केन्द्रीय आयुर्वेदीय विज्ञान अनुसंधान परिषद्
आयुष मन्त्रालय, भारत सरकार
जवाहर लाल नेहरू भारतीय चिकित्सा एवं होम्योपैथी अनुसंधान भवन
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CENTRAL COUNCIL FOR RESEARCH IN AYURVEDIC SCIENCES
Ministry of AYUSH, Govt. of India
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EXPRESSION OF INTEREST (EOI) FOR MANUFACTURING AND SUPPLY OF AYURVEDIC DRUGS FOR RESEARCH PURPOSE IN CCRAS

Advertisement No. TECH/EOI - 01 /2015

Central Council for Research in Ayurvedic Sciences (CCRAS) invites Expression of Interest from experienced and capable GMP certified Ayurvedic Pharmaceutical Industries in India.

The details are available at www.ccras.nic.in. The interested firms may submit Expression of Interest by 25th May 2015 as per guidelines given at above cited website. CCRAS reserves the right to accept or reject any or all applications without assigning any reason.

DIRECTOR GENERAL
CCRAS, NEW DELHI

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केंद्रीय आयुर्वेदीय विज्ञान अनुसंधान परिषद् में अनुसंधान के उद्देश्य हेतु आयुर्वेदिक औषधियों के विनिर्माण एवं आपूर्ति हेतु रुचि की अभिव्यक्ति

विज्ञापन सं. तकनीकी/रुचि की अभिव्यक्ति-01/2015

केंद्रीय आयुर्वेदीय विज्ञान अनुसंधान परिषद्(सीसीआरएएस) भारत में अनुभवी एवं योग्य जी.एम.पी. प्रमाणित आयुर्वेदिक भेषज निर्माण उद्योगों से रुचि की अभिव्यक्ति आमंत्रित करती है ।

विवरण www.ccras.nic.in. में उपलब्ध है। इच्छुक फर्में उक्त वेबसाईट में दिए गए दिशा- निर्देशों के अनुसार रुचि की अभिव्यक्ति को 25 मई, 2015 तक प्रस्तुत कर सकती हैं। केंद्रीय आयुर्वेदीय विज्ञान अनुसंधान परिषद् के पास किसी या सभी आवेदनों को बिना कोई कारण बताए स्वीकार या अस्वीकार करने का अधिकार सुरक्षित है।

महानिदेशक
केंद्रीय आयुर्वेदीय विज्ञान अनुसंधान परिषद्

CENTRAL COUNCIL FOR RESEARCH IN AYURVEDIC SCIENCES

Expression of Interest (EoI) for manufacturing and supply of Ayurvedic drugs for Research purpose in CCRAS

Background: The Central Council for Research in Ayurvedic sciences (CCRAS), Ministry of AYUSH, Government of India, is an apex body in India for undertaking, coordinating, formulating, developing and promoting research on scientific lines in Ayurvedic sciences. The activities are carried out through 30 Institutes/Centres/Units located all over India and also through collaborative studies with various Universities, Hospitals and Institutes. The research activities of the Council include Medicinal Plant Research (Medico-ethno botanical Survey, Pharmacognosy and Tissue Culture), Drug Standardization, Pharmacological Research, Clinical Research, Literary Research & Documentation, Reproductive & Child Health Care Programme and Tribal Health Care Research Programme.

Purpose: The council is focusing on validation of classical Ayurvedic formulations and also developing new drugs based on leads from Ayurvedic literature or claims. For clinical studies, quality and standardized medicines are required to be utilized which may be pharmacopoeial or non-pharmacopoeial. Council invites Expression of Interest (EoI) for manufacturing and supply of Ayurvedic drugs for Research purpose in CCRAS from GMP certified Ayurvedic pharmaceutical industries in India with following benchmark, terms and conditions.

The interested firms fulfilling the same may offer their consent for this purpose.

Terms and conditions:

Essential:

1. The manufacturer/Pharmaceutical Company should have valid GMP certificate for maximum varieties of dosage forms and fulfil other requirements as per Schedule T of D&C Act.
2. It should have an in house Quality Control Section and R&D facility and requisite expertise.
3. It should be having at least 10 years of experience in manufacturing and marketing of Ayurvedic drugs.
4. It should be able to prepare drugs as per SOP provided by the Council for manufacturing of classical formulations and Coded /Proprietary drugs developed by the Council. No manufacturing should be outsourced.
5. The pharmaceutical company must sign a Non Disclosure Agreement to maintain the confidentiality of New/coded drugs developed by the council through R&D and shall not claim any Intellectual Property Rights at any stage.

6. The manufacturer should comply with quality standards of raw materials, intermediate and finished products (viz. Pharmacognostic, Physico-chemical and Safety parameters) as per format provided by the Council/or able to develop in-house standard where ever required. The cost of these tests with codal formalities will be borne by the Council. If few parameters of Q.C. & standardization is outsourced the same should be from NABL accredited/Govt. laboratory.
7. It should also comply with the requirement of packing and labelling specifications provided by the Council.
8. It should have in house capability to produce different types of plant extracts as required for the manufacturing of Coded/Proprietary drugs developed by the Council or should be able to procure them from reputed Companies having capacity to produce export quality plant extracts along with SOPs of manufacture and COAs as will be specified by the Council.
9. It should be able to supply the drugs in a time bound manner within a maximum period of three months/or as specified from time to time in the supply order.
10. It should be able to supply the drugs directly to specified destinations through transportation.
11. The manufacturer should allow CCRAS team to oversee the manufacturing procedure as and when required.
12. The manufacturer has to provide samples of each raw drug and the finished product of required quantity as will be specified by the council.

Desirable:

1. WHO GMP compliance of the manufacturing unit.
2. Prior experience in manufacturing of Ayurvedic drugs for research purpose.
3. Experience of manufacturing export quality Ayurvedic products and exporting abroad.
4. They should be able to provide lab scale samples of Ayurvedic drugs as and when required, along with standards and COAs.
5. Inclination towards Research and active mutual interaction with CCRAS as and when required in the process of preparation and supply.

How to apply : The details, consent alongwith requisite documents substantiating the terms and conditions may be furnished to the following address by **25th May 2015** for further appraisal.

Director General
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