

Evolution de la RMN vers l'imagerie médicale

Les Origines : Laboratoire de spectroscopie et luminescence (Prof. J. Janin)

**Spectroscopie
hertzienne**

Spectrométrie
optique

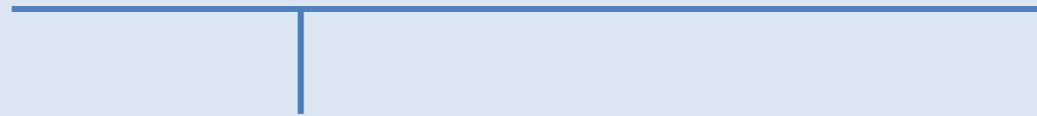
Matériaux
luminescents

Physico-Chimie
quantique

Lasers

RPE (Prof. A. Erbeia, Prof. J. Delmau)

RMN



Physico-Chimie

Spectroscopie RMN

Dynamique moléculaire

Chimie organique

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Spectroscopie RMN

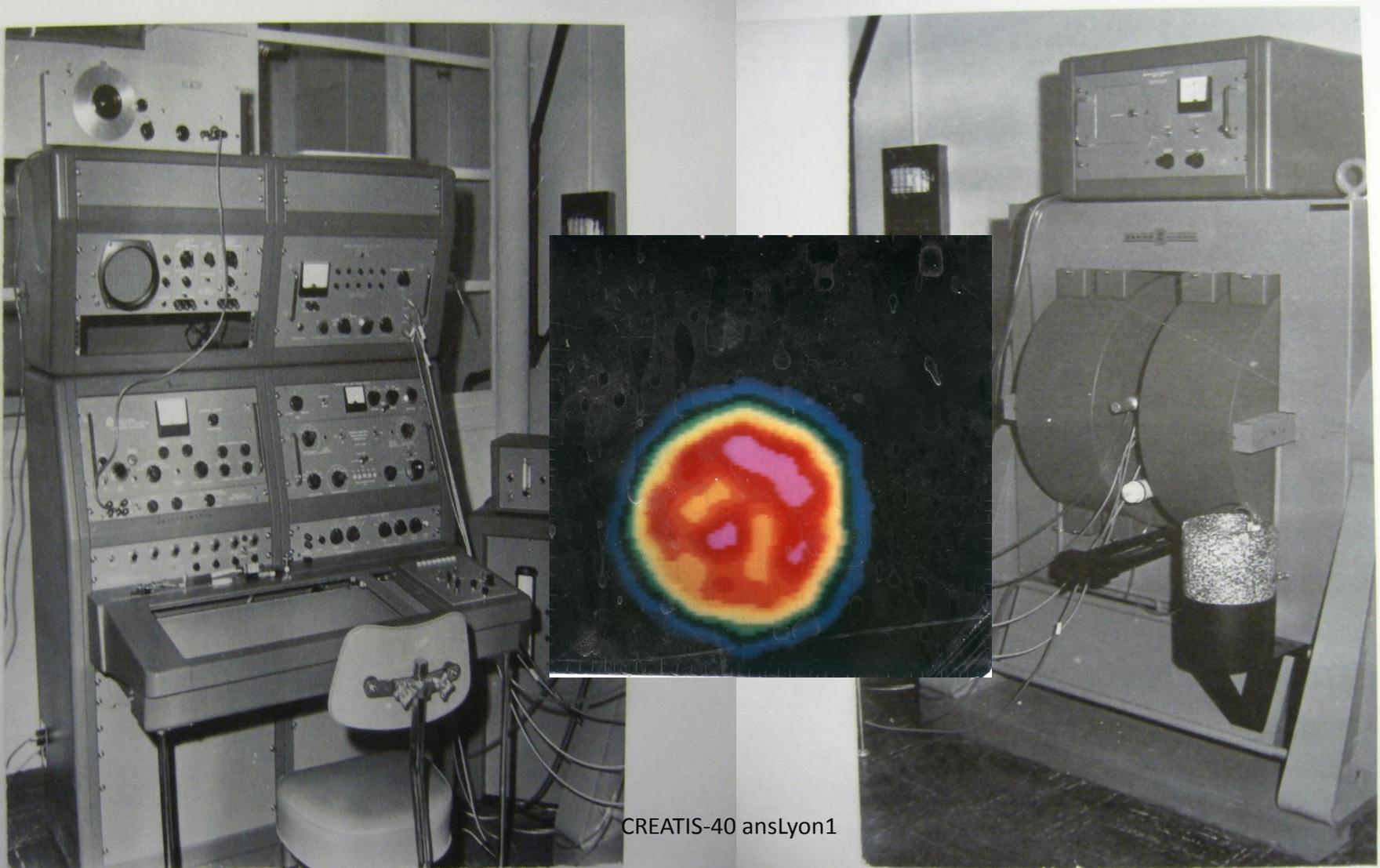
Dynamique moléculaire

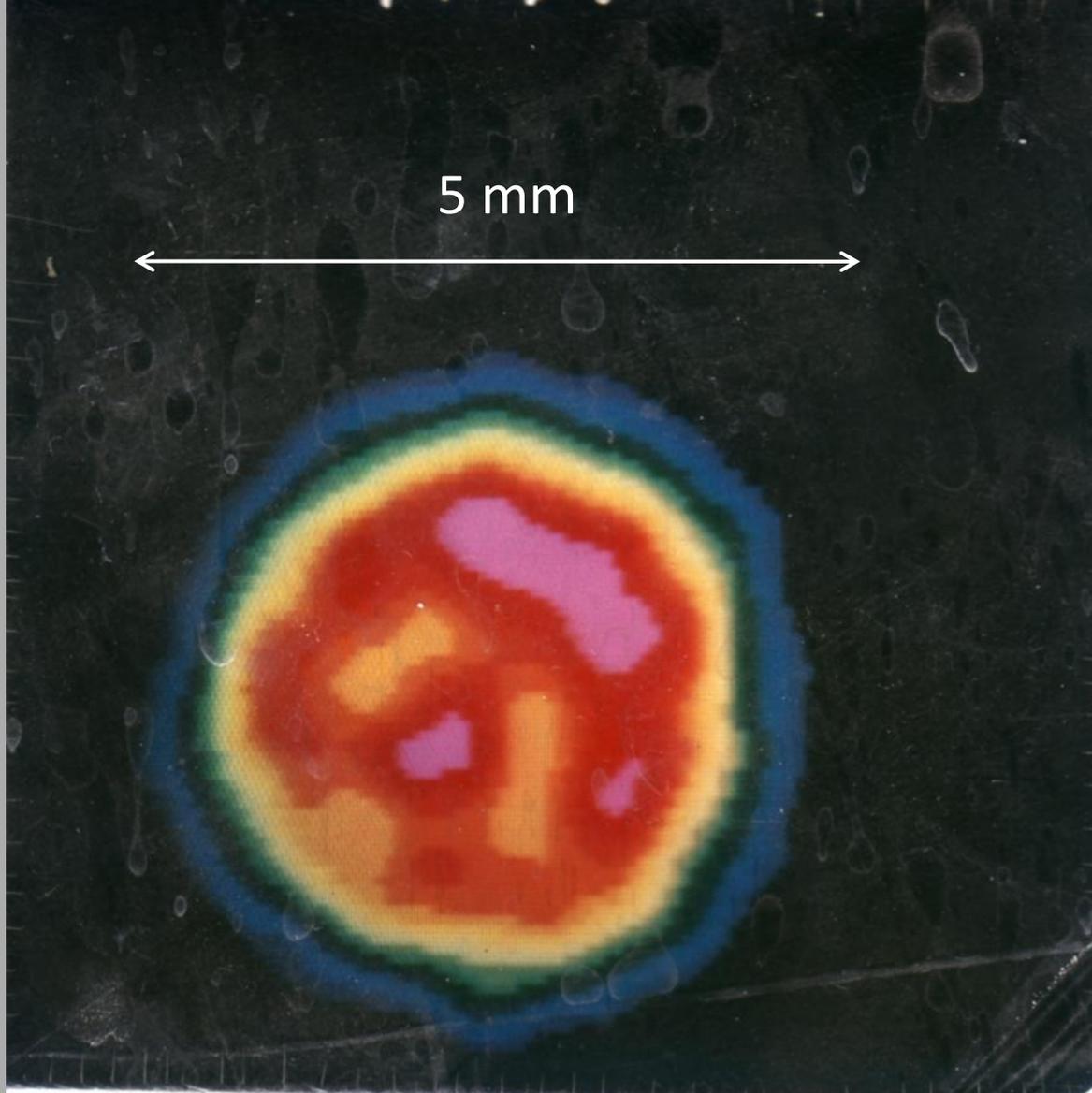
Chimie organique

Liens avec RMN biomédicale actuelle

Produits de contraste, Imagerie Moléculaire, Imagerie fonctionnelle, Imagerie métabolique
Imagerie de diffusion, Spectroscopie in vivo

SPECTROMETRE 100 MHz (Varian HA 100) à ondes continues





5 mm



2,35 teslas

100 MHz

Glec # 4,7 mT/m

BPFP (72 proj.)

J.J. MALLET
A.BONMARTIN
H. MEHIER

D. GRAVERON
C. LAPRAY
J.C. DUPLAN
A.BRIGUET

(Avril 1978)

Développement à partir de 1981

Labo RMN: B1*, UPRES,UMR CNRS

-Imageur résistif 20 MHz (Institut de Physique Nucléaire de Lyon 1980-1986)

-Imageur résistif corps entier Philips 5,5 MHz (Bâtiment Oméga 1986-1994)

**-Spectromètre 200 MHz Centre d'études du Métabolisme
(HEH/Inserm U80, 1987-1991)**

-Système SMIS-Oxford à 2 teslas (Bâtiment CPE, 1994...) + mise en place de la plateforme de spectroscopie commune UCB-CPE (4 spectromètres + 1 du centre des très hauts champs)

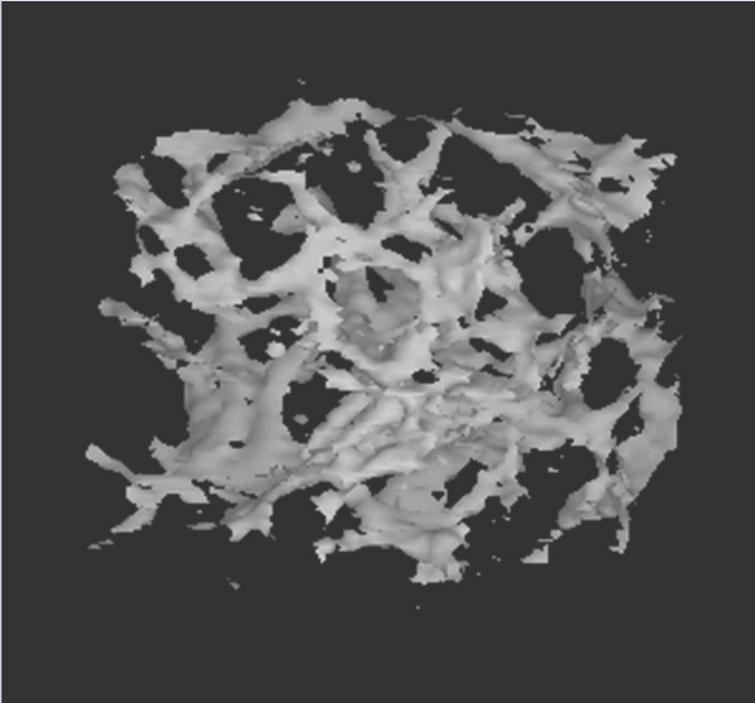
-Système (Bruker-Magnex) 7 teslas (Animage, CERMEP , 2003)

-Corps entier Siemens 1,5 teslas (CERMEP, 2004)

-Système Bruker-Magnex 4,7 teslas (Bâtiment CPE, 2005)

QUELQUES RESULTATS ou THEMES SIGNIFICATIFS

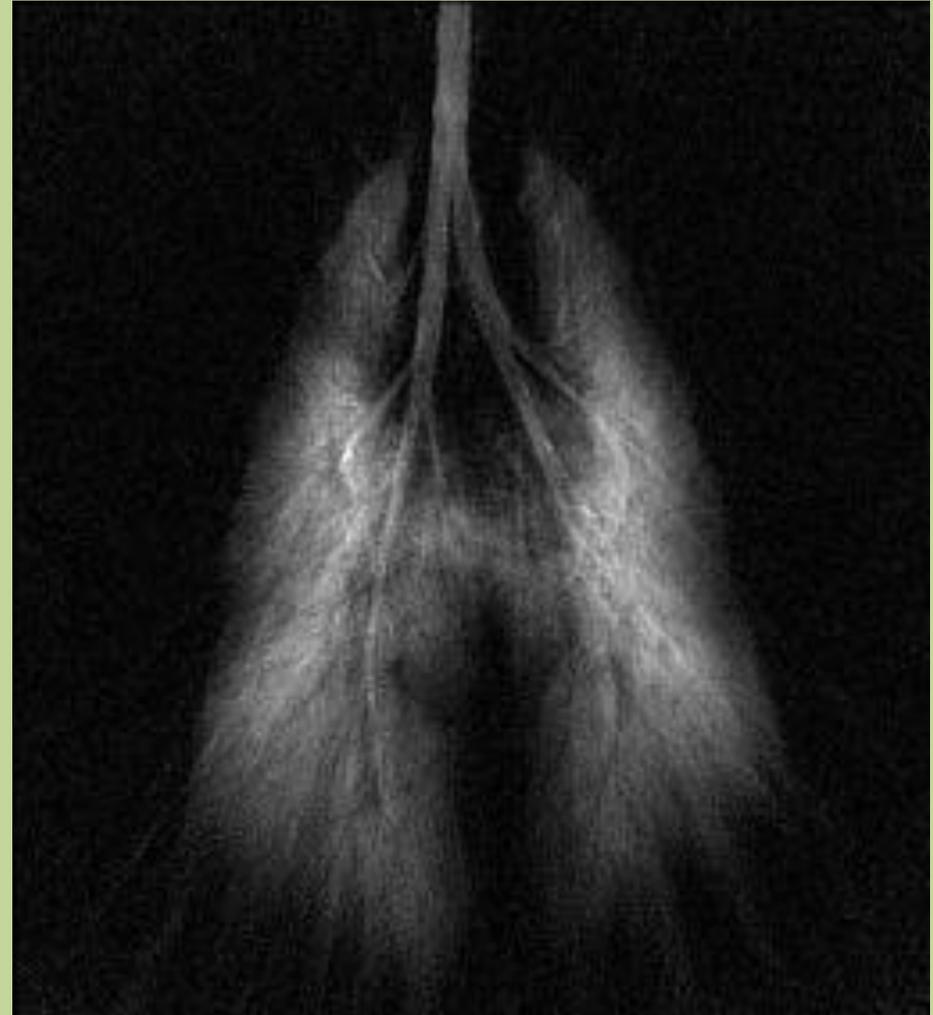
Imagerie des tissus denses (« Biomatériaux vivants »)



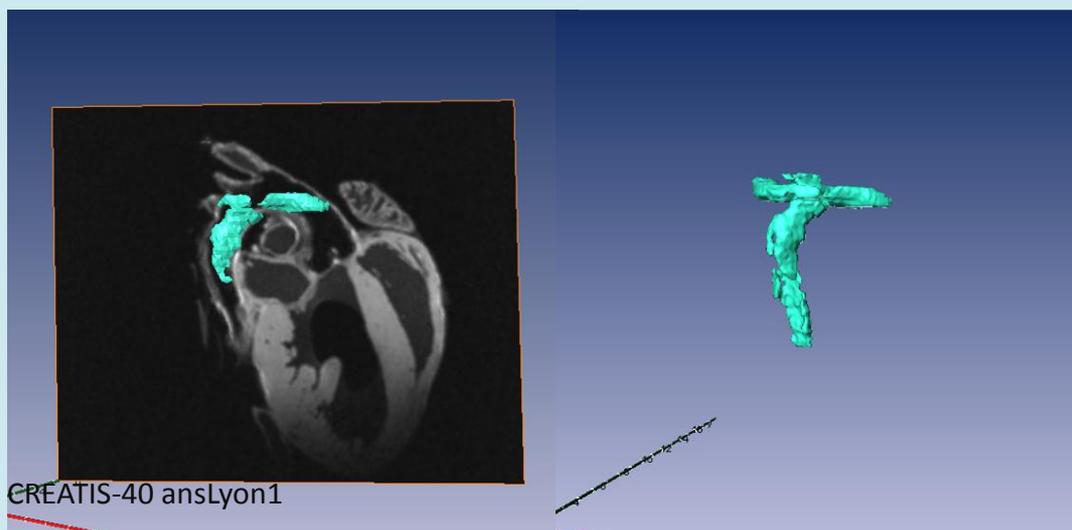
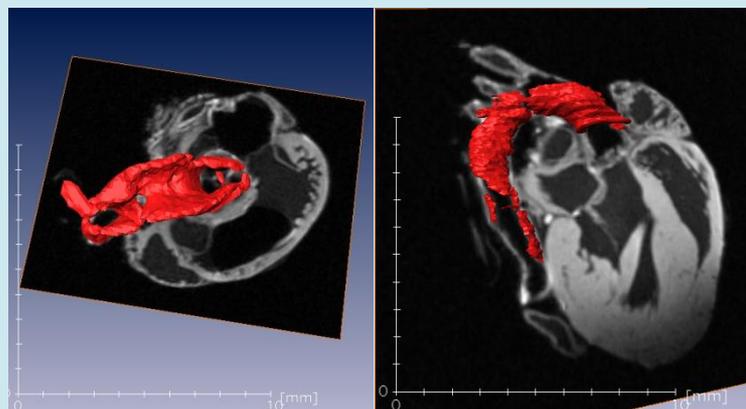
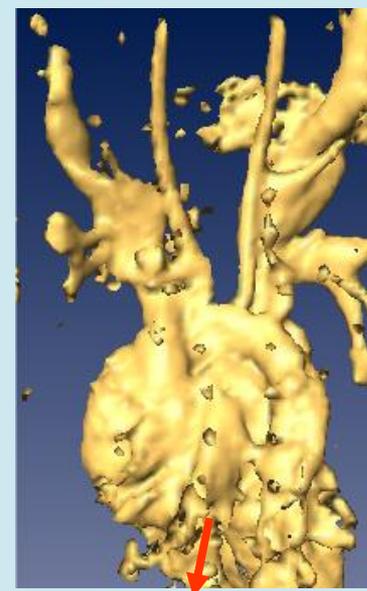
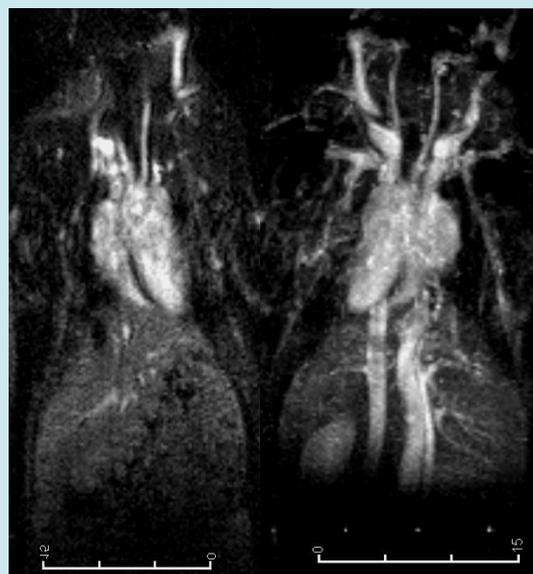
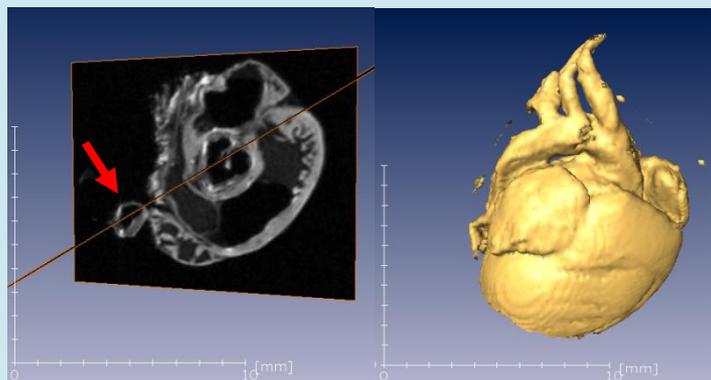
Ventilation pulmonaire avec hélium 3 hyperpolarisé (modèle animal)

RESOLUTION
temporelle : 100
ms
spatiale ~ 300 μ m

FOV = 42 mm
40³ respirations
120 cm of He3
durée acq.= 1mn

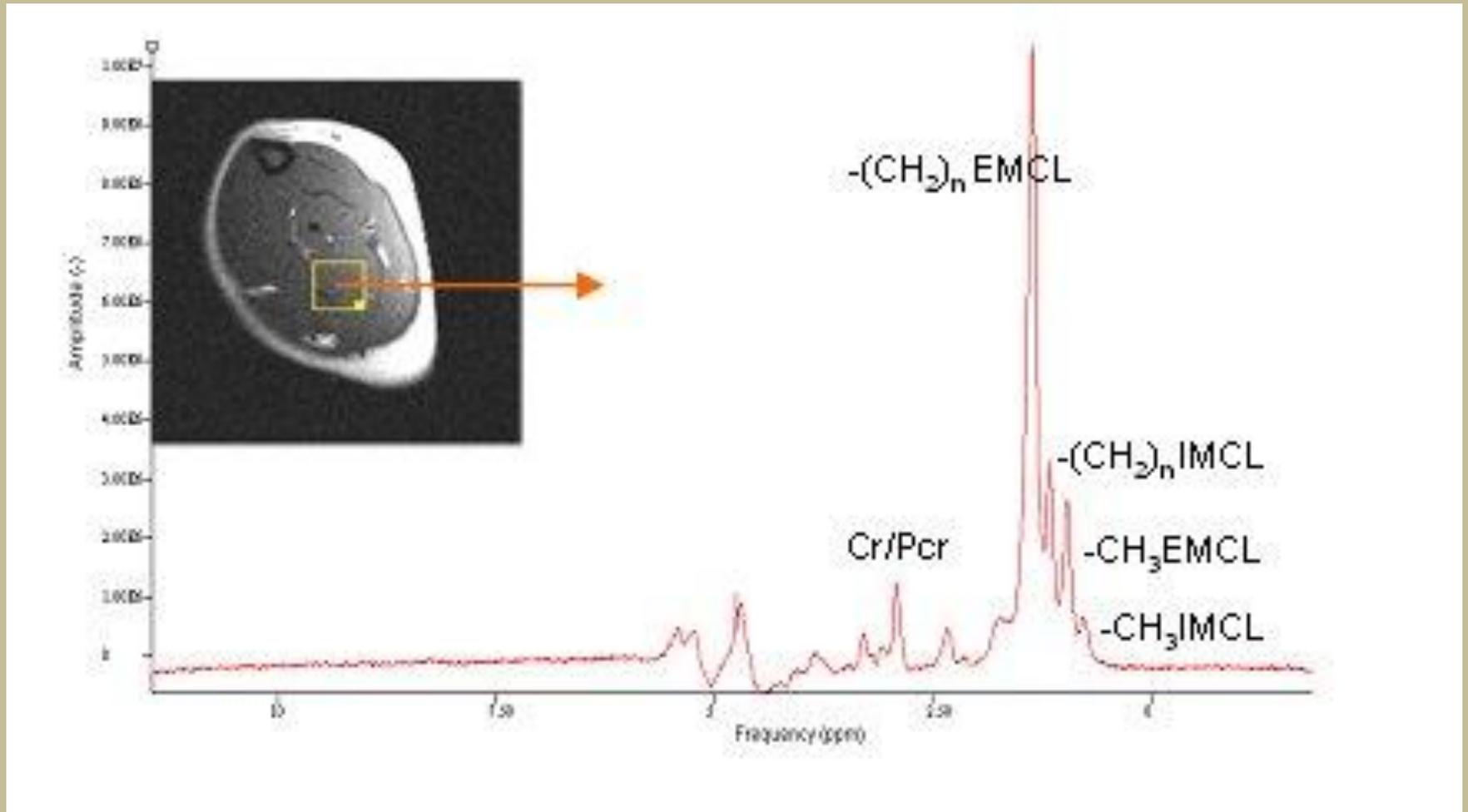


La Plaque d'Athérome (souris)



Données Emmanuelle CANET

Discrimination des lipides intra/extra-musculaires par spectroscopie proton localisée à l'angle magique

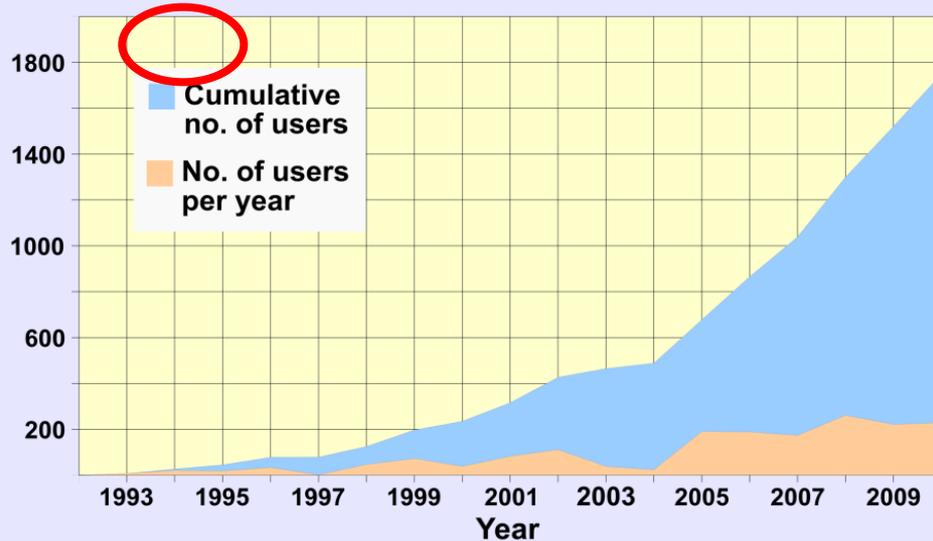


Travaux Rachida FISSOUNE et Hassem HIBA

jMRUI Software Package

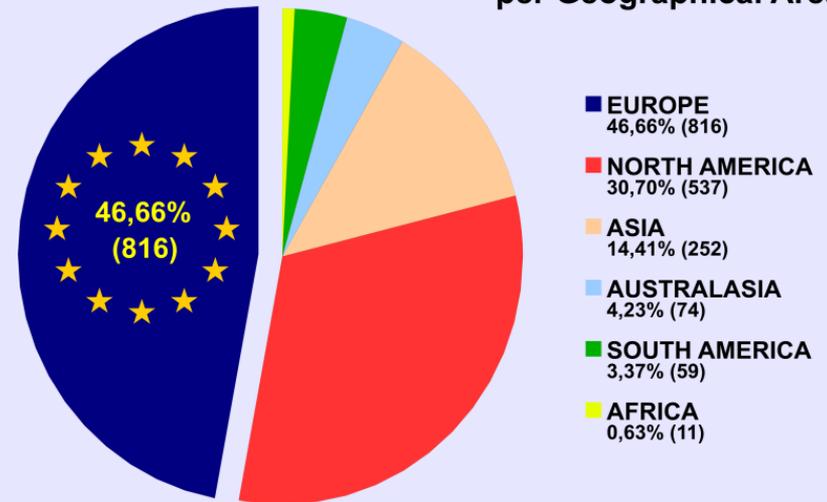
Advanced Signal Processing for Medical MRS

Number of MRUI Licensees



  FAST - MRTN-CT-2006-035801
<http://www.fast-mrs.eu>

Number of MRUI Licensees per Geographical Area



  FAST - MRTN-CT-2006-035801
<http://www.fast-mrs.eu>

Data as of 2010-10-27

- <http://www.mrui.uab.es/mrui/>
- Environments: Windows 2000, XP, Vista, Linux

- Danielle GRAVERON

EVOLUTION DE LA RMN BIOMEDICALE

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Palm NMR and 1-Chip NMR

Nan Sun, Member, IEEE, Tae-Jong Yoon, Hakho Lee, William Andress, Ralph Weissleder, and Donhee Ham, Member, IEEE

Abstract—In our earlier work, we developed a 2-kg NMR system, which was 60× lighter, 40× smaller, yet 60× more spin-mass sensitive than a 120-kg state-of-the-art commercial benchtop system. Here we report on two new nuclear magnetic resonance (NMR) systems that represent further orders-of-magnitude size reduction and lab-on-a-chip capability. The first system, which weighs 0.1 kg and can be held in the palm of a hand, is the smallest NMR system ever built, and is 1200× lighter, 1200× smaller, yet 150× more spin-mass sensitive than the commercial system. It is enabled by combining the physics of NMR with a CMOS RF transceiver. The second system, which even integrates a sample coil, directly interfaces the CMOS chip with a sample for lab-on-a-chip operation. The two systems detect biological objects such as avidin, human chorionic gonadotropin, and human bladder cancer cells.

Index Terms—Biosensors, CMOS, radio-frequency integrated circuits (RFICs), lab on a chip, nuclear magnetic resonance (NMR).

I. INTRODUCTION

PARTICULAR types of atomic nuclei, such as protons in hydrogen atoms, act like tiny bar magnets due to their spin. Nuclear magnetic resonance (NMR) is a resonant interaction between a radio-frequency (RF) magnetic field and the nuclei

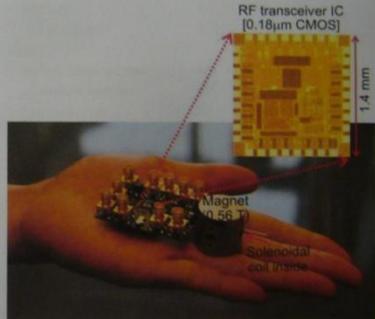


Fig. 1. 0.1-kg palm NMR system.

hospitals, testing facilities, and laboratories. A case in point is

J.B. Haun *et al* Sci Transl Med **3**: 71ra16 (2011)

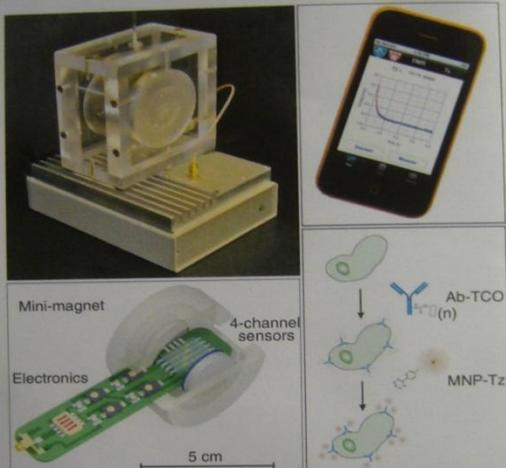
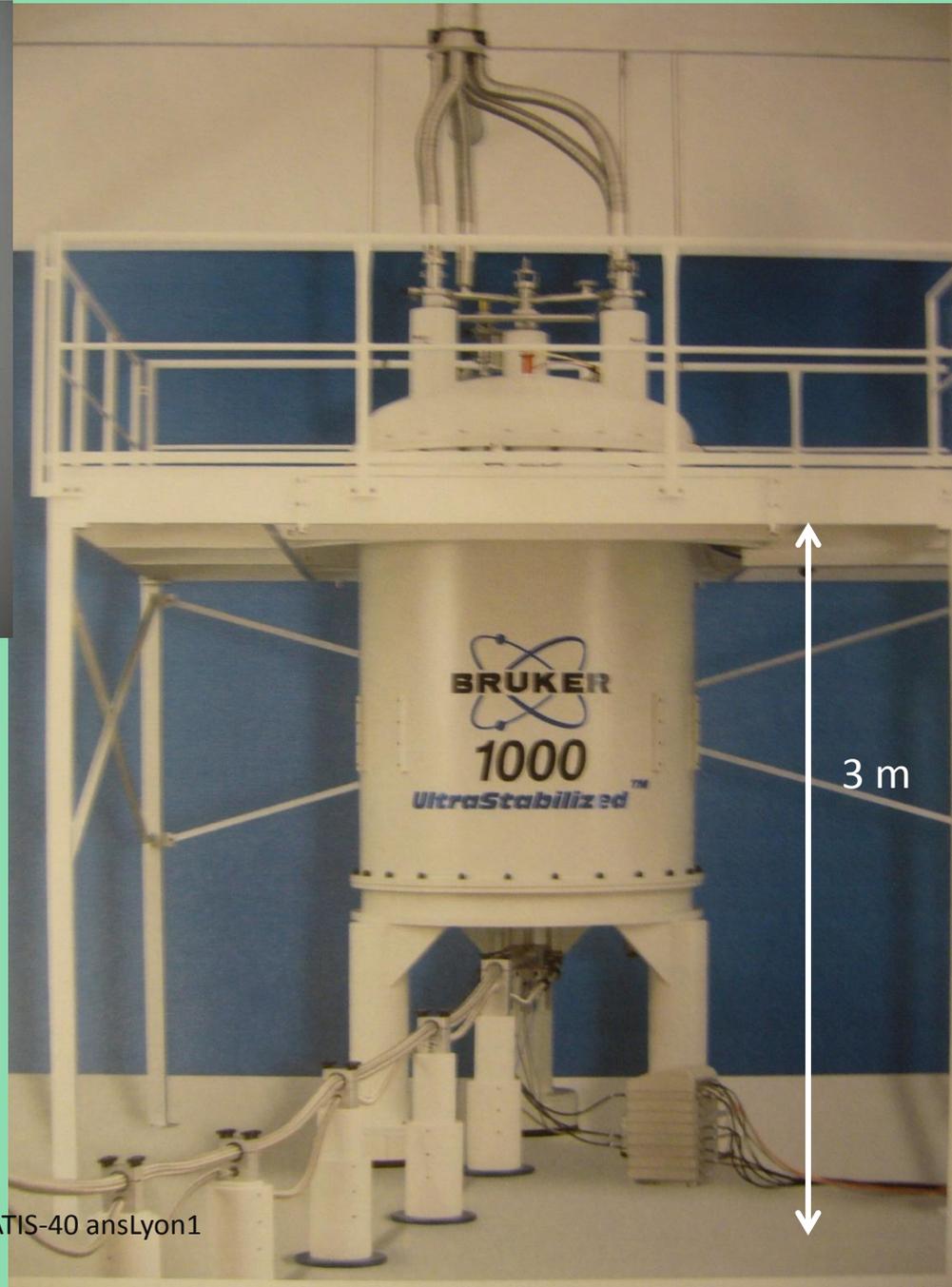


Fig. 1. The μ NMR clinical analysis system (DMR-3) and bioconjugation strategy (BOND-2). (Left, top) Complete μ NMR system for use at the patient's bedside. The bottom component contains all of the circuitry for NMR measurements, whereas the top enclosure holds a permanent magnet and chip-sized microliter-volume sensors. (Left, bottom) State-of-the-art μ NMR probe used for sensing within the mini magnet. (Right, top) Smart phone interface for operating the μ NMR system. (Right, bottom) Fine-needle aspirates from each patient sample were processed with a bio-orthogonal amplification strategy adapted for clinical samples. The bio-orthogonal amplification allows ultrasensitive detection of cells



CREATIS-40 ansLyon1