

### 3D Morphometry and Microdensitometry for Dental and Other Life Science Research







# Overview

- 1. Principles of Micro-Computed X-ray Tomography ("micro-CT")
  - Scanning Workflow of uCT
  - o Mechanical Setup
  - X-ray Attenuation
  - How reconstruction works
  - Analysis Workflow
- 2. SkyScan 1172
- **3**. Solutions for Life Science Applications
- 4. New Features in DataViewer, CTVox, CTAN

## What is tomography?



#### 1. **"Scan"**

A set of x-ray "projection" images are taken over a rotation of the imaging axis of 180 or 360 degrees

#### 2. "Reconstruction"

The "projection" images are processed by the filtered Feldkamp cone-beam method to create the stack of crossection slices

#### 3. "Analysis and visualisation"

The reconstructed crossection slices are processed into 3d models for morphometric measurements and virtual visual inspection







### The in vivo and ex vivo methods of microCT

In the *"ex vivo"* scanner type, the sample rotates on a stage around a vertical axis, allowing angular projection images over 360 degrees





In the **"in vivo"** scanner type, the sample – e.g. a live mouse or rat – lies still on a horizontal bed, while x-ray source and camera rotate around the sample bed, over 360 degrees







### Micro-CT analysis pathway



3D Morpho-

BMD calibrated densitometry

### X-ray attenuation

**Röntgen:** The attenuation of x-rays of wavelength  $\lambda$  is

 $I/I_0 = \exp(-\mu x)$ , (1)

where  $I_0$  is the intensity of the unattenuated x-ray beam, I is the beam's intensity after traversing x thickness of (homogeneous) material (in cm) with  $\mu$  being the linear attenuation coefficient (in cm<sup>-1</sup>).

In terms of the mass attenuation coefficient  $\mu/\rho$  (units cm<sup>2</sup>/g)

$$I/I_0 = \exp[(-\mu/\rho) \rho x)]$$
 . (2)

In terms of what happens in each thickness element dx

$$dI/I = -(\mu/\rho) \rho dx$$
 . (3)

Adding the increments of the attenuation along the direction of x-ray propagation yields the more general form

$$I = I_0 \exp[-\int \mu(s) ds]$$
 , (4)

where  $\mu(s)$  is the linear absorption coefficient at position s along ray s. The problem is assigning the correct value of  $\mu$  to each position along this ray (and along all the other rays traversing the sample) knowing only the values of the line integral for the various

orientations of s, i.e.,

```
\int \mu(s) ds = \ln(I_0/I) . (5)
```

For compounds or mixtures and wt. fractions  $w_i$ 

$$<\mu> = \sum w_i (\mu/\rho)_i <\rho>$$
 . (6)



Cartoon of back projection reconstruction

Absorption profiles from an idealized specimen.



Back projection (2)



Back projection (3)

### Third profile projected

### **Objects identified**



### Relative error in thickness measurements from partial volumes – threshold one-half of voxel



### Importance of Contrast Resolution



Reproduced from SR Stock, <u>MicroComputed Tomography Methodology</u> and <u>Applications</u>, In press 2008. © CRC Press/Taylor and Francis

# X-ray density contrast in biological samples

Photoelectric absorption ( $\tau$ ) is very strongly dependent on the atomic number Z of the absorbing material.

X-ray density therefore is primarily determined by a materials *elemental composition* and the position of these elements in the periodic table





# Atomic compositions of some biological tissues

	Н	С	Ν	0	Na	Mg	Р	S	Cl	К	Са
Atomic number Z	1	6	7	8	11	12	15	16	17	19	20
Tissue: percent content by mass											
Fat	11.4	59.8	0.7	27.8	0.1			0.1	0.1		
(Water)	11.2			88.8							
Blood	10.2	11.0	3.3	74.5	0.1		0.1	0.2	0.3	0.2	
Liver	10.2	13.9	3.0	71.6	0.3		0.2	0.3	0.2	0.3	
Brain	10.7	14.5	2.2	71.2	0.2		0.4	0.2	0.3	0.3	
Bone	3.4	15.5	4.2	43.5	0.1		10.3	0.3			22.5

### Four biological tissues can be resolved by x-ray micro-CT due to natural absorption contrast

- Bone
- Lean tissue
- Fat
- Lung







However contrast can be added by the use of contrast agents, such as resins containing heavy elements (e.g.

Microfil<sup>™</sup>)



OPEN OACCESS Freely available online



Three-Dimensional Characterization of the Vascular Bed in Bone Metastasis of the Rat by Microcomputed Tomography (MicroCT)

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### 2D histomorphometry requires **indirect model assumptions** to make 3D parameter estimates





1. w/D = Tan
$$\theta$$
, so w = D Tan $\theta$   
2. T/W = Cos $\theta$ , so W = T/Cos $\theta$   
3. w/W =  $\frac{D}{T}$  Tan $\theta$  Cos $\theta$  =  $\frac{D}{T}$  Sin $\theta$ 

The mean value of  $\sin\theta$  is given by:

4. 
$$\overline{\sin\theta} = \int \frac{\sin\theta \, d\theta}{\theta} = \frac{-\cos\theta}{\theta}$$

Assuming all values of  $\theta$  between 0 and 90 ( $\pi/2$  radians) to be equally likely, the mean value of w/W ( $\overline{w}/\overline{w}$ ) is given by:

$$\overline{w/W} = \frac{2}{\pi} \cdot \frac{D}{T} (-\cos \pi/2 + \cos \theta)$$
$$= \frac{2}{\pi} \cdot \frac{D}{T} = \frac{2}{\pi} R, \text{ where } R = \frac{D}{T}$$



TABLE 4. DERIVED INDICES IN BONE HISTOMORPHOMETRY

Type of index	Name of index <sup>a</sup>	Abbreviation <sup>a</sup>	Formula <sup>b</sup>		
I. Structural	Trabecular number	Tb.N	(BV/TV)/Tb.Th <sup>c</sup>		
	Trabecular separation	Tb.Sp	$(1/Tb.N) - Tb.Th^{c}$		
II. Kinetic	Mineralizing surface <sup>d</sup>	MS	(dLS + sLS/2)/BS <sup>e</sup>		
	Mineral apposition rate	MAR	Ir.L.Th/Ir.L.t		
	Adjusted apposition ratef	Aj.AR	MAR*(MS/OS)		
	Osteoid apposition rate	OAR	same <sup>g</sup>		
	Mineral formation rated	MFR	MAR*(MS/BS)		
	Bone formation rate <sup>d</sup>	BFR	same <sup>g</sup>		
	Bone resorption rate <sup>d</sup>	BRs.R	see text		
	Mineralization lag time	Mlt	O.Th/Aj. AR		
	Osteoid maturation time	Omt	O.Th/MAR <sup>h</sup>		
	Formation period	FP	W.Th/Aj.AR		
	Resorption period	Rs.P	FP*(Oc.S/OS) <sup>h</sup>		
	Reversal period	Rv.P	FP*(ES - Oc.S)/OS		
	Remodeling period <sup>i</sup>	Rm.P	FP*(ES + OS)/OS		
	BMU lifespan (sigma)	Sg (or $\sigma$ )	see text		
	Quiescent period	QP	FP*(QS/OS)		
	Total period <sup>i</sup>	Tt.P	FP*(BS/OS)		
	Activation frequency <sup>k</sup>	Ac.f <sup>1</sup>	(1/Tt.P)		

# By contrast, micro-CT morphometry of bone makes DIRECT MEASUREMENTS in 3D, no model assumptions





Fig. 1. Medial axis with circles.



Fig. 4. Balls inside the shape.



Fig. 7. Covering test on two balls restricted to  $\frac{1}{8}\mathbb{Z}^2$ .







### Color-coded thickness map in cortical bone Haversian canals (SkyScan1172 scan)



# Overview

- 1. Principles of micro-computed x-ray tomography ("micro-CT")
- 2. SkyScan 1172
  - Specifications
  - Adaptive Geometry
  - Thermal Correction
  - In Situ Stages
- **3**. Solutions for Life Science Applications
- 4. New Features in DataViewer, CTVox, CTAN



SkyScan 1172				
Min. voxel size	0.6 µm			
Max.sample FOV*	50 x 70 mm			
Max. X-ray voltage	100 kV			
X-ray camera mega-pixels	11 Mp (4000 x 2670)			
Key advantages	Resolution, speed (adaptive geometry), high throughput, versatility – wide range of applications			

#### Adaptive Geometry



#### The SkyScan 1172 High Resolution Micro-CT

#### **Adaptive Geometry**



### SkyScan 1172 / 1272 are the only scanner with Adaptive Geometry

### Image Clarity

### **Projection Pixel Shift Correction**



### In Situ Stages

### Materials Testing Stage



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  - o Dental (Scaffolds)
  - o Bone
  - o Bone (Biomechanics / Implants)
  - o Soft Tissue
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### Orthodontic analysis of the mouse mandible

• The molars can be automatically separated from the rest of the mandible by the image processing tools



SkyScan 1172, 6 micron voxel

Separation of molars from the mandible by size criteria. Note: periodontal ligament separates the molars from mandibular bone



1. Crossection



2. Binarised whole image



3. Mandible > molars: remove smaller part



4. Molars < mandible: remove larger part

• 3D animation of separated mandible and molars



# Imaging of human tooth samples

 Preliminary micro-CT results from analysis of MIH (molar incisor hypomineralisation) tooth samples

## MIH tooth sample 1



### MIH tooth sample 2



# Surface modeling of an MIH tooth (CTVol)



### Enamel crown with treatment applied





### Surface rendering

Volume rendering
# Tooth with implant material lining the pulp cavity



Three-dimensional Evaluation of Effectiveness of Hand and Rotary Instrumentation for Retreatment of Canals Filled with Different Materials

Mobammad Hammad, MSc, Alison Qualtrough, PhD, and Nick Silikas, PhD

#### (J Endod 2008;34:1370-1373)



# Endodontic fillings and treatments in pulp cavity



Within the pulp cavity



Connecting the pulp cavity with the outside

# Tooth fragment showing severe caries with some endodontic treatments



Binarisation and surface rendering of the caries region

#### In research from Jeddah, Saudi Arabia, SkyScan CT-Analyser software tools allow demineralisation to be quantified in selected dentine bands around the pulp canal:





#### Endodontic procedures: before and after



Endodontic research requires COMPARISON of a root canal before (gold) and after (blue) a treatment procedure

## SkyScan CTAn allows comparison of before and after datasets, using custom processing functions



Here the white mask is the root BEFORE the procedure

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Dental related and periodontal reconstruction involves the development of osteoconductive scaffold materials



Viewed by Guestion 5/9/2012

## Scaffold materials can be scanned ex vivo, such as PLA polymer type scaffolds





Poly LLA CO CL



Poly LLA CO DXO

Poly LLA July 25, 2017

## Analysis of scaffolds involves – as always – the definition and delineation of the **VOI** (volume of interest) for measurement



July 25, 2017

## Auto-wrapping or "shrink-wrap" of the VOI around an irregular object boundary (SkyScan CT-Analyser)







A PLA CO CL type scaffold – imaged by the SkyScan 1172 A woven type scaffold – imaged by the SkyScan 1172

In other, calcified scaffolds, detailed analysis of the porosity is required, to assess accessibility to bone cells





Acta Biomaterialia xxx (2011) xxx-xxx



- <sup>2</sup> Influence of the pore generator on the evolution of the mechanical properties
- and the porosity and interconnectivity of a calcium phosphate cement

<sup>4</sup> Q1 M.A. Lopez-Heredia<sup>a</sup>, K. Sariibrahimoglu<sup>a,1</sup>, W. Yang<sup>a,1</sup>, M. Bohner<sup>b</sup>, D. Yamashita<sup>c</sup>, A. Kunstar<sup>d</sup>,

- 5 A.A. van Apeldoorn<sup>d</sup>, E.M. Bronkhorst<sup>a</sup>, R.P. Félix Lanao<sup>a</sup>, S.C.G. Leeuwenburgh<sup>a</sup>, K. Itatani<sup>c</sup>, F. Yang<sup>a</sup>,
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## Pore size distribution



**Fig. 6.** Mean size of the open porosity for the different CPC–PLGA basic samples. S, small particles (~20  $\mu$ m); B, big particles (~40  $\mu$ m); SB, mixture of small and big particles. Numbers next to prefixes are the weight percentages of PLGA loaded into the CPC.

The analysis used here, developed by SkyScan, assesses how far a **virtual sphere** can penetrate a scaffold





...to simulate cell ingrowth

## The sphere analysis: curve displacement to right means the furthest access into scaffold





# Porosity analysis applied to starch fiber mesh scaffold research

TISSUE ENGINEERING: Part A Volume 14, Number 00, 2008 © Mary Ann Liebert, Inc. DOI: 10.1089/ten.tea.2008.0025

> The Role of Lipase and α-Amylase in the Degradation of Starch/Poly(ε-Caprolactone) Fiber Meshes and the Osteogenic Differentiation of Cultured Marrow Stromal Cells

Ana M. Martins, B.S.,<sup>1–3</sup> Quynh P. Pham, Ph.D.,<sup>3</sup> Patricia B. Malafaya, Ph.D.,<sup>1,2</sup> Rui A. Sousa, Ph.D.,<sup>1,2</sup> Manuela E. Gomes, Ph.D.,<sup>1,2</sup> Robert M. Raphael, Ph.D.,<sup>3</sup> F. Kurtis Kasper, Ph.D.,<sup>3</sup> Rui L. Reis, Ph.D., D.Sc.,<sup>1,2</sup> and Antonios G. Mikos, Ph.D.<sup>3</sup>



FIG. 2. Starch and poly( $\epsilon$ -caprolactone) fiber mesh images obtained using micro-computed tomography before (A, G) and after degradation with  $\alpha$ -amylase (150 U/L) under static (B–F) and dynamic (H–L) conditions up to 30 days. The scale bar is 2 mm and applies to all images.

FIG. 3. Starch and poly(ε-caprolactone) fiber mesh images obtained using micro-computed tomography before (A, G) and after degradation with lipase (110U/L) under static (B–F) and dynamic (H–L) conditions up to 30 days. The scale bar is 2 mm and applies to all images.

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#### Differential effects on bone of estrogen receptor $\alpha$ and androgen receptor activation in orchidectomized adult male mice

Sofia Movérare\*, Katrien Venken<sup>†</sup>, Anna-Lena Eriksson\*, Niklas Andersson\*, Stanko Skrtic\*, Jon Wergedal<sup>‡</sup>, Subburaman Mohan<sup>‡</sup>, Phil Salmon<sup>§</sup>, Roger Bouillon<sup>†</sup>, Jan-Åke Gustafsson<sup>¶</sup>, Dirk Vanderschueren<sup>†</sup>, and Claes Ohlsson\*<sup>∥</sup>

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Edited by Elwood V. Jensen, University of Cincinnati Medical Center, Cincinnati, OH, and approved September 5, 2003 (received for review May 22, 2003)

Moverare *et al.* showed that testosterone had its own pathway, its bone protective effect was not "piggy-backing" the estrogen pathway









WT sham vehicle

#### WT ORX vehicle



WT ORX estrogen



WT ORX DHT









**DERKO** sham vehicle

DERKO ORX vehicle

DERKO ORX estrogen

**DERKO ORX DHT** 



JOURNAL OF BONE AND MINERAL RESEARCH Volume 19, Number 11, 2004 Published online on August 23, 2004; doi: 10.1359/JBMR.040819 © 2004 American Society for Bone and Mineral Research

#### Additive Protective Effects of Estrogen and Androgen Treatment on Trabecular Bone in Ovariectomized Rats

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#### Experimental protocol:

Age at surgery: 12 weeks Group numbers (n): 7-8 Duration of treatment: 6 weeks

- Sham operated
- Ovariectomised
- OVX, estradiol (E2)
- OVX, DH testosterone (DHT)
- OVX, E2 + DHT



Sham-op



OVX



Testosterone DHT)

Estradiol



Estradiol + DHT







## Mechanical loading enhances the anabolic effects of intermittent parathyroid hormone (1–34) on trabecular and cortical bone in mice

Toshihiro Sugiyama \*, Leanne K. Saxon, Gul Zaman, Alaa Moustafa, Andrew Sunters, Joanna S. Price, Lance E. Lanyon

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#### Parathyroid hormone (1-34)



#### Imaging and analysis of bone fracture healing





#### **Otsu multi-level thresholding**

effectively delineated the mineralised callus from cortical bone. Upper images: 2 weeks post fracture; lower images, 6 weeks post fracture, showing the secondary "cortical" bone forming from the periphery of the callus.

## Adaptive segmentation of the bone plus callus.



Percent callus volume







While **callus volume** and thickness decreased monotonically over the 2-6 week post-fracture period, and the nonmineralised callus (chondrocytes) likewise,

the architectural parameters

of callus connectivity and complexity showed significant change only during 4-6 weeks post-fracture.

## **Thickness histogram** is informative of structural change





#### **Fracture callus**

remodeling shows a progression toward **higher callus structure thickness**, at the expense of depletion of thinner callus structures. From 2-6 weeks post-fracture.

#### Triphosphate biomaterial scaffold, SkyScan1172



#### Biomaterial implants into rabbit calvarium, SkyScan1172


## Arthritis models: assessment of the bone involvement











Paw with CIA

Paw with CIA, treated

Periosteal reaction around the tarsals is characteristic of CIA and quantifiable

Accurate semi-automated quantication is as important as visualising arthritis-related pathology.







Otsu multi-threshold – morphological close + open - despeckle

### Parameterization of arthritic changes





### Accurate imaging and parameterisation of bone disruption in arthritis models



### Mouse ankle, control

Mouse ankle, collageninduced arthritis (CIA)

# The rabbit subchondral bone: osteoarthritis and osteoporosis





Control, left; osteoporisis + osteoarthritis, right. SkyScan1272, 3 um voxel size





Above: osteocyte lacunae from canine cortical bone, SkyScan 1172



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## Biomechanics: head-banging woodpeckers

#### OPEN a ACCESS Freely available online



## Why Do Woodpeckers Resist Head Impact Injury: A Biomechanical Investigation

#### Lizhen Wang<sup>1,2</sup>, Jason Tak-Man Cheung<sup>3</sup>, Fang Pu<sup>1</sup>, Deyu Li<sup>1</sup>, Ming Zhang<sup>2\*</sup>, Yubo Fan<sup>1\*</sup>

1 Key Laboratory for Biomechanics and Mechanobiology of Ministry of Education, School of Biological Science and Medical Engineering, Beihang University, Beijing, People's Republic of China, 2 Department of Health Technology and Informatics, the Hong Kong Polytechnic University, Hong Kong, 3 Li Ning Sports Science Research Center, Beijing, People's Republic of China



**Figure 3. Anatomical structures of head and hyoid bone.** (a) Great Spotted woodpecker's head; (b) Great Spotted Woodpecker's hyoid bone; (c) Eurasian hoopoe's hyoid bone. doi:10.1371/journal.pone.0026490.g003



# Micro-CT images (SkyScan1076) provide the input material for FEA wood-peck simulations







**Figure 2. Micro-morphology of cranial bone.** (a) The micro-CT scanning images of Great Spotted woodpecker's head on the coronal plane; (b) The micro-CT scanning images of Eurasian hoopoe's head on the coronal plane; (c) The SEM image of Great Spotted woodpecker's cranial bone; (d) The SEM image of Eurasian hoopoe's cranial bone. doi:10.1371/journal.pone.0026490.g002

**Figure 4. Micro-CT image and the FE models of Great Spotted Woodpecker' head.** (a) Micro-CT image of Great Spotted Woodpecker' head; (b) Beak<sub>Lower</sub>>Beak<sub>Upper</sub> FE model; (c) Beak<sub>Lower</sub> = Beak<sub>Upper</sub> FE model; (d) Beak<sub>Lower</sub><Beak<sub>Upper</sub> FE model. doi:10.1371/journal.pone.0026490.g004

### Stress sequence of a wood-peck





**Figure 7.** The effective stress distribution of woodpecker's head during pecking. doi:10.1371/journal.pone.0026490.g007

# Metallic implants play a major role in orthopedic bone reconstruction and repair



Journal of Periodontology; Copyright 2011

DOI: 10.1902/jop.2011.110569

### Alveolar Bone Remodeling Around Immediate Implants Placed in Accordance With the Extraction Socket Classification — A Three-Dimensional Micro-Computed Tomography Analysis

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A series of reconstructed sagital and axial micro computed tomography images illustrating the measurements of the buccal bone thickness and volume of bone around immediate implant. a.1, b.1, and c.1) sagital sections show the buccal and lingual bone thickness around the implant placed in Group-1, -2 and -3, respectively. a.2, b.2, and c.2) Axial section at the level of the crestal bone showing the different buccal bone thickness around dental implants at the crestal bone level in each Group-1, -2, and -3, respectively. (Red line shows the border of the buccal and lingual bone).









TISSUE ENGINEERING: Part A Volume 00, Number 00, 2013 © Mary Ann Liebert, Inc. DOI: 10.1089/ten.tea.2013.0181 **Original Article** 



Enhanced Bone Regeneration of Cortical Segmental Bone Defects Using Porous Titanium Scaffolds Incorporated with Colloidal Gelatin Gels for Timeand Dose-Controlled Delivery of Dual Growth Factors

Johan van der Stok, MD,<sup>1,\*</sup> Huanan Wang, MSc,<sup>2,\*</sup> Saber Amin Yavari, MSc,<sup>3</sup> Michiel Siebelt, MD,<sup>1</sup> Marjan Sandker, MD,<sup>1</sup> Jan H. Waarsing, PhD,<sup>1</sup> Jan A.N. Verhaar, MD, PhD,<sup>1</sup> Holger Jahr, PhD,<sup>1</sup> Amir A. Zadpoor, PhD,<sup>3</sup> Sander C.G. Leeuwenburgh, PhD,<sup>2</sup> and Harrie Weinans, PhD<sup>1,3,4</sup>



**FIG. 1.** Photographs of porous titanium scaffolds in the shape of the 6-mm bone segment that was replaced during the animal experiment before **(A)** and after **(B)** incorporation with colloidal gelatin gels. Micro-CT images of perpendicular **(C)** and horizontal **(D)** cross sections of porous titanium scaffold (black) incorporated with colloidal gelatin gels containing iodine-based radiographic contrast agent ioxaglate (gray). Scale bar = 1 mm. Micro-CT, micro-computed tomography.





**FIG. 4.** Representative transversal micro-CT images of the porous titanium scaffolds containing unloaded **(A)**, fibroblast growth factor-2 (FGF-2) **(B)**, bone morphogenetic protein-2 (BMP-2) **(C)**, or BMP-2/FGF-2 **(D)** gels after 12 weeks. Porous titanium scaffolds and fixation screws appear in black, whereas bone appears in dark gray. Scale bar=1 mm.



FIG. 5. Representative 3D micro-CT images of bone bridging the porous titanium scaffolds containing unloaded (A), FGF-2 (B), BMP-2 (C), or BMP-2/FGF-2 (D) gels after 12 weeks. Porous titanium scaffolds are shown in transparent, whereas bone appears in dark grey. Color images available online at www.liebertpub.com/tea

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Contrast agents can confer contrast to soft tissue, opening the development of "micro-CT histology"





Mouse embryo 15 d, PTA

Mouse brain with Microfil infiltration of blood vessels: brain (left), femur (right)



Paddlefish (tetraodon) larva, stained by phosphotungstic acid (PTA)



Muscle, stained by phosphotungstic acid (PTA)

### Cartilage staining for microCT





PTA – baseline: calcified cartilage is visible, but not nonmineralised cartilage





PTA – 24 hours – successful cartilage staining but bone partly demineralised.











Staining of the mouse joint allows accurate imaging and quantiation of cartilage in arthritis models

## Lung imaging and analysis for pulmonary research: human lung autopsy at high resolution



Cross-section: 2 micron pixel

MIP (maximum intensity projection)

## Lung imaging and analysis for pulmonary research: Mouse lung sample at high resolution



## Quantification of lung tumour vasculature destruction

### Mouse lung tumour (yellow) and lung vasculature (red)

no treatment



#### Savai R. et.al. 2009, Neoplasia, 11, 48-56

Anti-VEGF treatment



## Brain vascularization imaged by blood vessel contrast agent: mouse brain with MicroFil



Mouse brain with Microfil™ vascular infiltration Bone vascularisation imaged by blood vessel contrast agent: mouse tibia and gastrocnemius muscle arterial structure



Micro-CT in animal anatomy and developmental biology: Head of embryo of Amia fish



Micro-CT in animal anatomy and developmental biology: Head of embryo of paddlefish



Micro-CT in animal anatomy and developmental biology: Whole embryo of paddlefish (with contrast agent for soft / neural tissues)



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  - Image Co-Registration
  - Link Datasets
  - Multi Dataset Loading in CTVox
  - Multi-threading Structure Thickness and Separation
  - Multi ROIs and Add/Subtract ROI
  - o Conditional Mean Filtering
  - Watershed Separation
  - Collections Tab
  - Reference "Upside Down"
  - Exclude Top/Bottom Surface Area

### New Features in "DataViewer"

#### Image Co-Registration of Datasets



Possibility to coregister two different dataset in the 3 orthogonal directions.

Important in comparative assessments: Pre- and Post Treatments Image Co-Registration of Datasets

# Links two datasets of same size into one screen. Ideal for changing the zoom and visual comparison.



New Features in "CTVox"

Load Multiple Datasets



Possibility to load up to 10 different datasets in CTVOX.

### New Features in "CT Analyzer"

### **Thickness Calculation**



Mapping of spatially resolved local thickness in 3D, illustrated by this thickness-color image, is now 5-8 times faster in CTAn.

- 3D thickness and separation calculation is now 5-8 times faster than the previous versions.
- Slightly more acccurate
- Tb.Th and Tb.Sp results will be 1-2% different than in previous versions

### New Features in "CT Analyzer"

### Multiple ROIs and Add/Sub ROIs



Multiple ROI shapes allow both molar tooth root canals to be delineated (above), and subtractive ROI shape drawing allows a hollow ROI for cortical bone to be drawn (below).

- 3D thickness and separation calculation is now 5-8 times faster than the previous versions.
- Slightly more acccurate
- Tb.Th and Tb.Sp results will be 1-2% different than in previous versions

New Features in "CT Analyzer"

Conditional Mean Filtering



### Conditional Mean Filtering


### **Conditional Mean Filtering**



"Watershed" Separation



"Watershed" Separation









#### "Collection" Tab







Figure 6. The collections tab allows often-used task lists (e.g. "root canal") to be available for quick access.

The new, fifth tab is added at the new first tab in the custom processing page. It is titled "collections" and its purpose is to allow the user to keep a list of frequently used task lists on hand for quick and easy call-up. Right-click "edit" on a task list in the collections tab to load it to the Task list tab – which now moves to second place just to the right of the collections tab

#### "Upside Down" Function for Reference Marker

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Direct Analyti	c		-
Z-position			
Reference	635	🚢 4.6992E+001 mm	Upside-Down
Oficet:	20	1.4901E+000 mm	Apply
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The button "Upside-Down" has been added in order to simplify use of the reference level where the landmark location is above, not below, the selected range of crosssections in the "Z" (height) axis. VOI for a region of a rat distal femur for trabecular bone analysis, and bone around a metal implant, are selected relative to reference points above the analyzed region (the clear, non-colored band).

#### **Excluding Top and Bottom Surfaces**



## Peripheral object area (exclude VOI boundaries)

In some analyses you need to measure object surface area but exclude surfaces that are artificially cut in the crosssectional plane by the top and bottom VOI boundaries. An example is the analysis of bone implant contact (BIC) around an orthopedic implant surface

The 2D slice-by-slice analysis not allows both intersection surface (BIC) and object surface to be measured while excluding these artificial cut surfaces.

# **THANK YOU**