



Medicines & Healthcare products  
Regulatory Agency

## Professional Posts available for the In Vitro Diagnostics Expert Advisory Group

### Advert

The Interim Devices Working Group (IDWG) is looking to appoint professional members to the In Vitro Diagnostics Expert Advisory Group (IVD EAG).

The IVD EAG is a subgroup of the IDWG that provides independent expert advice to the Medicines and Healthcare products Regulatory Agency (MHRA) on the development, implementation, and evaluation of IVD devices. The group's objective is to ensure that IVD regulations, guidance and regulatory practices support innovation, maintain high standards of safety and efficacy, and align with international and domestic leading practices.

IVD EAG is formed of experts in the field of in vitro diagnostics, including scientists, engineers, clinicians, pathologists, and diagnostics regulatory specialists. Representatives from relevant government departments and agencies, e.g. NICE, Devolved Administrations, NIHR, UK HSA, etc. Non-industry stakeholders, including representatives from academia and learned organisations such as the Royal Colleges and Biomedical faculties and patient or patient group representatives to ensure patient perspectives are considered.

The IVD EAG provides advice and recommendations to IDWG and MHRA on:

- The development and implementation of IVDs, including policies, regulations, and guidance.
- The research, development, and evaluation of certain IVDs, including their clinical evidence and performance.
- Applications for exceptional use authorisation of IVDs.
- Applications for Coronavirus test device approvals.
- The stakeholders, including professional organisations and patient groups, to gather diverse perspectives and insights.
- Emerging trends and advancements in IVD technologies to ensure the regulatory frameworks remain current with technological advancements.
- Matters arising from scientific, clinical, or regulatory evidence uncertainty and where leading practice guidance may be required.

IVD EAG also input into MHRA statements, advice, and recommendations.

IVD EAG members are not employed by MHRA. IVD EAG members are independent from MHRA and provide their time on a voluntary basis.

**The closing date for applications is 6 February 2026.**

# Role and responsibilities of the Professional Members of the IDWG's In Vitro Diagnostics Expert Advisory Group

As a member you will:

- Take full account of the evidence in making decisions and consider the analysis and interpretation of the evidence prepared by MHRA.
- Consider equality, diversity and inclusion in all of the EAG discussions.
- Have a flexible attitude, ability and enthusiasm to work as a team member
- Assessing benefit and risk in relation to evidence presented in order to contribute expertise to generate effective actions to mitigate risk
- Read papers in advance of meetings
- Occasionally comment on documents in between meetings by email
- Keep the work of the EAG confidential Declare conflicts of interest and adhere to the Code of Practice ([link](#))

## Essential Criteria

**The Member must be able to demonstrate the qualities, skills and experience to meet all the essential criteria for appointment.**

- Experience in IVD field;
- Current or previous experience of committee membership(s);
- Be recognised by their peers as a leader in their field e.g. national academies and Royal Colleges (for specialist posts only);
- Be able to assimilate and interpret complex scientific information and formulate evidence-based comments /advice at short notice;
- Be able and prepared to contribute actively to the work of the EAG, including on issues outside of own specialism;
- Be a skilled communicator;
- Be willing to develop a working knowledge and understanding of UK MDR and EU IVDR regulatory framework and procedures;
- Maintain strict confidentiality with respect to the work of the EAG;
- Be willing to declare conflicts of interest;
- Be committed to the values of selflessness, integrity, objectivity, accountability, professionalism, impartiality and consistency.

## Remuneration

- You may claim travel and subsistence expenses, which are properly and necessarily incurred in carrying out your role and responsibilities as Member of the IVD EAG, in line with travel and subsistence policy and rates for the MHRA. A copy of the policy and rates is available from the MHRA.

## Time Commitment

- 4 meetings per year, held virtually/in person.

## Tenure of office

- 3 years

## How to Apply

If you are interested in joining the **IVDEAG**, please email your application form, CV, completed declaration of interests form and monitoring form to [CSTrecruitment@mhra.gov.uk](mailto:CSTrecruitment@mhra.gov.uk), quoting the position and reference **IVDEAG25-2a** in the subject field. Please ensure that you provide evidence to support how you meet all of the essential criteria.

If you require an alternative format, please call the Appointments Team on 020 3080 6060 quoting the reference **IVDEAG25-2a**.

**The closing date for applications is 6 February 2026. The MHRA value and promote diversity and encourage applications from all sections of the community.**



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