|  |  |
| --- | --- |
| C:\Users\sn\OneDrive - IEC\coordination\template\iec_logo_100px.jpg**INTERNATIONAL****ELECTROTECHNICAL****COMMISSION** |  |

**Application form for international organizations wishing to create a liaison with an IEC technical committee**

**Technical Committee or Subcommittee concerned (TC number and title)**

**62:** **Medical equipment, software and systems**

**---------------**

**Applicant Organization:**

ITU/WHO FG-AI4H

**Liaison category (see the** [**ISO/IEC Directives**](https://www.iec.ch/members_experts/refdocs/) **Clause 1.17.2 for details):**

A (at the Committee level)

B (at the Committee level)

C (at the Working Group level)

**Background**

ITU/WHO Focus Group on AI for Health (FG-AI4H) is working towards international standards for trustworthy health AI technologies, with the important goal to develop an open, scalable benchmarking framework for the standardized quality assessment of these

See attachement

****

Detailed information concerning the general requirements applicable to liaisons, different categories of liaisons, eligibility criteria, rights and obligations of liaison organizations is contained in the [ISO/IEC Directives, Part 1 and IEC Supplement, Clause 1.17](https://www.iso.org/sites/directives/current/part1/index.xhtml#_idTextAnchor093).

## To be completed by the applicant organization

## Eligibility criteria

|  |
| --- |
| The applicant confirms that the organization: is not for profit is a legal entity (ITU is like ISO or IEC)(NOTE: this is a requirement only for Category A or B liaisons) is membership-based and open to members worldwide or over a broad region(NOTE: this is a requirement only for Category A or B liaisons) through its activities and membership demonstrates that it has the competence and expertise to contribute to the development of International Standards or the authority to promote their implementation has a process for stakeholder engagement and consensus decision-making to develop the input it provides |

|  |
| --- |
| ITU covers areas for AI at the international standardization level, which are important for the medical device community (e.g. conformity process, auditing) but yet not covered for AI by ISO and IEC. There are already close connections at the expert level which should be supported and intensified by the liaison. |

**Justification for liaison request**

|  |  |  |  |
| --- | --- | --- | --- |
| **Contact details of liaison representative(s)** | | | |
| **Title** | **First name** | **Last name** | **E-mail** |
| **Prof.** | **Thomas** | **Wiegand** | **thomas.wiegand@hhi.fraunhofer.de** |
|  |  |  |  |

|  |  |
| --- | --- |
| I accept the responsibilities and obligations of liaison organizations, as outlined in the [ISO/IEC Directives, Part 1 and IEC Supplement, Clause 1.17](https://www.iso.org/sites/directives/current/part1/index.xhtml#_idTextAnchor093), including the [IEC Copyright Policy and Implementation Guidelines](https://www.iec.ch/members_experts/refdocs/policy/IEC_copyright_policy_implementation_guidelines_2020.pdf). | |
| Name | Date |