

Research Article

A LabVIEW-Based Virtual Instrument System for Laser-Induced Fluorescence Spectroscopy

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We report the design and operation of a Virtual Instrument (VI) system based on LabVIEW 2009 for laser-induced fluorescence experiments. This system achieves synchronous control of equipment and acquisition of real-time fluorescence data communicating with a single computer via GPIB, USB, RS232, and parallel ports. The reported VI system can also accomplish data display, saving, and analysis, and printing the results. The VI system performs sequences of operations automatically, and this system has been successfully applied to obtain the excitation and dispersion spectra of α -methyl-naphthalene. The reported VI system opens up new possibilities for researchers and increases the efficiency and precision of experiments. The design and operation of the VI system are described in detail in this paper, and the advantages that this system can provide are highlighted.

1. Introduction

Laser-induced fluorescence (LIF) is a technique that relies on the measurement of the excited state fluorescence that results from photoexcitation. The excitation source for molecular LIF is typically a tunable dye laser so that selective excitation and high resolution can be achieved. Combined with the supersonic-jet expansion technique, LIF has been used to study the excited state structures of molecules and the ultra-fast dynamics of molecular systems [1]. LIF has also been successfully applied for quantitative measurement of concentrations in combustion, plasma, spray, and flow phenomena, in some cases visualizing concentrations down to nanomolar levels [2].

Unlike other analytical spectroscopy techniques, where ready-to-use commercial spectrometers have everything for the sample detection in a package, laboratories usually design and set up the LIF experimental instruments by themselves in order to satisfy individual research needs. In addition to hardware installation, LIF also requires synchronized and automated control software for laser function, data acquisition, and communication between instruments [3, 4]. Most laboratories develop homemade programs for their experiments using different programming languages, such as Visual C++, LabVIEW, Visual Basic, or Turbo Pascal, but

text-based languages have the shortcomings of long developing cycles and difficulties in maintenance and expansion [5]. NI LabVIEW 2009 (National Instrument Inc., USA) is a graphical programming tool which can be employed to develop sophisticated measurements [6–8] and control [9, 10] systems using intuitive graphical icons and wires that resemble a flowchart instead of written lines of text [11–16].

In this paper, a virtual instrument system based on LabVIEW 2009 was developed for LIF experiments. This system can simultaneously control the scanning of a dye laser and a monochromator, the operation of a molecular pump, and data acquisition. Furthermore, it achieves real-time data processing and monitoring of individual instruments. An application of this program in the LIF spectroscopy examination of α -methyl-naphthalene is shown in this paper. Simple modifications for adapting this program to different LIF setups are also illustrated.

2. Configuration of System Hardware

There are two different kinds of spectra in LIF spectroscopy, dispersed spectra, and excitation spectra. The dispersed spectra are obtained with fixed excitation laser wavelength and the fluorescence spectrum is analyzed. In excitation spectra, the laser wavelength exciting the molecules to a

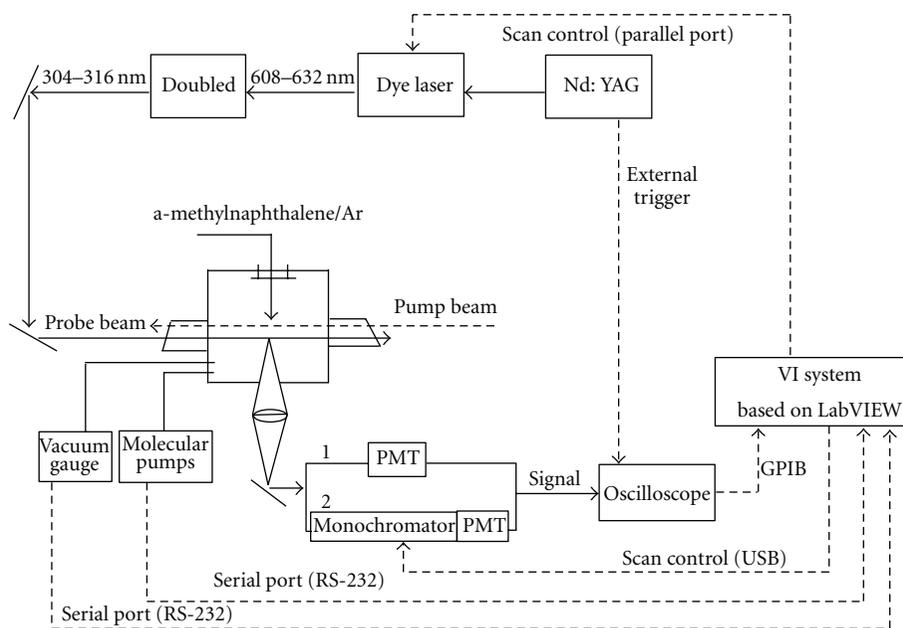


FIGURE 1: Hardware structure of VI System.

higher energy state is scanned and the total fluorescence is measured. Information on lower electronic states can be obtained by dispersed spectra, while the excitation spectra reveal parameters (energy levels, structure information, etc.) of upper level electronic states.

A conventional experimental setup for obtaining laser-induced fluorescence excitation spectra and dispersed spectra are depicted in Figure 1. The sample vapor is injected into a vacuum chamber, whose vacuum is maintained by a molecular pump and a mechanical pump, via a supersonic jet. The excitation laser beam (probe beam) is provided by a frequency doubled dye laser pumped by a Nd:YAG laser. The probe beam intercrosse the sample 10 to 15 mm downstream of the supersonic jet nozzle; when free radicals are studied, a pump laser beam propagates across the sample close to the nozzle and generates the free radicals. Fluorescence of the excited molecules is collected perpendicular to the probe beam by the optics and imaged onto the detection devices. For excitation spectra, total fluorescence is detected by PhotoMultiplier Tube (PMT) and converted into electronic signals, which are received and displayed in real time on an oscilloscope (synchronized by the triggers from the Nd:YAG laser). A General-Purpose Interface Bus (GPIB) card installed in the computer transfers data from the oscilloscope to the computer via the GPIB port, and data acquisition and processing are executed by the VI system. At the same time, the VI controls the dye laser scans by sending trigger pulses to the laser controller via the parallel port of the computer. The acquisition and control process for dispersed spectra are similar to those for excitation spectra. For dispersed spectra, the fluorescence signals are transmitted through a monochromator before being collected by the PMT. The VI system communicates with and controls the scan of the monochromator via the USB port in the monochromator. In

addition, the VI system also communicates with the vacuum gauge via an RS232 port, and the system displays the in situ values of the vacuum. The VI system also builds up an interface to communicate with the operating program of the individual instruments (e.g., the dye laser and the molecular pump); specific operating programs are usually included with the instruments at the time of purchase.

3. Programming

Figure 2 shows the flow chart of the VI system that performs the LIF experiments and data processing tasks. It incorporates several subVIs, each one of which carries out a specific function, including data acquisition and display, data processing, as well as generating and saving the results.

There are three icons, Excitation Spectrum, Dispersed Spectrum, and Data Processing, on the front page of the virtual instrument system, each is dedicated to a specific action or experimental technique. When an icon is selected, the corresponding subVI program will be activated, and it will open the panel where specific tasks can be executed. For programming, an *Event Structure* for incident response is used to call the corresponding subVI [17]. When the subVI starts, the *Event Structure* waits until the precompiled program code is executed and the task is finished, then the *Event Structure* returns to wait for another event.

3.1. Program for Excitation Spectrum. To fulfill the requirements of LIF excitation spectra, the Excitation Spectrum subVI consists of fluorescence data acquisition, data display and saving, vacuum monitoring, communication with other application modules, and synchronization of laser-wavelength-scan subVIs.

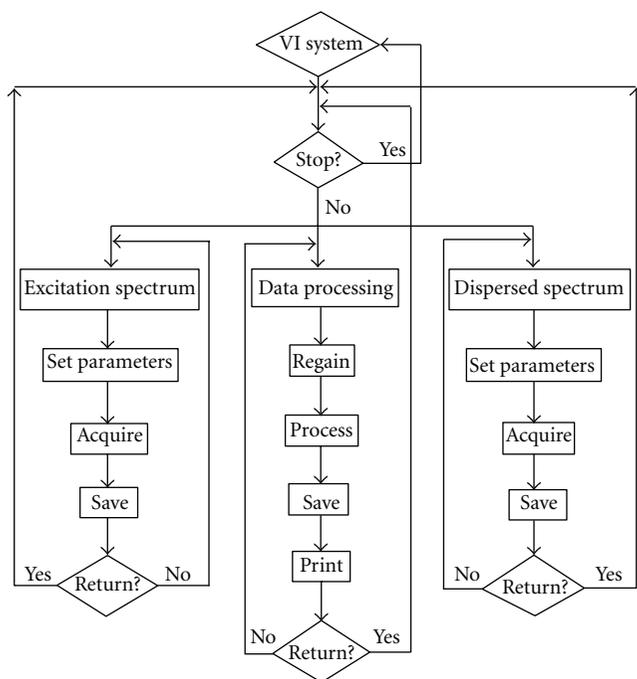


FIGURE 2: Flow chart of VI system.

3.1.1. Data Acquisition and Processing. In the experiment, the fluorescence signal is collected by the PMT and transferred to a TDS3032B oscilloscope. The TDS3032B oscilloscope supports the GPIB interface communication with a computer where the interface is an eight-bit parallel bus that provides high-speed data transfer and multiple-instrument control [18]. Detailed programming is shown in the upper left *Flat Sequence Structure* of Figure 3. We invoked the *GPIB Write* function from the LabVIEW *Instrument I/O* function to set the GPIB address of the oscilloscope and specify the data parameters to be transferred from the oscilloscope, including the portion of the waveform that users want to transfer, the waveform source, data format, and the number of bytes per data point. We then invoked the *GPIB Read* function and wired the corresponding GPIB address of the oscilloscope to the *GPIB Read*, and also specified the number of bytes the function read from the oscilloscope. Finally, we called the function nodes *Math Pattern*, *String Subset*, *Decimate 1D Array*, and *Join Numbers* to receive waveform data from the oscilloscope and display it in situ in the front panel waveform graph. Data from different oscilloscope channels (CH1, CH2, or Dual CHs) can be acquired by wiring selected terminals (in a combo box) to the case structure prior to data acquisition. For the end-user, the above process is easily achieved by choosing the channel from the pull-down list on the front panel (Figure 4) and clicking on the Acquire button.

There are two sequential subdiagrams in the upper right case structure of Figure 3. The left one performs the analysis and processing of the fluorescence data obtained from the TDS3032B oscilloscope at a particular laser wavelength. Four cursors were created, and property nodes were set to acquire the cursor's horizontal axis positions on a waveform graph and to define the gate width. We then used the *Array Subset*

to return two portions of fluorescence data between the two sets of cursors. On the front panel, users can directly drag the cursors to where they want, or they can simply set positions in the cursor legend to choose the gate on the graph. Integrals of data in both gates are computed separately using the *1D Numeric Integration.vi*, in which the integration method is the Trapezoidal Rule. The difference between the two gate integrals is calculated for the purpose of background subtraction. Finally, the fluorescence intensity and the current wavelength are sent to the XY Graph for real-time spectroscopy display. The front panel of the fluorescence Excitation Spectrum is shown in Figure 4, in which Choose Gates is shown in the upper portion and Spectroscopy Display in the lower part.

As soon as the data processing is complete, a TTL trigger is sent to step the dye laser to the next wavelength. TTL subVI, shown in the right sequential diagram in the case structure of Figure 3, calls the *Outport.vi* in *Port I/O*, which specifies a parallel port address for sending the trigger pulse and also defines the pulse width. A BNC cable is made with the inner core wired to the specified TTL pulse pin of the computer parallel port, and the outside shield wired to the ground pin. Thus, a TTL signal can be sent to the dye laser control unit from the computer parallel port when a data collection cycle is completed and the laser is stepped to the next scan wavelength. The settings for start position, end position, and step size of the laser wavelength scan must be identical in our VI program and in the laser control software. The TTL subVI is placed in a comparison case structure. If the current position is smaller than the end position, the comparison function will return a *False* value to initiate the program *TTL.vi* which will move the laser to the next wavelength to start a new data acquisition and processing cycle. If the current position is greater than or equal to the end position, the *True* case gives a message dialog box indicating that data acquisition (DAQ) is finished. Subsequently, data can be saved by employing the *Write to Spreadsheet* file and the *Build Array* and *Transpose 2D Array* nodes. This task is completed by clicking on the save button on the front panel and the fluorescence data will be saved according to the file name entered. The data acquisition process can be stopped at anytime by clicking the Acquire button again (in case the user wants to interrupt the experiment). An abort function is also programmed using the *Stop* node in the *Application Control* function, so users can terminate the whole VI system by clicking on the Abort-Execution button in the upper-right corner (shown in Figure 4). For convenience, the help function of the program can be shown by clicking on the Help button on the front panel.

3.1.2. Vacuum Monitoring and Accessing Other Application Software. The program for the real-time display of the vacuum gauge is shown in the lower-left parallel loop of Figure 3. To communicate with the vacuum gauge, the *Serial* function templates in the LabVIEW *Instrument I/O* can be selected. Parameters such as VISA resource name, baud rate, data bits, and parity in the *VISA Configure Serial Port* are set according to the vacuum gauge communication protocol [19, 20]. The *VISA write* function sends command characters

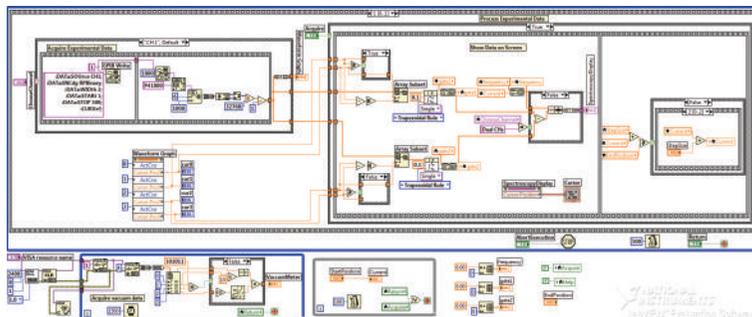


FIGURE 3: Program for excitation spectrum.

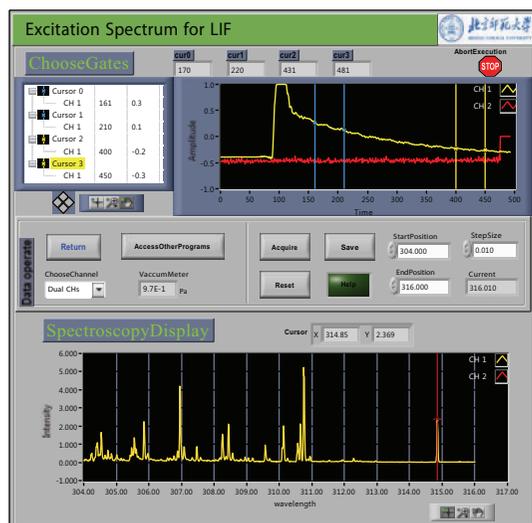


FIGURE 4: Front panel for excitation spectrum.

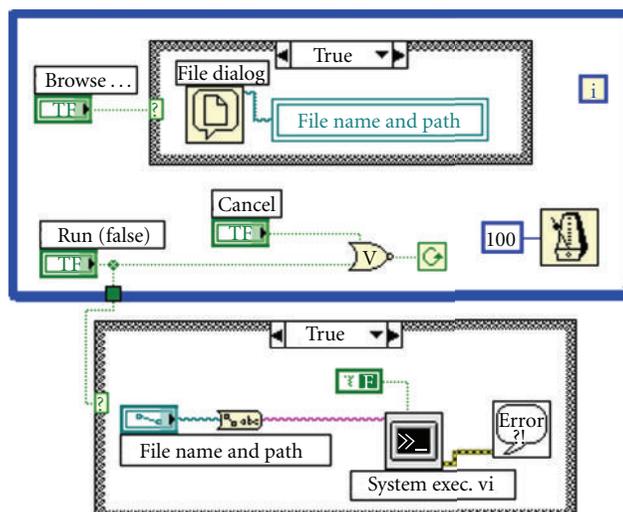


FIGURE 5: SubVI for Access Other Programs.

to the vacuum gauge and receives vacuum values in return. The nodes *String to Byte Array*, *String to Single Precision Float*, and *Index Array* transform the vacuum data into the required format and display the values on the front panel indicator.

For convenience, the VI system was designed to interface with the other applications that already exist for the individual instruments. By clicking on the *Access-Other-Programs* button on the excitation spectrum panel in Figure 4, users can select the application program by browsing the computer disks. Figure 5 shows the block diagram of the subVI for accessing other programs. The *Browse Boolean* command is wired to the upper case structure, which includes a *File Dialog Express VI*. The application can be selected when the file path is displayed on the front panel. When the user confirms the file selection, the application program is activated via the *System Exec.vi* shown in the lower case structure in Figure 5.

3.2. Program for Dispersed Spectra. A change in the dispersed spectrum program is the addition of synchronizing fluorescence DAQ with a monochromator wavelength scan instead of a dye laser scan. The 300i monochromator used in our dispersed LIF experiment was an Omni- λ 3008 spec-

trometer (Beijing Zolix Instrument Company). Communication between the Omni- λ 3008 spectrometer and the computer is achieved via a USB port on the Omni- λ 3008 spectrometer. The Omni- λ 3008 spectrometer is operated by a *ZolixOmniSpec ActiveX* (OmniSpec; provided by the manufacturer). In the LabVIEW VI system, we invoke an *ActiveX Container* and created a *ZolixOmniSpec ActiveX* object; an *Automation Open* in the *ActiveX* functions palette is called, which returns an automation reference number and allows the specific functions programmed in the *ZolixOmniSpec* object to be accessed. Property nodes in the *ActiveX* palette are used for reading the USB port information. The Omni- λ 3008 spectrometer USB serial string is found by calling the *SearchZolixUSBDevice* and *GetZolixUSBSerial* invoke methods, and then the string is wired to the *USBSerials* property node. The invoke commands (*Open* and *Connect*) are used for setting the connection status of the monochromator, and the status is displayed by the *Boolean* indicator on the front panel. The initial states of the Omni- λ 3008 monochromator components, such as the exit port, grating and current wavelength, are obtained by employing the *GetExitPort*, *GetCurrentGrating*, and *CurrentWave* methods, respectively. Next, different parameters are set according

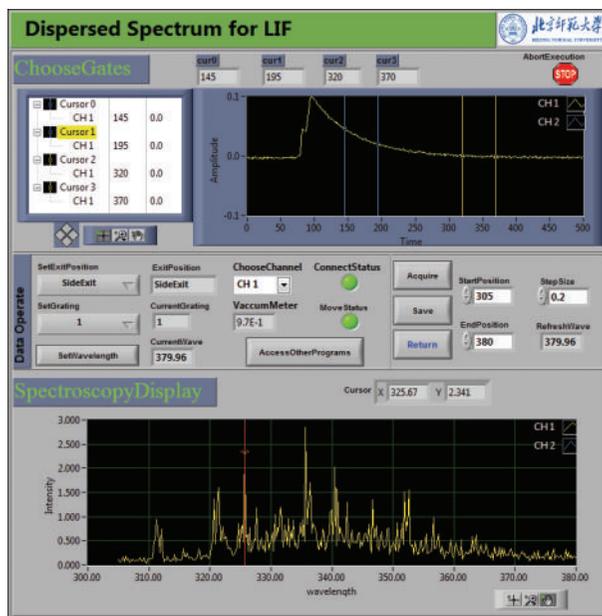


FIGURE 6: Front panel for dispersed spectrum.

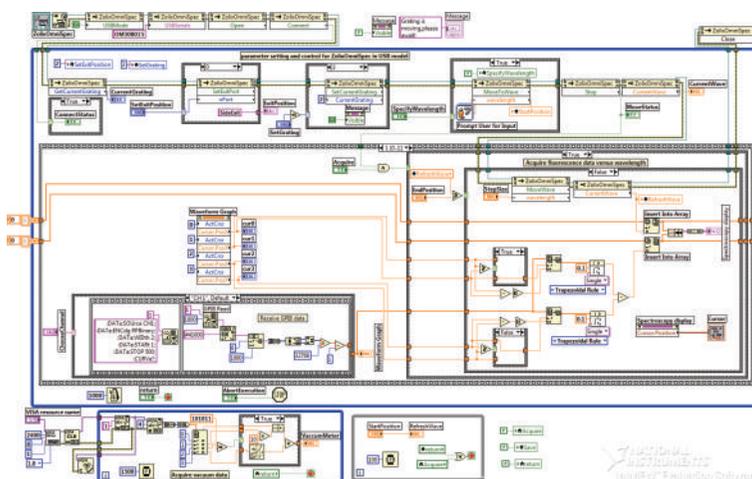


FIGURE 7: Program for dispersed spectrum.

to experimental requirements. This step involves the user's input of the experimental parameters by selecting from the pull-down menu on the front panel. The *SetExitPort* and *SetCurrentGrating* in the case structure return to the default state (which is set as *Do Nothing*) as soon as the commands are given to avoid sending looped control commands to the OmniSpec. The *Prompt User for Input* function in the *Dialog and User Interface* is called, and it prompts the user to input a specific wavelength, which is subsequently sent to the wavelength terminal in the *MoveToWave* method. After using the *MoveToWave* method, a *Stop* function is required to stop the grating from moving. The operation panel and program are shown in Figures 6 and 7.

A similar comparison case structure is applied here to step the monochromator scan wavelength as the one used in the Excitation LIF spectrum program. Identically to

the excitation experiment, the start position, end position, step size of the monochromator scan, and the gate for fluorescence signal integration will await user input on front panel. Executed from the front panel, data acquisition will begin, and the real-time spectrum will be displayed in the XY Graph. Using the *Save* icon on the front panel, the experimental data can be stored in the computer via *Write to Spreadsheet File.vi*. An *Automation Close* is programmed in the VI system to disconnect the USB interface to the Omni- λ 3008 and release the system resources when the Dispersed Spectrum subVI is closed.

3.3. Program for Data Processing. The data processing subVI (Figure 8) can conveniently redisplay and analyze previously saved data [21, 22]. This subVI can automatically accomplish wavelength calibration, unit conversion, and peak and

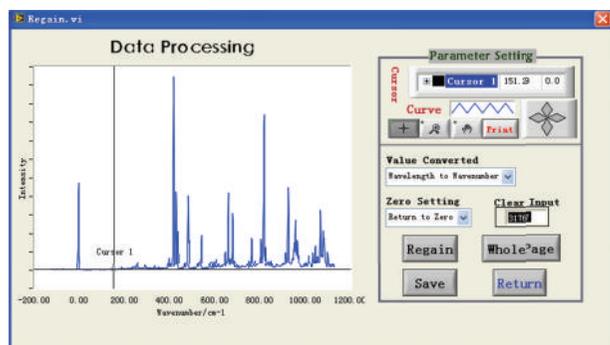


FIGURE 8: Front panel for data processing.

band-width reading of the spectrum curve. Users can also inspect each dataset using cursor move, data refresh, figure enlarge and diminish, and graph recover tools. In the VI programming, a *File Dialog* function will prompt the user to upload a data file. The *Read From Spreadsheet File.vi* and *Index Array* function will convert the wavelength and intensity data to x and y arrays, respectively. Wavelength unit conversion is realized by the case structures, and then the processed data will be sent to the XY Graph for display and saving by *Write To Spreadsheet File.vi*. Spectrum graph printing is achieved by using the relevant *Report Generation* functions.

4. Application

The VI system reported here has been used to take fluorescence spectra of molecules and free radicals in our laboratory. The excitation spectrum of *a*-methylanthalene is shown in Figure 4 (in wavelength, nm) and Figure 8 (in wavenumber, cm^{-1}). Dual CHs were used for data acquisition, one for studying the fluorescence signal and another for monitoring the laser power fluctuation using a photodiode. High resolution and signal-to-noise ratios were achieved in the spectra. The obtained excitation spectrum was consistent with previously reported results [23, 24] with spectral origin at 31767 cm^{-1} (314.80 nm). The dispersed fluorescence spectrum of *a*-methylanthalene, shown in Figure 6, was obtained by exciting the most prominent band at the wavelength of 310.70 nm. Dispersed spectra utilizing excitation of other bands (306.50 nm and 314.80 nm) were also obtained. Spectral analysis was conducted assisted by theoretical calculation and would be reported elsewhere [25].

5. Conclusions

A VI system for LIF spectroscopy based on LabVIEW 2009 has been developed. The VI system integrates a variety of laboratory equipment and controls the experiment data acquisition process using a single computer via GPIB, USB, RS232, and parallel ports. It achieves synchronous control of equipment and real-time fluorescence data acquisition. Data display, saving, and analysis can also be accomplished using the reported VI system. Furthermore, the VI system

can monitor running of individual instruments and changing of vacuum and laser power in real time. Therefore, the users can timely judge, analyze, and process on-site information, and high resolution and signal-to-noise ratios of molecular or free radicals can be easily achieved in the spectra. Our approach has been to develop the VI system for general use in LIF experiments. The instruments used in this paper, such as the Nd:YAG laser (SureliteIII, Continuum), dye laser (NarrowScan, Radiant Dye), digital delay generator (DG535, Stanford), oscilloscope (TDS3032B, Tektronix), and PMT (CR110, Hamamatsu) are commonly used in many laboratories. When different instruments are to be used in an experiment, the users simply replace the communication method according to the communication protocol for the corresponding replacement instrument or change the file path in the Access-Other-Program subVI by following the instructions provided in the VI. The VI system has user-friendly front panel windows and is easy to operate. Help and Instructions implanted in the program will assist the user in becoming familiar with each application step.

The VI system was tested extensively and applied in the determination of the excitation spectrum and the dispersed spectrum of *a*-methylanthalene with satisfactory experimental results. The VI system promotes and expands the capabilities of individual instruments, and it is an intelligent improving for existing hardware sources of laboratory. Lots of complicated functional operations are merely finished at the click of button on the VI system. The VI system not only avoids the manual error during the separate instrument control, data acquisition, and data processing, but ensures the accuracy and repeatability of experimental data and improves the efficiency of experiment. Furthermore, the VI system has a wide expandability of functions, minor modifications can be applied to a different LIF setups based on automatic measurement and control. The VI system will be provided upon request.

Acknowledgments

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