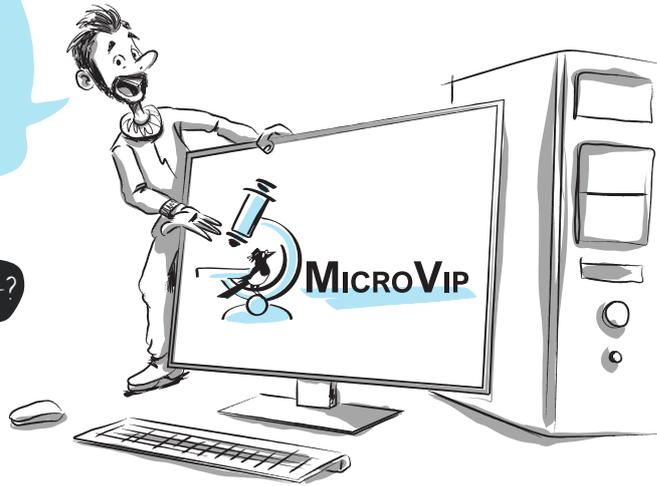


Hi, I'm Ali, Ph.D.
STUDENT AT CREATIS LYON.
LET ME INTRODUCE MICROVIP,
A MODERN MICROSCOPY SIMULATION
PLATFORM.

MODERN MICROSCOPY, WHAT'S THAT?



WE ALL REMEMBER
MICROSCOPES FROM HIGH
SCHOOL BIOLOGY CLASSES!



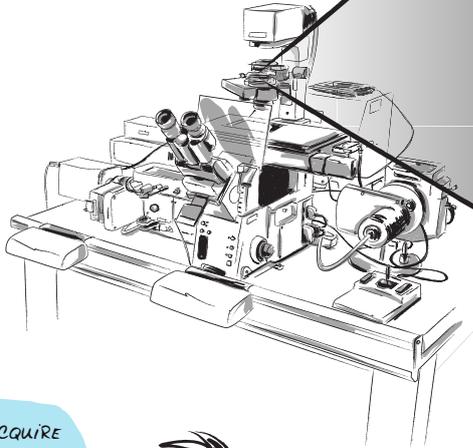
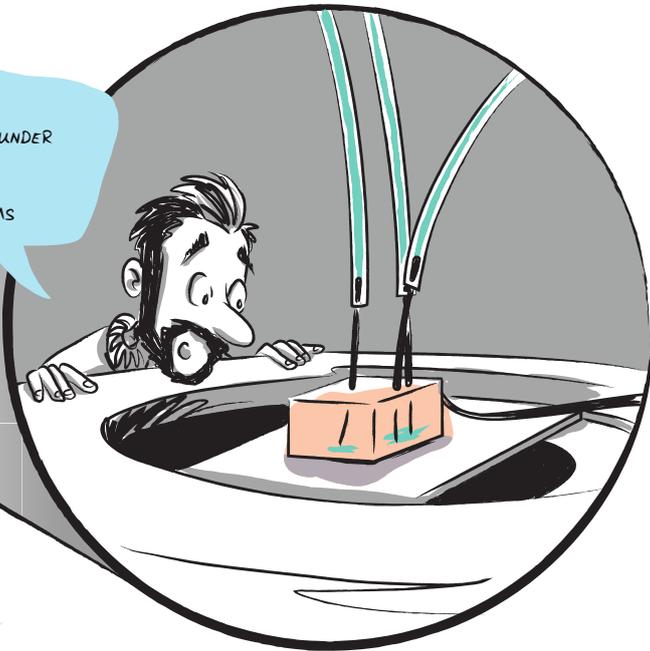
BUT HERE...



WE'RE TALKING
MODERN MICROSCOPY, BECAUSE
THERE'S BEEN NUMEROUS BREAKTHROUGHS
THESE PAST YEARS...



SAMPLES CAN BE AUTOMATICALLY CARRIED UNDER THE MICROSCOPE LENS VIA MICROFLUIDIC SYSTEMS



WE CAN ACQUIRE 3D IMAGES

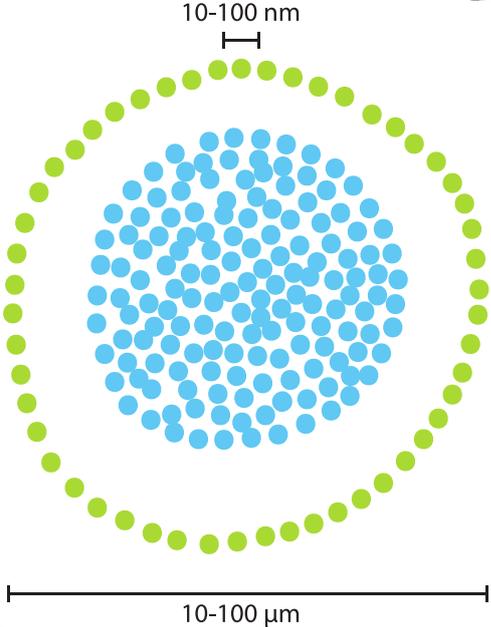


STRUCTURES OF INTEREST CAN BE HIGHLIGHTED USING FLUORESCENT MARKERS



WE CAN OBSERVE DETAILS WITH UNRIVALED RESOLUTION

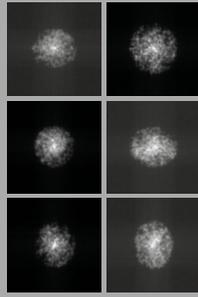
SUPER-RESOLUTION!!



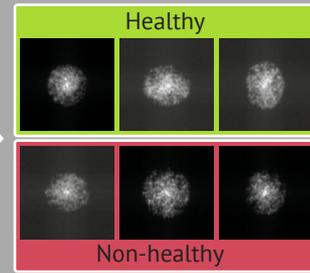
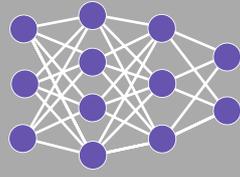
IMPRESSIVE! WHAT IS IT USED FOR?

UNDERSTANDING CELLS' METABOLISM, SEE WHAT HAPPENS WHEN THEY DYSFUNCTION...

MANY THINGS



Machine learning

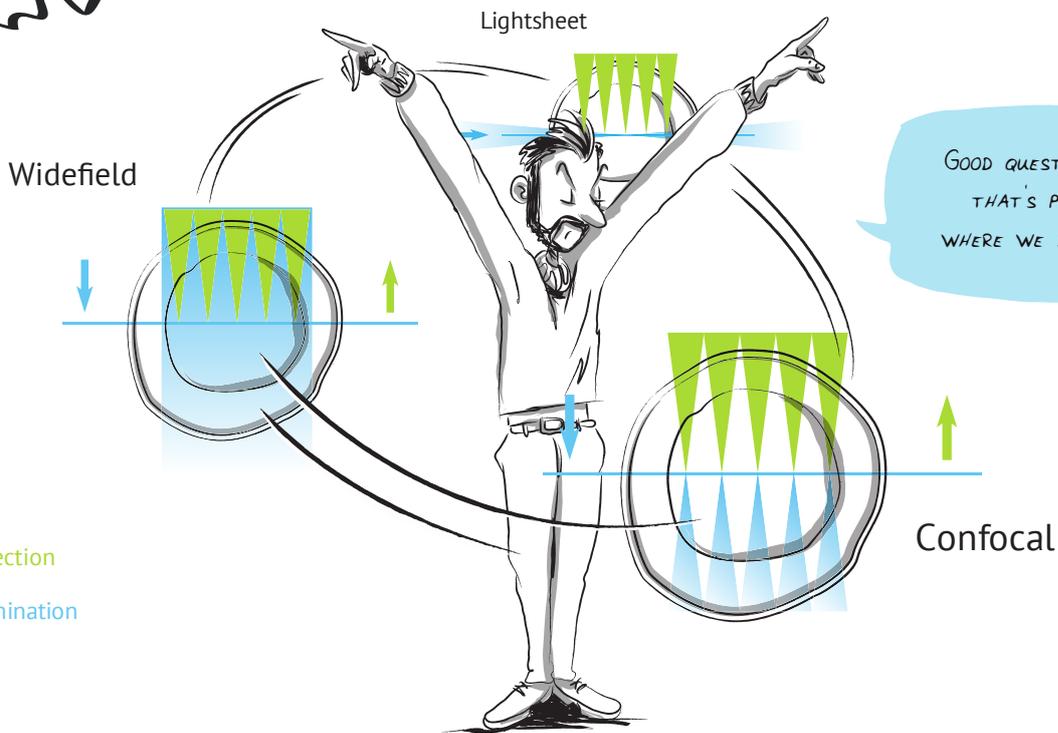


... DUE TO SICKNESS SUCH AS VIRUSES OR CANCER

... IT BECOMES POSSIBLE TO AUTOMATICALLY ANALYSE IMAGES AND THEN AUTOMATICALLY SORT THEM BY CELL TYPE.

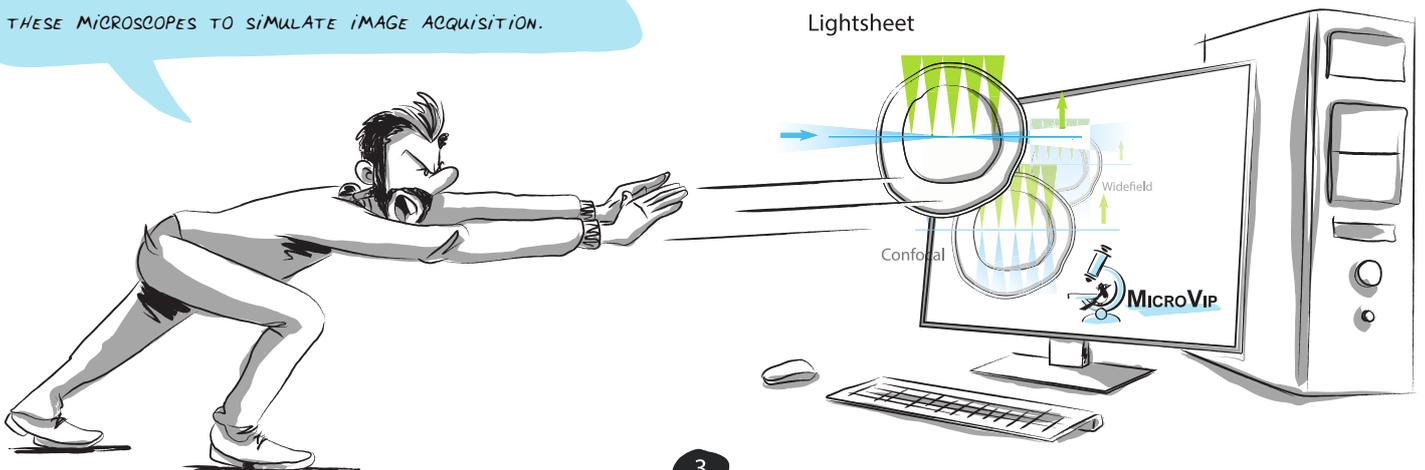
MOREOVER, USING ARTIFICIAL INTELLIGENCE...

BUT THERE ARE MANY MICROSCOPE TYPES, HOW DOES ONE KNOW WHICH ONE'S APPROPRIATE?



GOOD QUESTION, THAT'S PRECISELY WHERE WE STEP IN!

WE IMPLEMENTED A DIGITAL VERSION OF ALL THESE MICROSCOPES TO SIMULATE IMAGE ACQUISITION.



TELL ME MORE ABOUT IT!

FIRST, WE MODEL THE BIOLOGICAL SAMPLE OF INTEREST.

IN OUR CASE, IT'S CHROMATIN CHAINS

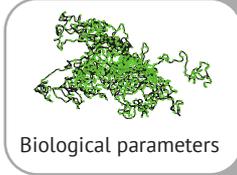
WHOSE ORGANISATION DEPENDS ON THE CELL'S HEALTH STATE.

AFTERWARDS, WE SIMULATE THESE CHAINS' IMAGES THROUGH DIFFERENT MICROSCOPES

WE FINALLY TRAIN ALGORITHMS TO SORT GENERATED CELL IMAGES.

THIS ALLOW US TO CHOOSE THE MOST APPROPRIATE

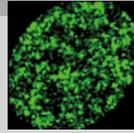
MICROSCOPE.



Biological parameters

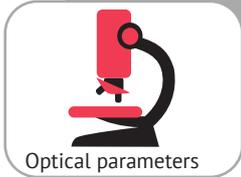


MICROVIP



Simulated image

Image analysis



Optical parameters



BUT WHY DON'T YOU PERFORM REAL MICROSCOPY EXPERIMENTS RATHER THAN GOING THROUGH ALL THESE SIMULATION STEPS?



ACTUALLY, IT'S A HUGE TIME GAIN BECAUSE

ARTIFICIAL INTELLIGENCE NEEDS ANNOTATED IMAGES

IN OTHER WORDS, WE NEED TO INDICATE WHETHER EACH CELL IS HEALTHY OR NOT. THIS IS VERY TIME-CONSUMING!

WITH SIMULATED SAMPLES WE AUTOMATICALLY HAVE THIS ANNOTATION, NO NEED TO PERFORM IT MANUALLY.

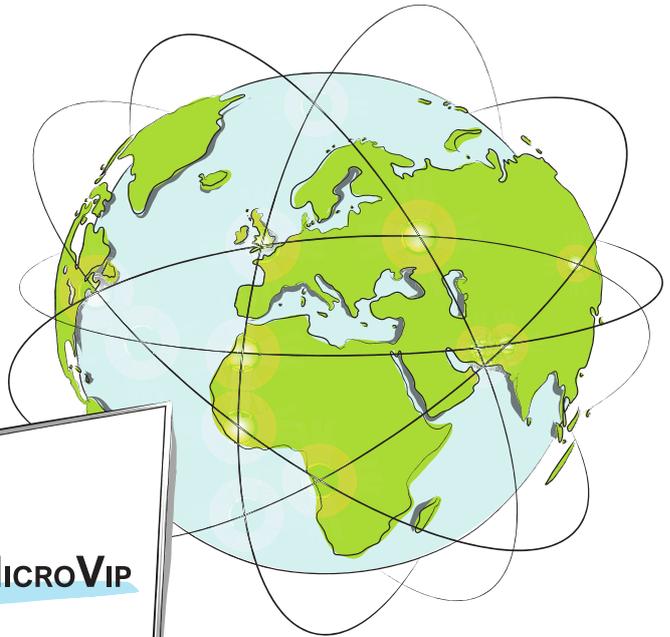
I SEE. THEN, WHAT DOES MICROVIP LOOK LIKE?

IT SIMPLY IS A WEB PORTAL
RUNNING ON A LARGE COMPUTATION GRID
SCATTERED ALL AROUND EUROPE.

IT ALLOWS RAPID GENERATION OF
A HIGH NUMBER OF SIMULATED IMAGES,
AUTOMATICALLY ANNOTATED.



THIS PORTAL IS AVAILABLE TO
RESEARCHERS ALL AROUND THE WORLD



DURING MY PH.D.

FOR INSTANCE, WE COLLABORATED
WITH IMPERIAL COLLEGE LONDON,
PLITECNICO MILANO, LARIS IN ANGERS,
AND EVEN THE COMPANY ELVESYS.



Present work is part of European project PROCHIP "Chromatin organization PROfiling with high-throughput super-resolution microscope on a CHIP".

It has been made possible by complementary expertise and collaboration of CREATIS (CNRS UMR 5220, INSERM U1294) and LARIS (EA 7315), as well as of their European partners CNR-IFN and University of Trento (Italy), ICL (UK), and Elvelys (France).

This project is funded by the European Union as part of HORIZON 2020 program with identifying number 801336.

Notably, this project would not have been possible without the contributions from Ali Ahmad, David Rousseau, Carole Frindel, David Sarrut, Frédéric Cervenansky, Sorina Pop, Axel Bonnet, Guillaume Vanel as well as Clémence Heller for the illustrations.

