**Recruitment Announcement**

**Global Pharma Key Opinion Leaders Program**

Korea Health Industry Development Institute (KHIDI) is a government-affiliated institution which provides professional and systematic support to develop domestic health industry and enhance health services since its establishment in 1999. As the nation’s only one organization in responsible for fostering health industry, KHIDI has been playing a major role in the industry development in fields of health services, pharmaceuticals, medical devices and beauty-cosmetic.

KHIDI is looking for “Global Pharma Key Opinion Leaders (GPKOL)” who can provide on-line consulting services to Korean pharmaceutical companies for new drug development and global commercialization.

**1. Job Description**

* Roles

1. Provide **online consulting** for Korean companies through e-mail, phone calls, etc
2. Participate in **GPKOL** **symposium** in South Korea and **one-on-one business meetings** with Korean companies

* Give presentations on topics related to new drug development, overseas clinical trial, approval, and global marketing and have open discussion (Q&A) with Korean companies
* Shall be invited to Korea at least once a year for GPKOL symposium (for less than a 7-day stay).

1. Publisha **manuscript (Trends paper)** that focuses on important pharmaceutical and related areas as well as the latest industry trends in Bio-Pharm (publish A4 10~15 sheets or less)

* The pharmaceutical Industry Information Portal, the trends paper and manuscript are being carried out (http://www.epharmakorea.or.kr)) in conjunction with the online consulting service
* Consulting Fields

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| **Fields** | **Related Sectors** |
| R&D  Planning | * Discovery of Drug Candidate, Preclinical, New Drug Value Evaluation, Data Output Analysis * Phased Approval, Clinical, Linkage between R&D and Marketing |
| Clinical Trial | * Preclinical and Clinical Trial (Data Analysis, Protocol Design, etc.) |
| GMP | * GMP audit (inspector) (EMEA, FDA, ICH Guideline, etc.) * Software validation and documentation for cGMP * Consultation on a new construction of pharmaceutical plants |
| Regulatory Affairs  (RA) | * + Approval and pricing of new drug, IMD, biosimilars and generics at multinational pharmaceutical company or FDA   + Consultation in WHO PQ process (ex. Work experience in USP) |
| Technical Marketing | * + To provide training and consulting on overseas expansion strategy and marketing channel development in such case of Korean companies’ market expansion into emerging countries   + To provide training and consulting on patent acquisition, marketing prediction, strategic alliance   + To establish partner network for each market and region   + To provide training and consulting on the provision of diverse commercialization models including financing |
| Project Management  (PM) | * + To provide training and consulting on the full cycle of new drug development from R&D planning, clinical trial and production (GMP) to regulatory affairs (RA) and technical marketing (BD) |

* Compensation

1. **Compensation for Customized Online Consulting**

Consultancy fee of KRW 500,000 (about USD 500) or so shall be paid per consulting

* Consulting report shall be submitted to get paid

1. **Compensation and Reimbursement for GPKOL Symposium**

Round-Trip Airfare, Staying Expenses, Instruction Fee

* Staying expenses shall be reimbursed in accordance with price standard of KHIDI.

1. **Compensation for Manuscript (Article)**

Manuscript fee of KRW 500,000(about USD 500) or so shall be paid per manuscript

* The payment of allowance regarding online consulting, manuscript etc. has to comply in accordance with KHIDI "GPKOL’s directions”.
* Qualification

At least **5 years working experience in new drug development, clinical trial, approval and global marketing** at multinational pharmaceutical companies or organizations such as FDA, NIH (US) or EMA (Europe) or NMPA(China)

* Applicants with Ph.D. degree in the applicable fields and have experience with RA institution preferred

**2. Selection Process**

Application period (June.7th ~ June.24th) → Qualifications review (June.30th) → Result announcement (July. 13th~)

* Selection schedule could be changed. Applicants shall be notified individually of the result

**3. Required Documents**

1. GPKOL Application form 1-4
2. A copy of CV
3. A copy of Certificate of the Highest level of Education
4. A copy of proof documents of present or past employment
5. A copy of documents of major achievements such as publication or accomplishments in R&D, technical transfer, approval and patent, if applicable
6. A copy of award in relevant field, if applicable

**4. How to Apply**

* The complete application package must be submitted by E-mail Only
* Application deadline: 11:59 PM (Korea Local time), June 24th, 2022
* Submitted documents shall not be returned and not be used for other purposes than recruiting process. These documents are protected under the Personal Information Protection Law.
* Inquiries
* Pharma & Bio-Pharma Global Team, KHIDI

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