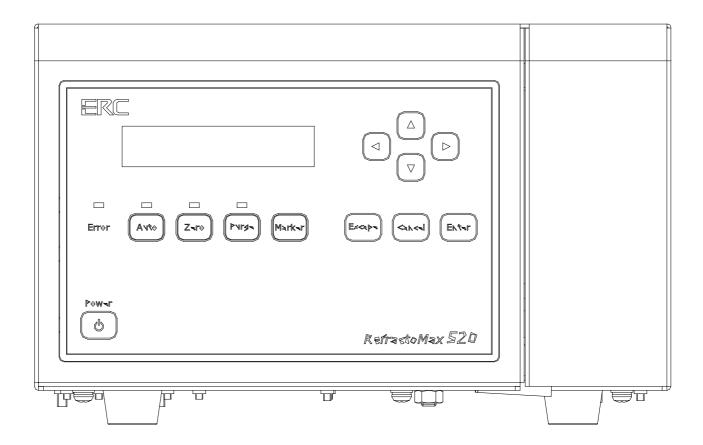
RefractoMax524 RI Detector Operator's Manual

High Sensitive Refractive Index Detector for High Performance Liquid Chromatography



Publication No. 77-5014-10

Important Notice

The information contained in this manual is subject to change without notice.

ERC Inc. (ERC) assumes no responsibility for any errors that may appear in this manual. This manual is believed to be complete and accurate at the time of publication.

In no event shall ERC be liable for incidental or consequential damages in connection with or arising from the use of this manual.

If the detector is used in a manner not specified by this Operator's Manual, the protection provided by the detector may be impaired.

Spare Parts Availability

It is the policy of ERC to provide maintenance spare parts for a period of seven (7) years after the final production of the instrument. Spare parts may be available after the seven (7) year period but only on an "as available" basis.

The following is a Federal Communication Commission advisory:

WARNING:

This equipment generates, uses, and can radiate radio frequency energy and if not installed and used in accordance with the instruction manual may cause interference to radio communications. It has been tested and found to comply with the limits for Class A computing device pursuant to Subpart J of Part 15 of FCC rules, which are designed to provide reasonable protection against such interference when operated in a commercial environment. Operation of this equipment in a residential area is likely to cause interference in which case the user at his own expense will be required to take whatever measures may be required to correct the interference.

Operating Instructions

This manual is provided to help you establish operating conditions that will make safe and efficient use of your equipment available.

Special considerations and precautions are also described in this manual that shall appear in the form of WARNING, CAUTION and NOTE as described below.

It is important that you operate your equipment in accordance with this manual and any additional information that may be provided by ERC from time to time.



Alert you to potentially hazardous situations that could result in serious injury, and how to avoid these situations.

Alert you to situations that may cause moderate injury and/or equipment damage, and how to avoid these situations.

note	Information to help you to achieve optimal performance from your equipment.
note	The following Trademarks and Registered Trademarks are found in this manual. Teflon® is Registered Trademark of E.I. du Pont de Nemours and Company.

Before Starting:

We thank you very much for your choosing RefractoMax524 Refractive Index Detector for your High Performance Liquid Chromatography System.

RefractoMax524 Refractive Index Detector is designed to be an analytical device for pure research purpose and may not be suitable for in vitro diagnostic analysis.

To operate RefractoMax524 Refractive Index Detector properly, you are strongly recommended to go through this Operator's Manual. Improper use of this detector shall be dangerous and may cause a hazardous result.

Limited Warranty Policy

ERC warrants its products against defects in materials and workmanship for the period of twelve (12) months from the date of shipment out of its factory.

ERC will, at its option, repair or replace products that are proved to be defective.

The aforementioned warranty policy shall not be applied to defects being caused by:

- (1) Improper or Inadequate maintenance, adjustment, calibration or operation by the user(s);
- (2) User-supplied software, hardware, interfacing or consumable;
- (3) Unauthorized modification or misuse;
- (4) Operation outside of the environmental and electrical specifications for the product;
- (5) Improper site preparation and maintenance; or
- (6) User induced contamination or leaks.

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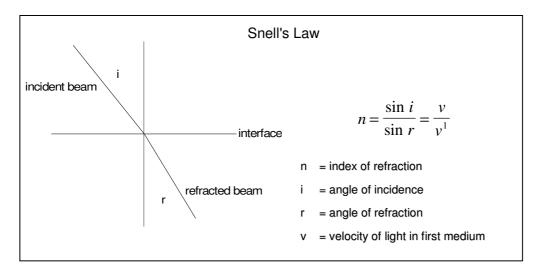
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Section 1. Introduction

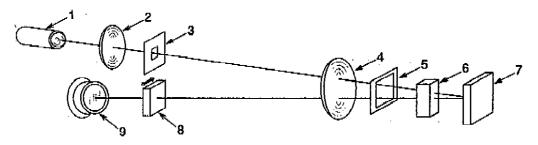
1-1. Principle of refractive index detection

RefractoMax524 Refractive Index Detector (called as "RefractoMax524" hereafter) is a high-performance universal detector designed for analyses requiring the continuous monitoring of the refractive index of a flowing liquid with respect to a reference.

RefractoMax524 is a deflection or Snell type refractive index detector. Snell's law states that a parallel light beam, when passing through a dielectric interface separating two media of different refractive index at an angle of incidence greater than zero, will be refracted as a function of the magnitude of difference of the refractive indices of the two media.



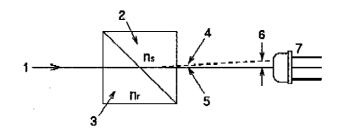
Optical System of RefractoMax524



[1] Tungsten lamp [2] condense lens [3] First Slit [4] Collimator Lens [5] Second Slit [6] Flow Cell [7] Mirror [8] Null Glass [9] Photodiode

Light from a low-power, long-lifetime tungsten lamp is collimated by a lens and slit and passed through reference and sample cells, reflected off a mirror, passed back through the optical cells, and focused by lenses onto a pair of photo sensing diodes (photo sensor).

During operation, RefractoMax524's reference and sample cells are filled with mobile phase. The reference cell is then isolated from the flow path and mobile phase flows through the sample cell only. As long as no difference exists between the refractive indices of the media of the two cells, there is no refraction of the light passing through them.



[1] Light Beam [2] Sample Cell [3] Reference Cell [4] Light Axis (Ns>Nr)
[5] Light Axis (Ns=Nr) [6] Distance between ...[4]&[5] at the photo sensor
[7] Photo sensor

Ns: Refractive Index of mobile phase in Sample Cell Nr: Refractive Index of mobile phase in Reference Cell

The light shines on a pair of photodiodes, each of which gives an electrical signal; these signals are amplified and the difference between the two signals is measured. Zero refraction should generate a zero-volt difference in these signals. An electrically controlled mechanical linkage allows the user to optimize the photodiodes' outputs for zero deflection via a refractive lens in the optical path. Additional circuitry enables the user to easily correct the signal output to electronic zero.



[1] photo sensor A [2] photo sensor B [3] light beam

When a change occurs in the refractive index of the mobile phase, the light passing through the interface between the sample and reference cells is refracted, causing the light intensity on one photodiode to increase and on the other to decrease. This difference gives a signal having both amplitude and polarity; the signal is amplified to drive a chart recorder or integrator.

1-2. Specifications

 Construction Refractive Index Range Range Linearity Noise Response Time Polarity Auto Zero Auto Zero Range Auto Zero Resolution Offset Range 	Deflection type $1.00 \sim 1.75$ $1/4 \sim 512 \mu$ RIU 600μ RIU $\leq 5 n$ RIU(Response : 3 seconds) 0.1, 0.25, 0.5, 1.0, 1.5, 2, 3, 6 sec. Positive/Negative Optical & Electrical Auto-Zero All Refractive Index Range $\leq 1 (@8mV/\mu RIU) / 4 (@2mV/\mu RIU) n$ RIU 0 to 500mV (same with Integrator output sensitivity)
 Offset Resolution Integrator Output Recorder Output 	10mV (same with Integrator output sensitivity) 0 to 1V/FS (Sensitivity: 2mV/μRIU, 8mV/μRIU) 0 to 10mV/FS
 Event Marker Temperature Control Operator Support External Communication Cell Volume Maximum Flow Rate Pressure Rating Internal Volume 	Marker out: $\geq 5\%$ of FSOFF, 30 to 55° C (1° C increment): 80° C Thermal ProtectorSpan Check/Validation DisplayRS232C2.5µl1 ml/min (mobile phase: pure water)50 kPa (0.5 kgf/cm^2)Inlet Port / Flow Cell:approx. 10μ lFlow Cell / Outlet Port:approx. 120μ lTotal:
 Wetted Material Power Requirement Power Consumption Dimensions Weight EMC Standards Safety Standards 	SST316, Teflon, Quarts Glass AC100 ~ 240V +/- 10%: 50-60Hz 120VA maximum H150 x W260 x D450 (mm) 12 kgs (26 lbs) EN61326-1,EN61326-2-1 EN61010-1
Conditions to secure safet	y (EN61010-1)

Indoor use Altitude up to 2000m Temperature 5°C to 40°C

Maximum relative humidity 80% for temperatures up to 31°C

decreasing linearly to 50% relative humidity at 40°C

Transient over voltages according to installation category II

Pollution Degree 2 in accordance with IEC 60664

Section 2. Unpacking and Installation

2-1. Installation Site Requirements

To install RefractoMax524, please make sure that any of followings shall be kept away from it to prevent interference on your analysis. Refractive Index Detector, in general, is very sensitive to the change in ambient temperature and airflow that results a drift of baseline.

- Air fan (air-conditioner-cooler, heating equipment and ventilation)
- Open doors or windows
- Direct sunlight
- Dart or dusts
- Vibration source
- Electro-Magnetic Wave or High Frequency

2-2. Power Requirements

RefractoMax524 requires 50-60 Hz single-phase power source capable of providing AC 100/240 + -10%.

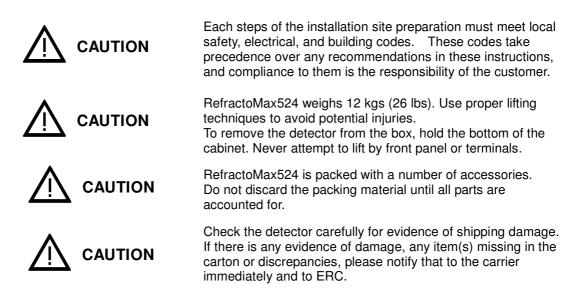


Running RefractoMax524 on a voltage other than the correct single-phase supply will void the warranty.

RefractoMax524 is designed to operate with single-phase (phase-neutral) power ONLY. If your facility provides only phase-phase (i.e., three-phase) power consult ERC.

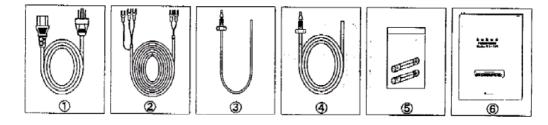
2-3. Unpacking and Inspection

This section describes in details how to install RefractoMax524. This detector is designed to be operator installed.



2-4. Standard Accessories

(1) P/N XXXXXX	Power Cord	1ea
(2) P/N 4303190	Output Signal Cable (with ferrite core)	1ea
(3) P/N XXXXXX	Inlet Tube (SST: 1.6 x 0.1 x 500mm with nut & ferrule)	1ea
(4) P/N 8809170	Outlet Tube (PTFE: 2.5 x 1.5 x 1500mm with nut & ferrule)	1ea
(5) P/N 2401120	Fuse 3.15A(T3.15AL/250V)	2ea
(6) P/N XXXXXX	Operator's Manual	1ea





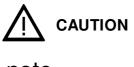
The power cord for the US and other AC 120V, 60 Hz applications is terminated in a 3-prong parallel blade plug. Outside of the 120V, 60 Hz areas, the detector shall be accompanied with a power cord (P/N 4301310). Be sure using appropriate power cord for your power source.

2-5. Loosing the locking screws

To prevent damage may occur during the shipment, the optical block is fixed by two (2) locking screws (5mm Allen bolts). As you are installing RefractoMax524, please make sure to loosen these screws.

- 1. Move RefractoMax524 to the side of a bench to make the locking screws accessible while supporting it to prevent it from falling.
- 2. Loosen the locking screws, but do not remove them, using the 5mm hexagonal key.
- 3. Move the detector securely back onto the bench.

Once the locking screws were loosened, the rubber insulator equipped with the optical block becomes functional and absorb an external shock or vibration.



Do not remove the screws. Do not run them out so far that the detector's weight rests on the screw heads instead of its feet.

note

Detector will not stabilize if locking screws are not loosened

2-6. Making a connection

2-6-1. Power Line

The power switch is located on the lower left of front panel as viewed in 4-1. The power should be OFF before connecting the power cord to the detector. The receptacle for the power cord is located on the rear of the detector. Refer to 4-2.

Connect the end of grounding cable to the ground terminals on the back panel and to a known ground.



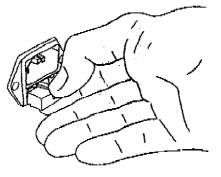
Electrically conducting spills can occur when conductive HPLC solvents are spilled on or in the instrument. Ground the instrument properly to protect the operator from electrical shock. Proper grounding also protects the instrument from power line noise. Verify that the instrument is properly grounded through the power line ground terminal. Do not remove or otherwise disable the power cord's ground prong.

2-6-2. Replacing Fuse



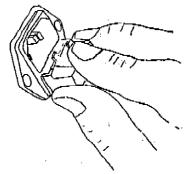
Replace only with same type and rating of fuse. Fuse for RefractMax524: P/N 2401190 3.15A(T3.15AL/250V)

To remove fuse holder



Turn off the power of detector. Remove the power cord. Push the lever located just above of Fuse holder cover. Pull the holder out as you hear clicking sound.

To re-install fuse holder



Replacing fuse. Push the fuse holder in until clicking Sound is heard. Make sure the holder is locked.

2-6-3. Tubing Connections

• Connection to the Inlet



RefractoMax524 must be the last component in your LC system. No detector or backpressure regulator can follow it. Do not use narrow-bore tubing for the outlet.

Hook up the narrow-bore stainless steel tubing of the Standard Accessory (0.25 mm ID, 1m long) with the inlet (IN) port of front panel.

• Connection on the Outlet Side

Hook up Teflon® tubing (1.5 mm ID, 1.5 m long) of the Standard Accessory with the outlet (OUT) port of front panel. Lead the opposite end of tubing into drain bottle.



Do not expose the purge valve against backpressures greater than 50 kPa (7 psi). Also, never subject the reference and sample cells to back pressures greater than 700 kPa (100 psi). High backpressures can break the cells.

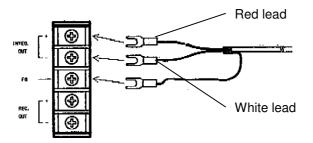
2-6-4. Cable Connections



The GROUND terminals, on the lower right side of the rear panel (see), are not the same as the recorder ground or Integrator ground terminals. Do not inter change these connections.

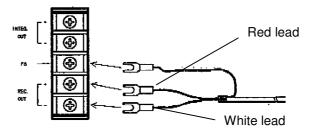
Integrator Connection

Using the output signal cable supplied in the Standard Accessory, connect one end to integrator, white lead to integrator negative (–) and red lead to integrator positive (+). Connect the third lead to the GROUND (FG).



• Recorder connection

Using the output signal cable supplied in the Standard Accessory, connect one end to the chart recorder, white lead to recorder negative (–) terminal and red lead to recorder positive (+). Connect the opposite end to the recorder's 10 mV terminal. Connect the third lead to the GROUND (FG).



2-7. Solvent Recommendations

Grade of Solvents

HPLC grade solvents are recommended for better instrument performance and chromatographic data.

Solvent Filtering

All solvents, including deionized water, should be filtered by in-line solvent filter.

Solvent Degassing

Solvents should be properly degassed prior to use. Solvents that are not properly degassed may cause bubble formation in pump heads and in the detector cells potentially causing pressure flow problems and a noisy chromatographic baseline. Degassing is especially critical when the column temperature is elevated above room temperature, or with solvents that have a high gas solubility, e.g., methanol. Analysis times that run greater than 5 to 6 hours will also require degassing of solvents.

On-line (continuous) solvent degassing is most ideal to have a constant state of mobile phase solvent over time, as the degassed solvent would start reintroducing the air.

Additional Precautions

Some solvents may corrode the wetted surfaces of the detector, if they are left in the detector after operation. The quartz cell window is easily etched by strong bases. It is recommended that some solvents be rinsed from the detector for overnight and weekend storage.

The solvents used with the detector are limited by the materials used for the wetted parts they come in contact with (e.g., quartz glass, Teflon® and 316 stainless steel). These limitations should be considered as you are choosing mobile phase solvents.

When an organic solvent that contains halogens, such as chloroform and methylene chloride, is used, flush out entire flow path with a solvent compatible with your chromatographic condition. (for example, hexane or another hydrocarbon)

For isocratic work with a normal phase column, the alcohols methanol or 2-propanol, a non-carcinogenic aromatic such as the xylenes, acetone, or an ether that is non-volatile and not a ready producer of peroxides can be used. DO NOT use ethyl ether.

If a solvent listed below is in use, flush out all the flow path sufficiently with an inert solvent that is compatible with your chromatographic system. Buffers, acids, and other highly ionic aqueous solutions should be flushed out with large amounts of water (5-10 times the volume of liquid from pump head to detector outlet If you neglect this flushing, the pump, injector, and column may become corroded and badly damaged.

Sulfuric Acid, Boric Acid, Citric Acid, Acetic Acid, Lactic Acid, Acetic Anhydride, KOH, NaOH, Hydrazine, Sodium formate Ammonium salts: -formate, -perchlorate, - nitrate, -citrate, -oxalate, -sulfate, -H₂PO₄, K₂CO₃ K, Na salts: -bicarbonate, -chlorate, -nitrite

Following solvents should be avoided.

```
Hydrohalogenic, Metal Halides >2M, KCI, Ammonium Halides,
Ammonium Formate, All hypochlorites, Tetrachloromethane
Acids: HCI, HF, etc.
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Fluorocarbon solvents will alter Teflon® over long exposure. Flush with pentane or another light hydrocarbon.

note

If you want to replace one solvent with an immiscible one, flush out the existing mobile phase with an intermediary solvent inter-miscible with your initial and final solvents. For example, you want to replace water in your HPLC with chloroform; water and chloroform are immiscible. Replace the water in your system with 2-propanol, which is freely miscible with water and chloroform. When you are certain all water is removed, replace the 2-propanol with chloroform. See the Miscibility Chart in Appendix.

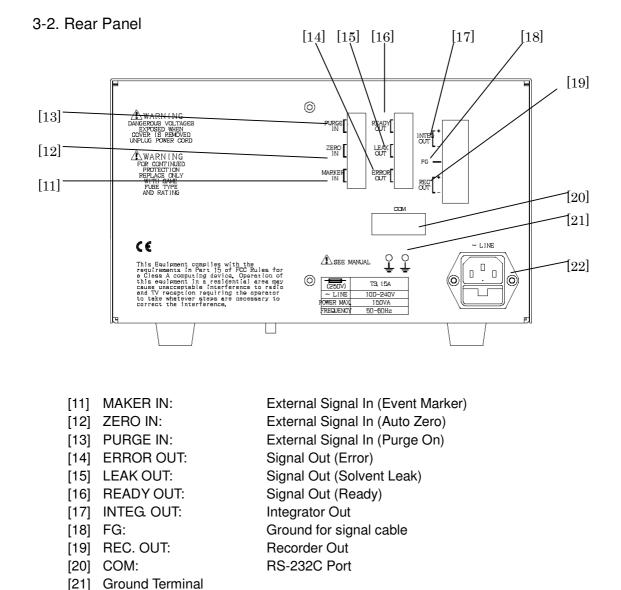
2-8. Characteristics of commonly used mobile phase

						(Those in bold should not be employed)					
Polar		Polarity Viscosity	Refractive	UV Cut-off	Flash Point	Fire Point			Vapor	Boiling Point	Gravity
	$E^2(Al_2O_3)$	_c P20 °C	Index	(nm)	(°C)	(°C)	Upper	Lower	Density	(°C)	
Fluoroalkanes	-0.25		1.25	()	()					()	
n-Pentane	0.00	0.23	1.358	210	<-40	308.9	1.5	7.8	2.5	36.1	0.6
Hexane	0.00	0.20	1.375	210	-21.7	233.9	1.2	7.5	3.0	68.9	0.0
Isooctane	0.00		1.404	210	-21.7	200.0	1.2	7.5	0.0	00.5	0.7
Petroleum ether	0.01	0.3	1.404	210							
n-Decan	0.01	0.92	1.412	210	46.1	207.8	0.8	5.4	4.9	173.9	0.7
Cyclohexane	0.04	1.00	1.427	210	-20	260	1.3	8	2.9	81.7	0.8
Cyclopentane	0.04	0.47	1.406	210	-20	200	1.5	0	2.3	01.7	0.0
Diisobutylene	0.05	0.47	1.411	210							
i-Pentene	0.08		1.371	210	-17.8	272.8	1.5	8.7	2.4	30	0.7
Carbon disulfide	0.08	0.37	1.626	380	-17.8	100	1.3	44	2.4	46.1	1.3
Carbon tetrachloride	0.15	0.37	1.466	265	-30	100	1.5	44	2.0	40.1	1.5
					10.0	040.0	1.0	0.0	0.7	100.1	0.0
Amyl chloride	0.26	0.43	1.413	225	12.8	343.3	1.6	8.6	3.7	106.1	0.9
Busy chloride	0.26		1.436	220	-9.4 o-17.2	460 463.9	1.8 1.6	10.1 6.0	3.2	144.4	
-		0.00.0	9 to 1 50					6.0 7.0	0 70		0.0
Xylene	0.26	0.62-0.	8 to 1.50	290	m-25	527.8	1.1		3.78	138.9	0.9
-		0.07	1.000		p-25	528.9	1.1	7.0	0.5	138.3	0.7
i-Propyl ether	0.28	0.37	1.368	220	-27.8	443.3	1.4	21	3.5	68.9	0.7
i-Propyl chloride	0.29	0.33	1.378	225	-32.2	593.3	2.8	1037	2.7	35	0.9
Tolune	0.29	0.59	1.496	285	4.4	536.1	1.4	6.7	3.1	110.6	0.9
n-Propyl-chloride	0.30	0.35	1.389	225	<-17.8		2.6	11.1	2.7	46.1	0.9
Chlorobenzene	0.30	0.80	1.525		32.2	637.8	1.3	7.1	3.9	132.2	1.1
Benzene	0.32	0.65	1.501	280	-11.1	562.2	1.4	7.1	2.8	80	0.9
Ethyl bromide	0.37		1.424			511.1	6.7	11.3	3.8	37.8	1.4
Ethyl ether	0.38	0.23	1.353	220	-45	180	1.9	48	2.6	35	0.7
Ethyl sulfide	0.38	0.45	1.442	290							
Chloroform	0.40	0.57	1.443	245							
Methylene-chlolride	0.42	0.44	1.424	245	-50	518.9	3.8	15.4	2.2	38.5	0.9
Methyl I-butyl ketone	0.43		1.394	330							
Tetrahydrofurane	0.45		1.408	220	-14.4	321.1	2	11.8	2.5	66.1	0.9
Ethylen dichloride	0.49	0.79	1.445	230	13.3	412.3	6.2	16	3.4	83.9	1.3
Methyl ethyl ketone	0.51		1.381	330	-6.1	515.6	1.8	10	2.5	80	0.8
i-Nitropropane	0.53		1.400	380	48.9	420.6	2.6		3.14	131.1	1.0
Acetone	0.56	0.32	1.359	220	-17.8	537.8	2.6	12.8	2.0	56.7	0.8
Dioxane	0.56	1.54	1.422	260	12.2	180	2.0	22	3.0	101.1	1.0
Ethyl acetate	0.58	0.45	1.370	260	4.4	460	1.8	8	3.5	90	0.9
Methyl acetate	0.60	0.37	1.362	210	-10	501.7	3.1	16	2.6	60	0.9
Amyl alcohol	0.61	4.1	1.410		32.8	300	1.2	10.0	3.0	137.8	0.8
Dimethyl sulfoxide	0.62	2.24									
Aniline	0.62	4.4	1.586		70	617.2	1.3		3.2	184.4	1.0
Dimethyl amine	0.63	0.38	1.387	275	<-17.8	312.2	1.8	10.1	2.5	56.7	0.7
Nitromethane	0.64	0.67	1.394	380	35	418.3	7.3		2.1	101.1	1.1
Acetonitrile	0.65	0.37	1.344	210	5.6				1.4	81.7	0.8
Pyridine	0.71	0.94	1.510	305	20		1.8	12.4	2.7	115	1.0
Butyl cellosolve	0.74			220							
i-Propanol		0.0	1.00		44 7	000.0	0.0	10	0.1	00.0	0.0
n-Propanol	0.82	2.3	1.38	210	11.7	398.9	2.0	12	2.1	82.8	0.8
Ethanol	0.88	1.20	1.361	210	12.8	422.8	4.3	19	1.6	78.3	0.8
Methanol	0.95	0.60	1.329	210	11.1	463.9	7.3	36	1.1	63.9	0.8
Ethylene glycol	1.11	19.9	1.427	210	111.1	412.8	3.2	-		197.2	1.1

(Those in bold should not be employed)

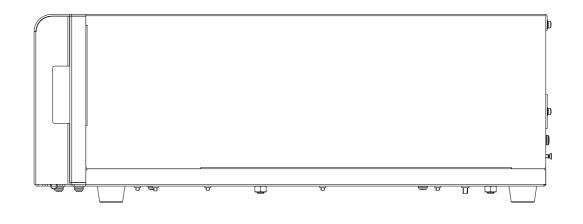
ERC [6]- \triangleright ·[9] 0 IN [8] Erro Aute Zero Pur∮ Marke Ente [13]-Powe [4] Q RefractoMax 52 [1] Т Ш st T म₹ Τ [5] [10] [3] [2] [12] [11] [1] Power Switch: Press this key once to turn on or off the unit. [2] Event Marker Key: Press to generate a marker signal. (5% of Full Scale) [3] Purge Key: Press to turn purge valve on or off to change flow path. When the valve is on, LED above the key illuminates and "PURGE" appears on the lower line of LCD display. Solvent flows through reference side of flow cell instead of sample side. [4] Auto Set Up Key: Press this key to do Auto Set Up. As "Auto Set Up" is going on, LED above the key illuminates and "AUTO" appears at the lower line of LCD display. Press this key to do Auto-Zero. As Auto-Zero is going on, [5] Auto-Zero Key: LED above is illuminated and "AUTO ZERO" appears on the lower line of LCD display. [6] LCD Display: Liquid Crystal Display [7] Arrow Keys: Press to change screen, to move cursor or to edit values. [8] **Outlet Port:** [9] Inlet Port: Press to save the edited data or to confirm the command. [10] Enter Key: [11] Cancel Key: Press this key to scrap the edited data. [12] Escape Key: Press this key once to return to Monitor Screen. [13] Error LED: Error message

Section 3. Function Keys and Displays 3-1. Front Panel



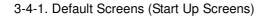
3-3. Side View

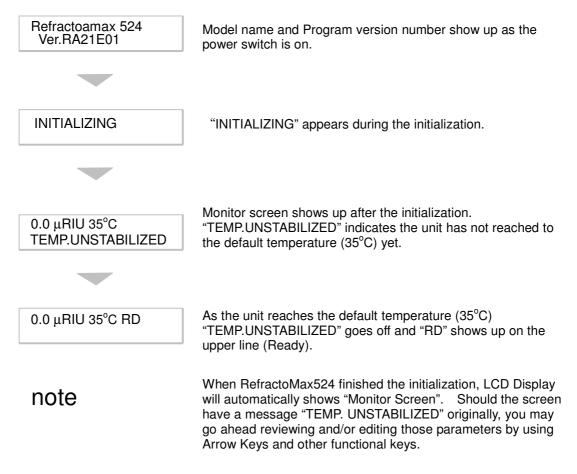
[22] Power Inlet



3-4. LCD Display (User Screens)

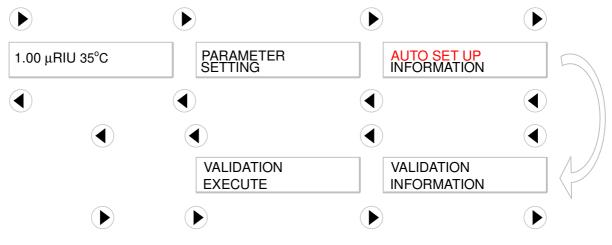
RefractoMax524 has a LCD display on the front panel as shown in 3-1. Front View here above. The display consists of following four screens and by solvent delivery pump, connect a tubing line directly from the pump to RefractoMax524 inlet port bypassing the column. Some cleaning solutions should be injected, however, directly into the flow cells by syringe due to their high corrosiveness or safety concern.



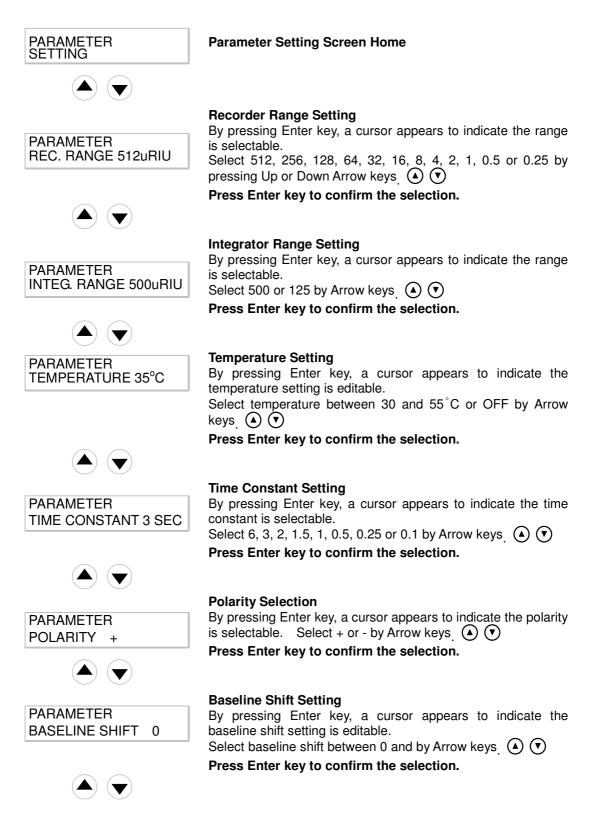


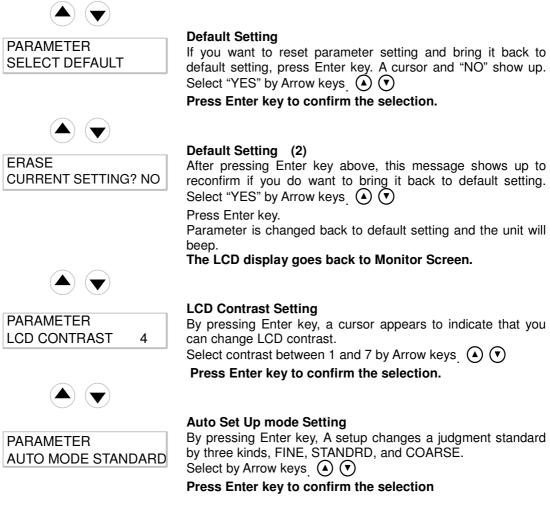
3-4-2. Screen Sequence

By pressing Arrow Keys, LCD Display will change from one screen to the other as follow



3-4-3. Parameter Setting Screen





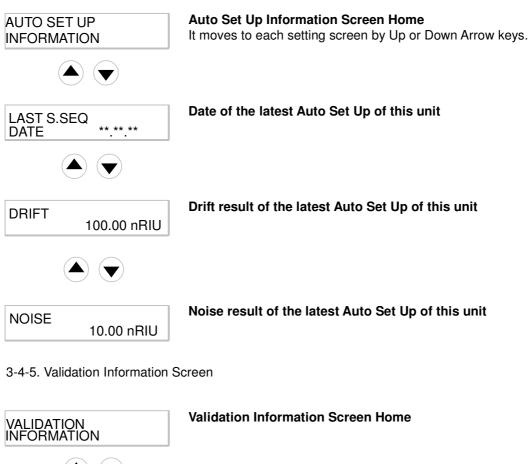
Auto Set Up mode

	Fine	Standard (Default)	Coarse			
Purge Cycle		30 sec.				
Number of Cycle		3				
Time to Auto Zero		240 sec.				
Equilibration Time	80 min.	60 min.	40 min.			
Measuring Time	80 min.	60 min.	40 min.			
Drift	100 nRIU/h	500 nRIU/h	2500 nRIU/h			
Noise		50 nRIU				

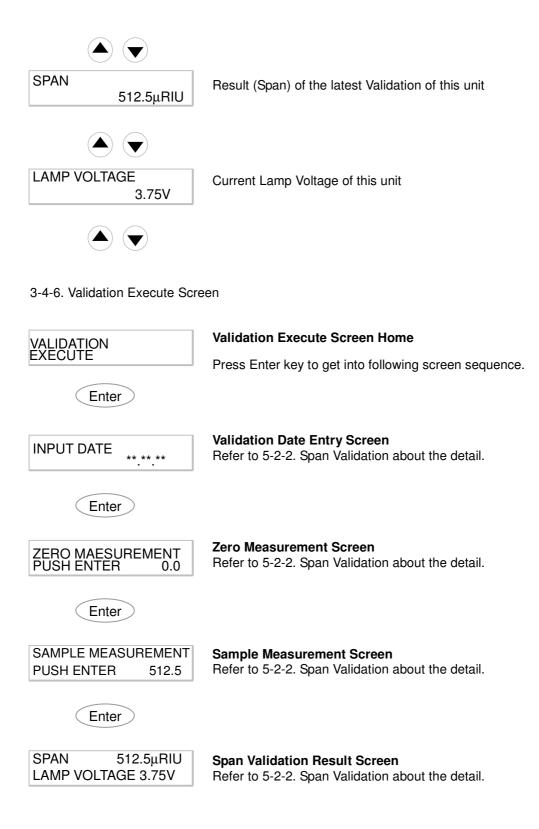
Parameter Default Setting

Parameter	Setting value	Unit	Default
REC. RANGE	0.25, 0.5, 1, 2, 4, 8, 16, 32, 64, 128, 256, 512 (12 steps)	μRIU/10mV	512
INTEG. RANGE	125 or 500	μRIU/1V	500
TEMPERATURE	OFF, 30 to 55 (increment: 1°C)	Celsius	35°C
TIME CONSTANT	0.1, 0.25, 0.5, 1, 1.5, 2, 3, 6 (8 steps)	Second	3
POLARITY	+ or -	N/A	+
BASELINE SHIFT	0 to 50 (increment: 10mV)	10mV	0
SELECT DEFAULT	YES or NO	N/A	NO
LCD CONTRAST	1 to 7 (increment: 1)	N/A	4

^{3-4-4.} Auto Set Up Information Screen







3-4-7. Error Messages

To alert you about the situation that is hazardous or may cause a deterioration of your analysis, RefractoMax524 flashes following five different error messages.

100.0 μRIU 35°C LEAKAGE ERROR **Solvent Leak Error** This indicates there is a solvent leakage inside of cabinet.



When this error message comes on, power off the detector at once.

Please double-check if there is any solvent leak. In case you cannot fix problems, please contact our local representatives in your area.

Null Glass Home Position Error

This indicates the null glass doesn't come back to its home position.

If you are commanding Auto-Zero via external control devise, refer to the procedure for "Optical Balance Error.

If you have this error as you are turning on RefractoMax524, please contact our local representatives in your area.

100.0 μRIU 70°C
OVERHEAT ERROR

600.0 μRIU 35°C

HOME POSITION ERROR

Overheating Error

This indicates overheating of optical block. Please contact our local representatives in your area.

Optical Balance Error

100.0 μRIU 70°C OPT.BALANCE ERROR	This indicates that Auto-Zero wasn't successful. Insufficient solvent exchange of reference flow path might cause this.
	Try purging procedure to fill reference flow path with fresh solvent at once.
100.0 μRIU 35°C	Parameter Error
PARAMETER ERROR	This indicates the units failed to store data.

3-4-8. Other Messages

There are four other messages as described below. Some of them are more like a warning message.

0.0 μRIU 35°C INTENSITY

Low Light Intensity

This indicates inadequate light intensity. There are several possible causes for this error as follow.

- 1. Different state of solvents between sample-side flow cell and reference-side flow cell. Purge reference-side flow path with fresh solvent.
- 2. Air bubble in flow cells. Repeat purge on and off with flow to remove the air bubble.
- 3. Optical axis is off from the center. Press Zero Key to do "Auto-Zero".

Unstable Temperature

0.0 μRIU 35°C **TEMP.UNSTABILIZED**

This indicates temperature of optical system hasn't met the preset temperature.

0.0 μRIU 35°C PURGE

Purge

This indicates the purge valve is on. Solvent flows through reference side of flow cell instead of sample side.

0.0 μRIU 35°C	
AUTO ZERO	

Auto-Zero

This indicates the purge valve is on. Solvent flows through reference side of flow cell instead of sample side.

Section 4. Getting Started

4-1. Just in Case !

Prior to operation, check the following one more time.

- Locking screws are loosened.
- Mobile phase solvent is freshly made and degassed well.
- All wetted parts are chemically compatible to the mobile phase solvent.
- Mobile phase flushed through the entire flow path; all incompatible or immiscible solvents have been flushed out.
- Cable connections are properly made to chart recorder, integrator, data system, or other external equipment.
- All tubing connections are properly made and checked for leaks.
- Power cord is plugged into appropriate power receptacle.
- Proper fuse is installed.
- Drain tube is installed.
- Power switch is ON.



Before activating the purge valve (LED light OFF), pump about 10 ml of liquid through the cell.

This will flush possible dust or particulate matter and reduce the possibility of damaging the valve seals.

- 4-2. Start Up
 - (1) Start pumping mobile phase solvent at flow rate 1 ml/min to reference cell (Purge On).
 - (2) Press Purge key in every 10 seconds to on/off the purge valve for few minutes.
 - (3) Keep pumping mobile phase solvent to reference cell for about 20 minutes from the above step (2).
 - (4) Press Purge key to turn off the valve. Mobile phase solvent flows to sample-side flow cell.
 - (5) Wait until the baseline is stabilized.
 - (6) Press Zero Key to do Auto Zero.



Wearing protective gloves and goggles is advised.

4-3. Parameter Setting

Set the parameter on Parameter-Setting Screen (Refer to 4-3-3).

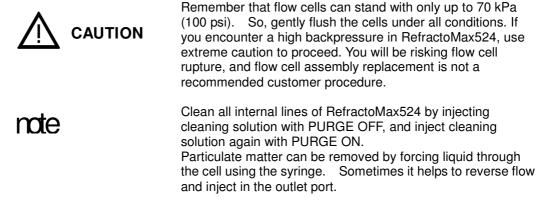
Section 5. Maintenance and Validation

5-1. Cell Cleaning

In many cases, performance degradation in sensitive instruments equipped with flow-through cells is caused by cell contamination. The use of filtered solvents with in-line solvent filters will protect the cell from contamination and reduce the amount of cleaning required. However contamination from trapped particulates or bubbles, from precipitates, or from thin films of residues can still occur.

5-1-1. Preparations

To introduce cleaning solution into RefractoMax524 by solvent delivery pump, connect a tubing line directly from the pump to RefractoMax524 inlet port bypassing the column. Some cleaning solutions should be injected, however, directly into the flow cells by syringe due to their high corrosiveness or safety concern.



5-1-2. Cleaning

Depends on the solvents in use, the cleaning procedure is varied. Following is a procedure for typical.

- (1) Inject cleaning solution (acetone) by syringe from the inlet port (5 ml).
- (2) Inject de-ionized water by syringe from the inlet port (5 ml).
- (3) Inject nitric acid solution (15%) by syringe from the inlet port (5 ml).
- (4) Expel nitric acid solution completely by flowing de-ionized water adequately.
- (5) Exchange de-ionized water with the mobile phase solvent.

If buffers or solutions of high salt content have been in use, the cells may be contaminated by precipitated salt. Pumping a large amount of distilled, de-ionized water, such as 1 ml/min, for up to several hours, is the simplest clean-up procedure.

An elevated cell temperature will speed dissolution. The water wash can be acidified, if the precipitated salt is more soluble in acidic solutions. However, do not use strongly basic (pH 10 or higher) solutions, as these will etch the refractive index cells.

If contamination is suspected when a non-aqueous solvent is in use, flush the cells with a solvent that is (1) miscible with your mobile phase, (2) a good solvent for the predicted contaminant, and (3) generally of greater polarity than your mobile phase.



Do not allow nitric acid to contact methanol. An explosion could result. Completely rinse the flow cell with water following cleaning with nitric acid.



Corrosive acids are used. Use extreme caution to avoid spillage on skin, clothing, or the instrument. Protective gloves are advised.



Never put hydrochloric acid in the cell. This acid in any concentration will corrode the cell. Diluted (10-20%) or concentrated nitric acid is a good cleaning solution. The sample and reference cells should be filled with water or air (blown dry) before proceeding.

Filled the syringe with cleaning solution and connect to the inlet port of the RefractoMax524. Carefully inject the cleaning solution. For safety, make sure that outlet tubing is led to drain bottle.

Flush acid cleaning solutions from the cells with large amounts of water, such as 1 ml/min. Flush for 15 to 30 minutes.

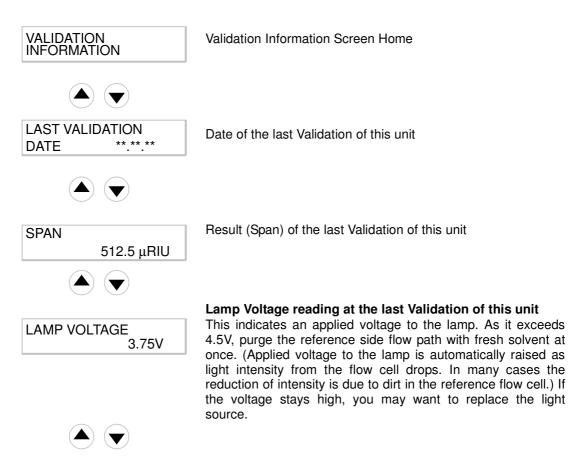


For shutdown and storage, please review Section 7. Shutdown Procedure.

5-2. Validation

From time to time, you are recommended to validate your HPLC system to keep an accuracy and credibility of your analysis.

5-2-1. Validation Information



5-2-2. Span Validation

To do Span Validation, follow the below procedure.

(1) Preparation of standard sucrose solution:

Weigh out 350 mg sucrose and transfer quantitatively to a volumetric flask. Dissolve in the 100 ml de-ionized, filtered, degassed water and dilute to flask's mark.

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note
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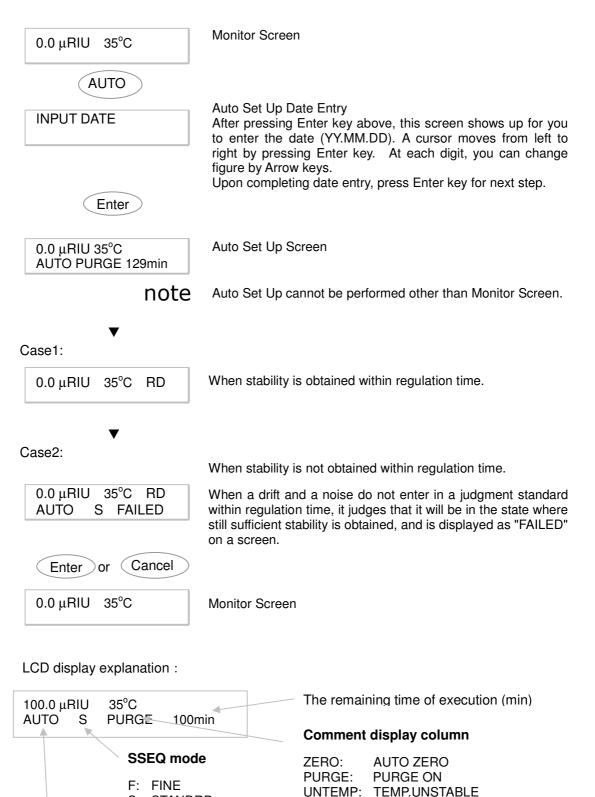
Use freshly made sucrose solution always.

- (2) Equilibrate RefractoMax524 by pumping de-ionized water through both reference and sample cells. Use the same de-ionized water as that used to prepare the sucrose standard solution. Start pumping at flow rate to 1 ml/min.
- (3) Make sure that the baseline is stabilized and the drift is equal or less than 500 nRIU/h by monitoring recorder chart.

VALIDATION EXECUTE					
Enter	Press Enter key to get into following screen sequence.				
INPUT DATE	Validation Date Entry After pressing Enter key above, this screen shows up for you enter the date (YY.MM.DD). A cursor moves from left to right by pressing Enter key. At each digit, you can change figure by Arrow keys. Upon completing date entry, press Enter key for next step.				
Enter					
ZERO MEASUREMENT PUSH ENTER 0.0	Zero Measurement Screen Press Auto-Zero key to do Auto Zero.				
Enter					
SAMPLE MEASUREMENT PUSH ENTER 512.5	Sample Measurement Screen				
(5) Make sure that the purg(6) Filled the syringe with s(7) Validation screen will co(8) The result should be with	tubing from the inlet port of RefractoMax524. The valve is off. Tandard sucrose solution and gently inject from the inlet port. The on with the measuring result. Thin 487 to 537 (512 μ RIU +/-5%) The vove the above sample measurement result.				
Enter					
SPAN 512.5 μRIU LAMP VOLTAGE 3.75V					
Enter	By pressing Enter key, the above validation results will be stored. The screen will go back to Validation Execute Home Screen				

NOte Make sure that you have put the tube back to inlet port.

5-2-3. Auto Set Up Execute



Indicates that RefractoMax524 is under Auto Set Up

S: STANDRD

C: COARSE

INTENS:

INTENSITY

note:

- 1) When an error occurs during Auto Set Up execution, execution is stopped and the contents of an error are displayed.
- 2) Communication (RS232C) with the exterior cannot be performed during Auto Set Up execution.
- 3) During Auto Set Up execution, Escape key is invalid.

Section 6. Shutdown Procedure

6-1. Corrosive Solvents

Some solvents may corrode the detector, if they are left in the detector and should be thoroughly flushed from the entire system, including the reference and sample flow cell.

The quartz flow cell window, in particular, is easily etched by strong bases. Do not turn power to RefractoMax524 off without rinsing these solvents from the detector.

Some solvents can be left in the cells at the end of an operation. For example, water, acetonitrile, 2-propanol, the xylenes, and paraffinic hydrocarbons are quite innocuous. They may be left in RefractoMax524 overnight or over a weekend.

6-2. No Flow Shutdown Versus Reduced Flow Shutdown

A continuous slow flow through RefractoMax524 is the preferable shutdown procedure if the situation permits. (Especially if buffers, tetrahydrofuran and organohalocarbons are in use.)

Reduced flow may be 0.5 ml/min to 0.01 ml/min but the HPLC pump must be able to stay primed at reduced flow.

Buffers:

Even if the buffer is non-corrosive, it is better to keep the solvent flowing at a reduced rate to eliminate the possibility of salt precipitation in the flow cells and tubing.

Tetrahydrofuran:

Because THF does oxidize, you may find that, if you keep solvent flowing at a reduced rate, the chromatographic system takes less time to re-stabilize upon start-up. Generally a reduced-flow shutdown procedure will minimize re-stabilization time; the time saved is noticeable with THF as the solvent.

Organohalocarbons, such as Methylene chloride and Chloroform:

Keep a small amount of flow to keep down the amount of corrosive chloride impurities in the cell.

6-3. Long-Term Storage

If RefractoMax524 will not be used for a week or more, the sample and reference cells should be blown dried.

If the detector is to be exposed to sub-freezing temperatures, an antifreeze flush, such as methanol must be used. Ideally, you are suggested to expel residual solution out of RefractoMax524 and to blown dry.

Section 7. Troubleshooting

7-1. Introduction

Malfunctions within RefractoMax524 can arise from three general sources:

- RefractoMax524 itself can be dirty, or operating non-optimally.
- The HPLC system can have a broken, dirty, or non-optimally operating component, but the problem is manifesting itself in RefractoMax524.
- A mobile phase and/or column problem, which by its very nature is spread throughout the HPLC system but appears as a malfunction of RefractoMax524.

To troubleshoot, you must be able to isolate the performance of RefractoMax524 within the HPLC system from its performance outside the HPLC system. Therefore, this section begins with guidelines for testing RefractoMax524 as a stand-alone.

Following is the Troubleshooting Table that lists the observed phenomenon with the possible cause and the suggested solution.

7-2. Make sure how the detector itself is working.

You must know how well the detector performs by itself before you can troubleshoot it in an HPLC system.

To perform the tests, disconnect all cables from the detector except one signal cable from recorder output to a calibrated, functioning recorder. The recorder should match input to the output of RefractoMax524 (10 mV).

Before proceeding, verify also that the locking screws on the optics module have been loosened; that the correct voltage is supplied and the correct fuse installed; and that degassed water is in the reference cell, the sample cell, and the entire flow path.

7-3. To isolate the cause of problem

As you go through the Troubleshooting Table, you will have occasional instructions for HPLC systems. Generally, however, the rule of thumb is to add one component at a time back into the HPLC system so that, should the condition arise again, the component causing the problem is indicated.

You will begin by adding the pump to RefractoMax524 first and you will add the column last.

If another type of detector is available, it is a good idea to use it before RefractoMax524 to aid in troubleshooting as reference. Let's assume that you have a UV detector before RefractoMax524 and only RefractoMax524 has a noisy baseline.

One possible implication is that the noise arises from pressure fluctuation, to which the refractive index detector is more sensitive. On the other hand, if both detectors are showing noise, a power line current may be indicated.

If both detectors show anomalous baseline performance, such as huge peaks that continue indefinitely, a bleed-off problem (material from the column or immiscible solvents trapped in the system) is more likely.

7-4. Troubleshooting (Diagnosis Chart)

Problem	Possible Cause	Solution						
Noise	Bubble in the pump.	Purge pump heads. Use only premixed/degassed mobile phase.						
	Bubble in the detector.	Elevate drain reservoir above the level of the flow cells to create slight backpressure. Premix and degas mobile phase well.						
	Dirty flow cell(s).	Clean flow cells. Refer to Cell Cleaning.						
	Weak lamp.	Check validation screen.						
	Ambient temperature fluctuations.	Move detector to a more stable environment.						
	Vapor pressure of the mobile phase is too high for the detector.	Reduce or turn temperature control off. Modify method to exclude or decrease concentration of						
		troublesome solvent.						
	Electrical transients from power line or radio frequency source.	Isolate the detector power source from other heavy equipment, motors, etc.						
		Ground the detector to earth.						
Noise appears in 8-10 hours of operation.	Formation of gases in the mobile phase reservoir.	Consider using on-line solvent degasser(R).						
Cyclic Noise	Ambient temperature fluctuation.	Move the detector to a more stable environment. Place a cover over the detector						
	Bubbles in reference cell.	Flush detector with Purge on/off						
Cyclic Noise matching pump stroke frequency	Drain tube is too small (narrow bore).	Verify that correct exit line is installed. Check outlet tubing for crimps						
Drift	Flow cell(s) dirty.	Clean reference and sample cells. Refer to Cell cleaning procedure.						
	Flow cell(s) damaged.	Check for liquid in rear drain tube indicating a broken cell						
	Contamination from HPLC System.	Flush HPLC system with a solvent stronger than the mobile phase (less polar for reverse phase, more polar for normal phase, etc.) until contaminant disappears.						

Problem	Possible Cause	Solution						
Drift (continued)	Contaminated or non-HPLC grade solvents.	Prepare fresh mobile phase (premixed/degassed)						
	Vapor pressure of mobile phase is too high for operating temperature causing bubble formation in reference cell.	Reduce or turn temperature control off. Modify method to exclude or decrease concentration of troublesome solvent.						
	Tetrahydrofuran (THF) in the mobile phase will oxidize in the reference cell.	Add an antioxidant to stabilize the THF, if compatible with other chromatographic requirements. Allow >2 hours stabilization time for oxidation in reference cell to reach a steady state condition.						
Baseline drift occurs in few hours of start.	Reference cell solvent has aged and deteriorated.	Flush reference cell with mobile phase.						
Baseline will not zero	Sample and reference cells do not contain identical solutions.	Flush sample and reference cells with mobile phase.						
	Reference cell contains air bubbles.	Flush sample and reference cells with mobile phase.						
	Flow cell(s) dirty.	Clean flow cell. Refer to Cell Cleaning procedure.						
	Flow cell(s) damaged.	Check for liquid in drain tube indicating a broken cell.						
	Deteriorating lamp or lamp out of adjustment.	Check validation screen.						
Intensity Alarm	Flow cell(s) dirty.	Clean flow cell. Refer to Cell cleaning procedure.						
	Lamp is burned out.							

Appendix. Miscibility Chart of Solvents

Acetic Acid	1															
Acetonitrile		1														
Chloroform			1													
Cyclohexane		Х]												
Methylene Chloride]				X =	= Imr	nisc	ible				
Dimethyl Formamide				Х]										
Dioxane								_								
Ethyl Ether																
Hexane		Х				Х				_						
Methanol				Х					Х							
Methyl t-Butyl Ether																
Trimethylpentane		Х				Х				Х						
Pentane		Х				Х				Х						
Propanol-2															_	
Tetrahydrofuran																
Water			Х	Х	Х			Х	Х		Х	Х	Х			
	Acetic Acid	Acetonitrile	Chloroform	Cyclohexane	Methylene Chloride	Dimethyl Formamide	Dioxane	Ethyl Ether ×	Hexane	Methano	Methyl t-Butyl Ether ×	Trimethylpentane	Pentane	Propanol-2	Tetrahydrofuran	Water

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