SCIENCE'S COMPASS

NET TIP

Mailbox: www.sciencemag.org/cgi/dmail?53884

Turning the Big Five-O

Spend a few minutes looking at World Wide Web pages and commonly used patterns emerge. You see similar colors used in Web pages, similar fonts, and similar layouts. Ten people viewing the same site will see identical images, because sites do not adapt to the user or retain preferences set during a previous visit. The current Web style seems to have reached a steady state over the past year. However, this is about to change.

Why? Because the core technology behind the building of Web pages will change dramatically with the introduction later this year of the next generation of browser software. In December 1994, Netscape's first browser was offered to the public. As of September 1998, the two dominant companies in the browser field, Microsoft and Netscape, had released fourth generation products. Now, by significantly enhancing the languages and the browser itself, the 5.0 versions of browsers will transform the Web into a sophisticated development platform. The sky is truly the limit with these new tools.

What features can we expect to see in the 5.0 browsers? For Netscape Communicator 5.0, there is little information to report. The company has decided to build this next version in an open way, by assembling contributions from individual developers around the world who will design components of the new browser. Netscape hints that the product will be available by late 1998.

Microsoft, however, has just released a working preview version of Internet Explorer 5.0 (IE5) to developers, and a final version is slated for late 1998. IE5 was constructed at Microsoft and has not been a public effort. IE5 is a powerful piece of software that will radically alter a Web page in three general areas: document structure, user preferences, and speed.

The document structure of today's Web pages is derived from Hypertext Markup Language (HTML has evolved to version 4.0). HTML will not be replaced, instead it will be augmented. IE5 will allow developers to position all items and figures on a page precisely, to construct one page from the content of multiple URLs, and to build a page quickly from data packets coded in Extensible Markup Language (XML). These changes describe a fraction of the possibilities that will become available.

The user experience will change with the increased use of Dynamic HTML (DHTML), first released in Internet Explorer 4.0. DHTML in IE5 will make all elements on a page (the graphics, text, buttons, forms, and so forth) dynamic. With this ability, Web developers can make pages that change to fit the preferences of the user. For example, an electronic journal could be designed in which the sections of articles could be rearranged based on viewer preferences. Would you want to see only abstracts? Click on a button to show only the abstract component of the articles. Would you want to see the results sections first? Click on a button to switch the order of the pieces of the article. Also, with the new technology of IE5, the browser will retain your layout preference for future visits.

The IE5 browser will also be faster. Each Web page must be processed for viewing, or rendered, by code within the heart of the browser. This code has been overhauled in IE5 and optimized for speed. For instance, for table structures used to position text and graphics, the time it takes to visualize a page after transfer to your computer could become 1/100 of what it is now.

Scientists will soon have desktop tools to distribute and format the most complex data and information. For more information on the next generation of browsers, see the links collected at www.mednav.com/zone/science.

-ROBERT SIKORSKI AND RICHARD PETERS

TECHSIGHTING

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The Art of Bubble Making

agnetic resonance imaging (MRI) reflects detection of the relaxation of ¹H nuclei after a biological specimen has been exposed to a magnetic field. Because such specimens are mainly water, MRI has had enormous utility. When the proton signal is weak, as it is in blood, contrast agents that magnify the relaxation of ¹H (such as gadolinium–diethylenetriamine pentaacetic acid) are usually added. For visualizing the lungs, which lack a significant proton signal source altogether, laserpolarized gases (³He and ¹²⁹Xe) are used. Then the gas itself becomes the source of the signal. These particular gases offer at least a 20-fold increase in signal-to-noise ratio compared with ¹H MRI images. In addition, these gases do not occur naturally in the body, and thus there is no background signal. Such contrast agents have been used with great success for visualizing anatomical air space. Furthermore, ¹²⁹Xe dissolves reasonably easily, so researchers have been able to view tissues such as blood, muscle, and brain, either by letting the gas dissolve in the pulmonary blood flow or by dissolving it in lipid vesicles. Helium-3 has a larger magnetic moment and currently a higher degree of polarization than ¹²⁹Xe and thus would offer a 10-fold improvement in signal over ¹²⁹Xe. Unfortunately, ³He has very poor solubility (one-tenth to one-hundredth that of ¹²⁹Xe), so it has been exclusively used for air space imaging of biological samples.

A report published last month in the Proceedings of the National Academy of Sciences U.S.A. may promote the use of ³He for visualizing fluid and solid tissues (1). In a preliminary feasibility study, researchers at the Duke University Medical Center were able to generate microbubbles of ³He and to view the vasculature of live rats. To generate these microbubbles, they used two 10-cm³ syringes connected via a three-way stopcock. The gas generated by the laser polarizer was collected in a holding cell. Then, 2 cm³ of it was quickly withdrawn from the cell into the first syringe, which had been evacuated to prevent paramagnetic oxygen in the syringe from depolarizing ³He. Next, 8 cm³ of fluid in the second syringe was manually flushed back and forth between the syringes, producing a suspension of ³He microbubbles. This was done rapidly because the microbubbles quickly rise out of the suspension. For this study, the time between agitation and image acquisition was reduced to about 10 s. The authors measured the suspended microbubbles with a Coulter counter. Mean bubble diameter was 31.8 um with a mean standard deviation of 10.4 µm. The large distribution of bubble size was due to the hand agitation technique, and more homogeneous distribution can be expected with other mechanical methods (such as sonication). The size of the bubbles was also significantly larger than the ideal size of 8 µm (the size of a red blood cell), but the authors decided this size was acceptable for this in vivo feasibility study. They also noted that the ³He suspension differed significantly in different fluids: in all, they tried six fluids and found that Hexabrix (Mallinckrodt) offered by far the most homogeneous distribution of bubbles and the best signal-to-noise ratio.

They injected this suspension of ³He into both the arterial and venous systems

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