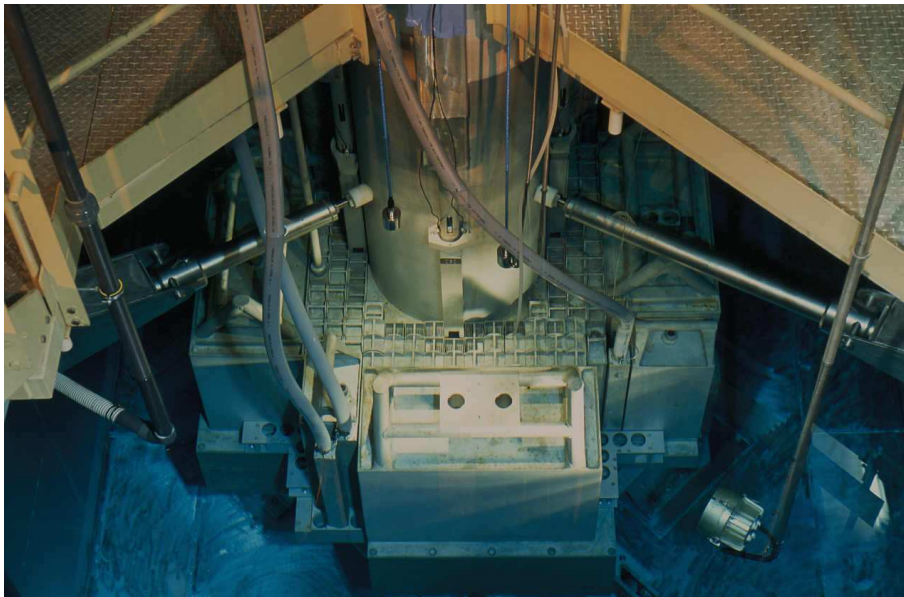




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CARMEN: an experimental configuration in the MINERVE critical facility for the qualification of neutron cross sections in epithermal spectrum



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Outline



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2. The MINERVE facility
3. Oscillation technique of measurement
4. OSMOSE and OCEAN programs
5. Conception of the CARMEN lattice
 - Main characteristics of the CARMEN configuration
 - Estimation of experimental signals
 - Optimization of the design
 - Mechanical design
6. Conclusion and perspectives

Introduction



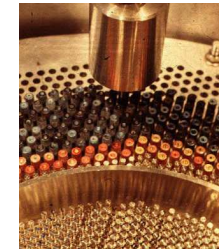
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To gain experimental data to under-moderated reactors, the

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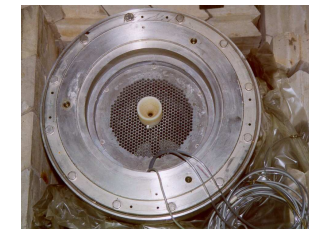
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Qualification of neutronic parameters
(ERASME program in the EOLE facility (1985))



Determination of capture rates (heavy nuclides, fission products)
(ICARE irradiations in the MELUSINE facility (1986-1988))

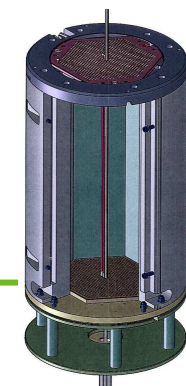
Measurement of the global capture of fission products
(oscillation of spent fuels)
(MORGANE program in the MINERVE facility (1986))



Complementary results were foreseen:

Improvement of cross sections for heavy nuclides and new neutron absorbers
(OSMOSE and OCEAN programs in the MINERVE facility)

A new configuration has been designed:
CARMEN (Core with An epitheRMAl nEutron moderation)



CADARACHE

IGORR September 19-23 2010 Knoxville, TN USA

The MINERVE facility

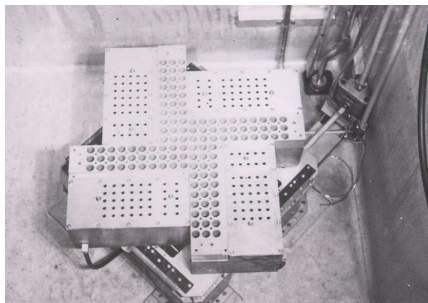


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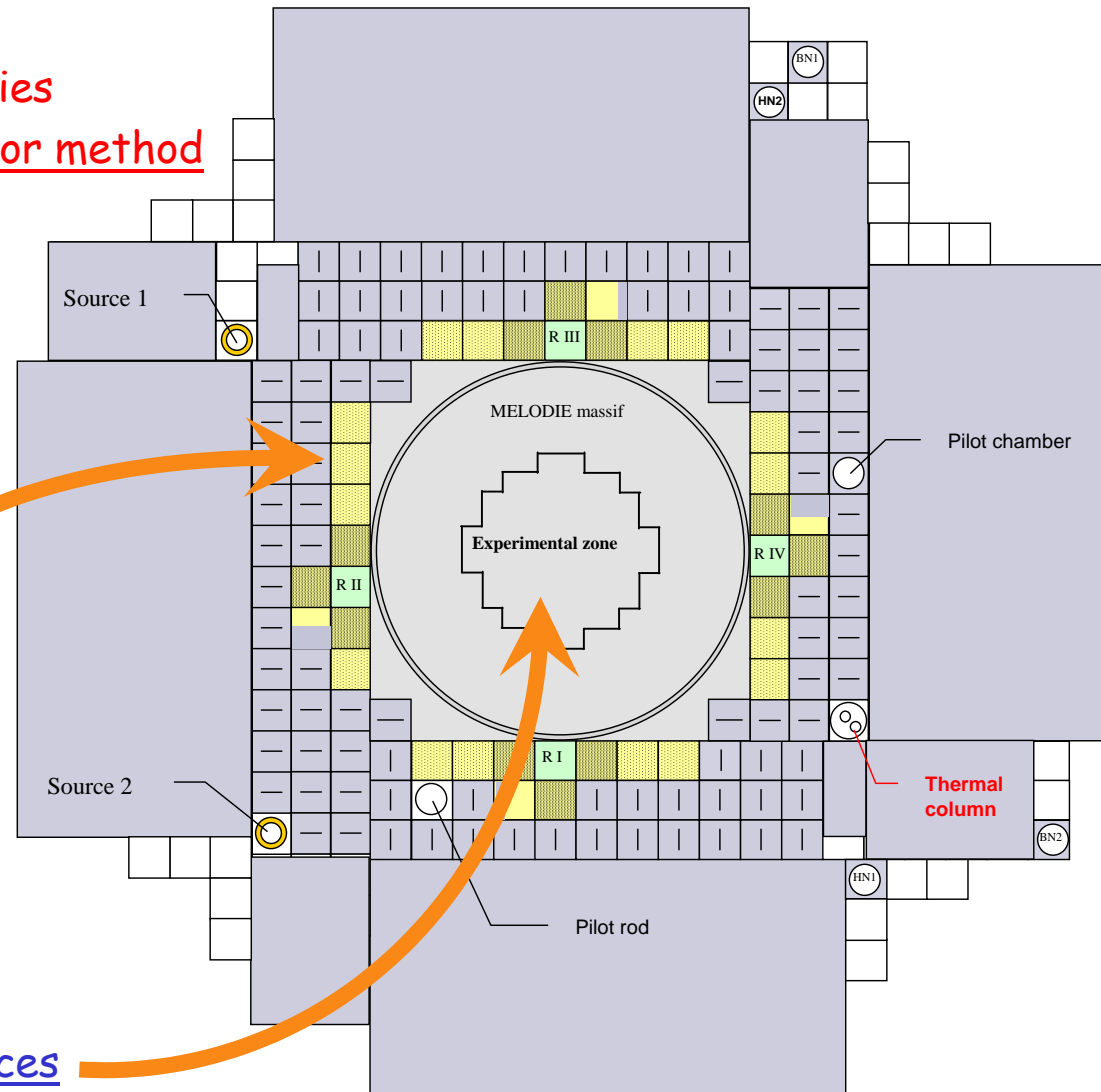
Devoted to neutronics studies
using the reactivity oscillator method

- Pool: 100 m³ of water
- Zero power (< 100 W)
- Thermal flux: 10⁷ n.cm⁻².s⁻¹.W⁻¹

- Driver zone on mobile grids

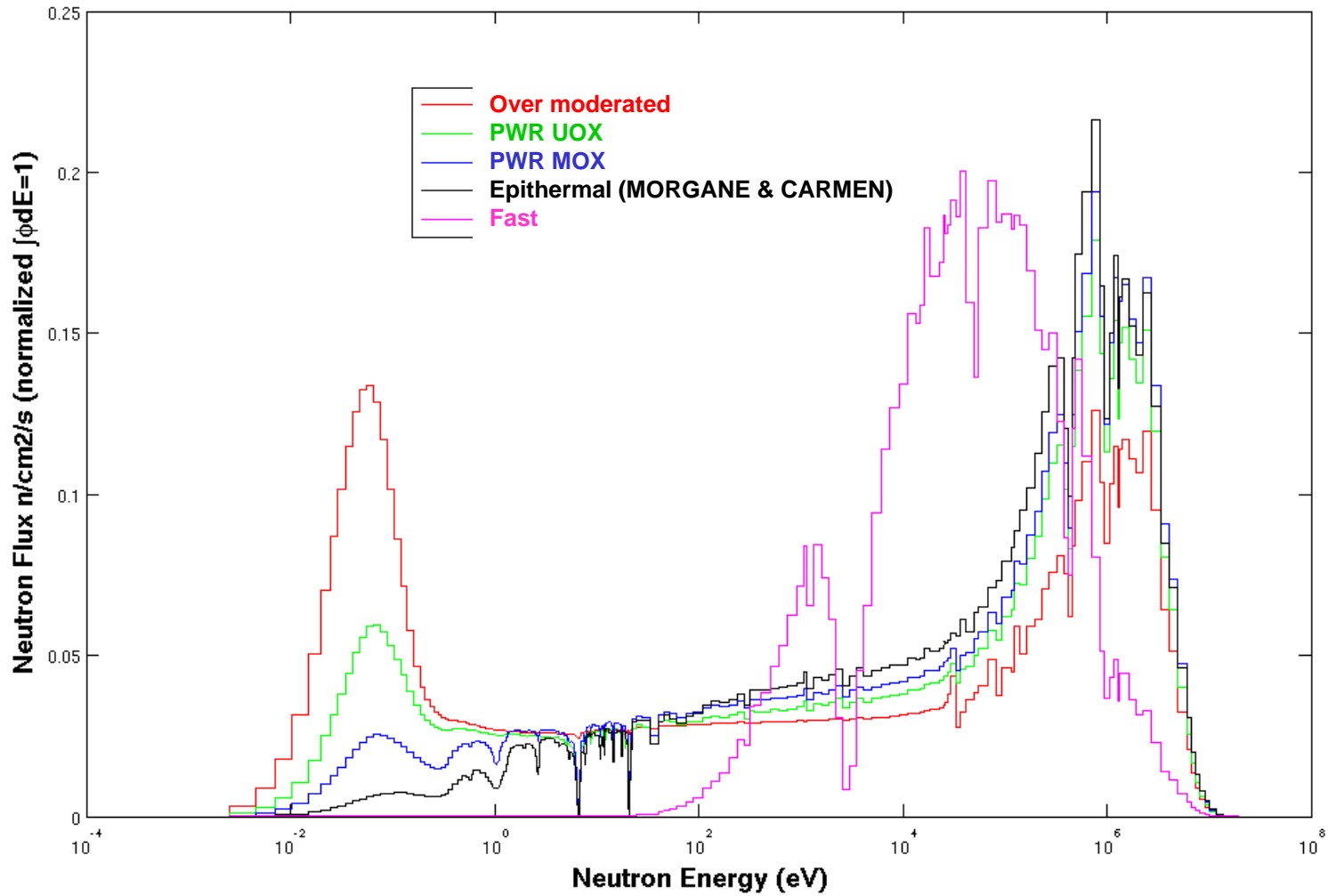


- Central cavity for experimental lattices

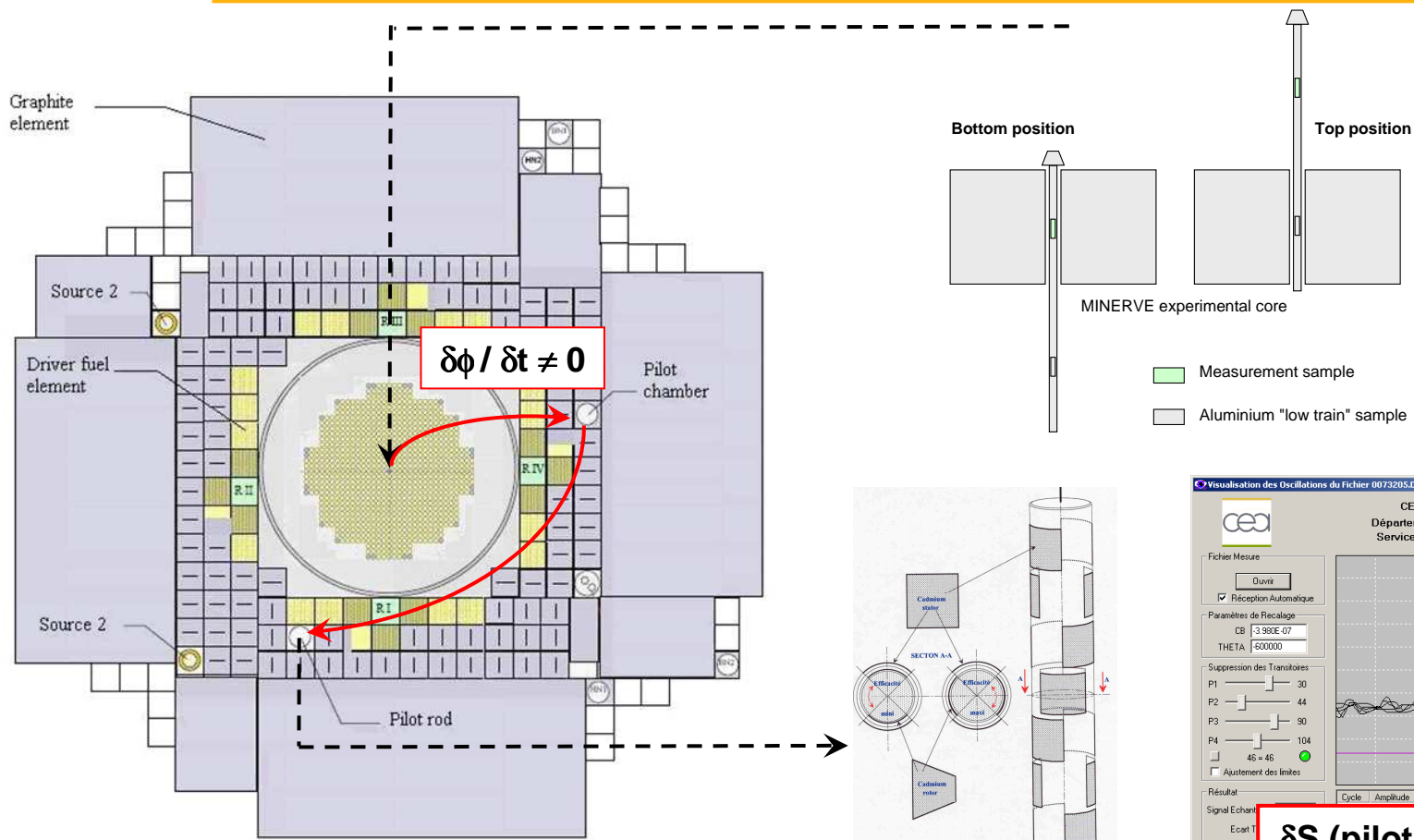


The MINERVE facility

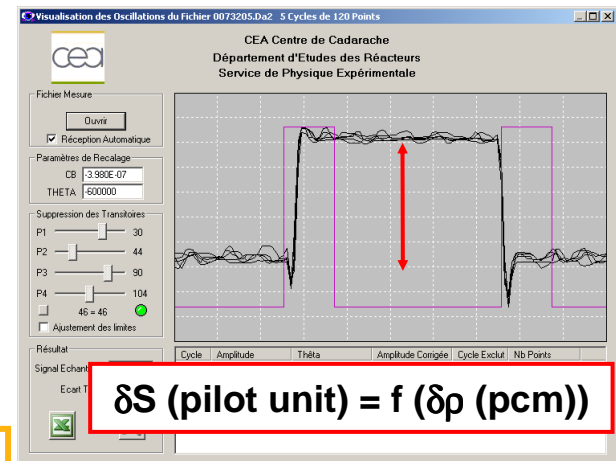
Neutronics spectra in the experimental zone :



Oscillation technique of measurement



Accuracy: about 3% for absolute reactivity worth (including the uncertainties on the material balance and on the calibration step)
 Reactivity effects of less than 2 cents can be measured



1 measurement = 5 cycles of 120 s
 1 sample = 5 measurements

The OSMOSE and OCEAN programs (2005 - 2012)

Partners:



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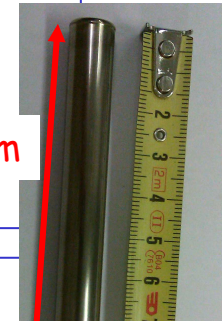


To improve the knowledge on the absorption cross sections of:

OSMOSE : Oscillation in Minerve of isotopes in "Eupractic" Spectra

<u>Actinides:</u>	Th-232	U-233	Np-237	Pu-238	Am-241	Cm-244
		U-234		Pu-239	Am-243	Cm-245
		URE		Pu-240		
				Pu-241		
				Pu-242		

L = 10.35 cm



OCEAN : Oscillation in Core of Samples of Neutron Absorbers

<u>Absorbers:</u>	Eu-151	Gd-155	Dy-160	Er-166	Hf-177
	Eu-153	Gd-157	Dy-161	Er-167	Hf-178
	Eu nat	Gd nat	Dy-162	Er-168	Hf-179
			Dy-163	Er-170	Hf-180
			Dy-164		

∅ = 1.06 cm

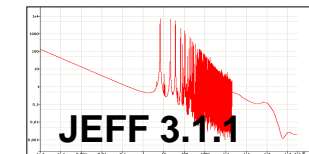


To be integrated into the library JEFF3.1.1

Since 2006 PWR UOX type spectrum (R1-UO2)

Measure in different neutron spectra for having a better decomposition in energy domains for the qualification of nuclear data

2012 Epithermal (High Conversion LWR) type spectrum (CARMEN)



Characteristics of the CARMEN configuration

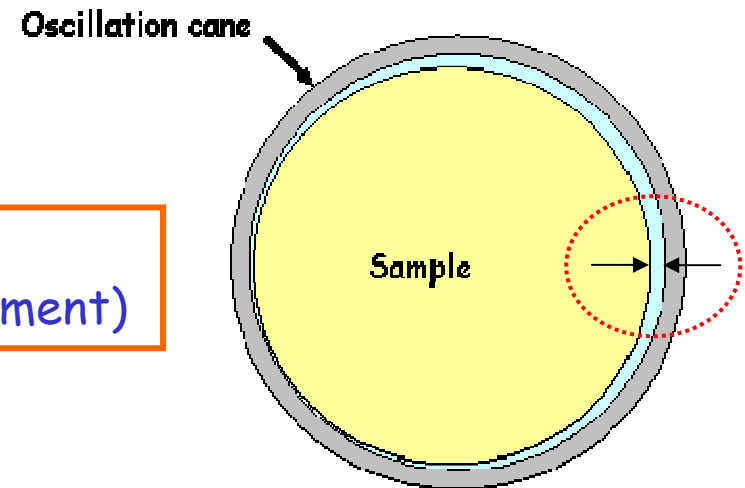
Main parameters required for the design:



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- Epithermal spectrum with a moderation ratio $V_m/V_f = 0.9$
- A high content in plutonium (representative of under-moderated concepts)
- Pins already available in the facility (7% in Pu and 3.7% in U-235)
- Several safety criteria to be respected (importance of the experimental zone compared to the driver zone)

Oscillation in a dry environment
(to improve the reproducibility of the measurement)



Estimation of experimental signals



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Experimental signals in R1-MOX lower than in R1-UO2

	R1-UO2 configuration	R1-MOX configuration	CARMEN configuration
ΔS_{H8-H1} (pilot unit)	410 400 \pm 1 000	119 600 \pm 1 000	expected signal \sim 50 000 \pm 1 000
Relative uncertainties	0.24%	0.84%	\sim 2%

To optimize relative uncertainties for CARMEN lattice, experimental signals have to be as high as possible

Whatever the experimental lattice:

$$\Delta S = \alpha^{\text{calib}} \Delta \rho$$

Estimation of experimental signals



$\alpha_{\text{CARMEN}}^{\text{calib}}$ can be estimated by a combination of:

- 3D Monte-Carlo calculations (MCNP5 code)
- 2D deterministic calculation (APOLLO2.8 code)
- results of previous measurement (R1-UO2)

Checking of this method with the well known R1-MOX lattice

	Estimation	Measurement
$\alpha_{\text{R1-MOX}}^{\text{calib}}$ (pilot unit)	815 ± 98	790 ± 15

Good agreement

Optimization of the design

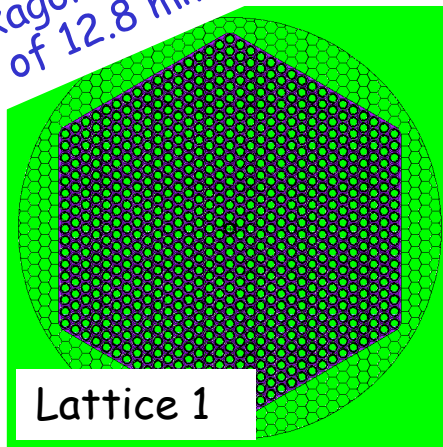


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Lattice number	Calibration factor α_{CARMEN}^{calib}	ΔS_{CARMEN} (pilot unit)
1	835 ± 89	$114\,178 \pm 12\,170$
2	1063 ± 90	$145\,355 \pm 12\,307$
3	1475 ± 103	$201\,692 \pm 14\,084$

ΔS_{R1-MOX}

Hexagonal lattice
of 12.8 mm

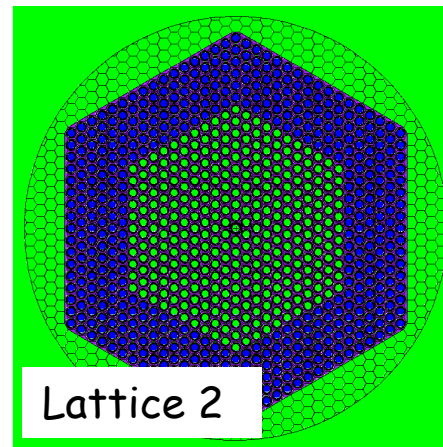


Lattice 1

Homogeneous

816 MOX 7% fuel pins

Overclad 11 mm of \varnothing_{ext}



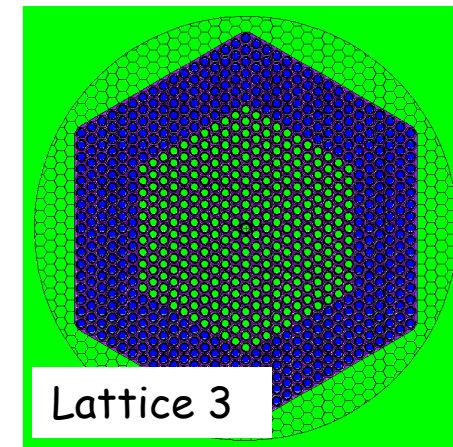
Lattice 2

Heterogeneous

330 MOX 7% fuel pins

486 UO2 (3.7% U-235) fuel pins (buffer zone)

Overclad $\varnothing_{ext}=11$ mm



Lattice 3

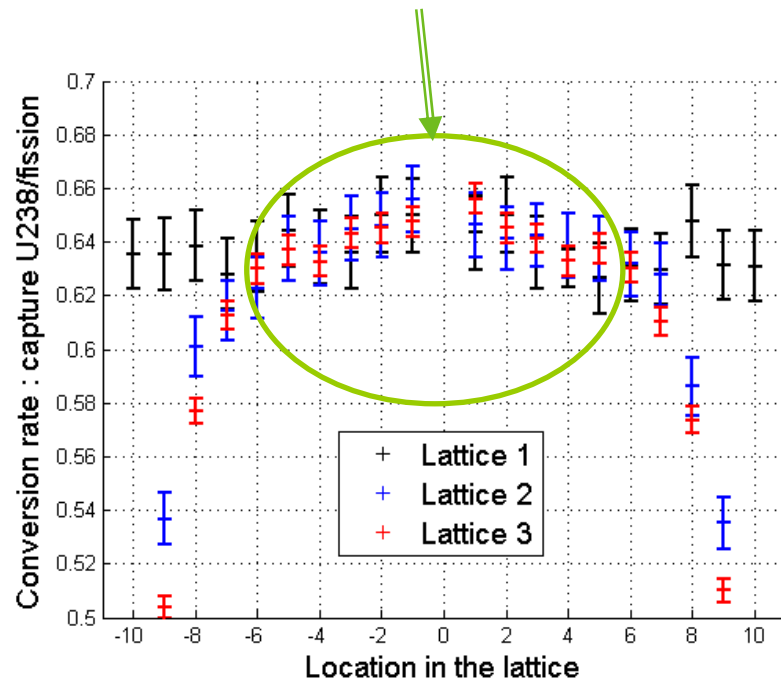
UO2 drilled overclad $\varnothing_{ext}=10.2$ mm

Optimization of the design

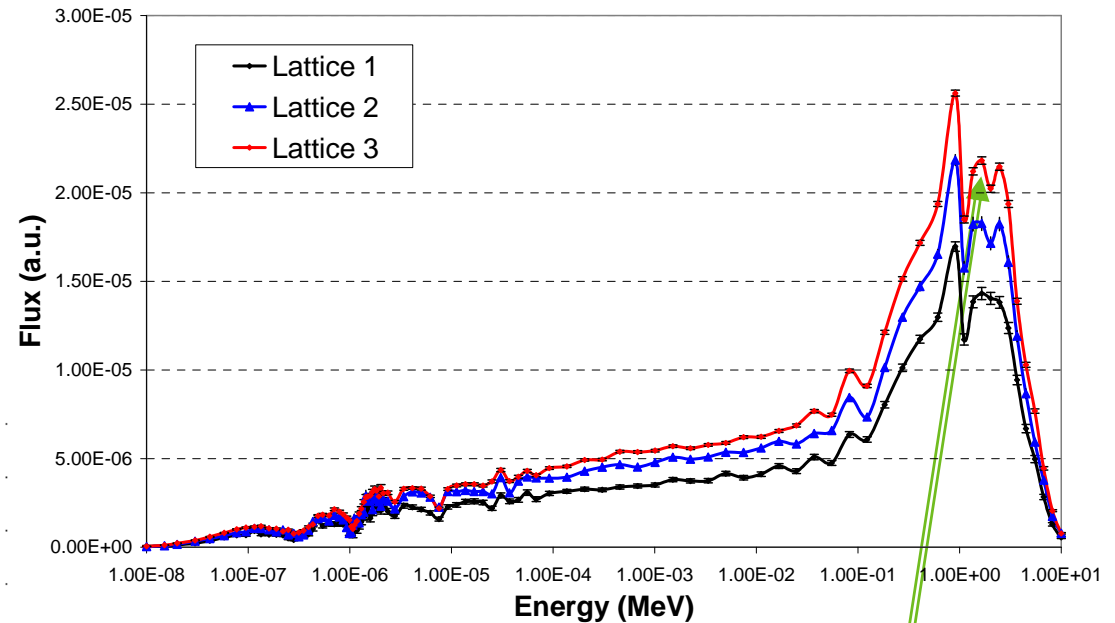


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- Same conversion ratio $\left(\frac{U_{238}}{F_{total}}\right)$ around the oscillation device



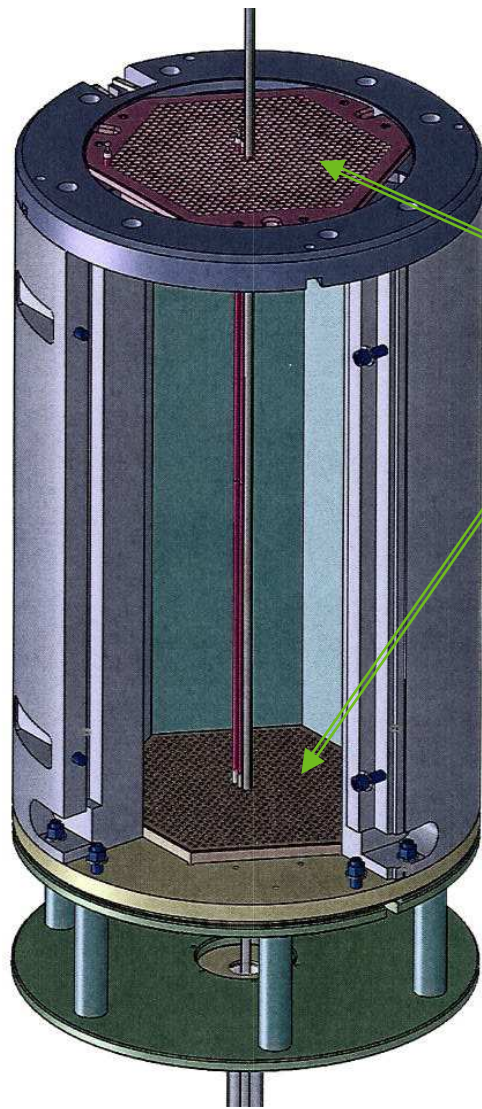
Neutron spectra in the experimental device



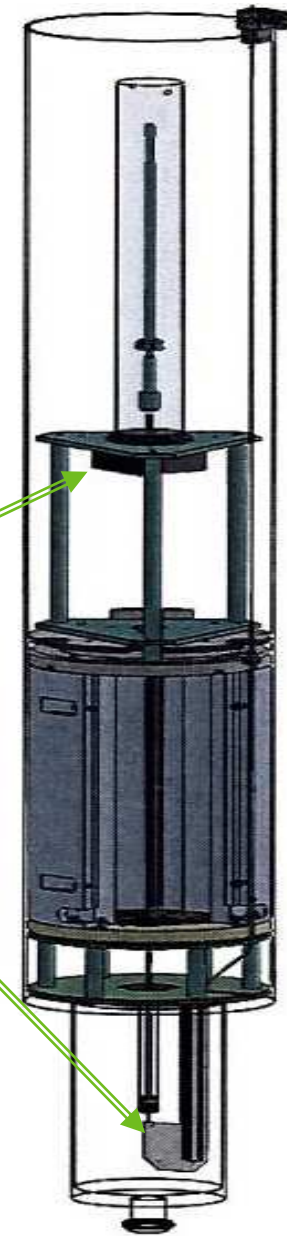
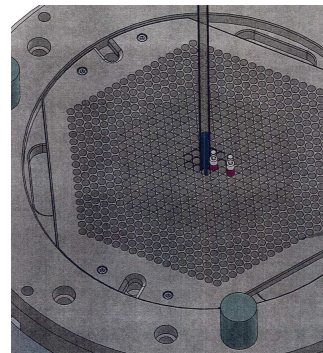
- Small increase of the flux level (lattice 3)

Third lattice will provide better results

Mechanical design



- 2 dedicated grid in an aluminum cask (versatility)
- Thick grid to drive the pins under 2 m of water
- Biological protection
- Dedicated device for extracting samples from the top of the pool



Conclusion and perspectives



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- Neutronic conception achieved
 - Mechanical building in progress
 - Reduction of experimental uncertainties
 - New calibration samples
 - Oscillations in CARMEN lattice should start in 2012
- ⇒ Improvements of nuclear data used for the JEFF3 library

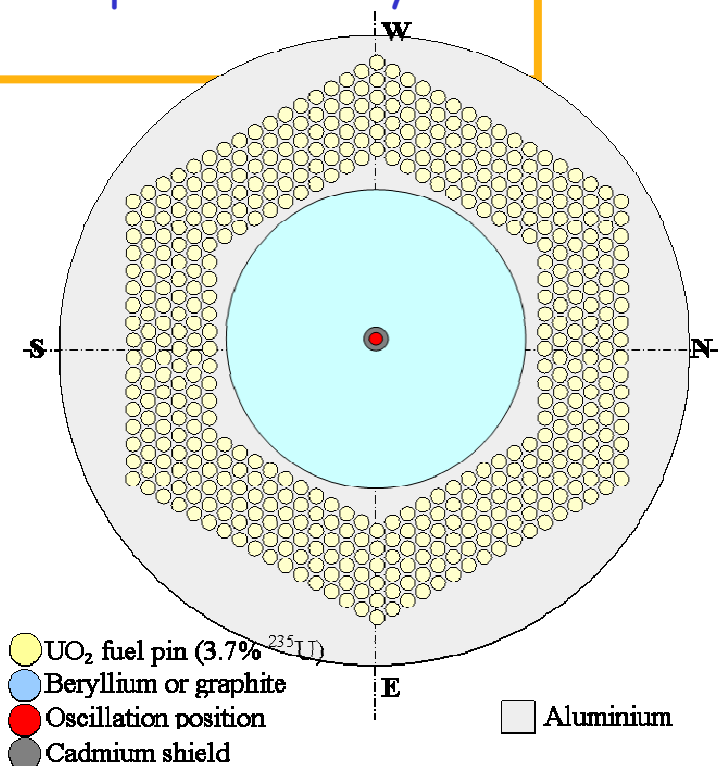
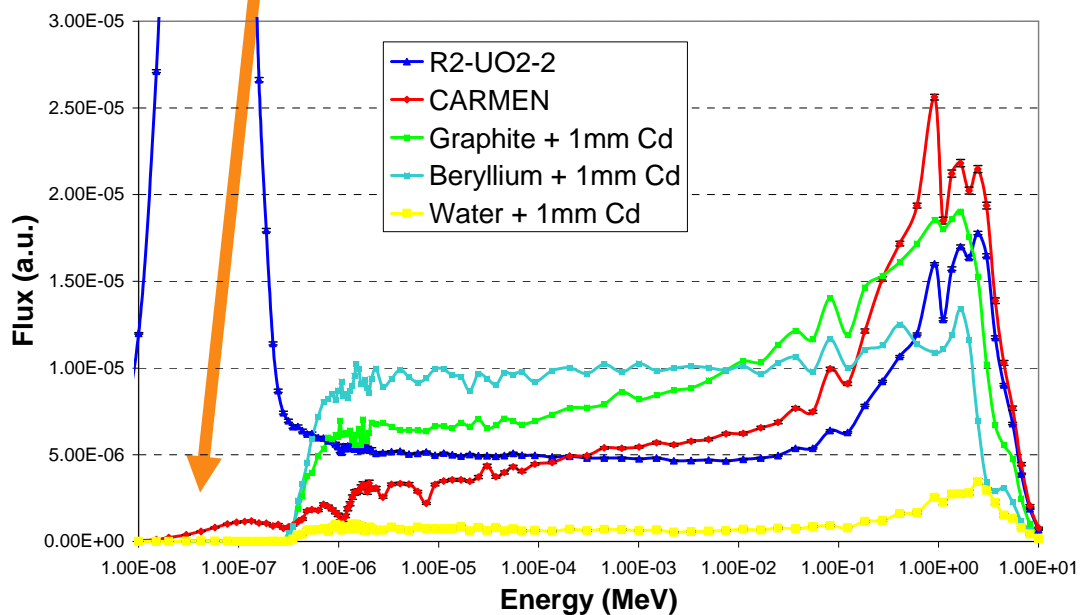
Conclusion and perspectives



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- Measurements under Cadmium shield to complete the decomposition in energy domains for the qualification of nuclear data
- MOX fuel pins can be replaced by Graphite or Beryllium cylinders

Neutron spectra in the experimental device





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Estimation of experimental signals

A reactivity effect introduced by a sample is exactly compensated by an automatic pilot rod, made of overlapping cadmium sectors:

$$\left. \begin{aligned} \Delta\rho &= \frac{\Delta N_{Cd} \int \sigma_{Cd}(E)\Phi(E)\Phi^*(E)dE}{I_f} \\ \Delta N_{Cd} &= c\Delta S \end{aligned} \right\} \Rightarrow \Delta S = \frac{1}{c} \frac{I_f}{\int \sigma_{Cd}(E)\Phi(E)\Phi^*(E)dE} \Delta\rho \quad \text{Eq. 1}$$

As the proportionality factor c depends only of the acquisition system, Eq 1 is rewritten for each core configuration:

$$\Delta S_C = \frac{\left(\int \sigma_{Cd}(E)\Phi(E)\Phi^*(E)dE \right)_R}{\left(\int \sigma_{Cd}(E)\Phi(E)\Phi^*(E)dE \right)_C} \frac{\Delta\rho_C}{\Delta\rho_R} \Delta S_R \quad \text{Eq. 2}$$

The integrals can be simplified:

the capture cross section of cadmium is essentially thermal,

and by assuming the same spectral variations for both the adjoint and direct neutron fluxes:

$$\Delta S_C = \frac{\left(\Phi_{th}^2 \right)_R}{\left(\Phi_{th}^2 \right)_C} \frac{\Delta\rho_C}{\Delta\rho_R} \Delta S_R \quad \text{Eq. 3}$$

The experimental signals in each configurations can be related to the reactivity effects calculated from 2D deterministic calculations though the same calibration process:

$$\Delta S = \alpha^{calib} \Delta\rho^{A2} \Rightarrow \alpha_C^{calib} = \frac{\Delta\rho_C}{\Delta\rho_R} \frac{\left(\Phi_{th}^2 \right)_R}{\left(\Phi_{th}^2 \right)_C} \frac{\Delta\rho_R^{A2}}{\Delta\rho_C^{A2}} \alpha_R^{calib}$$