



Restructuring the ATC L-Class: Antineoplastics and Immunomodulating Agents

Enhancing the classification structure for
improved categorization



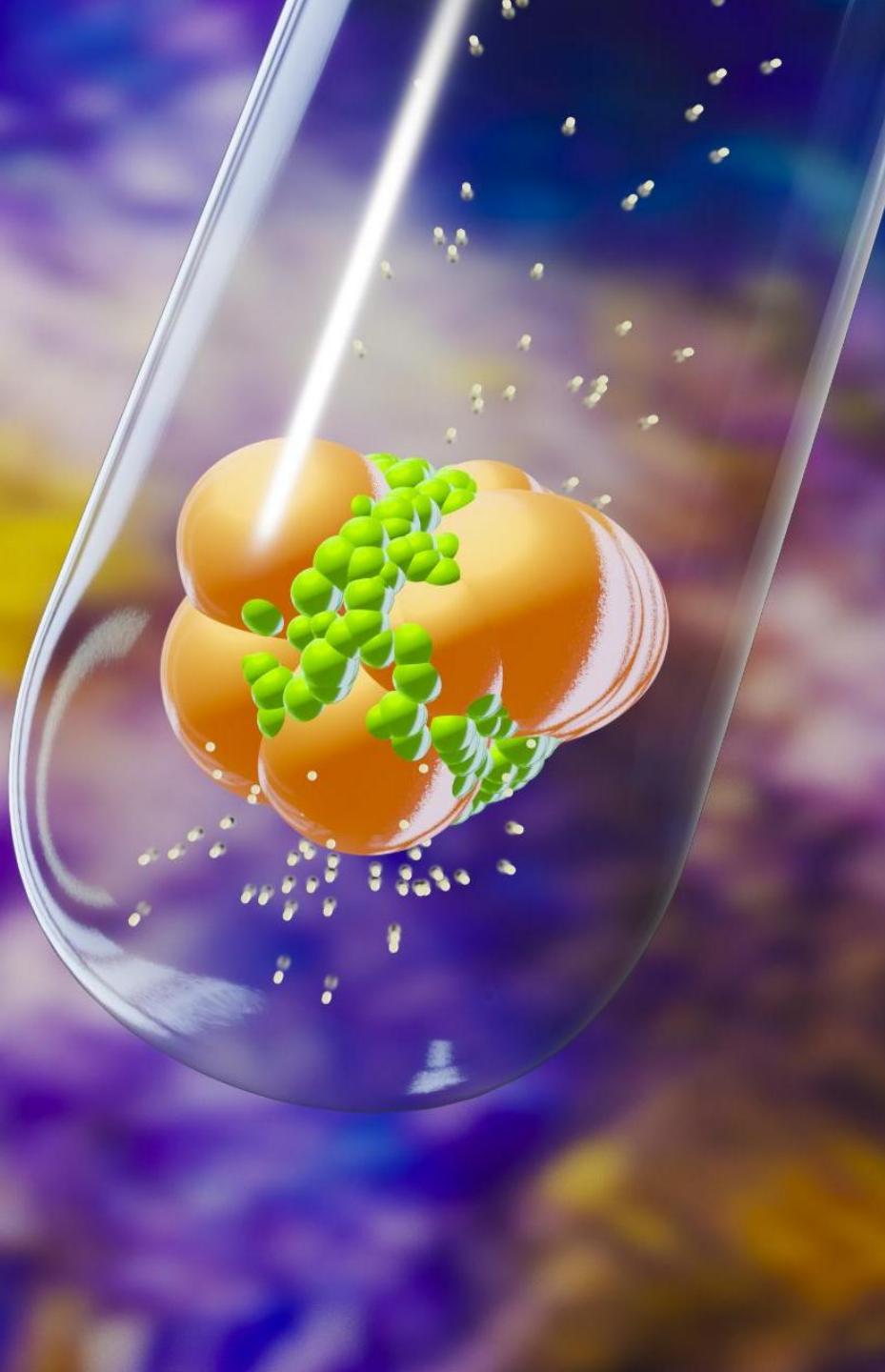
Today's Webinar Agenda

- Challenges in the Current ATC L-Class
- Proposed Changes
- Revised Classification Structure in Detail
- Benefits of Restructuring
- Impact of Revised Classification Structure
- Stakeholder Engagement and Support
- Implementation Timeline
- Q&A





Challenges in the Current ATC L-Class



Limitations of Combining Antineoplastic and Immunomodulating Agents

Combined Drug Classification

- The ATC L-Class currently groups antineoplastic and immunomodulating agents together

Challenges with Emerging Therapies

- New therapies have necessitated the introduction of new classes
- This has resulted in space issues in the current classification structure

Current L-Class Structure

L: ANTINEOPLASTIC AND IMMUNOMODULATING AGENTS

L1: ANTINEOPLASTICS		L2: CYTOSTATIC HORMONE THERAPY	L3: IMMUNOSTIMULATING AGENTS	L4: IMMUNOSUPPRESSANTS
L1A: ALKYLATING AGENTS	L1H: PROTEIN KINASE INHIBITOR ANTINEOPLASTICS	L2A: CYTOSTATIC HORMONES	L3A: IMMUNOSTIMULATING AGENTS EXCLUDING INTERFERONS	L4A: ANTI-TNF PRODUCTS
L1B: ANTIMETABOLITES	L1J: PROTEASOME INHIBITOR ANTINEOPLASTICS	L2B: CYTOSTATIC HORMONE ANTAGONISTS	L3B: INTERFERONS	L4B: INTERLEUKIN INHIBITORS
L1C: PLANT-BASED ANTINEOPLASTICS	L1K: LIDOMIDE ANTINEOPLASTICS			L4C: JAK INHIBITORS
L1D: ANTINEOPLASTIC ANTIBIOTICS	L1L: PARP INHIBITOR ANTINEOPLASTICS			L4D: OTHER IMMUNOSUPPRESSANTS
L1F: PLATINUM ANTINEOPLASTICS	L1X: ALL OTHER ANTINEOPLASTICS			
L1G: MONOCLONAL ANTIBODY ANTINEOPLASTICS				

Current Protein Kinase Inhibitor Structure Has Little Space for Future Growth

L	ANTINEOPLASTIC AND IMMUNOMODULATING AGENTS
L1	ANTINEOPLASTICS
L1H	PROTEIN KINASE INHIBITOR ANTINEOPLASTICS
L1H1	Protein kinase inhibitor antineoplastics, BCR-ABL
L1H2	Protein kinase inhibitor antineoplastics, EGFR
L1H3	Protein kinase inhibitor antineoplastics, ALK
L1H4	Protein kinase inhibitor antineoplastics, BRAF/MEK
L1H5	Protein kinase inhibitor antineoplastics, CDK 4/6
L1H6	Protein kinase inhibitor antineoplastics, BTK
L1H7	Protein kinase inhibitor antineoplastics, JAK
L1H9	Protein kinase inhibitor antineoplastics, other



Proposed Changes to the ATC L-Class Structure

Key Changes



New ATC1 for Immunological Products

- A dedicated 1ST level ATC established exclusively for immunological products

Promotion of antineoplastic classes from 3rd level to 2nd level

- This will future-proof the classification, allowing for innovation and emerging therapies

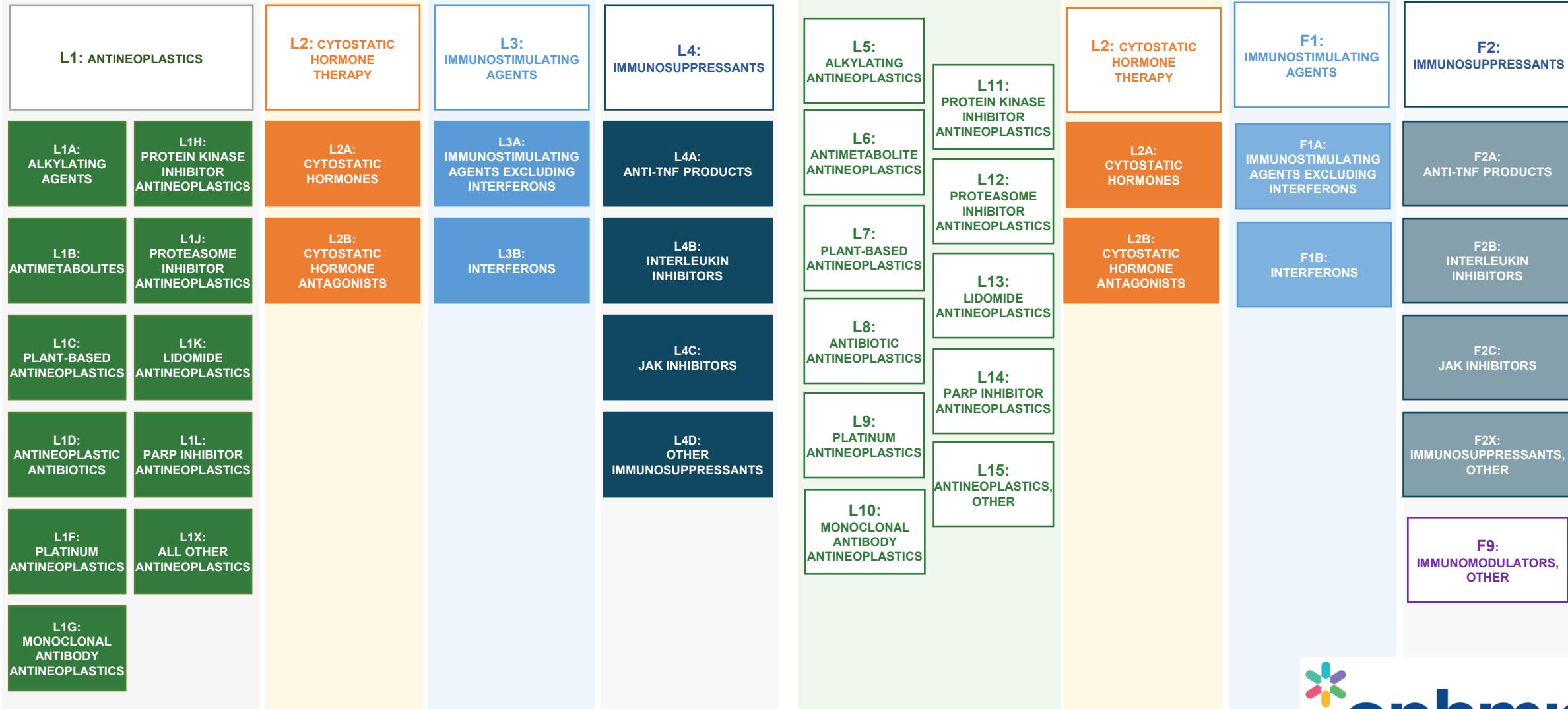
CURRENT

FUTURE

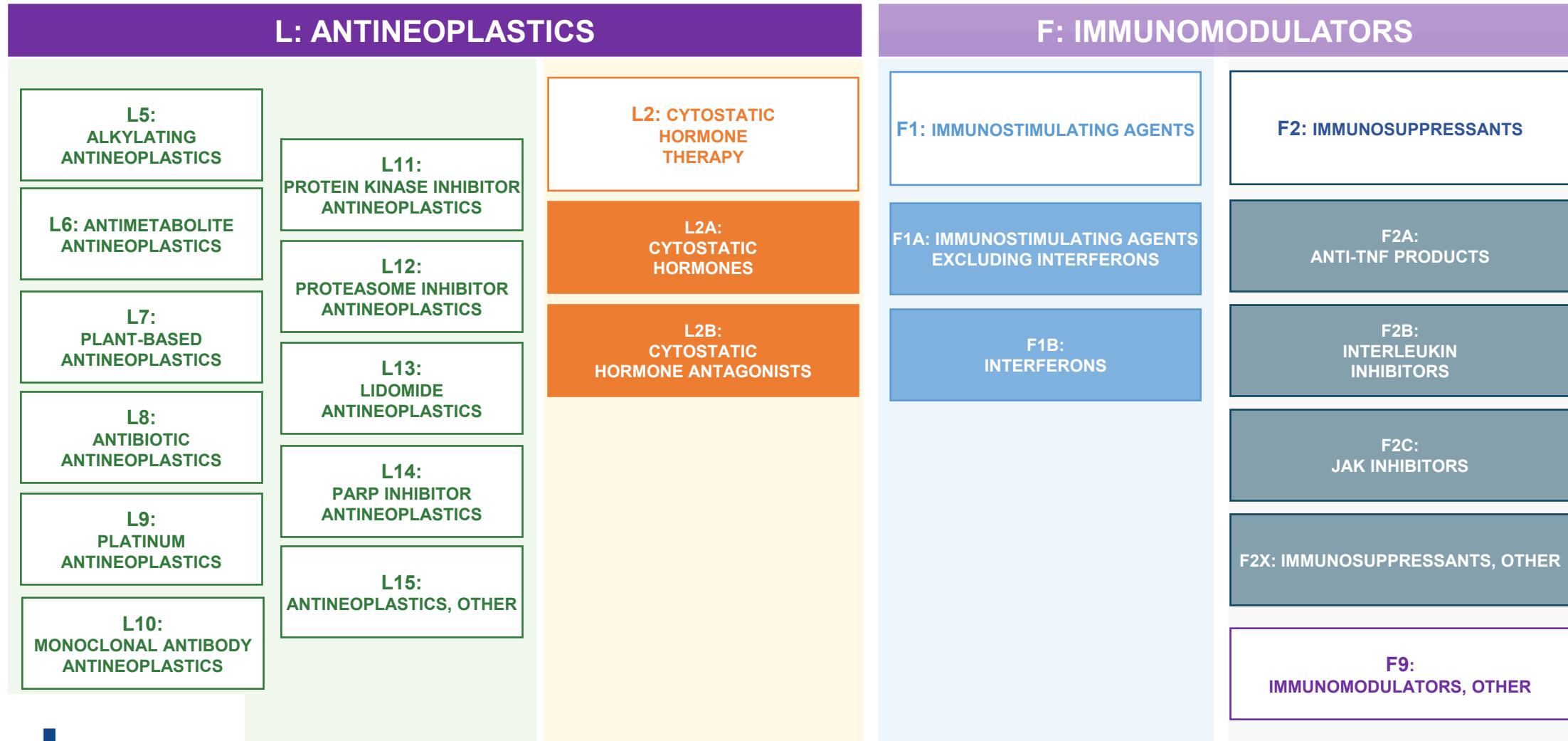
L: ANTOINEOPLASTIC AND IMMUNOMODULATING AGENTS

L: ANTOINEOPLASTICS

F: IMMUNOMODULATORS



Proposed L-Class and F-Class Structure



Proposed Protein Kinase Inhibitor Structure Allows for Approval of Novel Targeted Agents

L	ANTINEOPLASTICS
L11	PROTEIN KINASE INHIBITOR ANTINEOPLASTICS
L11A	Protein kinase inhibitor antineoplastics, BCR-ABL
L11B	Protein kinase inhibitor antineoplastics, EGFR
L11C	Protein kinase inhibitor antineoplastics, ALK
L11D	Protein kinase inhibitor antineoplastics, BRAF/MEK
L11E	Protein kinase inhibitor antineoplastics, CDK 4/6
L11F	Protein kinase inhibitor antineoplastics, BTK
L11G	Protein kinase inhibitor antineoplastics, JAK
L11X	Protein kinase inhibitor antineoplastics, other



Benefits of the ATC L-Class Restructuring

Future-Proofing the Classification System



Improved Clarity

- By giving oncology and immunology their own first level classes

Adaptability to Future

- By creating space so the classification can continue to adapt to future developments for many years to come



Impacts of the ATC L-Class Restructuring

Impacts of the Revised Classification Structure

Sales Ranking Changes

Based on IQVIA Q2/2025 data

Current ATC1	% Share
L: Antineoplastics and Immunomodulating Agents	26%
A: Alimentary Tract and Metabolism	19%
N: Nervous System	10%
J: General Systemic Antiinfectives	9%

Future ATC1	% Share
A: Alimentary Tract and Metabolism	19%
L: Antineoplastics	15%
N: Nervous System	10%
F: Immunomodulators	10%

Practical Considerations

- Internal systems, processes and analyses will need to be updated to accommodate this change



Stakeholder Engagement and Implementation Timeline

Stakeholder Feedback and Available Resources

Encouraging Feedback

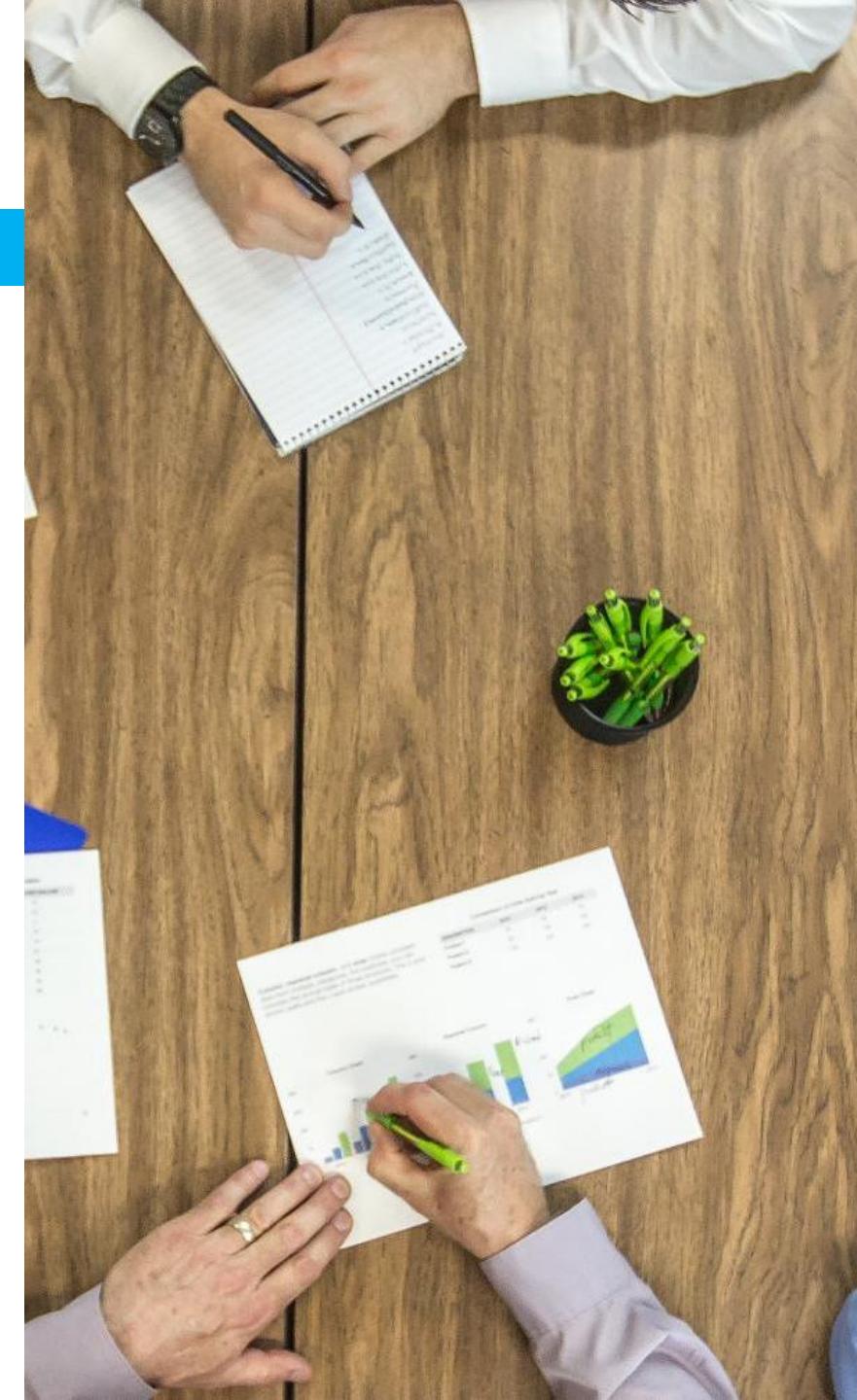
- Organizations, especially oncology teams, are encouraged to review proposals and provide meaningful feedback

Resources

- Webinar
- News articles and LinkedIn posts
- Available on the EPHMRA website

Any questions or feedback, please contact:

Bernadette Rogers (generalmanager@ephmra.org)



Key Dates for Voting and Implementation

Voting Schedule

- A vote by EPHMRA members to approve these changes is scheduled for May/June of 2026

Implementation Date

- Upon approval, the new structure will be implemented starting January 2027 when all products will be moved to their new classifications





Q&A



CURRENT

FUTURE

L: ANTOINEOPLASTIC AND IMMUNOMODULATING AGENTS

L: ANTOINEOPLASTICS

F: IMMUNOMODULATORS

