occur as a consequence of a qualitatively significant change in an independent parameter; and

--inferring that a dependent parameter has a qualitatively significant value.

Questions of the following type will build on the other structures, incorporating the kinds of mathematical descriptions of human physiology necessary to predict the effects of quantitative deviations from normal:

"What is the effect of infusing hyperosmotic saline? How much should be used to return plasma volume to normal? How fast will normalization occur?"

PSYCHOLOGICAL STUDIES OF EXPLANATION

Within the next several months, Mr. Teach will have completed all
data analysis for his research and will have completed most of his
dissertation writing. Thereafter we will work on analyzing the results in
light of our interest in the design of explanation capabilities for medical
expert systems. We expect to have several papers based on this work
completed or in preparation by this time next year.

EXPLANATIONS FOR A RULE-BASED EXPERT SYSTEM

As described in the GUIDON report elsewhere in this document, Dr.
Rennels and others from the NEOMYCIN project will continue in their effort
to implement a system to provide explanations for that program. This
project will allow us to learn about optimal techniques for encoding and
explaining the strategic tasking knowledge that is interfaced with that
system's domain knowledge.

B. Requirements for Continued SUMEX Use

All the work we are doing is totally dependent on the SUMEX resource.
As of yet, none of these young projects has been sufficiently mature to
justify their transfer to one of the SUMEX personal workstations, so the
KI-10's continue to be key elements in our research plan.

In addition, we have long appreciated the benefits of GUEST and
network access to the programs we are developing. SUMEX greatly enhances
our ability to obtain feedback from interested physicians and computer
scientists around the country. As our programs mature in the months ahead,
it will become increasingly important that we be able to make them
available for demonstration and for access by distant collaborators via the
SUMEX network.

C. Requirements for Additional Computing Resources

The mainframe machine should continue to provide a suitable
environment for our work in the months ahead. We have no plans to transfer
to other hardware soon. However, the response time on the main machine
continues to be a major problem, even though the SUMEX workstations have
provided additional cycles to permit off-loading of some users from the
PDP-10.

D. Recommendations for Future Community and Resource Development

We concur with the suggestion, expressed by some of the other project
reports, that consideration should be given to an upgrade of SUMEX to a
more modern central mainframe machine.

II.A.1.5   MOLGEN Project

MOLGEN - A Computer Science Application to Molecular Biology

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I. SUMMARY OF RESEARCH PROGRAM

    A. Project Rationale

    The MOLGEN project has focused on research into the applications of
symbolic computation and inference to the field of molecular biology. This
has taken the specific form of systems which provide assistance to the
experimental scientist in various tasks, the most important of which have
been the design of complex experiment plans and the analysis of nucleic
acid sequences. We plan to expand and improve these systems and build new
ones to meet the rapidly growing needs of the domain of recombinant DNA
technology. We do this with the view of including the widest possible
national user community through the facilities available on the SUMEX-AIM
computer resource.

    It is only within the last few years that the domain of molecular
biology has needed automated methods for experimental assistance. The
advent of rapid DNA cloning and sequencing methods has had an explosive
effect on the amount of data that can be most readily represented and
analyzed by computer. Moreover we have already reached a point where
progress in the analysis of the information in DNA sequences is being
limited by the combinatorics of the various types of analytical comparison
methods available. The application of judicious rules for the detection of
profitable directions of analysis and for pruning those which obviously
lack merit will have an autocatalytic effect on this field in the immediate
future.

    The MOLGEN project has continuing computer science goals of exploring
issues of knowledge representation, problem-solving, and planning within a
real and complex domain. The project operates in a framework of
collaboration between the Heuristic Programming Project (HPP) in the
Computer Science Department and various domain experts in the departments
of Biochemistry, Medicine, and Genetics. It draws from the experience of
several other projects in the HPP which deal with applications of
artificial intelligence to medicine, organic chemistry, and engineering.

We have begun a transition from being primarily a computer science research project to being an interdisciplinary project with a strong applications focus. The tools that we have already developed will be improved to the point where they make a significant contribution to both research and engineering in the domain of molecular biology.

B. Medical Relevance and Collaboration

The field of molecular biology is nearing the point where the results of current research will have immediate and important application to the pharmaceutical and chemical industries. Already, clinical testing has begun with synthetic interferon and human growth hormone produced by recombinant DNA technology. Governmental reports estimate that there are more than 200 new and established industrial firms already undertaking product development using these new genetic tools.

The programs being developed in the MOLGEN project have already proven useful and important to a considerable number of molecular biologists. Currently several dozen researchers in various laboratories at Stanford (Prof. Paul Berg's, Prof. Stanley Cohen's, Prof. Laurence Kedes', Prof. Douglas Brutlag's, Prof. Henry Kaplan's, and Prof. Douglas Wallace's) and over 400 others throughout the country are using MOLGEN programs over the SUMEX-AIM facility. We have exported some of our programs to users outside the range of our computer network (University of Geneva [Switzerland], Imperial Cancer Research Fund [England], and European Molecular Biology Institute [Heidelberg] are examples).

C. Highlights of Research Progress

Accomplishments:

The current year has seen intensive work on what might be termed a second-generation experiment design system. We have also begun to explore the capabilities of the SUMEX Dolphin computers for MOLGEN work. This section will summarize those projects as well as continuing work on other MOLGEN research.

1. Representation Research

The domain of molecular biology has proven a fruitful testbed in the development of a flexible software package, the Unit System, for symbolic representation of knowledge. The package is already in use by a variety of research projects both within the Heuristic Programming Project at Stanford and at other institutions. It provides for acquisition and storage of many different types of knowledge, ranging from simple declarative types like integers and strings to complex declarative types like nucleic acid restriction maps to procedural types like a rule language in a subset of English.

However, it has become clear that there are several bottlenecks in practical use of the Unit System on the main SUMEX computer system. First, there is the address space limitation. It is well known that Interlisp

programs on the DEC 10 and 20 series of computers have relatively little
user memory available. Several of the MOLGEN knowledge bases have already
exceeded that limitation. Second, it is clear that communication with the
Unit System via 1200 baud display terminals does not allow the presentation
of enough information at a fast enough speed. We are very optimistic that
both of these problems will be solved by use of the Xerox Dolphin
workstation computer. The Dolphin has a far larger address space than DEC
10 and 20 computers, as well as a large bit-map display with window and
graphic capabilities.

Our initial experiments on the Dolphin have been encouraging. We
were able to move the Unit System to the Dolphin with almost no
difficulties. We have implemented a simple display interface that allows
far more information to be directly accessible to the knowledge base
builder. We are now in the state of allowing several MOLGEN biologists to
begin using the Dolphins as practical knowledge entry tools.

2. Planning Research

The problem of designing laboratory experiments in molecular biology
has been fundamental to MOLGEN research. Previous work consisted of two
major subparts, each resulting in a doctoral thesis in computer science.
The two systems, developed by Peter Friedland and Mark Stefik, produce
reasonable experiment designs on test problems suggested by laboratory
scientists.

The last year has seen the design and construction of a hybrid
planning system combining the best of the previous two experiment planning
systems. This work was done by Yumi Iwasaki, a master's candidate in
computer science. The system, known as SPEX, uses Friedland's method of
experiment design by the stepwise refinement of abstracted or "skeletal"
plans, and Stefik's method for planning strategy control by "layers of
planning spaces." It also keeps extensive records of all decisions made
during the planning process; this will allow SPEX to be used as an
experimental laboratory for work on explanation, verification,
optimization, and debugging of plans.

In addition to the above general planning work, Rene' Bach has
designed and implemented a small expert system for the problem of
suggesting sequencing strategies for large nucleic acid molecules. The
system, MAXAMIZE, was built entirely within the Unit System. It takes
information about a given molecule (restriction map, already sequenced
regions, etc.) along with information specific to the laboratory (how many
bases may be comfortably sequenced by that lab, enzymes available, etc.)
and provides an optimal or near-optimal strategy.

3. Knowledge Base Construction

Over seven man-years have now been spent in constructing knowledge
bases for various fields of molecular biology. A new project was initiated
this year with Professors Stanley Falkow and Esther Lederberg of the
Stanford Medical Microbiology Department. Also, an entire series of

knowledge-base development projects has begun at the Imperial Cancer
Research Fund in London, England, using the Unit System which was exported
to a DEC 2060 at the ICRF.

The knowledge bases developed under the MOLGEN grant have begun to
find their way into the daily laboratory practice of many of the scientists
associated with the project. They have a provided a mechanism for managing
the explosive growth of data and strategies in many areas of molecular
biology without the necessity of building special purpose systems for each
area. Also, the expert scientists themselves have been able to design and
build their own systems, avoiding the time and reliability problem of a
knowledge base passing through the filter of a computer scientist
intermediary. The knowledge bases have served as "intelligent
encyclopedias," as simulation systems, and as training vehicles.

It should also be noted that the Unit System allows for the easy
transfer of knowledge from one knowledge base to another, and indeed the
various expert molecular biologists have freely shared information as they
work on related knowledge bases.

4. Other Applications of Symbolic Computation to Molecular Biology

MOLGEN work in the previous year has concentrated on basic research,
there now being adequate commercial avenues for applied development of
symbolic computation tools for molecular biology. However, we have
continued to support large-scale guest access to the SEQ program for
nucleotide sequence analysis and the Los Alamos Nucleic Acid Sequence
Library. The GENET guest community has continued to grow. Because of this
growth, a committee, headed by Professor Allan Maxam of Harvard University,
has been formed to set policy for future guest access via SUMEX or other
possible nationally networked systems.

We believe that GENET has been a clear demonstration that SUMEX is a
model for building national communities of research scientists. We hope
that the GENET committee will find adequate mechanisms for supporting this
community without detracting from the primary research goals of SUMEX-AIM.

Research in Progress:

The next year will be spent in testing the SPEX experiment design
system in the domain of cloning experiments, continuing the testing of the
Dolphins for knowledge acquisition, and starting to explore the areas of
plan debugging and explanation. We are also beginning work on automatic
acquisition of experiment design strategies and discovery of gene
regulation signals in the nucleic acid sequence library.

1. Cloning Advisory System

The SPEX experiment design system is now operational. In parallel
with its development, we have begun to construct a large knowledge base
specific to tools and techniques used during cloning experiments. The
basic strategy or skeletal plan of all cloning experiments is quite

straight-forward: First, isolate the piece of DNA you wish to clone,
second, select a vector to carry the clone, third, insert the DNA into the
vector, fourth, select a host for expression of the hybrid molecule, fifth,
insert the hybrid into the host, and sixth, select for the protein or
nucleic acid product that was the eventual goal of the cloning experiment.
Following this skeletal plan, the cloning knowledge base will contain
information on DNA isolation methods, cloning vectors, insertion methods,
hosts, host insertion methods, and selection methods. Professor Stanley
Falkow and several others are assisting us in the building of this expert
knowledge base.

We expect the cloning knowledge base to be essentially complete by
the end of this year. We will also have accumulated substantial experience
with running SPEX on that knowledge base. Sometime during this year, we
will release the cloning advisory system to several of our molecular
biologist colleagues for further testing and improvement. We also hope to
gain more information about experiment design heuristics in general, using
SPEX as a laboratory for this testing. The issues to be studied include
optimal methods for plan expansion: depth-first breadth-first, and
heuristic are among the possibilities, and how abstract or detailed to make
the skeletal plans which drive the experiment design system.

2.   Research on Plan Debugging and Explanation

SPEX keeps complete records of all decisions made during the course
of designing an experiment. These include strategic decisions as to which
general planning heuristics to employ and which domain-specific skeletal
plans to use, as well as tactical decisions made in the course of choosing
specific operators to instantiate a plan step. In addition, SPEX keeps a
dynamic model of the world state as assumed after the execution of each
plan step.

We will now make use of this information to add plan explanation and
debugging facilities to the SPEX system. Users of SPEX will be able to
easily determine which strategies and rules led to each planning decision.
A limited querying interface will be added, probably more graphical through
the Dolphin facilities than natural language.

Plan debugging can take two general forms. First, the debugging of
plans generated originally by SPEX. In that case the debugging process
will consist of attempting to ascertain the point at which the plan failed
by comparing the actual state of the experiment at various points with
SPEX's model of the world at those points. Presumably this will lead to
identifying the instantiation step which failed; if no step failed, then
the skeletal plan which generated the experiment design must have been
faulty. In the former case, an alternate instantiation will be suggested
(and possibly the knowledge base updated to reflect the error). In the
latter case, an alternate skeletal plan will be used and the faulty one
will be corrected (initially manually, but we hope by some automated means
in the future).

The second general form of plan debugging occurs when the plan was generated outside of SPEX. In this case a plan verification step must first take place to ensure that the original plan design and model correspond to the knowledge base. If errors are found, then alternatives will be suggested, if not then debugging will proceed as above.

3. Further Developments of the Unit System

Considerable effort will be spent in taking full advantage of the many capabilities for graphical interaction of the Dolphin. We expect improvements to be opportunistic as our molecular biologist collaborators begin to do their routine knowledge base development on the Dolphin.

4. Work in Plan and Model Discovery

Initial exploration has begun in two new projects in what might be termed automated knowledge acquisition. The first has to do with acquiring skeletal plans for the SPEX knowledge base. Currently skeletal plans are described directly by molecular biologists. We would like to explore means of automatically extracting these abstracted plans from successful experiment designs, either from the literature or in an interactive mode from scientists. One of the major questions to be resolved concerns determining how specific or abstract to make the skeletal plans; specificity leads to efficiency, abstractness leads to generality--a balance is needed.

The second project is in the automated discovery of rules for the regulation of genes from the data stored in the nucleotide sequence library. It is well known that close to 90% of all DNA in higher organisms does not code directly for proteins, but has to do with the regulation of those genes that do code for proteins. There is currently no widely believed mechanism which explains even an appreciable fraction of regulatory processes. The information used for gene regulation is stored somewhere within the nucleotide sequence itself. We have access to a large collection of such sequence data (over 600000 bases) and to the best currently available sequence analysis system (SEQ). We plan to add semantic pattern finding procedures to SEQ, probably through a Unit System interface, that will allow proposed theories to be tested on the data base.

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E. Funding Support

Computer Science, Laurence H. Kedes, Investigator, Howard Hughes Medical
Institute and Associate Professor of Medicine, and Douglas L. Brutlag,
Associate Professor of Biochemistry. MOLGEN is currently funded from 10/81
to 9/82 at $159,785 including indirect costs as the second year of a three
year renewal.

II.   INTERACTIONS WITH THE SUMEX-AIM RESOURCE

       SUMEX-AIM continues to provide the bulk of our computing resources.
The facility has not only provided excellent support for our programming
efforts but has served as a major communication link among members of the
project. Systems available on SUMEX-AIM such as INTERLISP, TV-EDIT, and
BULLETIN BOARD have made possible the project's programming, documentation
and communication efforts. The interactive environment of the facility is
especially important in this type of project development.

       Unfortunately, the computing environment at SUMEX has suffered in the
recent past from heavy demands on cycle time creating serious real-time
delays for programmers and knowledge-base building especially. The Units
Editor is especially sensitive because of its relatively large demands on
cpu-resources. Accordingly, a significant fraction of the MOLGEN group
activity has been transferred to the SCORE computer in the Department of
Computer Science at Stanford. When SUMEX hardware is updated, we
anticipate that its response time will improve and the MOLGEN computing
will return full time to SUMEX. It is clear, however, that the MOLGEN
project continues to thrive and prosper because of the computing
environment only available at SUMEX: the interactive environment including
instantaneous communications among collaborators who are physically distant
(even on the Stanford campus), and especially the unique telecommunications
facilities that have allowed the development of the GENET community with
its access to MOLGEN applications tools are two clear examples.

       We have taken advantage of the collective expertise on medically-
oriented knowledge-based systems of the other SUMEX-AIM projects. In
addition to especially close ties with other projects at Stanford, we have
greatly benefitted by interaction with other projects at yearly meetings
and through exchange of working papers and ideas over the system.

       The ability for instant communication with a large number of experts
in this field has been a determining factor in the success of the MOLGEN
project. It has made possible the near instantaneous dissemination of
MOLGEN systems to a host of experimental users in laboratories across the
country. The wide-ranging input from these users has greatly improved the
general utility of our project.

       We find it very difficult to find fault with any aspect of the SUMEX
resource management. It has made it easy for us to expand our user group,
to give demonstrations (through the 20/20 adjunct system), and to
disseminate software to non-SUMEX users overseas.

III. RESEARCH PLANS

A. Justification and Requirements for Continued SUMEX Use

The MOLGEN project depends heavily on the SUMEX facility. We have already developed several useful tools on the facility and are continuing research toward applying the methods of artificial intelligence to the field of molecular biology. The community of potential users is growing nearly exponentially as researchers from most of the bio-medical fields become interested in the technology of recombinant DNA. We believe the MOLGEN work is already important to this growing community and will continue to be important. The evidence for this is an already large list of pilot exo-MOLGEN users on SUMEX.

SUMEX is currently having difficulty meeting the research needs of the MOLGEN project adequately. We expect to need more file space as our knowledge bases grow; perhaps an additional 5000 disk blocks in the next few years for that work. Our real difficulties will come in the applications testing of MOLGEN tools. We support with great enthusiasm the acquisition of satellite computers for technology transfer and hope that the SUMEX staff continue to develop and support these systems. One of the oft-mentioned problems of artificial intelligence research is exactly the problem of taking prototypical systems and applying them to real problems. SUMEX gives the MOLGEN project a chance to conquer that problem and potentially supply scientific computing resources to a national audience of bio-medical research scientists.

II.A.1.6   MYCIN Projects Group

MYCIN Projects

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I.   SUMMARY OF RESEARCH PROGRAM

A. Project Rationale

The MYCIN Projects are a set of related research programs, each
devoted to the development of knowledge-based expert systems for
application to medicine and the allied sciences. The name was derived from
our first system, the MYCIN program. That research has now given way to
three sub-projects (EMYCIN, GUIDON, and ONCOCIN), each of which is
discussed in the sections that appear below. The key issue for all sub-
projects has been to develop programs that can provide advice similar in
quality to that given by human experts, and to develop systems that are
easy to use and acceptable to physicians and medical students.

The success of the original MYCIN infectious disease consultation
program has led us to try to generalize and expand the methods employed in
that program to a number of ends:

(1) to develop consultation systems for other domains (our generalized
    system-building tool is known as "Essential MYCIN", or EMYCIN, and
    has been applied in several new areas; ONCOCIN is our a more recent
    consultation system and was inspired by EMYCIN although it is an
    entirely new program);

(2) to explore other uses of the MYCIN knowledge base (our tutoring
    system, GUIDON, uses the infectious disease knowledge in MYCIN to
    teach medical students about diagnosis and management of infections);

(3) to continue to improve the interactive process, both for the
    developer of a knowledge-based system, and for the user of such a
    system (both EMYCIN and ONCOCIN have stressed simplified techniques
    for interacting with a knowledge base and entering data); and

(4) to experiment with alternate techniques for knowledge representation,
    recognizing that the pure production rule method used in MYCIN was
    inadequate at times and frequently led to confusion regarding the
    separation of strategy or control processes from domain knowledge
    (ONCOCIN uses production rules as only one of several knowledge

representation techniques, and the work on GUIDON has led to a more robust revised version of MYCIN known as NEOMYCIN).

B. Medical Relevance and Collaboration

By utilizing our EMYCIN system to collaborate on building the PUFF program, we learned that it is possible in a short period of time to develop a clinically useful consultation system using the domain-independent parts of MYCIN. EMYCIN has since been applied in a number of additional medical domains. With each successive application we learn more about the representation of medical knowledge and the scope and limitations of the production rule formalism used in EMYCIN. For example, it has become clear that "shallow" rules relating signs and symptoms to diagnoses through a few intermediate concepts can be sufficient for high performance in medical diagnosis. On the other hand, such shallow rules are not always sufficient for teaching medical students because they lack the deeper causal links needed for justifying and remembering the more shallow associations (see GUIDON discussion below).

Although EMYCIN was not used to build our new ONCOCIN program, the lessons learned in building prior production rule systems have allowed us to create a large oncology protocol management system much more rapidly than was the case when we started to build MYCIN. We introduced ONCOCIN for use by Stanford oncologists in May 1981. This would not have been possible without the active collaboration of Stanford oncologists who helped with the construction of the knowledge base and also kept project computer scientists aware of the psychological and logistical issues related to the operation of a busy outpatient clinic.

In addition, there is a growing realization that medical knowledge, originally codified for the purpose of computer-based consultations, may be utilized in additional ways that are medically relevant. Using the knowledge to teach medical students is perhaps foremost among these, and GUIDON continues to focus on methods for augmenting clinical knowledge in order to facilitate its use in a tutorial setting. A particularly important aspect of this work is the insight that has been gained regarding the need to structure knowledge differently, and in more detail, when it is being used for different purposes (e.g., teaching as opposed to clinical decision making). This aspect of the GUIDON research has led to the development of a modified version of MYCIN, NEOMYCIN, which is an evolving computational model of medical diagnostic reasoning that we hope will enable us to better understand and teach diagnosis to students.

C. Highlights of Research Progress

1. Accomplishments This Past Year

EMYCIN

In the last year, work on EMYCIN was brought to a conclusion. The program was "frozen" and documentation completed. The program remains a

vehicle for computer science research, but there is no funding for
continued development of the tool. Our work centered around the following
activities:

(1) One of the most significant experiments with the program is the
     development of ROGET, a knowledge acquisition program that aids a
     system builder with the definition and creation of a new knowledge
     base for an EMYCIN system. ROGET itself is written in EMYCIN.
     Design was completed in this year and programming started, as
     described in more detail below.

(2) We continued to help users in the SUMEX community with new
     applications of EMYCIN and with questions about on-going
     applications. These are described below.

(3) Buchanan and Shortliffe have begun writing up the results of many
     years' work on MYCIN and EMYCIN. Because we view AI as an
     experimental science, we believe there is much to be learned in a
     detailed analysis of the experiments performed with rule-based
     consultation systems by the MYCIN group. This work is also briefly
     described below.

ROGET: A KNOWLEDGE-BASED SYSTEM FOR THE DESIGN OF EXPERT SYSTEMS

ROGET is a program under construction which is designed to assist an
expert with the initial formulation of an EMYCIN consultant. The ROGET
system is intended (1) to help the expert properly delimit the scope and
function of a consultant, (2) to identify the major types of inferences it
will make and their requisite facts, and (3) to educate the expert about
the use of the particular representation system being used to construct the
consultant.

Current expert-system building tools provide little or no guidance to
an expert during the initial formulation of a consultant system. A
knowledge engineer provides the expert with advice about what tasks the
consultant might have to perform and, more importantly, how to employ a
particular representation system to achieve this performance. Previous work
by Davis, Reboh, and van Melle have emphasized tools and techniques that
are appropriate only considerably after the basic "inference structure" of
the consultant has already been designed and placed on-line. These tools
are meant to assist the refinement of a knowledge base and they make the
implicit assumption that the expert is already fairly familiar with (i.e.,
has at least reading knowledge of) the representation system.

In contrast, the primary function of ROGET will be to guide and
instruct the expert as he selects the relevant concepts and inferences
required to solve a problem, indicate how they have been typically
represented within a specific representational framework, and provide
hands-on instruction regarding the tool's actual use. ROGET will provide an
interactive environment in which the expert can construct a prototypical
inference structure for a consultant and some of its "connective tissue".
In this manner, the expert can become familiar enough with the

representation system to use the existing knowledge acquisition tools to
further refine the knowledge base.

This research is predicated on the observation that several computer-
based consultants, particularly of the diagnostic variety, employ similar
types of inference, provide similar types of advice, and require similar
types of data. Furthermore, the actual representation of the domain-
specific counterparts of this knowledge within a particular representation
system has also taken rather stereotypic forms. This research will attempt
to capitalize on these "standard" forms by representing this type of
knowledge and developing a system that will use it to guide the acquisition
of new expertise in different domains.

Helping the expert identify an appropriate task for the consultation
system is a major function of any knowledge engineer. ROGET will assist
the expert with the proper scoping of the consultant by identifying the
applicable inferences for this problem and by suggesting the most
appropriate inferences for the current consultant to make. These
suggestions can be made on the basis of information about the experts
background, his experience with system construction, any constraints on his
time and other resources, and, most importantly, a description of the
"consultation setting", where and how the consultant will interact with
prospective clients.

The education of the expert about the intricacies of a particular
representation system is currently accomplished primarily through the use
of examples taken from other expert systems constructed with this tool.
ROGET will mimic this behavior as well. In addition to providing domain-
specific examples of each of the inference, advice, and data types
mentioned above, ROGET will also give examples of how the inference engine
interprets example knowledge bases. Again, knowing about the consultation
setting will help ROGET focus attention on specific portions of complete
knowledge bases that demonstrate how certain types of knowledge are needed
to make certain features of the inference engine run.

ROGET will be developed to assist an expert design an EMYCIN
consultant. To represent the consultation setting a simple, script-like
representation of the major events, people, and information flow for
various consultation scenarios will be developed. In the case of diagnostic
consultants this representation must be adequate to capture most "clinical
settings" at some fairly high level of abstraction. The expert will then
identify specific types of people, information, and events that are
applicable to his domain and use this perspective to identify points in the
information flow where a consultant would be most appropriate and
effective.

This same script-like representation will also be used to capture the
stereotypic inference steps that EMYCIN diagnostic consultants employ, to
identify what types of data are expected at certain points in the
consultation, and to understand how the advice the consultant gives will be
used by the client. The current inference engine of EMYCIN will be modified
to operate on the basis of this new representation of control knowledge, in

addition to the normal backward-chaining of rules. Furthermore, the
explanation machinery will be updated to provide alternative explanations
of the consultant's line of reasoning about a case based on this
representation of control.

Representation of this scriptal knowledge, the hierarchies of
inferences, data, and advice for each task, and the examples themselves
will employ a simple UNITS-style representation system embedded within the
current EMYCIN system. A trace of the design process for individual
consultants will be kept and used as the basis for "developmental
explanations" that could be used to indicate why the consultant is scoped
to perform certain tasks to the exclusion of others and to identify parts
of the knowledge base that are under-developed.

The initial ROGET system will be considered "done" when it can
successfully scope and design the basic inference structures for the
following EMYCIN consultants: MYCIN, GRAVIDA, PUFF, and SACON. At least
one general consultant task must be represented; currently this task is
diagnosis and simple therapy selection. Another potential task would be the
design of objects or plans by configuring schematic designs (ROGET itself,
R1, and Friedland's MOLGEN are examples of this type of "configuration"
consultant).

In summary, then, this project seeks to make explicit some important
aspects of knowledge-engineering expertise. It will identify a taxonomy of
the kinds of expert systems in terms of common tasks and how their place in
a larger information context effects their final design. This knowledge
will be used to help scope the initial design of a consultant and to
educate the expert about a representation system. It is concerned with
identifying expertise required for initial expert system design and this,
in turn, will require an analysis of the nature of expertise itself and the
current limitations on its representation, especially within the EMYCIN
system.

EMYCIN Applications

We spoke with many physicians around the country about potential
applications of EMYCIN in their own work. We encouraged several to submit
applications to SUMEX for pilot status. We collaborated with one physician
at length on a prototype system named COAGS, described below.

COAGS

During January and February of 1982 James Bennett and Robert T. Wall,
M.D., a hematologist at the Stanford Medical School, collaborated to
construct a prototype hematology consultant using the EMYCIN system. The
consultant, named COAGS, deals with patients about to undergo surgery when
routine coagulation screening tests indicate a potential bleeding problem.
The surgeon will typically refer the patient to a hematologist for a more
complete set of tests to confirm the disorder and to receive a
recommendation about any treatment that might allow the operation to
proceed.

The consultant requests that various collections of tests be performed in an attempt to confirm that a problem does exist. Most of the coagulation tests are not particularly accurate and repetition of critical tests with appropriate controls is an essential part of hematologists' attempts at confirmation. The tests can be expensive to perform and the consultant has been constructed to minimize the cost of diagnosing the most specific bleeding disorder possible.

The EMYCIN system did not require any improvements or modifications to represent Dr. Wall's clinical expertise. The consultant was constructed in a two month time period on weekly meetings between expert and knowledge engineer. We believe that this represented approximately 6 person-days.

ANALYSIS OF EXPERIMENTS WITH MYCIN & EMYCIN

Artificial Intelligence, or AI, is largely an experimental science in that at least as much progress has been made by building and analyzing programs as by examining theoretical questions. MYCIN is one of several well-known programs that embody some intelligence and provide data on the extent to which intelligent behavior can be programmed. As with other AI programs, its development was slow and not always in the forward direction. But we feel we learned some lessons in the course of over eight years of work on MYCIN, and related programs. The purpose of this analysis, then, is to share the results of many experiments performed in that time, and to paint a coherent picture of the work.

The MYCIN project is one of the clearest representatives of the experimental side of AI. It was begun in the spring of 1972 with a set of discussions among medical school and computer science researchers interested in applying more intelligence to computer programs that interpret medical data. Shortliffe's PhD dissertation in 1974 discusses the problem context and the MYCIN program that implemented a solution.

In itself, the 1974 version of MYCIN represents an experiment. We were testing the hypothesis, tentatively advanced in the DENDRAL work, that a production rule formalism was sufficient for the high performance, flexibility, and understandability that we demanded in an expert system. The positive answer to this question is one of the best-known lessons in the history of AI.

In addition to, or rather because of, the original MYCIN program and the knowledge of blood infections and meningitis that was accumulated for that work, many derivative projects explored variations on the original design. In a book-length analysis currently underway, we are examining many of the experiments that evolved in the period from 1972 to 1980 based on the 1972-74 design effort. We have chosen those pieces of work which, at least in retrospect, can be seen as posing clear questions and producing clear results that were documented in the AI or medical literature, or occasionally in technical reports.

Some of those other projects are discussed in this annual report, for example GUIDON and NEOMYCIN. Others have come to a logical conclusion:

TEIRESIAS, VM, CENTAUR, and now EMYCIN. In those major pieces of work, and several others as well, the questions posed and the answers we derived from program construction and testing are important scientific results. Documenting these results will, we hope, benefit the entire AI community.

Because almost all of the computing work was done on the SUMEX-AIM computer resource at Stanford, we recognize that this set of experiments would not have been possible without funding from the Division of Research Resources of the National Institutes of Health.

GUIDON

The purpose of the GUIDON project is to develop a tutorial program that uses a knowledge-based system for teaching material. Research over the past five years has shown that a knowledge representation that is adequate for developing a consultation system with good performance may be inadequate for teaching. Our study of MYCIN's meningitis rule base (1979-1980) led us to develop a new representation in which kinds of knowledge important to teaching are represented separately and explicitly. Using this new system, which we call NEOMYCIN, we proceeded in 1981 to develop a student modelling program that will be used in a tutorial program for teaching diagnostic strategies to students.

CONTINUING DEVELOPMENT OF NEOMYCIN

NEOMYCIN is distinguished from other AI consultation programs by its uses of an explicit set of domain-independent meta-rules for controlling all reasoning. These rules constitute the diagnostic procedure that we want to teach to students: the stages of diagnosis, how to focus on new hypotheses, and how to evaluate hypotheses. It has been a major undertaking, separate from the problem of representing disease knowledge, to design and test this diagnostic procedure. For example, this year we extracted the forward-directed reasoning that had previously been embedded in NEOMYCIN's interpreter and represented this as a "task" with 4 meta-rules (one meta-rule each for: follow-up questions, antecedent rules, trigger (hypothesis-suggesting) rules, and forward application of rules that confirm hypotheses already under consideration). By the end of the year, attention turned to extending NEOMYCIN's disease knowledge to include more diseases that present similarly to, or are confused with, meningitis.

THE IMAGE STUDENT MODELLING PROGRAM

Teaching diagnosis involves recognizing the intent behind a student's behavior, so that missing knowledge can be distinguished from inappropriate strategies. The teacher interprets behavior, critiques it, and provides advice about other approaches. To do this successfully and efficiently in a complex domain, the teacher benefits from multiple, complementary modeling strategies. IMAGE is a student modelling program that uses NEOMYCIN's meta-rules and disease knowledge to understands student diagnostic plans by a dual search strategy. IMAGE first produces multiple predictions of student behavior by a model-driven simulation of NEOMYCIN. Focused, data-driven searches then explain incongruities. By supplementing

each other, these methods lead to an efficient and robust plan understander.

A model of student strategies in medical diagnosis must disambiguate the possible purposes and knowledge underlying the student's actions. The approaches followed by other plan recognizers and student modelers are not sufficient here because:

(1) the complex domain makes thorough searches impractical, whether top-down or bottom-up;

(2) we are not modeling only facts and rules used in isolation, but also the procedures for applying them;

(3) every one of the student's actions must be monitored in case the teaching module decides to interrupt;

(4) his behavior must be evaluated and not just explained; and

(5) we might not have any explicit goal statements from the student, so we expect to rely only on his queries for problem data as evidence for his thinking.

The IMAGE program is now operational and will soon be tested in experiments with medical students.

SHORT-TERM PLANS FOR NEOMYCIN AND GUIDON

The next stages in the development of the GUIDON2 tutorial program (using IMAGE and NEOMYCIN) are:

(1) additions to NEOMYCIN's disease knowledge so we can fairly evaluate the program's focussing strategies;

(2) experimentation to determine the capabilities of IMAGE for explaining student (non-expert) behavior;

(3) development of an explanation system for NEOMYCIN, to serve as a testbed for explanations to be generated by the tutorial program (including use of a student/user model and condensation methods for presenting explanations at the appropriate level of detail) (Drs. Clancey and Shortliffe, and their graduate students, are collaborating on this project);

(4) design of various teaching scenarios for teaching NEOMYCIN's meta-rules, and representation of this teaching knowledge in a prototype version of GUIDON2.

ONCOCIN

The oncology chemotherapy consultation system, named ONCOCIN, has achieved many of its goals since work on the project began in July 1979. We are developing an interactive system to be used by oncology faculty and fellows in the Debbie Probst Oncology Day Care Center at Stanford University Medical Center.

The ONCOCIN research goals are directed both towards the basic science of artificial intelligence and towards the development of clinically useful oncology consultation tools.

We have undertaken AI research with the following aims:

(1) to implement and evaluate recently developed techniques designed to make computer technology more natural and acceptable to physicians [formal studies are underway to assess the success of our prototype system in achieving these goals];

(2) to extend the methods of rule-based consultation systems to interact with a large database of clinical information; and

(3) to continue basic research into the following problem areas: mechanisms for handling time relationships, techniques for quantifying uncertainty and interfacing such measures with a production rule methodology, approaches to acquiring knowledge interactively from clinical experts, assessment of knowledge base completeness and consistency.

The clinic system itself is intended to have a number of performance capabilities when it is completed. We outline those goals here, citing those which have been completely or partially accomplished and indicating those that have yet to be achieved:

(1) to assist with identification of current protocols that may apply to a given patient [not yet undertaken; will not be relevant until more protocols than lymphomas have been encoded]

(2) to assist with determining a patient's eligibility for a given protocol [not yet undertaken because lymphoma patients have been fully screened before they come to chemotherapy clinic];

(3) to provide detailed information on protocols in response to questions from clinic personnel [a query system is just now being started; it is being designed to assist both physicians and those building a protocol knowledge base];

(4) to assist with chemotherapy dose selection and attenuation for a given patient [fully implemented and being evaluated for patients under treatment for lymphoma];

(5) to provide reminders, at appropriate intervals, of follow-up tests and films required by the protocol in which a given patient is enrolled [fully implemented and being evaluated for patients under treatment for lymphoma];

(6) to reason about managing current patients in light of stored data from previous visits of (a) the individual patients [partially achieved, but much work remains], or (b) the aggregate of all "similar" patients [not yet attempted].

In the early years of the project, we developed a prototype of the ONCOCIN consultation system, drawing from programs and capabilities developed for the EMYCIN system-building project. We also undertook a detailed analysis of the day-to-day activities of the Stanford Oncology Clinic in order to determine how to introduce ONCOCIN with minimal disruption of an operation which is already running smoothly. Subsequently we completed the development of a special interface program that responds to commands from a customized keypad. Software protocols were developed for achieving communication between the interface program and the reasoning program, and we coordinated the printing routines needed to produce hardcopy flowsheets, patient summaries, and encounter sheets. Finally, lines were installed in the Stanford Oncology Day Care Center, and, beginning in May 1981, eight fellows in oncology began using the system three mornings per week for management of their patients enrolled in lymphoma chemotherapy protocols.

The introduction of the system last year has largely determined the nature of our subsequent research effort. That experience has proved to be a constant source of new ideas and challenges for system improvement. Much of our work in the last year has been in response to the lessons learned by observing the strengths and weaknesses of the computing system now that it is in routine use by physicians. Formal evaluation studies have continued in parallel with the ongoing development effort, and the data collection for them is within three months of completion; we expect to begin data analysis before the end of June.

IMPLEMENTATION OF ONCOCIN IN THE STANFORD ONCOLOGY CLINIC

ONCOCIN was introduced smoothly into the Oncology Clinic in May and June of 1981. Each oncology fellow was given approximately 45 minutes of training in the system's organization and in use of the terminal interface. They found it to be extremely easy to learn, and additional help was seldom needed after the training session. During the first several months our data manager was routinely present in the clinic to assist with questions, but it became clear that this was not necessary and since January of this year we have simply asked the physicians to call our project office if a problem arises.

Our Clinical Specialist Dr. Carlson, himself a fellow in oncology, has regularly reported to us the informal reactions of the physicians to the system. We will soon have formal data regarding their attitudes (see Study 1 below), but the informal feedback and suggestions were important

because they led to the design and implementation of several additional
features that have required a good deal of programmer time during the past
year. For example, the communications between the Reasoner and Interviewer
were streamlined for efficiency, and major changes were made to a special
category of data entries known as "special questions". These frequently
annoyed the physicians because they tended to interrupt the smooth flow of
the dialogue with the machine. The revised version of the interface
inserts these questions more benignly into the flowsheet data form.

Two major problems were identified that relate to knowledge
representation in ONCOCIN and control of the reasoning process. One of
these is the inability of the program to manage complex inferences that are
dependent upon analyzing a patient's response to therapy over all previous
visits to the clinic. This kind of analysis is typically ignored in the
formal protocols but is routinely used by experienced oncologists and is,
of course, a desirable capability for a consultation program in this area.
The development of techniques for managing temporal reasoning is a major
interest of our group and is of particular importance in dealing with
expert decision making in a domain such as cancer chemotherapy. Our
oncology collaborators are finding this aspect of their knowledge
particularly difficult to distill, and we expect that our attempts to
develop computer-based approaches to the problem will provide particularly
stimulating challenges in the months ahead.

The second problem was unanticipated until the system was implemented
but is having a major impact on our future plans. ONCOCIN provides a
therapy recommendation for the physician but allows those recommendations
to be overridden if the physician prefers an alternate treatment. If the
fellows change the therapy, they are asked to indicate the reason for
preferring their therapy to what ONCOCIN suggested. When there is a major
disagreement between the fellow and the computer, the physicians seldom
object to indicating the reasons for the change, and their entries have
been useful in further debugging the program's knowledge of lymphoma
treatment. However, the changes they make are frequently minor ones that
are more stylistic than substantive (e.g., some physicians prefer always to
give prednisone in multiples of 20 mgs when they are dealing with large
doses; others prefer to be more precise in their dosing and will ask
patients to take combinations of 5 and 20 mg tablets). In these cases, the
physicians tend to be annoyed by requests for reasons that they have
disagreed with ONCOCIN; they do not see the slight adjustments as being
real disagreements. We have attempted to respond to this problem by
developing criteria for determining when a physician's preferred treatment
is a clinically significant difference from what ONCOCIN has recommended.
The knowledge for determining what changes are "important" is different
from the kind of knowledge in the protocols themselves. It is more
"judgmental" and imprecise, but seems to be an important addition to the
program's knowledge base. Furthermore, this issue, and the observation
that the physicians frequently believe they know what to do and would
rather not be bothered by the program's suggestions, has led us to propose
a modified version of ONCOCIN that will verify the physician's plan rather
than offer a therapy recommendation for every case. We have referred to
this new work as "hypothesis assessment" (see below).

EVALUATIONS OF THE ONCOCIN SYSTEM

All three ONCOCIN evaluations are approaching completion, and we expect to have data analyzed and formal reports written by this time next year.

Study I is an evaluation of the program's impact on the attitude of the oncology fellows towards computers in general and ONCOCIN in particular. All physicians were administered questionnaires and structured interviews in the Spring of 1981 before ONCOCIN was introduced. The same questionnaires have recently been distributed to them once again, and follow-up interviews are just now underway. Pre- and post-ONCOCIN results will be analyzed as described in last year's annual report.

Study II is an evaluation of the program's impact on the completeness and accuracy of flowsheet data recorded with and without ONCOCIN. Research programmers have written routines to formally analyze on-line flowsheets for completeness and accuracy. Pre-ONCOCIN flowsheets have been entered into the system exactly as they were originally recorded by the physician. The same analytic routines will analyze these pre-ONCOCIN flowsheets and we will compare the pre- and post-ONCOCIN results. We expect to begin analysis of the post-ONCOCIN data within the next two months when a sufficient number of patient visits have been recorded to permit a statistically significant comparison.

Finally, Study III is examining the comparison between ONCOCIN's therapeutic advice and the treatment decisions made by oncology fellows in the same setting. As described in last year's report, expert evaluators are rating treatment plans without knowing whether the recommendation is that of ONCOCIN or one of the clinic physicians. Over 200 flowsheets are currently being evaluated by Stanford lymphoma experts, and we expect to have their ratings returned for analysis by the end of May.

THE INTERVIEWER/REASONER MODEL

We have continued to refine ONCOCIN's communication system for permitting the physician to interact with a rapid interface program while a slower reasoning program considers treatment options in a second process. After the system was implemented, several unexpected problems resulting from the use of asynchronous processes were discovered and required resolution. For example, the reasoning program has no straightforward way of knowing whether a piece of information that it needs has been skipped on the flowsheet by the physician or whether the doctor has simply not reached that part of the flowsheet yet. Similarly, the Reasoner needs techniques for dealing with changes to flowsheet entries that are made after the initial entry has been transmitted by the Interviewer and has been used by the Reasoner in its decision making process.

HYPOTHESIS ASSESSMENT

As mentioned above, we have decided to consider modifications to ONCOCIN that would permit it to function as an "observer" of the physician's own decisions rather than as a primary source of advice. By permitting the physician to enter his or her own therapy plan on the flowsheet, we can acknowledge the oncologist's ability to reach appropriate therapeutic decisions for most patients. ONCOCIN will simply compare the physician's plan with what it believes is the proper therapy. If the system agrees with the physician, or determines that small differences are clinically insignificant, no advice from the computer will be necessary. If significant disagreements occur, on the other hand, ONCOCIN will need to respond with warnings and explanations for why it feels that an alternate therapy plan may be preferable. Our early experience with ONCOCIN since its clinic implementation suggests that this mode of interaction may be preferred by the clinic physicians. It will require minimal changes to ONCOCIN's decision making approach, but the determination of what differences are clinically significant, and the optimal method for explaining their importance to the physician, will be exciting challenges and important theoretical problems. This work is just getting underway and will receive much attention during the coming year.

QUERY SYSTEM

We have also recently started work on the development of a query system that will permit easy access to the large ONCOCIN knowledge base. Now that we have encoded several hundred rules, it is becoming unwieldy for system builders to work from large hard-copy listings of the knowledge base, and physicians will also require direct access to the program's knowledge. The query system is being designed to permit this kind of access. Rather than dealing with natural language understanding by computer, we are designing ways that menu selection and the high-speed interface can be used to permit access to the information that is needed by a physician or system builder.

VERIFYING THE COMPLETENESS AND CONSISTENCY OF THE KNOWLEDGE BASE

An important question for AI researchers involved with the development of expert systems is how to ascertain that a knowledge base for a consultation program is complete and consistent. Dr. Motoi Suwa, a visitor to Stanford from Japan two years ago, became fascinated with this question and collaborated with us on a formal analysis of the developing ONCOCIN knowledge base. However, the programs that he wrote were never formally linked to our system for writing rules and modifying other parts of the knowledge base. As a result, we have recently spent time modifying his code so that it will operate as an integral part of ONCOCIN. The development of an improved interface to permit this program to be used by our expert collaborators is also planned for the coming year.

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