SIPRA Labs Limited 7-2-1813/5/A, Adjacent to post office Sanath Nagar Industrial Estate Hyderabad

SUB: Two Decades of Regulatory Services to Pharmaceutical Industry

Dear Sir/Madam,

We are glad to inform you that Sipra is completing **Two Decades** of fruitful association with the pharmaceutical industry by providing the Regulatory Research services. Sipra always believed in the policy of "Compliance First" and "Customer is our Boss" and strived all these years to fulfill the wide ranging aspirations of its esteemed clientele.

Date: 21/04/14

We are deeply indebted for the continued support provided by you, your organisation and industry as a whole. All these years, your organisation has appreciated our technical strengths and allowed us to continue our services.

On this auspicious occasion of completing **Two Decades** in customer service, we are launching a program called **"SIPRA FIRST"**, means **SIPRA** has to receive the first call from their clients for their future requirements.

In order to achieve these and to improve our services further, we are creating a data base of our esteemed clients, and their current and future outsourcing needs. We would like to have an interaction with you to get your honest and worthy feedback on our services as part of Two Decades in customer service. Also we will be in regular touch with you through Client Relation Management for every enquiry and guery you have on our services.

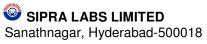
We thank you very much for your kind support and the honest feedback.

Thanking you,

From

Sipra labs

"SIPRA FIRST" FORM



Form No:

) Chairman:) Managing Director:) Head Office Address: Contact No:) Manufacturing / R & D Plants (Please provide the name of the locations): S. No (1) (2) (3) (4) R & D API Formulations Regulatory Status (Ex: USFDA, TGA, & etc)) Kindly specify your future expansions under planning? I Kindly specify your future products under planning and are you going to launch these products ationally or internationally? (if possible) I Kindly specify regulatory planning's of your Organisation?	1) Name of Organisati	on:			
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	nationally or internation	nally? (if possible)			
	3) Kindly specify regul	atory planning's of yo	ur Organisation?		
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) What are your expectations from an Outsourcing Partner?	3) What are your expe	ctations from an Outs	ourcing Partner?		

10) Kindly suggest some outsourcing services you are feeling difficult to manage?

SECTION II: Details of the Person Filling the Form and his Unit

1) Name of the Person:	
2) Designation:	
3) Unit & Division:	Buoto
4) Job Description:	PHOTO
5) Number of People Working in Your Team:	
6) Regulatory Compliance of Your Facility:	Phone:
, .g , ,	Email:
7) Kindly specify major test facilities/instruments available in your QC/organis	<u>l</u> ation:
8) Do you have any specific requirement of Instrument or Instrument based s	tudies?
Please specify the pharmacopieal studies being outsourced by your organics	isation:
of thouse speemy the prialinasspied stadies somig substances by year engain	oano
10) Does your facility/organisation require any additional studies for outsource	ing (for procent & uncoming
products)?	ing (for present & apcoming
11) Facility monitoring services (particle count, bio-load, flow and pressures)	are launched by SIPRA
recently. Does your GMP facility require it?	
12) Do you have any requirement of CRO services?	
1.27 2.3 year mane any requirement of error continues.	
13) Please name the services your organisation is outsourcing to others (other	er than to SIPRA)?
14) Can you give details of your associates who require our services?	
14) Carr you give details or your associates who require our services?	
15) How long you are associated with SIPRA?	
13) How long you are associated with SIFHA!	
40) K'adh assa' talan assay (assahada a OIDDA	
16) Kindly specify the reasons for selecting SIPRA:	

SECTION III: About SIPRA

1) Analytical

1) Present Service Group of SIPRA

4) Packaging Material

			O)1 Tecimical Studies	
	7) Invitro Studies	8) Facility Monitoring	9) Microbiology	
	10) Biotechnology	11) Environmental Studies		
		•		
2) R	legulatory Status of SIPRA is			
	1) US FDA	2) TGA Australia	3) Health Canada	
	4) MCC South Africa	5) NABL 17025	6) Medical ISO 15189: 2007	
	7) DCGI	8) DST approved R & D	9) CPCSEA registered animals facility	
3) N	lew Services added in the last		0) 0	
	1) HPLC – ELSD & ECD	2) Air jet Sieve	3) Scanning Electron Microscope	
	4) GC-MS/MS	5) IC- Amperometry	6) GC – Gas Analysis	
4\ C	tions to a suite and suitable also seems	was and in a second of LICE. Dance		
4) S			rour organisation interested in any of the these	
	1) Leachables and Extractat	oles 2) ICP-MS studies	3) Invitro Assays	
5) S	ipra is planning to purchase fo	allowing instruments this year		
٥, ٥	1) NMR 500 MHz	2) USP-4 Dissolution	3) BIA-CORE	
	.,	2) 331 1 21330141011	O/ Birt GOTTE	
Val	may Tick the above service	es of vour interest or service	s used by your facility in SIPRA	
2) K	indly aposity any other regula	tory requirement, which help t	as industry (other than appoified shove):	
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3) H	low SIPBA can participate in v			
-, -	on the contract of the contrac	our future activities as an out	sourcing partner?	
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		our future activities as an out	sourcing partner?	
		our future activities as an out	sourcing partner?	
4) C	an you suggest the improvem	your future activities as an outs		
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2) Stability Studies

5) BA/BE Studies

3) Method Development

6)Preclinical Studies