

the group aims to harness the benefits of Artificial Intelligence (AI) in incology and haematology pharmacy while ensuring its safe and iffective integration into clinical practice.  Deficives:  Eviewing Current Literature:  Examine evidence and emerging trends in AI applications relevant to incology and haematology pharmacy.  Evaluating AI Technologies:  Essess potential benefits, risks, and challenges associated with AI doption in pharmacy practice.  Eveloping Guidelines:  Formulate best practices and recommendations for ethical and esponsible AI use in pharmacy, focusing on patient safety and data
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dvising Regulatory Bodies:
rovide guidance to regulatory bodies, professional organizations, and ealthcare institutions on AI implementation policies and standards.
acilitating Education:
romote education, training, and awareness to enhance understanding nd proficiency in AI applications among pharmacists and stakeholders.
he Specialist Advisory Group comprises multidisciplinary experts, acluding but not limited to:
Clinical pharmacists specializing in oncology and haematology. Pharmacy informaticists with AI experience. Oncologists and haematologists interested in AI applications. Data scientists specializing in healthcare analytics and machine earning. Ethicists and legal experts on AI ethics, privacy, and regulatory ompliance. Representatives from regulatory agencies, professional ssociations, and industry stakeholders.
ctive Participation:



- Engage in meetings, discussions, and collaborative activities to achieve objectives.
- Attend at least 50% of meetings annually and actively contribute to the group's progress.

### **Expert Recommendations:**

- Provide subject matter expertise and insights based on the latest evidence-based practices and ethical principles.
- Offer recommendations for integrating AI into clinical practice, ensuring patient safety and data security.

## **Collaborative Development:**

- Work with stakeholders to develop comprehensive guidelines, best practices, and tools for AI integration in oncology and haematology pharmacy.
- Create educational resources and organize training sessions to support the professional development of pharmacists and healthcare providers in AI applications.

#### **Review of Al Technologies:**

- Critically evaluate emerging AI technologies, algorithms, and applications relevant to pharmacy practice.
- Identify opportunities for innovation as well as potential risks and challenges, ensuring that AI tools are safe, effective, and ethically sound.

#### **Communication of Findings:**

- Disseminate findings, recommendations, and best practices through various channels such as publications, presentations, and educational materials.
- Communicate regularly with relevant stakeholders, including pharmacists, healthcare providers, policymakers, and the public, to keep them informed about the group's activities and advancements in AI technology.

#### **Policy and Standards Development:**

- Advise regulatory bodies, professional organizations, and healthcare institutions on the development of policies and standards related to AI implementation in pharmacy practice.
- Ensure that these policies and standards promote the ethical and responsible use of AI technologies, protecting patient privacy and data security.



Facilitating Research and Innovation:						
	<ul> <li>Promote and support research initiatives that explore the application of AI in oncology and haematology pharmacy.</li> <li>Encourage collaboration between pharmacists, data scientists, and other healthcare professionals to foster innovation and improve patient outcomes through AI.</li> </ul>					
	Monitoring and Evaluation:					
	<ul> <li>Continuously monitor the impact of AI technologies on clinical practice and patient outcomes.</li> <li>Evaluate the effectiveness of implemented AI solutions and provide feedback for improvement.</li> </ul>					
	Ethical Oversight:					
	<ul> <li>Ensure that the integration of AI in oncology and haematology pharmacy adheres to high ethical standards.</li> <li>Address ethical concerns related to AI, such as bias, transparency, and accountability, and develop strategies to mitigate these issues.</li> </ul>					
Accountability	☐ The Chairperson ensures meetings are properly documented, with minutes presented to the BOPA Executive Committee.					
	☐ Members must attend at least 50% of meetings annually and actively					
	contribute to the group's progress.  ☐ The group reports to and is accountable to the BOPA Executive Committee, which reviews and approves the work plan annually.					
Frequency of Meetings:	<ul> <li>□ Meetings held regularly, with one face-to-face meeting per year.</li> <li>□ Teleconference and email discussions for interim communication.</li> <li>□ Formal decisions ratified and documented in meetings.</li> </ul>					
Quorum:	A minimum requirement for quorate to be achieved is attendance by 75% of core members.					
Ownership of Group Projects and Initiatives	All projects, initiatives and outcomes will be owned by BOPA.					
Reimbursement and financial management	BOPA will not re-imburse Group members for their time but will pay travel expenses and venue hire for the annual face to face meeting as appropriate. If additional face to face meeting are planned the executive committee must agree if travel expenses can be funded.  The group will prepare an annual estimated budget covering proposed projects, travel and other expenses for submission to treasurer to enter into annual financial prioritisation process for approval by Exec Committee.  The group is not profit generating in line with the BOPA and its charitable status.					
Communication Arrangements:	<ul> <li>☐ Minutes forwarded within three weeks, agendas and minutes posted online.</li> <li>☐ Agenda items submitted seven days before meetings.</li> <li>☐ Inter-meeting communication via email by the Chairperson or Secretary.</li> </ul>					



Declaration of	All potential or perceived conflicts of interest should be declared.
Interest:	

## **Document Control**

Document Title:	BOPA AI Specialist Advisory Group TERMS OF REFERENCE				
Author:			Current Version:		
Approved by:			Date Approved:		
Due for Review:					
Summary of Changes:		New Document			