

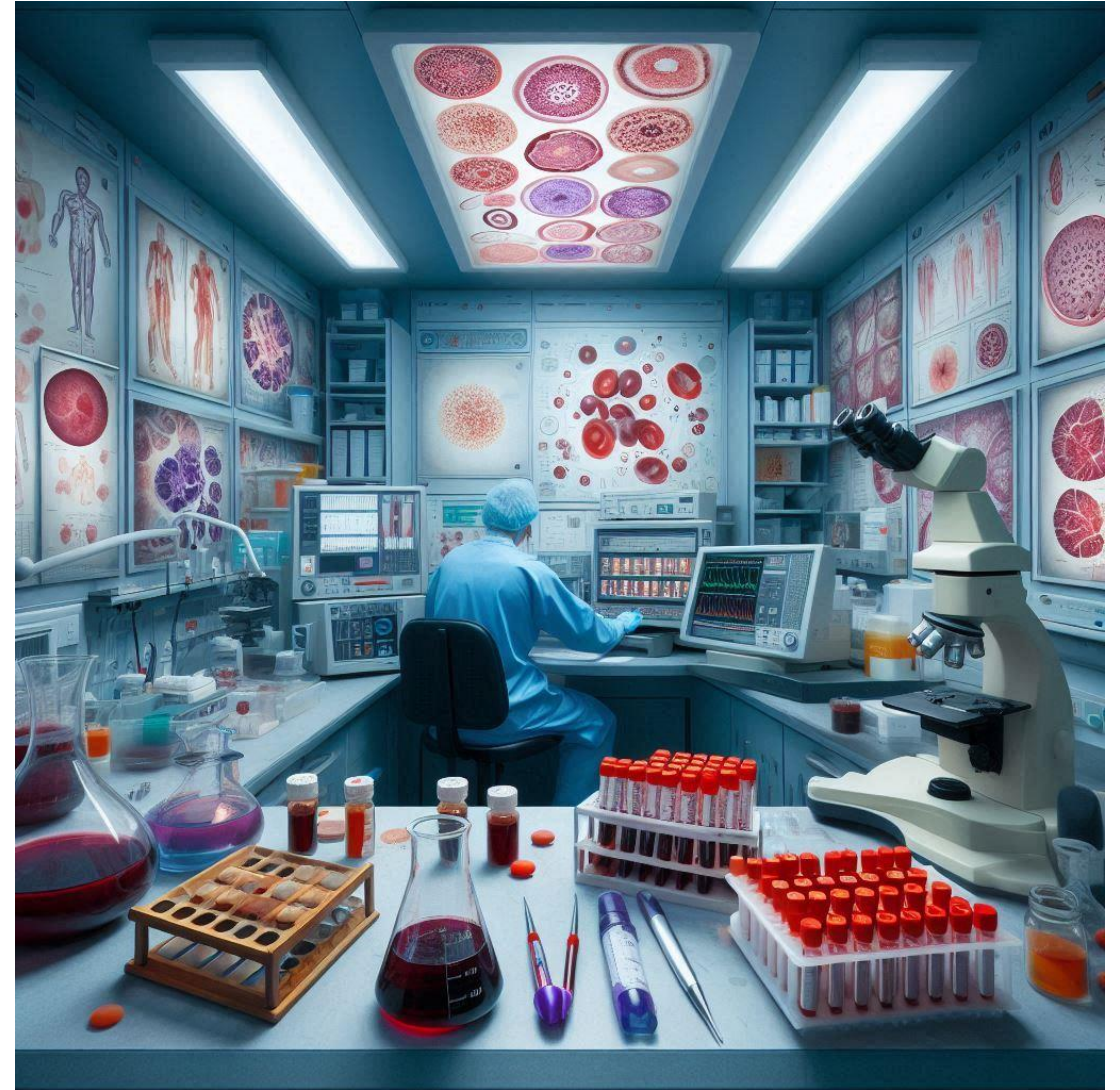
# Digital Haematology Task Force

Advancing Digital Pathology  
in Haematology within the UK

**Dr Timothy Farren**

On behalf of the BSH Lab SIG

and Dr Guy Hannah Co-Chair of DHTF



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- 1590 – Zacharias Jassen
- 1609 – Galileo Galilei
- 1625 – “Microscope” Faber
- 1665 – “Cells”
- 1667 – Robert Hooke
- 1676 - Leeuwenhoek



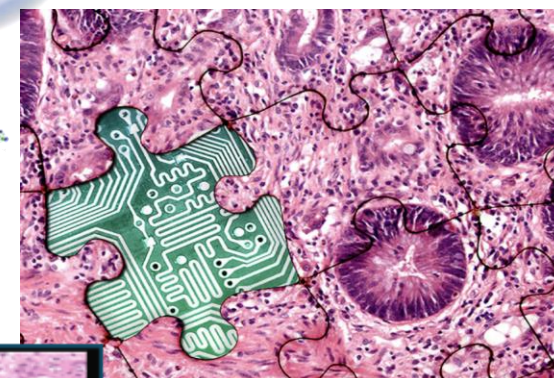
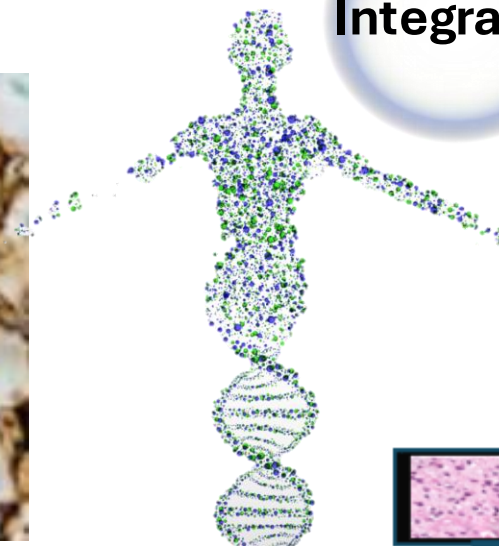
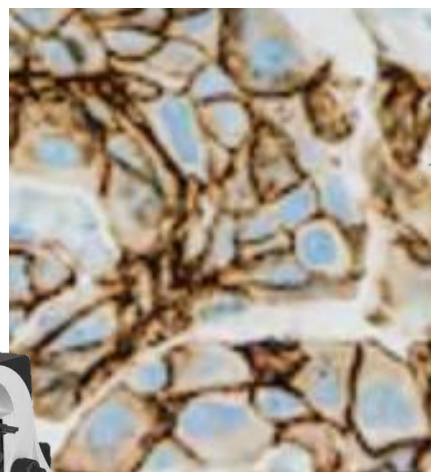
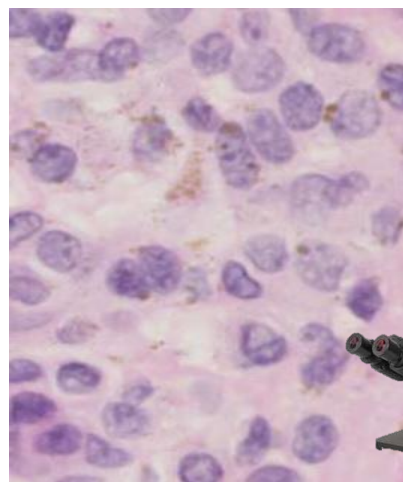


# Why Digital?

**LOW VALUE**  
PROCESS DRIVEN

**HIGH VALUE**

DATA DRIVEN  
PRECISION MEDICINE



Macroscopic  
1700-now

Microscopic  
1850-now

Antibodies/EM  
1970 - now

Molecular  
2000-now

Digital  
2018- now

**First Revolution**  
(Dissection/Microscope)

**Second Revolution**  
(Antibodies/Electron Microscope)

**Third revolution** (Computational)

- The microscope remains a haematologists best friend.

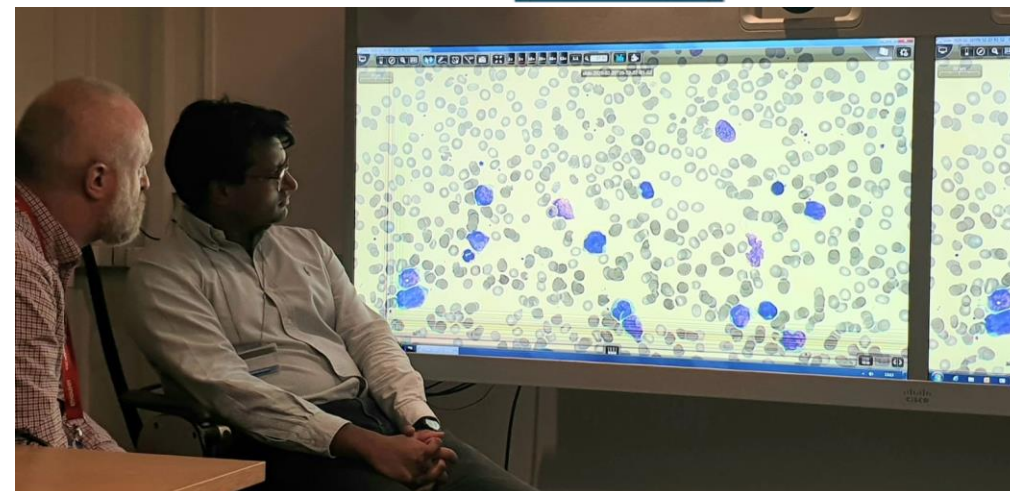
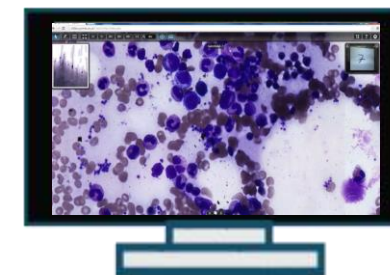
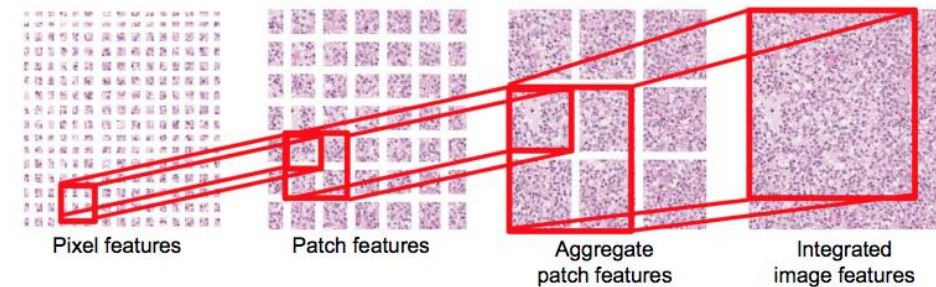
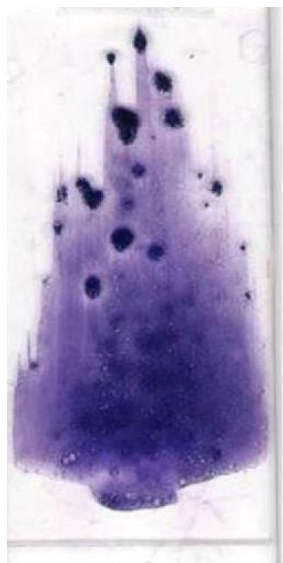
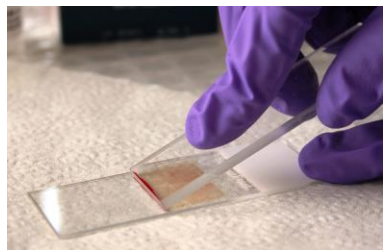


- Limitations:
  - Consistency / reproducibility
  - Specialist referral / multisite
  - Training / Education
  - Capacity and efficiency

# Current situation








**Human Tissue**  
(Microscope based interpretation)

**Data**  
(Available for Analytics)

- Already “embedded” by our histopathology colleagues.
- Remote reporting / telepathology
- Collaboration
- Standardisation of image analysis
- AI integration
- Training tools
  - Case libraries
  - Digital teaching sets
- Scalable
- Long Term Plan

# Why Digital Haematology?

 The Royal College of Pathologists  
Pathology: the science behind the cure

**Best practice recommendations for implementing digital pathology**  
January 2018



Authors: Simon Cross, Peter Furness, Laszlo Igali, David Snead, Darren Treanor

Unique document number	G162
Document name	Best practice recommendations for implementing digital pathology
Version number	1
Produced by	Simon Cross, Peter Furness, Laszlo Igali, David Snead and Darren Treanor on behalf of the Specialty Advisory Committee on Cellular Pathology.
Date active	January 2018
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Comments	In accordance with the College's pre-publication policy, this document was on The Royal College of Pathologists' website for consultation from 12 May 2017 to 12 June 2017. Responses and authors' comments are available to view on request.  <b>Dr Lorna Williamson</b> Director of Publishing and Engagement

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<https://www.rcpath.org/static/f465d1b3-797b-4297-b7fedc00b4d77e51/Best-practice-recommendations-for-implementing-digital-pathology.pdf>

 The Royal College of Pathologists  
Pathology: the science behind the cure

**Digital pathology strategy 2019**

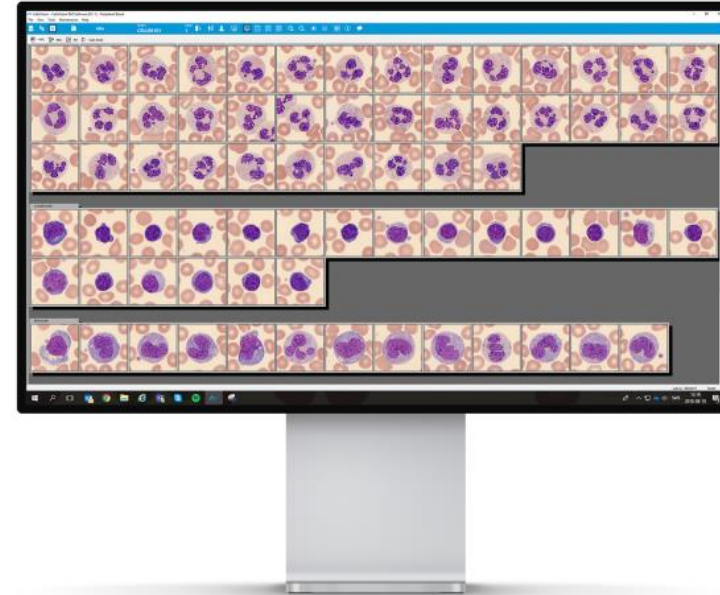


<https://www.rcpath.org/static/2248bb71-b773-4693-945bffda593f2f2f/cf251e84-f7d0-415d-bb67217219203066/Digital-Pathology-Strategy.pdf>



**Will it work?**

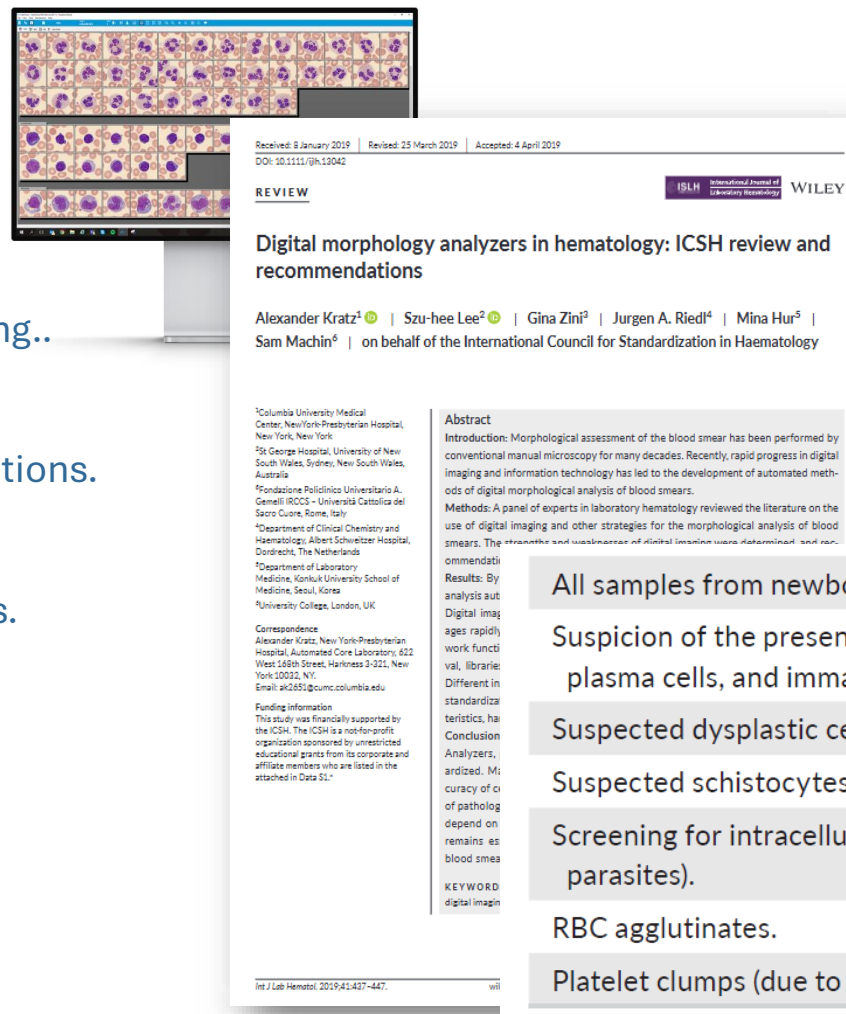
- Already works in Clinical Setting..





# Will it work?

- Already works in Clinical Setting..
- Increasing “Research” publications.
- Limited “Clinical” publications.
  - DI60
  - Cellavision
  - Vision Hema
  - EasyCell
  - NextSlide
  - HemaCAM
- Peripheral Blood focus.



All samples from newborns and from patients with leukemia.

Suspicion of the presence of pathological cell types, including blasts, plasma cells, and immature granulocytes.

Suspected dysplastic cells.

Suspected schistocytes.

Screening for intracellular parasites (eg, Malaria, Babesia, other parasites).

RBC agglutinates.

Platelet clumps (due to location in feather and lateral edges).

Kratz A, Lee S-H, Zini G, Riedl JA, Hur M, Machin S; on behalf of the International Council for Standardization in Haematology. Digital morphology analyzers in hematology: ICSH review and recommendations. *Int J Lab Hematol*. 2019; 41: 437–447. <https://doi.org/10.1111/ijlh.13042>





Received: 26 February 2022 | Accepted: 19 May 2022  
DOI: 10.1111/ijlh.13908

## REVIEW

ISLM International Journal of Laboratory Hematology WILEY

# Digital morphology in hematology diagnosis and education: The experience of the European LeukemiaNet WP10

Gina Zini<sup>1,2</sup> | Ombretta Barbagallo<sup>2</sup> | Fernando Scavone<sup>2</sup> | Marie C. Béné<sup>3</sup>

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<sup>2</sup>Transfusion Service, Fondazione Policlinico Universitario Agostino Gemelli IRCCS, Rome, Italy

<sup>3</sup>Hematology Biology, Nantes University Hospital and CRCINA, Nantes, France

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Gina Zini, Hematology, Catholic University of Sacred Heart, Rome, Italy.  
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## Abstract

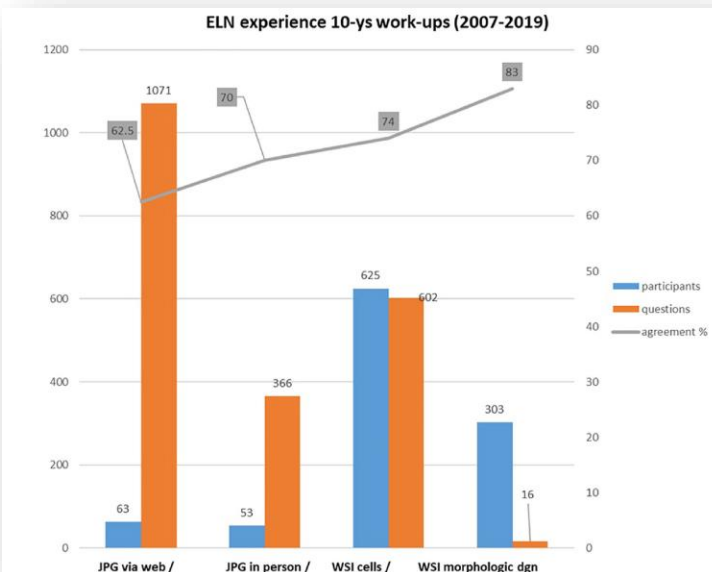
Hematological diagnostics is based on increasingly precise techniques of cellular and molecular analysis. The correct interpretation of the blood and bone marrow smears observed under an optical microscope still represents a cornerstone. Precise quantitative and qualitative cytomorphological criteria have recently been codified by up-to-date guidelines for diagnosing hematopoietic neoplasms. Morphological analysis has found formidable support in digital reproduction techniques, which have simplified the circulation of images for educational or consultation purposes. From 2007 to 2019, the Working Group WP10 of European LeukemiaNet (ELN) used, in annual exercises, digital images to support training in cytomorphology and verify harmonization and comparability in the interpretation of blood and bone marrow smears. We describe the design, development, and results of this program, which had 741 participants in-person or remotely, to which 2055 questions were submitted regarding the interpretation of cytomorphological images. We initially used circulation and presentation of digital microphotographs and then introduced a virtual microscopy (VM). Virtual slides were obtained using a whole slide imaging technique, similar to the one largely used in histopathology, to produce digitized scans of consecutive microscopic fields and reassemble them to obtain a complete virtual smear by stitching. Participants were required to identify cells in labeled fields of view of the virtual slides to obtain a morphological diagnosis. This work has demonstrated substantial improvements in diagnostic accuracy and harmonization with the VM technique. Between-observer concordance increased from 62.5% to 83.0%. The integrity of the digitalized film image, which provides a general context for cell abnormalities, was the main factor for this outcome.

**KEYWORDS**  
cytomorphology, digital imaging, European LeukemiaNet, leukemia, virtual microscopy, whole-slide-imaging

## 1 | INTRODUCTION

with particular relevance in myeloid forms.<sup>1,2</sup> The quantitative and qualitative microscopic diagnostic criteria of the FAB classification

- 2007-2019.
- JPEG and TIF – 2007 to 2011
- “Virtual Microscopy” utilising WSI.
  - From 2012
  - 62.5% to 83%
  - Enhanced dysplasia and blasts.



Zini G, Barbagallo O, Scavone F, Béné MC. Digital morphology in hematology diagnosis and education: The experience of the European LeukemiaNet WP10. *Int J Lab Hematol*. 2022 Sep;44 Suppl 1:37-44. doi: 10.1111/ijlh.13908. PMID: 36074713.

# Will it work?

## Phase 2 diagnostic evaluation of the 3DHISTECH Slide Scanning Technology for Digitising Bone Marrow Slides

Timothy Farren<sup>1</sup>, Samuel Machin<sup>2</sup>, Tom Butler<sup>3</sup>  
<sup>1</sup>Immunophenotyping Department (SIHMDS), Barts Health NHS Trust, London, UK <sup>2</sup>Department of Haematology, University College Hospital, London, UK <sup>3</sup>Department of Haematology, Barts Health NHS Trust, London, UK

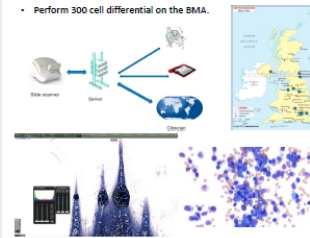
**NHS**  
**Barts Health**  
**NHS Trust**

## Introduction

Bone marrow aspirates and trephine biopsies are crucial to the diagnosis of haematological malignancies. Quantitative assessment of bone marrow aspirates by performing differential cell counts is fundamental in the diagnosis, stratification and prognosis of these malignancies. There has been much debate around the use of digital morphology. The use of digital technology to capture high resolution images of stained tissue sections continues to increase within the histopathology arena, and enables a diagnosis to be carried out remotely. Some progress has been made in histopathology and bone film analysis but studies on the use of digital morphology of bone marrow aspirates are still in their infancy.

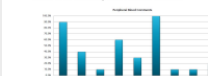
## Study Design

- Phase 2 study involving 10 UK Centres
- 7 real life clinical case scenarios
- Provide a differential diagnosis based on a brief case history, peripheral blood (PB) parameters, bone marrow aspirate (BMA) +/- trephine (BMT) digital morphology scanned using the bench top 3DHISTECH (provided through Sysmex UK Ltd), immunophenotyping and genetic testing.
- Comment on PB, BMA and BMT in terms of cellularity and abnormal cells.
- Perform 300 cell differential on the BMA.



## PB Review

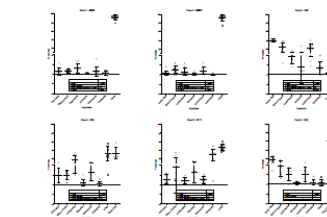
Case 1 - ABMI  
Peripheral blood film review comments  
Note free choice, not forced choice



## BMA/T review



## BMA differential



## WHO Classification

Case	Case	Case	Case
Case 1	Case 2	Case 3	Case 4
Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)
100%	100%	100%	100%
Case 5	Case 6	Case 7	Case 8
Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)	Acute myeloid leukaemia (AML)
100%	100%	100%	100%

## Conclusions

- Excellent concordance on PB comments
- Good concordance on BMA morphological descriptions of cellularity, overall haematopoiesis and abnormalities.
- ANOVA analysis demonstrated highly significant correlation between results from 300 cell differentials  $R^2 = 0.70$  to  $0.99$
- Overall a good concordance in reporting the differential diagnosis to WHO 2008.
- Glass and digital is comparable

Acknowledgements: Helen Barker (Sheffield), John Burthorn (Manchester), Peter Carey (Newcastle), Robert Cuthbert (Belfast), Angela Hamblin (Oxford), Simon Kimber (Sysmex UK Ltd), Khalid Sajja (BHRUT), Tracey Smith-Straney (Liverpool), Geoffrey Summerfield (Gateshead).



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# Purpose of the DHTF

*“To review and develop the use of digital pathology (and AI) in haematology in the UK to advance the integration of cutting-edge digital technology within haematology to improve diagnostic accuracy, efficiency and patient outcome.”*

## Proposed aims of the Task Force are:

1. Appraisal of Current Technology and Evidence
2. Data Collection and Experience Sharing
3. Development of research into Digital Pathology in Haematology
4. Development of National Strategy and Guidelines
5. Enhancement of Educational Resources





REVIEW

# Role of artificial intelligence in haematolymphoid diagnostics

Charlotte Syrykh,<sup>1</sup> Michiel van den Brand,<sup>2,3</sup> Jakob Nikolas Kather<sup>4,5</sup> & Camille Laurent<sup>1,6</sup>

<sup>1</sup>Department of Pathology, Institut Universitaire du Cancer-Oncopole de Toulouse CHU Toulouse, Toulouse, France, <sup>2</sup>Department of Pathology, Radboud University Medical Center, Nijmegen, <sup>3</sup>Pathology-DNA, Arnhem, The Netherlands, <sup>4</sup>Else Kroener Fresenius Center for Digital Health, Faculty of Medicine and University Hospital Carl Gustav Carus, TUD Dresden University of Technology, Dresden, <sup>5</sup>Medical Oncology, National Center for Tumor Diseases (NCT), University Hospital Heidelberg, Heidelberg, Germany and <sup>6</sup>INSERM UMR1037, CNRS UMR5071, Université Toulouse III-Paul Sabatier, Centre de Recherches en Cancérologie de Toulouse, Toulouse, France

Syrykh C, van den Brand M, Kather J N & Laurent C

(2025) Histopathology 86, 58–68. <https://doi.org/10.1111/his.15327>

## Role of artificial intelligence in haematolymphoid diagnostics

The advent of digital pathology and the deployment of high-throughput molecular techniques are generating an unprecedented mass of data. Thanks to advances in computational sciences, artificial intelligence (AI) approaches represent a promising avenue for extracting relevant information from complex data structures. From diagnostic assistance to powerful research tools, the potential fields of application of

machine learning techniques in pathology are vast and constitute the subject of considerable research work. The aim of this article is to provide an overview of the potential applications of AI in the field of haematopathology and to define the role that these emerging technologies could play in our laboratories in the short to medium term.

Keywords: artificial intelligence, haematopathology, lymphoma diagnosis

## Introduction

Lymphomas are among the ten most common cancers worldwide,<sup>1</sup> and are characterized by considerable clinical and biological heterogeneity, with variable prognosis and therapeutic response.<sup>2,3</sup> Lymphoma diagnosis requires in-depth histological analysis by expert pathologists, and relies on ancillary tissue staining techniques, now increasingly combined with

molecular analyses (fluorescence *in situ* hybridization [FISH], clonality, high-throughput sequencing). However, access to these sophisticated technologies is limited, and the risk of misdiagnosis by nonexpert pathologists remains high, as demonstrated in a French nationwide study (Lymphopath Network), which showed that 20% of the diagnoses are inaccurate, with a direct impact on patients' treatment.<sup>4,5</sup> Recent major breakthroughs in deep-learning

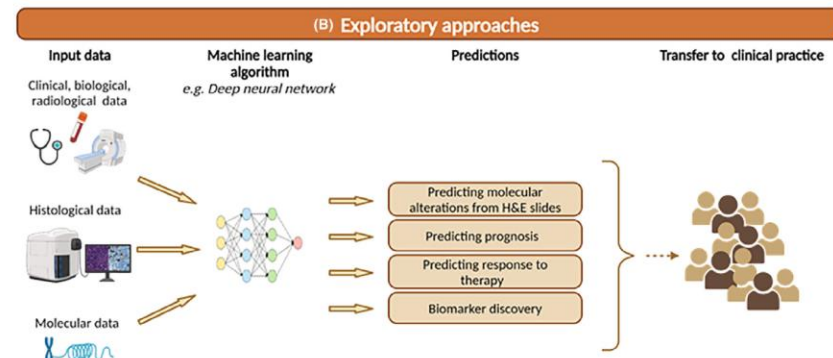
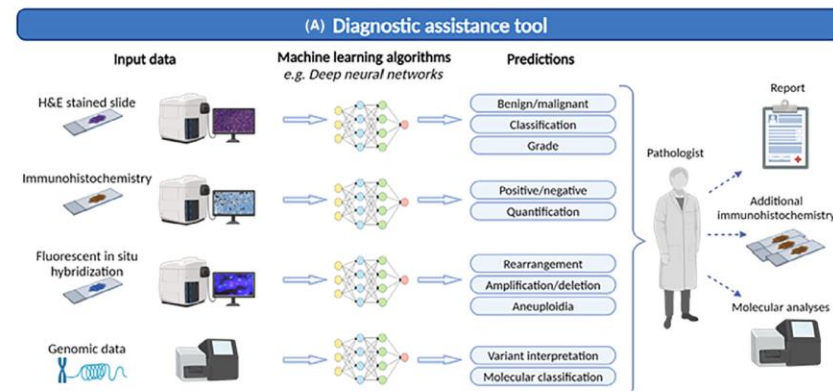
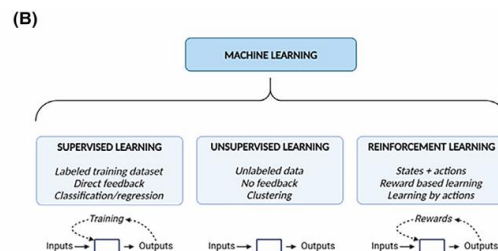
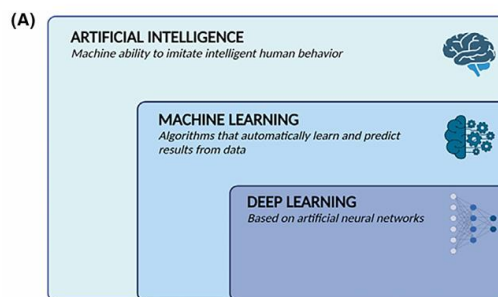
Address for correspondence: C Syrykh and C Laurent, Department of Pathology, Institut Universitaire du Cancer-Oncopole de Toulouse, CHU Toulouse, Toulouse, France; e-mail: [syrykh.charlotte@iuct-oncopole.fr](mailto:syrykh.charlotte@iuct-oncopole.fr) and [laurent.camille@iuct-oncopole.fr](mailto:laurent.camille@iuct-oncopole.fr)

**Abbreviations:** AI, artificial intelligence; AUCROC, area under the receiver operating characteristic curve; CLL, chronic lymphocytic leukaemia/lymphocytic lymphoma; COO, Cell Of Origin; DLBCL, diffuse large B-cell lymphoma; FISH, fluorescence *in situ* hybridization; FL, follicular lymphoma; H&E, haematoxylin-eosin; IPI, international prognostic index; MIP-b, biologic-Mantle cell lymphoma international prognostic index; ML, machine learning; NLP, natural language processing; WSI, whole slide image.

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# The role of Artificial Intelligence



- Cell classification.
- Anomaly detection.
- Workflow efficiency.
- Remote diagnostics.
- Challenges.
- Faster, more accurate diagnosis.



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