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COMPUTER RECREATIONS

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COMPUTER RECREATIONS

Exploring the field of genetic algorithms in a primordial computer sea full of flibs

by A. K. Dewdney

Imagine an abstract sea inhabited by abstract organisms called finite living blobs, or flibs. Each flib is equipped with the simplest decision-making apparatus possible. This is the biological equivalent of what computer scientists call a finite automaton. Each flib also contains a single chromosome consisting of a string of symbols that encodes the automaton. The flibs inhabit a primordial, digital soup in constant flux. These changes must be predicted accurately by the flib if it is to survive.

In the primordial soup I recently set simmering in my computer, flibs that predicted poorly died out. The best predictors left progeny that sometimes improved on ancestral performance. Eventually a line of perfect predictors evolved.

Flibs and their evolutionary tendencies illustrate nicely a form of programming known as the genetic algorithm. Pioneered by John H. Holland of the University of Michigan in the 1960's, the technique is sometimes able to solve difficult problems by evolving a sequence of approximate solutions. New solutions are produced by mating the best of the old solutions with one another. Before long a new solution that is superior to its parents appears and joins the list of preferred breeders. Genetic algorithms have been applied with some success to pattern recognition, classifier systems, pipeline operation, symbolic layout and a small number of other problems. In my computer soup the technique yielded superior flibs. Was this success due to the general efficacy of the genetic-algorithm method or to the simplicity of the predictive task facing the flibs? The question is hard to answer. It can be pondered and the underlying phenomenon can be reproduced by any interested reader who has a computer within reach.

A finite automaton has a finite number of states; an input signal causes

it to change automatically from one state to another. The kind of automaton used in a flib also generates signals. Incoming and outgoing signals are represented within the automaton by symbols. When a signal is received, the automaton changes state and emits a second signal.

A state-transition table is useful for representing the process. For example, a finite automaton that is capable of assuming three states, *A*, *B* and *C*, and that can handle afferent and efferent 0's and 1's fits nicely into a 3-by-4 table. For each state the automaton might find itself in, and for each symbol it might receive, there are two entries. The first entry gives the corresponding output symbol; the second entry gives the state that the automaton next assumes:

	0		1	
A	1	B	1	C
B	0	C	0	B
C	1	A	0	A

The automaton represented by this table might well find itself in state *C* at some time. If the automaton receives a 1, the table tells us the automaton will generate a 0 and enter state *A*.

Another representation, easier for

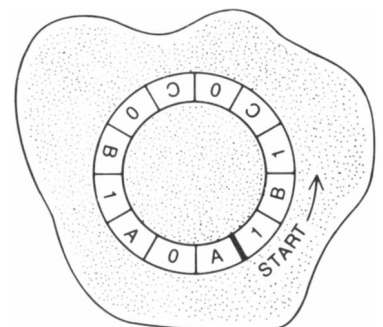
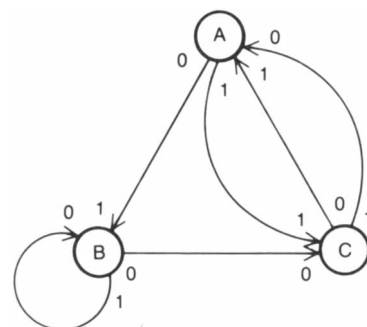
humans to read, is the state-transition diagram, in which circles represent states and arrows represent transitions. If an automaton goes from one state to another when it receives a specific symbol, an arrow should be drawn from one state circle to another. The arrow should be labeled both with the input symbol that caused the transition and with the resulting output symbol [see illustration below].

A finite automaton always begins its operations in a specific state, called the initial state. At each tick of an imaginary clock a new symbol arrives, a new symbol leaves, and a new state is entered. The automata used in my flibs all send and receive the same two symbols, 0 and 1.

How is one to interpret the behavior of a flib if so little is known about the creature's biology? Therein lies the joy of abstraction. The symbols received by the automaton are merely sensory messages from the environment. In corresponding fashion, an output symbol can be viewed as a response by the organism to the environment's most recent condition.

The concept of a flib is so flexible that input and output can represent a great variety of specific biological phenomena. For example, an input signal could represent a chemical or temperature gradient. The corresponding output symbol could be a command to an effector that controls cilia, or a spore-forming mechanism. A task of great importance to a creature wishing to evolve to some minimally acceptable level (say that of a university professor) is to predict the environment. To a flib the environment is a seemingly unending sequence of 0's and 1's. Insofar as symbols received indicate significant events, there is clearly some advantage in the ability of a flib to predict the next symbol, particularly if under some more specific interpretation of flib functioning the flib's survival were enhanced.

Most flibs are rather poor at predicting their environment in this sense. For example, the flib described by the



A state-transition diagram (left) and a corresponding flib with its chromosome (right)

state-transition table given above responds to the environmental sequence

0111000010110...

with the outputs

1000011001000...

At each stage of its operation the flib's output is its prediction of the next symbol to arrive from the environment. To find the number of correct predictions shift the output sequence one symbol to the right and compare it bit by bit with the input sequence. Count the number of matching symbols. In this case the flib predicted correctly only six of the 12 incoming symbols, a score that is no better than might result from random guessing.

One can easily demand too much from a finite automaton. Indeed, it is unfair to ask a flib to predict any non-periodic environment. Readers might like to ponder this point for a moment. Why must a perfectly predicted sequence of input symbols consist of the same basic string endlessly repeated?

For example, the 3-state flib that failed the prediction test just set for it succeeds brilliantly on the following environmental sequence:

010011010011010011...

Here the environment marches to the beat of a simple repetition, 010011.

There are several dozen 3-state flibs, but only a few of them can predict this sequence perfectly. Among flibs that have more than three states perfect predictors for a given environmental sequence are rare and become more so as the number of states increases. Predictability depends heavily on the period of the sequence: no n -state flib will ever be able to predict the sequence that results from continued repetition if the basic string of symbols is too long. There is evidently a relation between the number of states a flib can have and the largest period in a sequence that it predicts perfectly. Readers might enjoy discovering the relation for themselves. What is the longest period an n -state flib can predict?

A flib is more than a finite automa-

ton trying to predict its environment; it has a chromosome. Flibs periodically breed (by some unknown method). An examination of the chromosome in its relation to a flib's finite automaton shows how the inherited genes determine the behavior of the offspring. Start with the state-transition table and strip away the rows, one at a time, from top to bottom. Join the rows together end to end and then join the beginning of the string to its end. The result is a circular chromosome.

Before the final joining operation, the chromosome of our 3-state exemplar appears as a string of 12 genes:

1B1C0C0B1A0A

Strictly speaking, the symbols in this string are alleles. An allele is a specific form of a gene that appears at a given locus. As such, a gene can be specified either by its name or by its locus. Thus the seventh symbol from the left controls a flib's output symbol when it is in state B and a 1 is received from the environment. The locus here is 7.

I recently set up a primordial soup containing 10 4-state flibs in my personal computer. Before 1,000 of the time units I call chronons had passed none of the original flibs was alive. All had been replaced by superior predictors. The display screen showed the highest and lowest scores attained in the current population. The lowest score fluctuated a good deal; the highest score crept slowly upward [see illustration at left]. Just when I was beginning to give up hope that a perfect predictor would evolve, one suddenly appeared, whereupon the highest score jumped to 100.

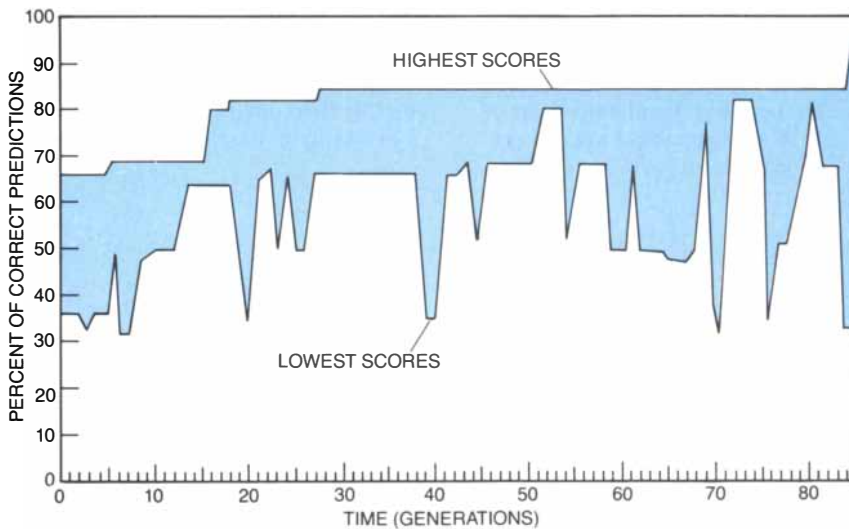
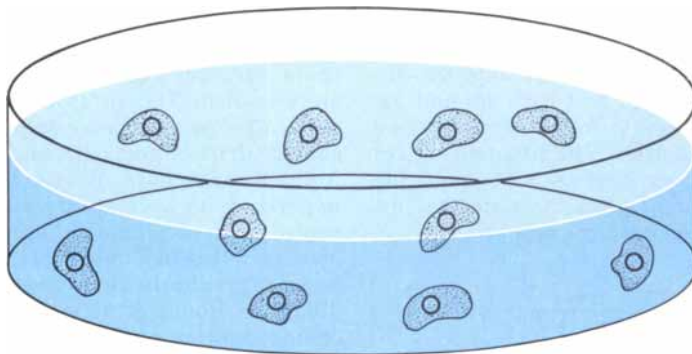
All of this raises the question of just how flibs evolve in my computer soup. Periodically a cosmic ray zips through the broth and strikes a random chromosome at a random locus; the result is that a specific gene is changed from one allele to another. For example, in the following 4-state flib chromosome the gene at locus 3 controls the output symbol for the transition from state A , when the creature receives a 1:

0D1C0D0B1A0C1B1A

A cosmic ray striking this gene changes the chromosome slightly:

0D0C0D0B1A0C1B1A

Mating is the other source of variation in the flib gene pool. During the mating season the highest-scoring flib shuffles genes with a randomly selected flib. The offspring bears a composite chromosome. One part comes from the superior parent, the other from



A soup of 10 flibs (top) evolves a perfect predictor (bottom)

Silent partners in world health

Schistosomiasis affects as many as 200 million people in Africa, Asia, the Middle East, Puerto Rico and Latin America. It is often called "snail fever" because at one stage of their life cycle, *Schistosoma* worms infect snails that live on the bottom of rivers and streams. These parasites invade the skin of humans who drink, wash or swim in contaminated waters. They can cause severe itching, fever, diarrhea, and eventually irreversible damage to the liver. For 16 years, researchers visiting the island of St. Lucia in the Caribbean have been testing the practicality of various methods of control. Three approaches have proven to be most effective.

First, a public health team sprayed the rivers and streams of St. Lucia to get rid of infested snails. New plumbing facilities were constructed to assure a supply of uncontaminated water. Finally, treatment of people carrying the parasite was greatly facilitated by a drug developed and supplied by Pfizer. While previous treatments had to be given by injection, this drug was given orally only once, making it much simpler to reach a large number of people. The total control and elimination of the parasite is not yet a reality, but this combined medical and environmental program has done much to make life better for the people of the island.

Developing a drug such as this is a significant task that takes a decade or more and tens of millions of dollars. It generally involves the synthesis of hundreds of compounds in the organic chemistry laboratory. These compounds are then screened for antiparasitic activity. If one or more of them shows promise, the next step is to do toxicity studies and learn all about how the potential new drugs behave in laboratory animals. Only after completion of extensive, time-consuming animal studies can the drug be tested for safety and effectiveness in humans. And clinical trials in human patients can last for several years. If the clinical trials indicate that the drug should be made

available, new technology must be developed to produce it on a mass basis, and in cases like this, with little if any profitability for the developer.

Drug research and development isn't always "good theater." And it's largely a team endeavor generally without charismatic heroes. The days of Paul Ehrlich and his "magic bullet" are long past. The work of the pharmaceutical industry isn't usually the stuff of TV documentaries. More often, the industry has been the silent partner of government agencies, physicians, nurses and their associates working together to improve public health in St. Lucia and other developing countries.

In the Third World, pharmaceuticals are perhaps even more important than in advanced industrial countries. Often they are the only form of advanced medical technology which is practicable. Other forms of care, such as surgery, are often too cumbersome and too demanding of scarce resources. Drugs, by comparison, are portable, relatively inexpensive and comparatively simple to use.

The vast majority of drugs for the Third World and also for developed countries originate in the pharmaceutical industry. The government agencies do not have the broad expertise or resources for drug development, and medical schools and universities have different missions. Only the major research/pharmaceutical companies have the necessary skills and resources. Most manufacturers of generic drugs lack the research capabilities to create new drugs and test them for safety and efficacy. And that's only one reason an economically viable research-based pharmaceutical industry is important to all of us.

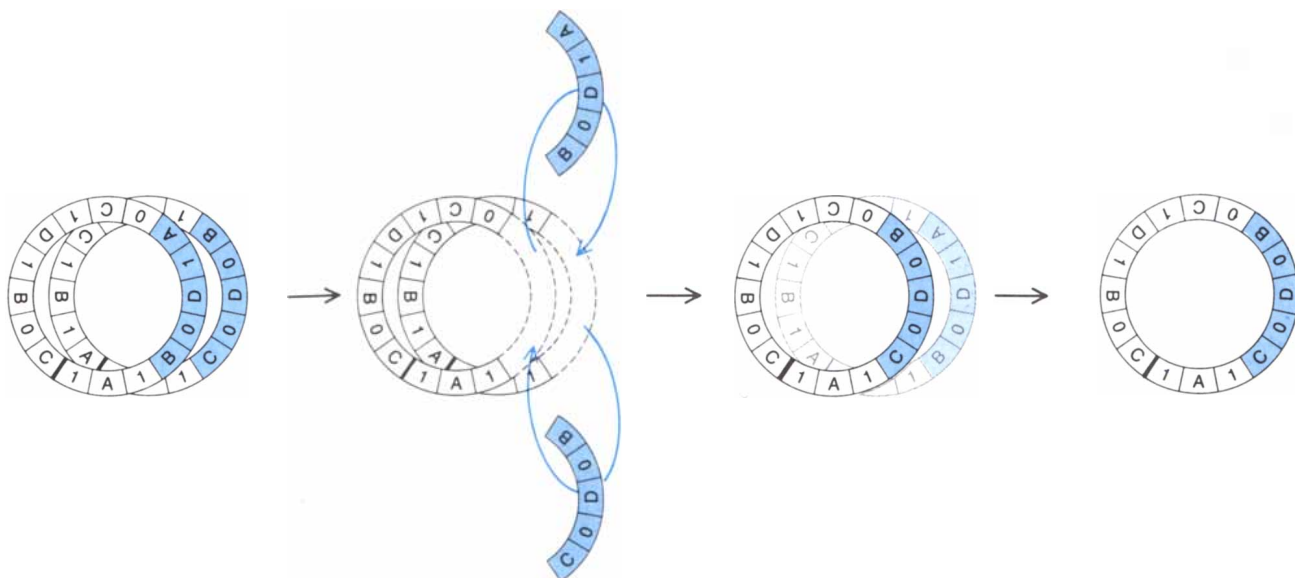
Pfizer is pleased to have been a partner in helping to reduce the hazards of one of the world's more widespread health problems. Pfizer is also pleased to be working on other solutions to similar health problems around the world.



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Crossover of two flib chromosomes, and the resulting chromosome of the offspring (right)

the winner of the mating lottery. The composition resembles a phenomenon called crossover that takes place in real chromosomes. In flib chromosomes crossover can be illustrated by combining the first (unaltered) chromosome listed above with another:

1A1B0D1A0C1D1B0C
 ↑ ↑

Arrows indicate randomly selected crossover points. The offspring's chromosome is identical with that of the second parent as far as the first crossover point. Between points it is identical with the first parent's chromosome. After the second point it is again identical with the chromosome of the second parent [see illustration above]:

1A1C0D0B0C1D1B0C

Before actually writing and testing the primordial program, I was somewhat skeptical of the value of crossover breeding. I was surprised to find, however, that if the first parent is a reasonably good predictor, the offspring tends to be as well.

Readers may judge the issue themselves by writing a program called AUTOSUP. Listed, the program does not extend much beyond a single page. It consists of four modules embedded in a loop. A limit that defines the top score should be set. As long as the top score is less than the limit the program should continue to run through the four modules.

In the first module the 10 flibs are scored on a sequence of 100 environmental symbols. The second module identifies the flibs with the highest and lowest scores that result. In the

third module the top-scoring flib is bred with a randomly selected mate. The offspring of this union replaces the bottom-scoring flib. In the fourth and last module a cosmic ray arrives, strikes a random flib and causes a mutation. Just before the program invokes the third (breeding) module a random number is selected. If the number falls below a certain threshold, the program will skip around the breeding module and execute the mutation module immediately. The threshold can be set to any level. Certain settings, however, are better than others; if the breeding module is executed too often, the small population quickly becomes dominated by the genes of the top-scoring flib. The gene pool loses diversity and evolution slows to a painful crawl if not to a downright standstill. Evolution slows down, in any event, as the scores get higher. The top-scoring flib remains on the scene for a lengthening period because it becomes increasingly unlikely that flibs superior to it will evolve.

Four arrays are useful in AUTOSUP. They are called *chrom*, *state*, *score* and *e*. *Chrom* is a two-dimensional array of 10 flibs and 16 genes. *Chrom*(*i*,*j*) refers to the *j*th gene of the *i*th flib. *State* and *score* contain the current state and score of the 10 flibs. The fourth array, *e*, contains the basic string used to generate environmental symbols. This array is received from the keyboard at the beginning of the program.

Flibs are evaluated by means of a double loop. The outer loop generates 100 environmental symbols and the inner loop increases the score of each flib if it correctly predicts the next symbol. One can test 4-state flibs adequately on environments of period 6, a chal-

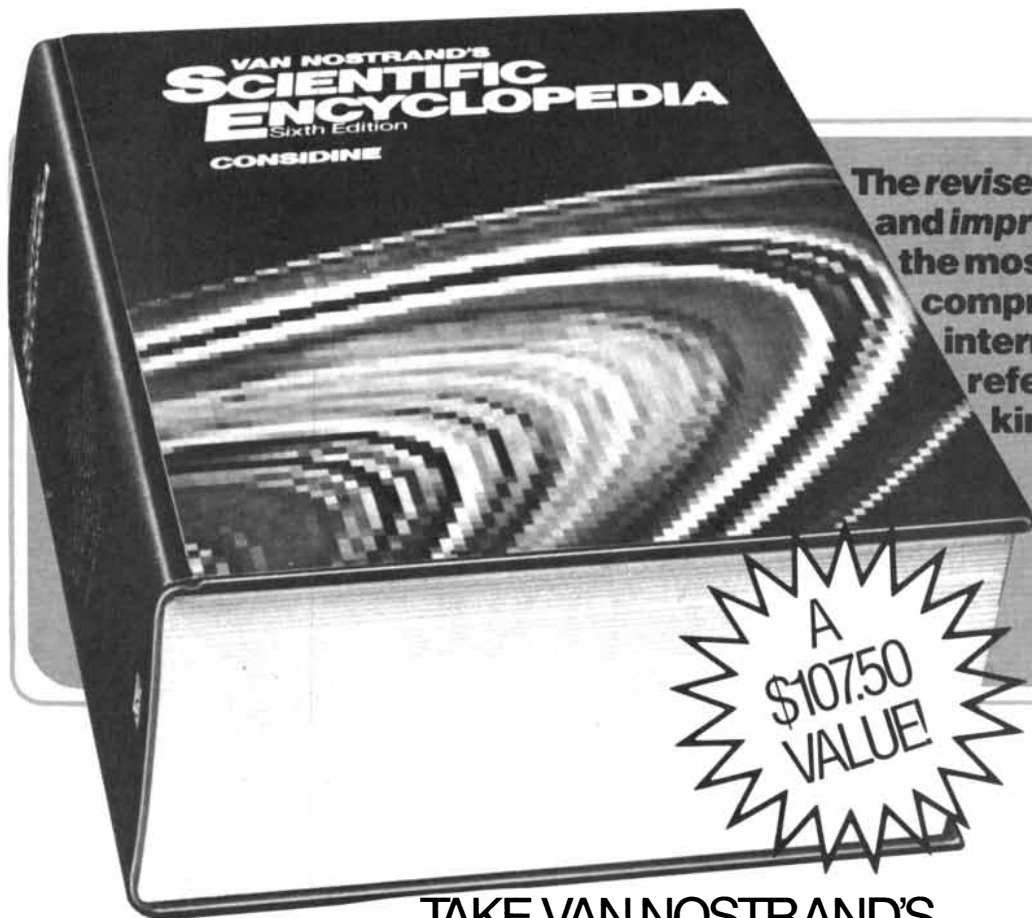
lenge of intermediate difficulty. Perfect predictors may require a day's run to evolve in a period-8 environment, whereas period-4 environments are almost no challenge at all. Two tricks are useful in this module. The first trick retrieves the next environmental symbol from the outer-loop index *i* by computing *i* modulo 6; the result is the remainder of *i* on division by 6. The number can be used to index the array *e*. As *i* runs from 1 to 100 the computed index runs through the array repeatedly, producing the proper sequence of environmental symbols. Given the index of the current symbol, the next symbol is easy to compute and look up. This symbol is compared with the prediction made by each flib in turn.

The second trick enables the program to find the flib's next state quickly and determine its output merely by scanning the chromosome. Instead of representing the four states by *A*, *B*, *C* and *D*, the numbers 0, 1, 2 and 3 are used as entries in the array *state*. If the environmental symbol is called *symp*, the output of the *i*th flib can be found by first using a simple formula:

$$l = 4 \times \text{state}(i) + 2 \times \text{symp}.$$

Then *chrom*(*i*,*l*) should be identified. The locus *l* on the *i*th flib's chromosome yields its output when the creature is in state *i* and is receiving input *symp*. The next state occupies the locus *l* + 1.

The module that determines the top and bottom flibs uses an exercise common in elementary programming courses: given an array of *n* numbers, write a program that finds the largest number. The solution involves setting a variable called *top* to 0 and then scan-



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ning the array within a simple loop. Each entry is then compared with *top*. If it is larger than *top*, *top* should be replaced by that entry. The index should be saved as well. The same procedure can be inverted and used to find the bottom score. This time a variable, *bot*, should be set at 100 and replaced by entries that are smaller.

The third module breeds the top-scoring flib with a randomly selected member of the population. The only difficulty in writing this segment arises in the selection of the two crossover points. I think it is easiest to select two random integers c_1 and c_2 from the range 1 through 16. If c_1 is greater than c_2 , their values should be exchanged. With just a little finesse readers will discover how three loops that range from 1 to c_1 , c_1 to c_2 and c_2 to 16 supply the machinery to move elements of *chrom* on the breeding rows into the destination row, which is occupied by the doomed flib with the lowest score.

In the fourth module a random flib index and a random *locus* should be selected. The parity of *locus* determines whether a state gene or an output gene is to mutate. If the value is 0, then 1 modulo 2 should be added to the number already stored there. This flips the bit, so to speak. If the *locus* value modulo 2 is 1, then 1 modulo 4 should be added to the array entry. This changes the state stored there.

Have I cheated? Surely a systematic change of state from 0 to 1 to 2 to 3 and back again is hardly a random effect. My answer is that it is random enough: the number of states is small enough so that one cannot expect the final outcome of the program to be much different from the outcome when more randomly selected states prevail. Indeed, I also cheated in a mild way by choosing c_1 and c_2 so carelessly: the method guarantees an advantage for certain substrings in relation to others. Again, I think differences between AUTOSUP and a statistically corrected crossover selection procedure would be slight. Either way there is so much juggling of genes and cracking of chromosomes that the top flib is hard put to recognize its own grandchildren.

The only parts of AUTOSUP as yet unspecified are its beginning and its end. The flibs initially occupying the soup should be selected randomly. For each gene in each flib an integer should be selected from the appropriate range and assigned to that gene. Finally, when a flib first exceeds the limit set in the outer loop, AUTOSUP should print it.

Readers embarking on this genetic adventure are warned that there is much exploring to do. Perhaps some explorers will become addicted. Ques-

tions to be answered concern the presence of evolution and its speed. When an environment period is too long for a 4-state perfect predictor to evolve, how fit do the flibs get? How do changes in the length of the period affect the amount of time it takes a perfect predictor to evolve? Nothing in the AUTOSUP description prevents extending the program to 5- and 6-state flibs. One can even alter the program to explore nonperiodic environments or ones that occasionally change their basic string.

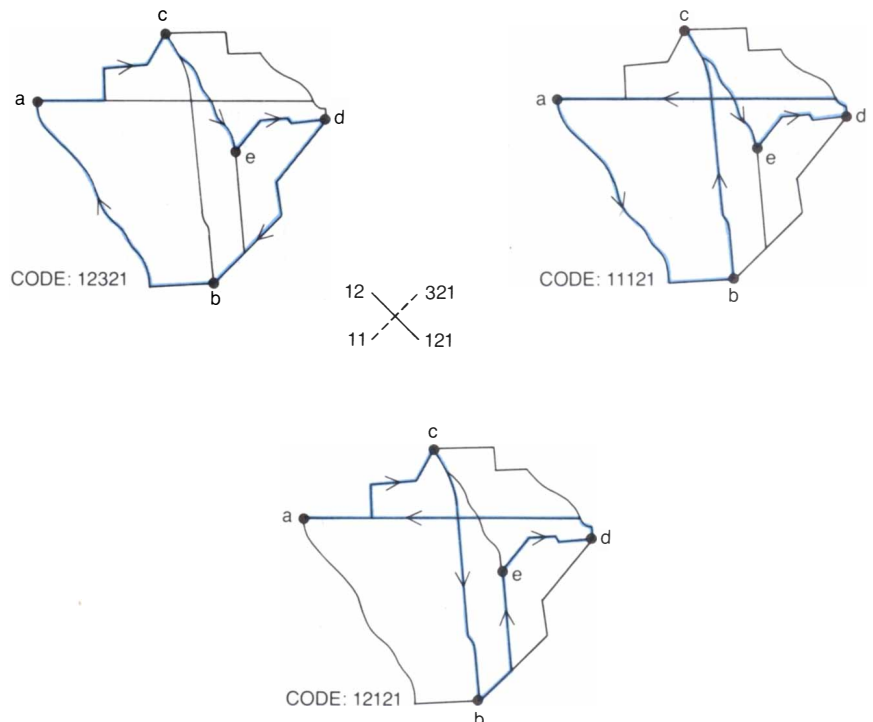
Automaton soup was inspired by a book that appeared in the early 1960's. Titled *Artificial Intelligence through Simulated Evolution*; the book describes a series of experiments in the evolution of automata by Lawrence J. Fogel, Alvin J. Owens and Michael J. Walsh. Automata were asked to predict periodic sequences and were allowed to evolve much the same way as our flibs. No breeding or crossover was allowed in this austere study, however.

It was Holland who suggested that I add the crossover facility to the automaton soup. As noted above, Holland is the acknowledged father of the genetic algorithm. Practitioners of the discipline, growing steadily in number, met at their first large-scale, funded conference, held at Carnegie-Mellon University. They discussed a wide range of theory and applications. A problem explored in several papers serves as an interesting introduction to the subject of genetic programming.

Called the traveling salesman problem, it poses the following challenge: Given a map of n cities connected by a network of roads, find the shortest tour of the n cities. Such a tour can then be used by a salesman or saleswoman to minimize travel expenses. In spite of my inclusion of salespeople of both sexes, the foregoing description has a 1940's ring to it. But more modern methods of travel and actual costs are easily incorporated without changing the mathematical skeleton implicit in the statement.

It is entirely possible that a tour of minimum length can be made to evolve just as perfect flib predictors evolved from lesser flibs. Each tour should be encoded in a chromosome. The shortest tours should be bred in the hope of obtaining yet shorter offspring. Crossover yields the chromosomes of the progeny.

It is a beguiling task to choose a good representation for a tour. For example, if one simply uses a list of the cities in the order visited, the offspring may not even be a tour. To get around this difficulty the authors of one paper, John Grefenstette, Rajeev Gopal, Brian Rosmaita and Dirk Van Gucht of Vanderbilt University, propose an ingenious chromosome. The representation for a five-city tour such as a, c, e, d, b turns out to be 12321. To obtain such a numerical string reference is made to some standard order for the cities, say a, b, c, d, e . Given a tour such as a, c, e, d, b , systematically remove cities from



Two parent salesman tours (top), and an offspring (bottom) resulting from genetic crossover

A THOROUGH EXPLANATION OF THE 16-VALVE SAAB 900 ENGINE.

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Hence the rule: Make the valves bigger and you increase what's called the *volumetric* efficiency. (In plainer English, the larger the valve, the easier it is for gas to come in and exhaust to go out.)

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they simply doubled the number of valves.

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Power output: 125 HP/92 kw @ 5500 rpm
Max. torque: . . . 123 ft. lbs./166 NM @ 3000 rpm
Compression ratio: 10.1:1
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Ignition: Bosch electronic with knock detector

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Max. torque: . . . 188 ft. lbs./255 NM @ 3000 rpm
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Ignition: Bosch electronic, Hall effect

car. If it has a four-cylinder engine, it probably has eight valves.

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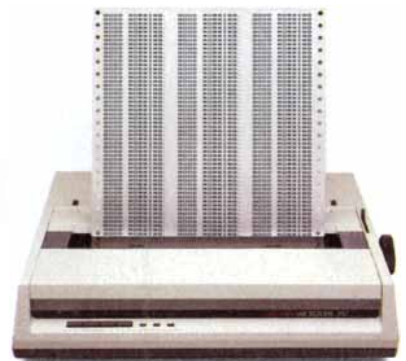
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the standard list in the order of the given tour: remove *a*, then *c*, *e* and so on. As each city is removed from the special list, note its position just before removal: *a* is first, *c* is second, *e* is third, *d* is second and, finally, *b* is first. Hence the chromosome 12321 emerges. Interestingly, when two such chromosomes are crossed over, the result is always a tour [see illustration on page 27]. With such a representation tours can now be bred, so to speak, for fitness.

Most practitioners of the art of genetic algorithms readily admit that the traveling salesman problem is one of their greatest challenges. Although experiments with the representation just described were not very encouraging, there are other genetic algorithms that perform better on the problem.

Still, no genetic algorithm has ever been able to conquer the traveling salesman problem in any well-accepted sense. This is undoubtedly owing to the difficulty of the problem itself. Because it is what theorists call NP-complete, it may be doomed to eternal insolubility in the practical sense.

Since its appearance in this column last August the MANDELZOOM program has been incarnated in hundreds of homes, schools and workplaces. Although the program has apparently awed adults, intrigued teenagers and frightened a few small children, to my surprise the mail on iteration diagrams, the secondary topic, nearly equals that on the Mandelbrot set.

Many readers have lost themselves in the colored intricacies of the Mandelbrot set by zooming ever deeper. Some readers who are determined to have their own color images but who lack color-display equipment have been ordering pictures from John H. Hubbard of Cornell University. I have been told by Heinz-Otto Peitgen, the Mandelbrot explorer whose images graced the August pages, that those images too are for sale; they are included with dozens of others in a color catalogue from Design-Büro-Weisser, Lothringer Strasse 23, D-2800 Bremen 1, West Germany. Hubbard and Peitgen together learned the art of image generation from Mandelbrot.

For readers whose equipment is limited to black and white, I should have thought of shades of gray. Such pictures can be nearly as inspiring as their colored counterparts. The best gray images were produced by David W. Brooks, who works with equipment at Prime Computer, Inc., in Framingham, Mass., to compute and plot his pictures. In his fabulous and delicate riots of halftones each shade of gray is rendered by tiny black squares of a

certain size; the squares are made by a laser printer. Brooks has been searching for the tiny filaments that are thought to connect miniature Mandelbrot to the main set. So far they have not appeared at any magnification used by Brooks. Mandelbrot has advised him that they are probably infinitesimal.

Those with less sophisticated equipment can still work with shades of gray on a black-and-white monitor. John B. Halleck of Salt Lake City varies the density of points per pixel to indicate different shades.

Another approach depends on black and white contours. Yekta Gursel of Cambridge, Mass., has generated views of the Mandelbrot set that rival the ones Brooks generates. Gursel replaces a discrete spectrum of colors with alternating bands of black and white. Gary J. Shannon of Grants Pass, Ore., suggested the same technique and Victor Andersen of Santa Clara, Calif., took it to an extreme. He suggested changing from black to white (or the converse) whenever the *count* variable changes from one pixel to its neighbor.

Two other explorations are worth mentioning. James A. Thigpen IV of Pearland, Tex., uses height instead of color. The Mandelbrot set becomes an immense plateau seen from an angle, with a complicated arrangement of spiky hills approaching the plateau in various places. Richard J. Palmaccio of Fort Lauderdale dispenses with the set altogether. His interest is in tracking individual complex numbers in the course of iteration. Their choreography near the boundary can result in spiral ballets or circular jigs.

The function $z^2 + c$ gives rise to the Mandelbrot set. Naturally other functions are possible, but they produce other sets. For example, Bruce Ikenaga of Case Western Reserve University has been exploring what appears to be a cubic cactus. The function $z^3 + (c-1)z - c$ produces a prickly and uncomfortable-looking set (at least in stark black and white) surrounded by mysterious miniature spiral galaxies.

There are mysteries in iteration diagrams as well: when the integers modulo n are squared, each number migrates to another, in effect. The iteration diagram appears when each number is replaced by a point and each migration is replaced by an arrow. I raised several questions about such diagrams. How many components do they have? Readers sent diagrams documenting their explorations for various values of n .

The largest diagrams were completed by Rosalind B. Marimont of Silver Spring, Md. She examined the integers

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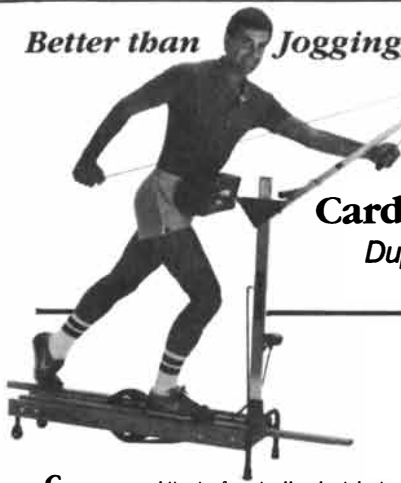
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modulo 1000 and reported four pairs of components in the resulting iteration diagram. Each component sported a single attractor, as usual, and the largest attractors had 20 numbers. As a mathematician Marimont is allowed to conjecture that the integers modulo 10^k will produce $k + 1$ pairs of components and that the largest attractors will have $4 \times 5^{k-2}$ numbers.

Stephen Eberhart of Reseda, Calif., investigated the case where n is a Fermat prime (a prime number of the form $2^{2^k} + 1$). Here the number 0 forms an attractor by itself and the remaining numbers all lie in one single, grand tree. A number-theorist friend affirms that this will always be the case for Fermat primes and that the tree is binary: each internal point has two incoming arrows.

Iteration diagrams, like numbers, can be multiplied. If n is the product of two relatively prime numbers, say p and q , the iteration diagram for the integers modulo n is the product respectively of the diagrams for p and q . This interesting observation was made by Stephen C. Locke of Florida Atlantic University. Locke has also described a fascinating relation between the n th iteration diagram and a diagram of a seemingly different kind, one in which the numbers, instead of being squared, are merely doubled. When n is a prime, the latter diagram for the integers modulo $n - 1$ is the same as our n th iteration diagram, except for a single isolated number forming an attractor by itself. Much the same observation was made in number-theory terms by Noam Elkies of New York City.

A powerful tool for analyzing the (squared) iteration diagrams was developed by Frank Palmer of Chicago. Apparently all the trees attached to a given attractor are isomorphic. This means essentially that they have precisely the same form.

Finally, Bruce R. Gilson of Silver Spring, Md., and Molly W. Williams of Kalamazoo, Mich., examined a quite different generalization of the numbers from 0 through 99. These may be regarded as numbers to different bases. As numbers to the base 3, for example, one would count 00, 01, 02, 10, 11, 12, 20, 21, 22 before arriving once more at 00. Such numbers also produce iteration diagrams that look like those arising for integers modulo n . Gilson proved the diagrams always have paired components when n is even but not a multiple of 4.

There was an error in the iteration diagram presented in the August column for the integers from 0 through 99. Two arrows were missing from two components and the direction of one attractor was reversed.

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