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STLENE

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Experiment Number	Time to Peak of Pulse ^a (sec)	Sol Height at Peak Power (cm)	ution Volume at Peak Power (liter)	Rate of Reactivity Addition (dollars/sec)	Minimum Doubling Time (sec)	Inverse Period (sec ⁻¹)	Specific Energy in Pulse (10 ¹² fissions/cm ³)
CRAC 01 CRAC 02 CRAC 03 CRAC 04 CRAC 05 CRAC 06 CRAC 07	232 72 427 197 21.6 22.8 3.9	329 255 274 331 82.2 82.4 29.7	226 173 186 225 56.0 56.2 20.5	0.00341 0.00141 0.00391 0.0667 0.0740 0.786	2.9 0.18 5.0 3.2 0.060 0.050 0.00157	0.24 3.9 0.138 0.216 11.6 13.9 442	1.2 1.0 0.93 1.0 1.1 1.2 2.0
CRAC 08 CRAC 09 CRAC 10	3.1 6.4 6.6	29.3 47.1 44.0	20.3 32.3 30.2	0.746 0.247 0.0772	0.00069 0.015 0.0176	1004 46 39.4	4.0 1.4 1.4
CRAC 11 CRAC 12 CRAC 13 CRAC 14 CRAC 15 CRAC 16 CRAC 16 CRAC 17 CRAC 18 CRAC 19	65 7.5 12.0 4.0 11.2 11.7 43.8 16.2	46.3 53.3 45.4 43.6 44.1 44.3 43.8 44.9	31.8 36.5 31.1 29.9 30.3 30.4 30.1 30.8	0.0156 0.157 0.0992 0.253 0.0755 0.0820 0.0168 0.0870	0.275 0.012 0.049 0.033 0.242 0.177 0.52 0.036	2.52 57.7 14.1 20.8 2.86 3.92 1.33 19.2	1.2 1.4 1.3 1.2 1.2 1.2 1.2 1.2 1.2
CRAC 20.1 CRAC 20.2 CRAC 20.3 CRAC 20.4 CRAC 20.5	2.2 2.2 2.4 3.4 2.5	28.4 28.4 28.6 29.2 28.5	19.7 19.7 19.8 20.2 19.7	0.674 0.684 0.691 0.685 0.616	0.0061 0.0063 0.0066 0.00118 0.0058	114 110 105 587 120	1.0 1.1 1.0 2.9 1.1
CRAC 21 CRAC 22 CRAC 23 CRAC 24 CRAC 25 CRAC 25 CRAC 26 CRAC 27 CRAC 28 CRAC 29	$ \begin{array}{r} 17.0 \\ 4.6 \\ 4.6 \\ \hline 7.1 \\ 8.3 \\ 4.7 \\ 8.8 \\ 12.3 \\ \end{array} $	45.4 28.9 39.6 33.0 33.6 41.3 42.3 44.9	31.1 20.0 27.2 22.2 23.1 28.4 29.1 30.8	0.0833 0.501 0.310 0.871 0.638 0.315 0.226 0.0808	0.032 0.00147 0.0058 0.00153 0.0027 0.012 0.011 0.032	21.6 471 120 453 252 58 63 21.7	1.0 2.1 1.6 $$ 1.7 1.7 1.2 1.3 1.1

Table 8. Characteristics of the First Pulse of the Experiments in the 300-mm-diam Cylinder.



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Report SRSC nº 223- September 1994

SILENE Reactor

Results of selected typical experiment

Francis BARBRY

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SUMMARY OF REFERENCE DATA AND LIST OF THE MOST REPRESENTATIVE EXPERIMENTS

SUMMARY

This document recapitulates the most representative experimental results found on the Silene reactor in a configuration without shielding in studies on solution criticality accidents. Most of this data is relative to the nominal operating concentration of the reactor, i.e. 71 g/l, but in the appendix we have also included the results of a test series at 220 g/l as well as the results of the reactor calibration campaign.

This set of reference data can of course be used by the operator, but was initially compiled for validating computing models of accidental criticality excursions.

Two documents reviewing the lessons learned from the SILENE experiments are given in appendix.

Il y a eu quelques erreurs de pages entre le français et l'anglais lors d'un 1^{er} tirage du rapport. Pour éviter un gaspillage ces exemplaires ont été conservés pour la diffusion interne SRSC, les documents partis à l'extérieur ayant été rectifiés. Les pages inversées concernent les tableaux, symboles, ... (pages entre le texte principal et les figures illustrant les expériences SILENE)

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1 - GENERAL INFORMATION AND SILENE OPERATING PRINCIPLE

2 - THE REACTOR CORE AND ITS VARIOUS MODES OF OPERATION

2 - 1 - Description of the core

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APPENDIX 1 - Results obtained on SILENE reactor with a uranium concentration of 220 g.l-1

APPENDIX 2 - Results of the SILENE reactor calibration campaign

APPENDIX 3 - A review of the SILENE criticality excursions experiments

<u>SILENE</u>

1 - GENERAL INFORMATION AND SILENE OPERATING PRINCIPLE

Silene is a homogeneous experimental reactor using a fissile solution of uranyl nitrate (U enriched in 235 U at 92.7%) as fuel.

The core is in the form of a small annular tank located in the middle of a large concrete room referred to as a "cell". The fissile solution required for operation of the reactor is prepared in a laboratory located in the cell basement (ref. 1 153).

The general operating principle of the reactor is as follows:

- The fissile solution, previously adjusted to the Silene operation concentration in a special large capacity tank, is pumped into the core up to predetermined supercritical level. During this phase, a control rod is present in the core to avoid divergence.
- The "divergence" power excursion is then produced by removing the rod from the core using a procedure which depends on the operating mode selected for the planned experiment.
- When the experiment has been completed, the fissile solution containing the radioactive fission products is dumped into a special tank located in a shielded room, allowing rapid access into the cell.

A ventilation circuit blows continuously through the upper part of the core to dilute radiolysis gases formed. After the time necessary for their radioactive decay, these gases are removed through ad hoc filtration systems.

2 - THE REACTOR CORE AND ITS VARIOUS MODES OF OPERATION

2.1 - Description of the core

The Silene core is in the form of a 1 m high, 360 mm diameter stainless steel core with a 70 mm diameter internal channel (Figure 3

- Main dimensions

External cylinder 360/368 mm Internal channel 70/76 mm Tank bottom thickness 36 mm Cover thickness 30 mm

- Composition of the stainless steel used (grade ZZ-CN 18.10)

Isotope	Number of atoms x 10^{-24} cm ⁻³
Carbon Silicon	1.200 10 ⁻⁴ 1.709 10 ⁻³
Iron Nickel	$\begin{array}{c} 1.661 & 10^{-2} \\ 5.922 & 10^{-2} \\ 8.178 & 10^{-3} \end{array}$

2.2 - The cell

The core is located in a concrete room with dimensions of 19 x 12 x 10 m and walls 1.5 m thick (Figure 4).

Isotope	Number of atoms x 10^{-24} cm ⁻³
Hydrogen	1.035 10-2
Boron-10	1.602 10-6
Oxygen	4.347 10-2
Aluminum	1.563 10-3
Silicon	$1.417 \ 10^{-2}$
Calcium	$6.424 \ 10^{-3}$
Iron	7.621 10-4
Iron	7.621 10-4

The concrete composition is as follows:

2.3 - The fissile solution at nominal operating concentration

The fissile solution used is uranyl nitrate, which, at the nominal concentration of 71 g.1⁻¹ chosen for reactor operation, has the following mean properties:

Density	1.161 g.cm ⁻³
Total U concentration	71 g.1 ⁻¹
Acidity H+	2 N
²³⁵ U enrichment	92.7%

giving the following composition :

Isotope	Number of atoms x 10^{-24} cm ⁻³
Hydrogen	6.258 10 ⁻²
Nitrogen	$1.569 \ 10^{-3}$
Oxygen	$3.576 \ 10^{-2}$
Uranium-234	1.060 10 ⁻⁶
Uranium-235	1.686 10-4
Uranium-236	4.350 10-7
Uranium-238	1.170 10 ⁻⁵

This is a mean composition used as reference in the calculations, since it is obvious that the value of these parameters must have varied during the experiments, reprocessing and various adjustments.

The reactor calibration experiments (Ref.3) were carried out at various concentrations. A series of specific experiments was performed at a concentration of about 200 g.1⁻¹. However, the mean nominal concentration during reactor operation is 71 g.1⁻¹.

2.4 - Control rods and the corresponding control devices

Divergence in accordance with the selected procedure is initiated by rapidly or slowly displacing a rod at a predetermined speed within the central core channel. The rods are annular, allowing for the possibility of adding test capsules in the central channel which then act as an irradiation channel. There are several types of rods, wrongly called control rods, but which are actually reactivity insertion rods. The two most commonly used are made of cadmium and boron carbide. A special rod was designed for "boiling" type experiments which require reactivities of up to 5800 pcm (7.3 \$).

The reactivity values of these rods for the 71 $g.1^{-1}$ concentration are as follows:

Cadmium	rod		∆kp	<	32 2 0	pcm	(4.1	\$)
B ₄ C rod			∆kp	<	45 6 0	рст	(5.8	\$)
Special	"boiling"	rod	∆kp	<	5800	рст	(7.3	\$)

Only the "Cadmium" rod was calibrated with precision (Ref. 4) since it was the only rod authorized for high reactivity insertions ("free evolution" mode).

Figures 5 show the rate of reactivity insertion with the C. rod and the "critical" position of the rod respectively, for different heights of the fissile solution within the core.

There are two types of control mechanisms for removing the rod from the core:

. a mechanical device allowing rod displacements $\leq 20 \text{ cm}.\text{s}^{-1}$;

. a pneumatic device which can eject the rod at about 1.5 m.s^{-1} .

It should be understood that when constructed, Silene had a single mechanical device operating as follows :

- in "pulse" ejection of the rod at 20 cm.s⁻¹

- in "free evolution" rod output at 2 cm.s^{-1}

- in "steady state" rod displacements at 2 mm.s⁻¹

Due to the fact that a significant percentage of "pulses" resulted in premature initiation of the chain reaction and therefore did not allow pure reactivity "steps", it was decided two years later that a pneumatic device would be added to allow faster rod ejections to avoid any initiation of divergence before the rod was totally removed from the core. Since this period, no "pulses" have "failed".

2.5 - Operating modes

Depending on the reactivity present within the core, the rod ejection speed, and the presence or absence of an auxiliary neutron source, Silene can operate in one of three different modes called "PULSE", "FREE EVOLUTION" and "STEADY STATE".

- Operation in "Pulse" mode is obtained by ejection of the rod at high speed (0.2 or approx. 2 m.s^{-1}) with or without the presence of an auxiliary neutron source, allowing it to obtain a very high power peak in a short time (up to 1000 megawatts in a few milliseconds). This is therefore a fast transient. The reactivity in this mode of operation is limited to 2400 pcm due to a pressure wave generated in the liquid (value limited to 8 bars absolute pressure in the Silene configuration).

- "Free evolution" operation takes place by removal of the rod at low speed (< 2 cm.s⁻¹) in the presence of an auxiliary neutron source¹. The reactivity engaged under normal operation cannot exceed 4000 pcm, but under exceptional circumstances can be increased to 5800 pcm if it is required to bring the fissile solution to boiling state.

- "Steady state" operation is obtained by having the rod controlled by a control sequence. Rod displacements take place very slowly (about

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¹ Reminder of the role of the external neutron source:

When the presence of an external neutron source is indicated in the specifications of the experiment, this refers to a 100 mCi Am-Be source located at the bottom of the core tank (Figure 3). Its role is to generate a deterministic initiation of the chain reaction by increasing the initial neutron population in the system (Ref. 5). It is obvious that to obtain a high reactivity "step", it is better to operate without an external neutron source and with the highest possible rod ejection speed so that all reactivity is inserted before the first persistent neutron chain is initiated.

2 mm.s⁻¹) in the presence of an auxiliary neutron source. In this mode of operation, Silene is brought to a predetermined stable power level.

The performances of Silene in the three operating modes described above are illustrated and specified in Figure No. $\boldsymbol{\delta}$.

3 - REACTOR DIAGNOSTICS

3.1 - Total number of fissions

The main objective of an experimental reactor is to have the closest possible control over the total released energy; in our case, this means determining the total number of fissions during the experiment.

Two methods can be used for this purpose, namely gamma spectrometry on the fission products formed, and thermal balance.

- Spectrometry

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The total number of fissions carried out during an experiment can be determined by taking a sample of the fissile solution after the experiment and analyzing judiciously selected fission products by gamma spectrometry. In the case of Silene, the fission products used are generally as follows:

Radionuclide	Energy KeV	Period	% emission	Fission yield
⁹⁹ TC	140.52	66.02 h	89.4 X	6.16 X
¹³² Te	228	78.8 h	90 X	4.26 X
¹⁴³ Ce	293.1	33 h	46 X	5.91 %
¹⁰³ Ru	497.1	39.35 dy	89.7 X	3 X
⁹⁵ Zr	756.7	6,5.2 dy	54 %	6.2 %
^{9 9} Mo	739.4	66.02 h	11.82 %	6.16 %

Applied strictly, this method provides an accuracy of 5%, taking into account all statistical errors as well as errors due to the data libraries used.

- Thermal balance

It was observed on Silene that during short duration experiments (<2 minutes), the system was significantly adiabatic due to extremely low thermal exchange coefficients with the environment. Heating the fissile solution can therefore be used as a method of diagnosis provided that the temperature of the solution is uniform, in other words if there is enough mixing by radiolysis gases, and also provided that the thermal characteristics of the solution are well known.

In the case of uranyl nitrate solutions, we can establish the thermal balance by using the following relationship determined from experiments carried out in the Valduc laboratory:

 $Cp = 0.0008154 \times C_{U_{+}} = 0.06 H +$

At a concentration of 71 g.1⁻¹, C_p is approximately 0.82 Cal.g⁻¹ °C⁻¹.

3.2 - Monitoring power and radiation emission rate

In order to monitor the evolution of reactor power, different types of detectors were used (fission chambers, scintillators, ionization chambers) which were power-calibrated based on a comparison between their signal and the number of fissions resulting from the previous diagnosis.

Certain precautions must be taken, however, to avoid errors due to:

- the time-of-flight of neutrons if measuring in thermal neutrons,

- radiation diffused by the concrete walls which, in particular, could significantly distort monitoring of the sudden power drop after the first peak.

On Silene, special collimated detectors placed behind shields were designed to overcome these difficulties.

Dose-calibrated detectors were used for measuring neutron and gamma radiation dose rates (kerma rads for the neutron radiation and tissue rads for the gamma radiation).

3.3 - Routine diagnosis

Since gamma spectrometry is long and complicated, routine diagnosis of the reactor is performed using:

- 1. Activation detectors (S, Au, Cu, Mg) calibrated during calibration experiments,
- 2. Detectors to monitor the power of the reactor, whose information is sent to a data acquisition and processing system.

3.4 - Temperature and pressure measurements

Aside from the nuclear detectors mentioned above, the only nuclear instruments installed permanently on the reactor are two thermocouples (chromel-alumel) placed in the fissile solution at 20 and 30 cm from the bottom of the tank, and a pressure sensor (CEC BELL tight-wire type sensor) placed on the bottom of the core tank in the sole aim of ensuring that the pressure wave generated at the first power peak does not exceed 8 bars, the maximum limit allowed under the safety conditions of this facility.

Specific pressure measurements were also taken inside the fissile solution (Ref. $\boldsymbol{\xi}$).

3.5 - Measuring the fissile solution level

Two independent devices are used to measure the height of the fissile solution pumped into the core tank.

The first device is based on an "electromagnetic" principle and consists of making an electrical contact on the surface of the liquid. The corresponding assembly is thus installed on the upper part of the core.

There is another device at the bottom of the core tank that measures the level ultrasonically, consisting of two detectors immersed in the solution. The first detector sends a signal to the surface of the liquid and processes the information through the second detector referred to as the "reference" detector. The reference detector sends a signal to a fixed reference surface, making it possible to account for variations in wave velocity as a function of the density and temperature of the fissile solution.

4 - NEUTRON DATA FOR REACTIVITY CALCULATION WITH 71 g.1⁻¹ CONCENTRATION

4.1 - Neutron data

In order to obtain the reactivity values that appear in the result tables. we used the following neutron data determined by CEA computing codes (Apollo, Moret).

. Lifetime of prompt neutrons at 71 g/l $\overline{2}$ = 36 μ s . Delayed neutrons

GROUP	1	2	3	4	5	6
ci	1 28	1.24	1.26	1.23	1.26	1.20
ßi	0 021	0.139	0.126	0.252	0.074	0.026
li s	0 0124	0.0305	0.111	0.301	1.13	3.0

ci represents the efficiency of each group of delayed neutrons.

 $\beta_{\text{off}} = \sum_{i=1}^{6} \epsilon_i \times \beta_i = 794 \text{ pcm}$

4.2 - Reactivity calculations

In the result tables, a distinction is made between two reactivity values Δk .

 Δk_1 is defined as the reactivity present at the time of the first peak, corresponding to application of the Nordheim formula starting from the <u>minimum</u> period measured in the rise of the first peak.

$$\rho(\text{pcm}) = \frac{\frac{1}{2}/T_{\bullet}}{\frac{1}{1+\frac{1}{2}/T_{\bullet}}} \qquad \frac{1}{1+\frac{1}{2}/T_{\bullet}} \qquad \frac{i-6}{\Sigma} \qquad \frac{\epsilon_{i} \beta_{i}}{1+\lambda_{i} T_{\bullet}}$$

 Δk_p is defined as the total potential reactivity to be introduced in the core. In the case of a reactivity "step", implying an experiment where total reactivity is reached even before the first peak occurs, then $\Delta k_p = \Delta k_1$.

In the case of a reactivity ramp occurring in the presence of an external neutron source (this is often the case in "free evolution" experiments), then total reactivity is not reached in the core at the time the first peak occurs, and $\Delta k_n \neq \Delta k_1$.

For pure reactivity steps, an experimental relationship was established between Δk^p and ΔH (height of the additional solution above the delayed critical height).

For high reactivity levels (in "boiling" type experiments), which are not allowed in the "Pulse" mode in order to avoid steps that are too abrupt. the reactivity could only be calculated. Figure 8 illustrates the <u>calculated</u> evolution of k_{eff} as a function of the height of the solution The "fit" below represents this variation of k_{eff} on the Silene reacter.

Keff =
$$0.68276 + 1.194 10^{-2} \text{ x H} = 9.905 10^{-5} \text{ x H}^2$$

5 - RESULTS OBTAINED WITH THE 71 g.1⁻¹ CONCENTRATION

- Presentation of results

The most representative results of the Silene experiments are given in the form of:

- tables summarizing the main characteristics, the meaning of the symbols used being given beforehand,

- graphs illustrating the evolution in time, in experiments on the following parameters:

. power, energy

- . temperature of the fissile solution
- . pressure at the first power peak

. neutron and gamma dose rate (in general at 4 m from the source center).

To simplify presentation, however, the experiments have been classified into different catagories:

1 - experiments with slow kinetics: referring to all experiments carried out with a reactivity insertion of less than β (β being the prompt critical state \approx 794 pcm) $\rho_{CM} = /0^{-5}$, $k_{PC} = /.00794/$

2 - experiments at the prompt critical state ($\rho \neq \beta$)

3 and 4 - experiments with fast kinetics corresponding to reactivity steps $(\rho >> \beta \text{ and } \Delta k_p - \Delta k_1)$ with and without external neutron source.

5 - experiments corresponding to reactivity ramps: referring to all experiments carried out in the "free evolution" mode.

6 - "boiling" experiments. These are the experiments carried out in the "free evolution" mode but using a special reactivity rod allowing the insertion of more than 7 \$ of reactivity, thus bringing the reactor fissile solution to a boiling state. 7. Enfin nous avons isolé une dernière catégorie d'expériences pour leur intérêt sur le plan de la qualification des modèles de calcul : il s'agit d'une série d'expériences réalisées à une réactivité constante ($\Delta k_p = 2350$ pcm) mais avec des vitesses d'introduction de réactivité différentes obtenues en jouant sur la vitesse d'éjection de la barre cadmium.

Il est à noter qu'un dossier complet sur chaque expérience est archivé à Valduc et qu'il est possible d'y trouver des informations plus détaillées telles que le listing de l'évolution du nombre total de fissions en fonction du temps ou la position précise de la barre au moment du l^{er} pic de puissance.

Une synthèse des résultats de SILENE est donnée dans les références 7 et 8.

Références

Réf. 1 Brochure SILENE : document CEA/IPSN

Ref.2 F.BARBRY, "Fuel solution Criticality Accident studies with the SILENE reactor : Phenomenology, consequences and simulated intervention" International Seminar on Criticality studies programs and needs. DIJON (France) SEPT - 1983

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- Ref. 4 F. BARBRY, Etalonnage de la barre cadmium du réacteur SILENE Note technique SRSC n° 92.11, Sept. 1992
- Ref. 5 G. HANSEN, "Assembly of fissionable material in the presence of a weak neutron source", NSE 8-709-719, 1960
- Ref. 6 F. BARBRY, J.P. ROZAIN, "Formation of radiolysis gas and the appearance of a pressure increase during a criticality excursion in a fissile solution" Trans. Am. Nucl. Soc., 59, 181, (1989)
- Ref. 7 F. BARBRY, "A review of the SILENE criticality excursion experiments". Topical meeting on Physics and Methods in Criticality Safety, 34, 40, Sept 993, NASHVILLE, TN
- Ref. 8 F. BARBRY, "Dosimetry of neutron and gamma radiation in the CRAC ans SILENE criticality accident study program Rapport SRSC n° 205, 1991

Liste des tableaux de résultats des expériences SILENE

Tableau 1Expériences à cinétique lente $\Delta k \ll \beta$

Tableau 2Expériences en mode "SALVE" $\Delta k >> \beta$

 Tableau 3
 Expériences en mode "LIBRE EVOLUTION"

Tableau 4 Expérienes du type "Ebullition"

Annexe I Résultats des expériences à la concentration de solution de 220 g.l⁻¹

Annexe II Résultats des expériences de la campagne d'étalonnage du premier "coeur" de SILENE

Figures

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Fig. 1	Vue générale de l'installation
Fig. 2	Schéma de principe de fonctionnement
Fig. 3	Schéma de coeur
Fig. 4	Cellule
Fig. 5	Etalonnage de la barre cadmium
Fig. 6	Performances de SILENE dans les différents modes de fonctionnement
Fig. 7	Evolution du K _{eff} en fonction de la hauteur de solution fissile
Fig. 8	Relation entre la puissance spécifique maximale au 1^{er} pic E/V et l'inverse de la période ω

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+ Catalogue des résultats des expériences

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41.1 2 (11) 10 1 Cœur 2.Ecrans 2.Econs 3.Stockages solution avant experience 4.Stockage solution après experience 6.Cuve de confinement des gaz 6.Boite à gants de prise d'échantillons 7.Boite à gants d'extraction (retraitement) 8.Stockages des produits de fission 9.Cellule 10.Sous cellule nf4 11.Sous cellule nº3 12.Salle de contrôle 13.Saile calculateur 14.Dispositif d'exercices d'intervention







Fig:4 - SILENE Reactor diagram building







SILENE OPERATING MODES Fig 6 _







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Unshielded configuration

 $C_{Ut} \cong 71 \text{ g.l}^{-1}$

List of reference experiments

SYMBOLS USED

- CU_t Total uranium concentration (in g.l⁻¹) (93 % en ²³⁵U enriched uranium)
- H_f Final height of solution (cm)
- V_f Final volume (in liters)
- ΔH Solution addition beyond critical height
- N_f Total number of fissions
- Δk_p Total potential reactivity addition in the core

 $\Delta keff_1$ Effective reactivity present at the first peak

- $\Delta \theta$ Mean rise in temperature of the fissile solution at the end of the experiment (in °C), θ_i designating the initial temperature and θ_f the temperature reached at the equilibrium stage t_e
- T₂ Doubling time of power rise (in s)
- ω Reciprocal of period (in s⁻¹)

Λ

E Maximum power at the top of the first peak (in fissions/s)

- E Energy integrated up to the maximum power of the 1st peak
- Δp Dynamic pressure wave on the core tank bottom during the first peak (relative value in bars)
- E_{p1} Energy integrated up to the botton of the first peak (in fissions)
- E_{p2} Energy integrated during the 1st peak (in fissions)
- teq Time when equilibrim of solution temperature is reached
- E_{eq} Energy integrated up to t_{eq}

Duration : duration of the experiment

Neutron source : oui = yes no = non indicates the presence of an external neutronsource (100 mCi Am-be)



SLOW KINETICS EXPERIMENTS $\Delta k_p \leq \beta$

						1 st PEAK								
MALSER	C _{#1g} /1	H _c cm	∆H _{mm}	H _{fcm}	V _{fl}	T2,	^ω s ⁻¹	î.	É fissions s-1	É fissions	E PI Assistant	∆P _{bar}	.\.k.	AR.
LEJ-229	113	. 37.2	0.5	37.17	36.1	235	0.003	3900	1.4 1012	1.1 1015	<u> </u>	+	28	0.03
12.214	213	-37.9	1	37.85	36.8	98.8	0.007	2100	1.21013	4.3 1015			59	0.07
LEI-229	71.5	37.2	1	37.27	36.2	80	0.0087	1900	1.5 1013	5.0 1015			70	0.08
LEI-214	71.3	37.9	2	38.08	36.9	43.8	0.016	980	4.5 1013	1.0 1016			111	0.14
LE2-229	71.5	37.2	6.5	37.75	36.6	6.2	0.112	220	7.1 1014	1.9 1016			333	0.42
LEJ-214	71.3	37.9	7	38.56	37.4	4.2	0.165	140	1.2 1015	1.8 1016			390	0.49
S1-300	70.5	37.36	8.0	38.16	37.2	3.8	0.182	110	1.3 1015	2.2 1016			405	0.51
S1-J29	71.9	37.17	13.3	38.49	37.34	0.430	1.6	14.5	8.4 10 ¹⁵	1.6 1016	4.8 1016	<u> </u>	682	0.86
\$2-346	70.9	36.82	13.3	38.15	37.00	0.345	2.0	12.8	8.7 1015	1.7 1016	3.7 1016		700	0.88
\$2.300	70.8 .	37.36	15.5	38.91	37.74	0.130	5.3	9.5	1.7 1016	1.3 1016	4.2 1016	┼┼	769	0.97
\$1-258	70.5	37.74	15.5	39.29	38.11	0.110	6.3	6.3	1.9 1016	9.9 1015	3.9 1016	<u> </u>	779	0.98

			TEMPI	ERATURE	To	tal]
NUMBER	⁽ eq,	E Cl Junior	θί•ε	^{ک0} ° د	^{Ak} p pcm	^{At} _{PS}	DURATION	N fissions	SOURCE NEUTRONS	CATEGORY	f-/l
LE3-229	14000	3.1 1015	20	1	28	0.035	14400	3.1 1015	OUI	1	8.5953
LE2-214	9000	1.1 1016	19	2.2	59	0.074	16200	1.1 1016	OUI	1	2,4464
LE1-229	2800	1.2 1016	20	2.5	70	0.088	2880	1.2 1016	OUI	1	3,31E1-
LE1-214	8000	2.0 1016	19	4	111	0.14	16200	2.4 1016	OUI	1	6.5
LE2-229	6000	5.6 1016	20	11.5	333	0.42	14400	7.5 1016	OUI	1	I. 35 316
LE3-214	780	6.3 1016	19	13.5	390	0.49	780	6.3 1016	OUI	1	
S1-300	1000	6.0 1016	22.2	13.7	405	0.51	1100	6.5 1016	OUI	1	
\$1-329	37	6.0 1016	21.0	17.0	682	0.98	62	7.7 1016	OUI	2	
\$2-346	60	6.4 1016	19.0	19.0	700	0.88	170	8.8 1016	ουι	2	Real in the
\$2-300	60	7.0 1016	21.7	21.7	769	0.97	265	1.1 1017	NON	2	2,9151
\$1-258	60	7.7 1016	21.4	21.4	779	0.98	380	1.1 1017	NON	2	

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 $\overline{C}_{3} = 71,15$; $\overline{H}_{c} = 37,37$ $\overline{V}_{s} =$

	SILENE
"PULSE"	OPERATION EXPERIMENTS $\Delta k >> \beta$

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Table -2-

	·			}	I ST PEAK										··
NUMBER	^C utg/1	^H °cm	∆H _{mm}	^H f _{cm}	V _{f1}	Т _Э ,	ωs-1	t,	Ecusiona 1-1	É fissions	<i>E p</i> ₁	E _{p1}	^{∆P} bar	∆k ₁ _{PCM}	^{Δł} 1s
S2-258	70.5	37.74	21.0	39.84	38.6	0.010	68.6	1.526	7.1 1017	1.8 1018	4.5 1016	4.7 1016	<u> </u>	1046	1.32
S3-258	70.5	37.74	28.1	40.55	39.3	0.0038	181	1.432	4.9 1018	5.4 1016	8.1 1016	8.4 1016	1	1458	1.84
S1-362	69.9	37.57	46.0	42.17	40.9	0.00391	177	3.405	5.3 1018	5.6 1016	8.5 1016	8.5 1016	1.05	1420	1.79
S4-258	70.5	37.74	32.0	40.94	39.7	0.0028	249	1.798	9.1 1018	7.0 1016	1.0 1017	1.0 1017	1.9	1696	2.14
S3-300	70.8	37.36	36.0	40.96	39.7	0.00243	285	1.314	1.1 1019	6.6 1017	1.0 1017	1.0 1017	2.55	1832	2.31
S2-259	70.5	37.38	40.0	41.38	40.1	0.00198	350	1.238	1.7 1019	8.2 1016	1.3 1017	1.3 1017	3.7	2068	2.6
S4-259	70.5	37.38	44.0	41.78	40.5	0.00192	361	1.193	1.8 1019	8.6 1016	1.3 1017	1.3 1017	4.4	2108	2.65
\$3-259	70.5	37.38	44.0	41.78	40.5	0.00171	405	1.365	2.1 1019	7.9 1016	1.4 1017	1.4 1017	4.8	2269	2.86
S4-346	70.9	36.81	46.0	41.41	40.2	0.00162	428	1.234	2.4 1019	9.8 1016	1.5 1017	1.5 1017	5.6	2350	2.96
S1-346	70.9	36.82	46.0	41.42	40.2	0.00162	428	1.170	2.5 1019	9.8 1016	1.5 1017	1.5 1017	5.6	2350	2.96

			TEMP	ERATURE	TOTAL REACT	IVITY ADDITION		1	1			י ר
NUMBER	teq,	E _{CQ Justines}	θι.,	ΔΘ. c	^{∆k} _P _{pcm}	At PS	DURATION	N _f fissions	SOURCE	OBJECTIVES	CATEGORY	
S2-258	50	8.8 1016	20.4	28.3	1046	1.32	11220	1.5 1017	non	Cinétique +rentrée de la barre à t = 7 min	3	
S3-258	90	1.42 1017	20.4	35.7	1458	1.84	420	1.9 1017	non	Cinétique	3	4
S1-362	160	2.6 1017	20	50	2350	2.96	210	2.8 1017	oui	Salve avec source éjection barre Cd : 20 cm.s ⁻¹	4.7	4
S4-258	90	1.7 1017	20.5	40.2	1696	2.14	420	2.3 1017	лоп	Cinétique	3	
S3-300	· 80	1.8 1017	22.3	39.9	1832	2.31	120	2.1 1017	non	Cinétique	3	
S2-259	90	2.1 1017	20.6	47.4	2068	2.6	420	2.7 1017	non	Cinétique	3	
S4-259	90	2.2 1017	20.8	50.9	2108	2.65	10920	2.9 1017	oui	Cinétique + rentrée de la barre à t = 7 min	4	
\$3-259	70	2.1 1017	20.7	51.1	2269	2.86	420	2.9 1017	non	Cinétique	3	T tal
S4-346	180	2.9 1017	21.5	53	2350	2.96	180	2.9 1017	non	Ejection barre Cd rapide 1.5 m.s ⁻¹	3.7	5 21E15
S1-346	180	2.9 1017	21	53	2350	2.96	180	2.9 1017	oui	Ejection barre Cd rapide 1.5 m.s ⁻¹	4.7	1,

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"FREE EVOLUTION" MODE EXPERIMENTS

	Ι	[ſ			Ι		<u> </u>								
NUMBER	C _u _{Ig} /1	H _c cm	∆H _{mm}	H _{fcm}	v _{fl}	T2,	^ω s ⁻¹	• Efizsions , s ⁻¹	Êfissions	E _p	E _{p1}	∆P _{bar}	$\Delta k_{i_{PCM}}$	^{Δk} 1s	t eq,	E _{eq fissions}
LE1-362	69.9	37.57	46.0	42.17	40.90	0.136	5.1	2.0 10 ¹⁶	9.6 10 ¹⁵	4.1 1016	4.4 10 ¹⁶		766	0.95	300	2.9 1017
LE2-362	69.9	37.57	46.0	42.17	40.90	0.0245	28.3	1.8 10 ¹⁷	1.2 10 ¹⁶	4.5 1016	4.7 10 ¹⁶		890	1.12	160	2.6 1017
LE1-311	70.5	37.27	60.0	43.27	42.00	0.38	1.8	1.3 1010	2.0 10 ¹⁶	3.7 1016	4.6 10 ¹⁶	1	693	0.87	600	4.0 1017
LE1-258	70.5	37.74	60.0	43.74	42.43	0.0239	29.0	1.8 10 ¹⁷	1.1 10 ¹⁶	4.2 1016	4.6 1016		893	1.12	150	3.0 1017
LE1-273	70.7	37.54	62.0	43.74	42.43	0.025	27.7	1.7 1017	1.1 1016	4.4 10 ¹⁶	4.6 10 ¹⁶		890	1.12	140	3.0 1017

TEMPERA		ERATURE	TOT REACT	TAL TIVITY					I	Ţ
NUMBER	Θ _{i °c}	ΔΘ° c _{max}	$\Delta k_{p_{pcm}}$	^{Δk} _p s	DURATION	URATION N _{f flssions} Source NEUTRONS PARTICULARITY		CATEGORY		
LE1-362	20	52	2350	2.96	700	3.1 10 ¹⁷	OUI	vitesse sortie barre Cd 2 mm/s	5.7	
LE2-362	20	50	2350	2.96	210	2.8 10 ¹⁷	OUI	vitesse sortie barre Cd 9.5 mm/s	5.7	
LE1-311	20.0	63.4	2770	3.50	660	4.0 10 ¹⁷	OUI	vitesse sortie barre Cd 2.7 mm/s	5	
LE1-258	20.2 .	62.9	2720	3.42	500	3.9 1017	OUI	vitesse sortie barre Cd 1 cm/s	5.	
LE1-273	21	63	2820	3.55	1300	4.6 10 ¹⁷	OUI	Exercice Intervention/Phase post-accidentelle + redémarrage	5	- 1:08E1

Table -3-

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"BOILING" TYPE EXPERIMENTS

							1 st PEAK										
NUMBER	C _u tg/l	H _c cm	∆H _{mm}	^H f _{cm}	v _{fl}	T25	^ω s ⁻¹	Éfissions s ⁻¹	Ê _{fissions}	E P1 fissions	E P1 fissions	ΔP_{bar}	$\Delta k_{1_{PCM}}$	^{∆k} ¹s			
LE1-175	71.4	36.79	90	45.79	44.41	0.015	46	4.2 1017	1.7 1017	4.2 1016	4.6 1016		960	1.21			
LE1-176	69.8	38.07	114.9	49.56	48.07	0.016	43	4.5 1017	1.8 1016	3.7 1016	5.1 1016		950	1.20			
LE2-176	69.8	38.07	139.9	52.06	50.49	0.018	38	4.1 1017	1.7 1016	3.7 1016	5.6 1016		930	1.17			
LE2-343	70.6	37.1	140	51.1	49.57	0.016	42.5	3.8 1017	1.6 1016	3.2 1016	5.1 1016		950	1.20			
LE1-281	70.9	37.42	170	54.42	52.79	0.017	41	4.2 1017	1.7 1016	3.8 1016	5.4 1016		940	1.18			

	Temperature ^Θ ^j ° c ΔΘ° c _{mex}		TOT REACT	'AL IVITY				CATEGORY	
NUMBER			∆k _p pcm	∆k _{ps}	DURATION	N f fissions	SOURCE		
LE1-175	19.6		4000	5.0	540	5.4 1017	OUI		
LE1-176	18.8		4100	5.2	720	6.9 10 ¹⁷	OUI	5.6	
LE2-176	18.8		4800	6.0	900	7.4 1017	OUI	5.6	
LE2-343	22.4		5100	6.4	700	8.7 1017	OUI	5.6	
LE1-281	21		5700	7.2	600	8.6 1017	OUI	5.6	

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SLOW KINETICS EXPERIMENTS $\Delta k_p \leq \beta$

	Γ			A.12	All	AH						1st]	PEAK	<u> </u>			
NAGEER	C _{#Ig} /I	H _c cm	AH man	H f _{cm}	<i>v_{J1}</i>	72 ₅	°°s⁻i	î,	E fissions s-1	É fissions	E _{P2 feature}	^{∆P} bar	AR IPCN	ΔA1.			
LE3-229	71.5	37.2	0.5	37.17	36.1	235	0.003	3900	1.4 1012	1.1 1015			28	0.03			
LE2-214	71.3	37.9	1	37.85	36.8	98.8	0.007	2100	1.21013	4.3 1015			59	0.07			
LEI-229	71.5	37.2	-1	37.27	36.2	80	0.0087	1900	1.5 1013	5.0 1015			70	0.08			
LEI-214	71.3	37.9	2	38.08	36.9	43.8	0.016	980	4.5 1013	1.0 1016			111	0.14			
LE2-229	71.5	37.2	6.5	37.75	36.6	6.2	0.112	220	7.1 1014	1.9 1016			333	0.42			
LE3-214	71.3	37.9	7	38.56	37.4	4.2	0.165	140	1.2 1015	1.8 1016			390	0.49			
SI-300	70.8	37.36	8.0	38.16	37.2	3.8	0.182	110	1.3 1015	2.2 1016			405	0.51			
\$1-329	71.9	37.17	13.3	38.49	37.34	0.430	1.6	14.5	8.4 1015	1.6 1016	4.8 1016		682	0.86			
\$2-346	70.9	36.82	13.3	38.15	37.00	0.345	2.0	12.8	8.7 1015	1.7 1016	3.7 1016		700	0.88			
\$2-300	70.8	37.36	15.5	38.91	37.74	0.130	5.3	9.5	1.7 1016	1.3 1016	4.2 1016		769	0.97			
\$1-258	70.5	37.74	15.5	39.29	38.11	0.110	6.3	6.3	1.9 1016	9.9 1015	3.9 10 ¹⁶		779	0.98			

			TEMPI	RATURE	To	tal tivity				
NUMBER	^l eq;	E _{eff parties}	θί.ε	^{∆⊖} °	At post	^{Δ4} P ₅	DURATION	N fasions	SOURCE NEUTRONS	CATEGORY
LE3-229	14000	3.1 1015	20	1	28	0.035	14400	3.1 1015	OUI	1
LE2-214	9000	1.1 1016	19	2.2	59	0.074	16200	1.1 1016	OUI	1
LE1-229	2800	1.2 1016	20	2.5	70	0.088	2880	1.2 1016	OUI	1
LEI-214	8000	2.0 1016	19	4	111	0.14	16200	2.4 1016	OUI	1
LE2-229	6000	5.6 1016	20	11.5	333	0.42	14400	7.5 1016	OUI	1
LE3-214	780	6.3 1016	19	13.5	390	0.49	780	6.3 1016	OUI	1
\$1-300	1000	6.0 1016	22.2	13.7	405	0.51	1100	6.5 1016	ουι	1
\$1-329	37	6.0 1016	21.0	17.0	682	0.98	62	7.7 1016	OUI	2
\$2-346	60	6.4 1016	19.0	19.0	700	0.88	170	8.8 1016	OUI	2
\$2-300	60	7.0 1016	21.7	21.7	769	0.97	265	1.1 1017	NON	2
\$1-258	60	7.7 1016	21.4	21.4	779	0.98	380	1.1 1017	NON	2


Evolution de la puissance É, de l'énergie E et de la température solution.

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- SILENE 71 g/l configuration sans écran. Débits de dose neutrons et gamma à 4 m de l'axe du réacteur.















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Puissance, énergie et température

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Puissance, énergie et température (détail ler pic)



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: PUISSANCE (FISSIONS/SEC.) TEMPERATURE (C) 1.8 SILENE S1.258 ENERDIE PUISSANCE TEMPERATURE 4 0 ... TEMPS (SEC.) EVOLUTION DE LA PUISSANCE, DE L'ENERGIE ET DE SILENE LA TEMPERATURE AS SE . **n** · · · + ·





NUMBER .		H can	∆H _{mm}	H f _{cm}	v _{fl}	I ^M PTAK									
						T2,	⁶⁰ s ⁻¹	t,	Étuine s'	É fissions	E _{P1}	E _{P1}	^{∆P} bar	∆k _i , _{rov}	^{∆k} is
S2-258	70.5	37.74	21.0	39.84	38.6	0.010	68.6	1.526	7.1 1017	1.8 1016	4.5 1016	4.7 1016		1046	1.32
S3-258	70.5	37.74	28.1	40.55	39.3	0.0038	181	1.432	4.9 1018	5.4 1016	8.1 10 ¹⁶	8.4 1016	1	1458	1.84
S1-362	69.9	37.57	46.0	42.17	40.9	0.00391	177	3.405	5.3 1018	5.6 1016	8.5 1016	8.5 1016	1.05	1420	1.79
S4-258	70.5	37.74	32.0	40.94	39.7	0.0028	249	1.798	9.1 1018	7.0 1016	1.0 1017	1.0 1017	1.9	1696	2.14
S3-300	70.8	37.36	36.0	40.96	39.7	0.00243	285	1.314	1.1 1019	6.6 1017	1.0 1017	1.0 1017	2.55	1832	2.31
S2-259	70.5	37.38	40.0	41.38	40.1	0.00198	350	1.238	1.7 1019	8.2 1016	1.3 1017	1.3 1017	3.7	2068	2.6
S4-259	70.5	37.38	44.0	41.78	40.5	0.00192	361	1.193	1.8 1019	8.6 1018	1.3 1017	1.3 1017	4.4	2108	2.65
S3-259	70.5.	37.38	44.0	41.78	40.5	0.00171	405	1.365	2.1 1019	7.9 1016	1.4 1017	1.4 1017	4.8	2269	2.86
S4-346	70.9	36.81	46.0	41.41	40.2	0.00162	428	1.234	2.4 1019	9.8 1016	1.5 1017	1.5 1017	5.6	2350	2.96
S1-346	70.9	36.82	46.0	41.42	40.2	0.00162	428	1.170	2.5 1019	9.8 1016	1.5 1017	1.5 1017	5.6	2350	2.96

1			TEMPI	RATURE	TOTAL REACTIVITY ADDITION			T	1	1	T
NUMBER	leq;	E _{el} _{Autor}	^θ /•e ΔΘ•e _{max}		Ak p cm Ak ps		DURATION	N _f fissions	SOURCE	OBJECTIVES	CATEGORY
S2-258	50	8.8 1016	20.4	28.3	1046	1.32	11220	1.5 1017	non	Cinétique +rentrée de la barre à t = 7 min	3
S3-258	90	1.42 1017	20.4	35.7	1458	1.84	420	1.9 1017	non	Cinétique	3
S1-362	160	2.6 1017	20	50	2350	2.96	210	2.8 1017	oui	Salve avec source éjection barre Cd : 20 cm.s-1	4.7
S4-258	90	1.7 1017	20.5	40.2	1696	2.14	420	2.3 1017	non	Cinétique	3
S3-300	· 80	1.8 1017	22.3	39.9	1832	2.31	120	2.1 1017	non	Cinétique	3
S2-259	90	2.1 1017	20.6	47.4	2068	2.6	420	2.7 1017	non	Cinétique	3
S4-259	90	2.2 1017	20.8	50.9	2108	2.65	10920	2.9 1017	oui	Cinétique + rentrée de la barre à t = 7 min	4
S3-259	70	2.1 1017	20.7	51.1	2269	2.86	420	2.9 1017	non	Cinétique	3
S4-346	180	2.9 1017	21.5	53	2350	2.96	180	2.9 1017	non	Ejection barre Cd rapide 1.5 m.s ⁻¹	3.7
S1-346	180	2.9 1017	21	53	2350	2.96	180	2.9 1017	oui	Ejection barre Cd rapide 1.5 m.s ⁻¹	4.7

"FULSE" OPERATION EXPERIMENTS AX >> \$




































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SILENE

EXPERIENCES DU TYPES "EBULITION"

		·				1er PIC								
EXPERIENCE SILENE	C _u tg/l	H _{ccm}	∆H _{mm}	H _{fcm}	V _{f1}	<i>T</i> 2 _s	^ω s ⁻¹	• Efissions s ⁻¹	Ê _{fissions}	E Pi _{fissions}	E . P2 fissions	∆P _{bar}	∆k _{ipcm}	۵¢1s
LE1-175	71.4	36.79	90	45.79	44.41	0.015	46	4.2 10 ¹⁷	1.7 1017	4.2 1016	4.6 1016		960	1.21
LE1-176	69.8 ⁻	38.07	114.9	49.56	48.07	0.016	43	4.5 10 ¹⁷	1.8 1016	3.7 1016	5.1 1016		950	1.20
LE2-176	69.8	38.07	139.9	52.06	50.49	0.018	38	4.1 1017	1.7 1016	3.7 1016	5.6 1016		930	1.17
LE2-343	70.6	37.1	140	51.1	49.57	0.016	42.5	3.8 1017	1.6 10 ¹⁶	3.2 1016	5.1 1016		950	1.20
LE1-281	70.9	37.42	170	54.42	52.79	0.017	41	4.2 10 ¹⁷	1.7 1016	3.8 1016	5.4 1016		940	1.18

	Темі	PERATURE	REACT POTENT	IVITE MELLE				
EXPERIENCE SILENE	Θ _{i•c}	ΔΘ° c _{max}	^{∆k} _p _{pcm}	∆k _{ps}	DUREE S	N _f fissions	SOURCE NEUTRONS	CATEGORIE
LE1-175	19.6		4000	5.0	540	5.4 10 ¹⁷	OUI	
LE1-176	18.8		4100	5.2	720	6.9 1017	OUI	5.6
LE2-176	18.8		4800	6.0	900	7.4 1017	OUI	5.6
LE2-343	22.4		5100	6.4	700	8.7 1017	OUI	5.6
LE1-281	21		5700	7.2	600	8.6 1017	OUI	5.6











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SILENE LE2.362

















SILENE LE1.311 Débit de dose(nety) à 4m (rod xh⁻¹) 10⁶ Débits de dose neutron et gamma à 4 m. 105 ۵Ö٤ \equiv in 10⁰ 10-1 10-2 0.0000 TETIPS SEC 50.0000 100.0000 150.0000 250.0000 200.0000







LA TEMPERATURE AN SETN DU PEACTEUR





TEMPERATURE AU SEIN DU REACTEUR

DEBIT N 4HCRFH3 PUISSANCE(F18./SEC.) DEBIT ¥ 4H E 1 9 SILENE LE1-258 618 E17 PUISSANCE C 1 6 615 E14 C I 3 E 1 2 E 1 1 £ 1 0

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SILENE-EVOLUTION DE LA PUISSANCE AU SEIN DU REACTEUR Et decits de cose n'entre de company de

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LA TEMPERATURE AU SETN DU PEACTEUR



SILENE - Puissance, débits de dose neutrons et gamma à 4 m du réacteur.

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SILENE - Puissance, débits de dose neutrons et gamma à 4 m du réacteur.

SILENE

"BOILING" TYPE EXPERIMENTS

								-	1st]	PEAK		<u></u>		
NUMBER	C _u tg/l	H _c cm	∆H _{mm}	H _{fcm}	v _{fl}	^T 2 _s	^ω s ⁻¹	Éfissions s ⁻¹	Ê _{fissions}	E P1 fissions	E P2 fissions	^{∆P} bar	∆k ₁ _{PCM}	^{∆k} ¹s
LE1-175	71.4	36.79	90	45.79	44.41	0.015	46	4.2 1017	1.7 1017	4.2 1016	4.6 1016		960	1.21
LE1-176	69.8	38.07	114.9	49.56	48.07	0.016	43	4.5 10 ¹⁷	1.8 1016	3.7 10 ¹⁶	5.1 1016		950	1.20
LE2-176	69.8	38.07	139.9	52.06	50.49	0.018	38	4.1 10 ¹⁷	1.7 1016	3.7 1016	5.6 1016		930	1.17
LE2-343	70.6	37.1	140	51.1	49.57	0.016	42.5	3.8 1017	1.6 1016	3.2 1016	5.1 1016		950	1.20
LE1-281	70.9	37.42	170	54.42	52.79	0.017	41	4.2 10 ¹⁷	1.7 1016	3.8 1016	5.4 1016		940	1.18

	Тем	PERATURE	Тот React	'AL IVITY				
NUMBER	^Θ /•c	∆⊖°°c _{max}	^{∆k} _p _{pcm}	∆* _{PS}	DURATION	N f _{flssions}	SOURCE	CATEGORY
LE1-175	19.6		4000	5.0	540	5.4 10 ¹⁷	OUI	
LE1-176	18.8		4100	5.2	720	6.9 10 ¹⁷	OUI	5.6
LE2-176	18.8		4800	6.0	900	7.4 1017	OUI	5.6
LE2-343	22.4		5100	6.4	700	8.7 1017	OUI	5.6
LE1-281	21		5700	7.2	600	8.6 1017	OUI	5.6

Evolution de la puissance (toute l'expérience)

Fig: - Expérience du type "Ebullition"

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(toute l'expérience)

Evolution de la puissance (premiers pics)

5 _Expérience du type "Ebullition"

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Evolution de la puissance (toute l'exnérience)

Fig: 7 _Expérience du type # Ebullition"

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Evolution de la puissance (premiers nice)

Fig: 8 _Expérience du type z Ebullition"

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Evolution de la puissance (toute l'expérience)

13 _Expérience du type * Ebullition*

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APPENDIX 1

RESULTS OBTAINED ON SILENE REACTOR WITH A URANIUM CONCENTRATION OF 220 g.1⁻¹

1 - <u>Preamble</u>

A series of experiments was carried out in 1981 and 1982 using a uranium concentration of 220 g.1⁻¹ on the Silene reactor within the framework of studies on solution criticality accidents, in order to acquire experimental data in the following areas:

- 1. Maximum release of gaseous fission products when concentrated uranium solutions are in the boiling state.
- 2. Neutron and gamma radiation dosimetry associated with this type of configuration (concentrated medium, geometry different from an orthogonal cylinder).
- 3. Characterization of the power level of the boiling "steps".

The results presented here cover mainly kinetics (the effect in terms of reactivity of 1 millimeter of solution, period and power of the first peak), since the rest of the information was summarized previously in other documents.

2 - Presentation of results

Because the experiments were carried out on tank no. 1 at Silene during general calibration of the reactor (Ref. 1 and 2), and more recently on core no. 2 using a slightly different configuration (different thickness in the tank bottom and different fissile solution acidity), it is quite normal that **certain di**fferences will appear in critical characteristics of Silene with the 220 g.1⁻¹ concentration.

The results are presented in the following manner :

1. Neutron calculations,

2. Critical characteristics and effects of one millimeter,

3. Kinetic experiments on core no. 2,

4. Dosimetry.

3 - <u>Neutron calculations</u>

These calculations were made by the DAS/SEC at the time of the first calibration of Silene using the Moret code.

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- Characterisics of the fissile solution:

C_U = 220 g.1⁻¹ ²³⁵U enrichment = 92.7% Acidity = 2.84 M NO₃ - = 4.71 M Density = 1.386

In terms of number of atoms per cm³, this gives:

Number	of	²³⁵ U atoms	5.253	10 ²⁰
Number	of	²³⁸ U atoms	3.821	1019
Number	of	H atoms	5.802	10 ^{2 2}
Number	of	N atoms	2.838	1021
Number	of	0 atoms	3.780	10 ^{2 2}

Calculation of k_{eff} (Moret results)

Solution height	20	22	22.4	24	26.4	30	36.4	40
k.ff	0.931	0.961	0.965	0.999	1.025	1.065	0.110	1.127

The evolution of k_{eff} as a function of the solution height is illustrated in figure 1.

Lifetime of prompt neutrons $1 = 12.5 \ \mu s$ for $H = 24 \ cm$.

Taking into account the fact that the fissile solution heights used for the kinetic experiments were higher, let $L = 13 \ \mu s$ for the Nordheim calculations.

- Characteristics of delayed neutrons:

GROUP	1	2	3	4	5	6
$\begin{cases} \xi i \\ \beta i \chi \\ \lambda i s^{-1} \end{cases}$	1.38	1.33	1.35	1.34	1.35	1.20
	0.021	0.139	0.126	0.252	0.074	0.026
	0.0124	0.0305	0.111	0.301	1.13	3.00

 $\beta_{\text{off}} = \sum_{i=1}^{6} \xi_i \times \beta_i = 853 \text{ pcm}$

Graph no. 2 illustrates the Nordheim relationship between reactivity and doubling time T_2 for the Silene configuration at 220 g.1⁻¹.

4 - <u>Critical characteristics and effects on reactivity of one millimeter</u> of solution

<u>Core No. 1</u>:

fissile solution $C_U = 220 \text{ g.}1^{-1}$ $H^{+t} = 2.84 \text{ M}$ d = 1.3860

fissile solution height $H_c = 25.00$ cm (core tank bottom thickness 20 mm)

Effect of one millimeter $\Delta \rho$ ΔH

Experiment	-		Δρ		<u>Др</u> Дн	
NO.	s	∆H mm	p.c.m.	\$	pcm/mm	\$/mm
D4 - 01	45.6	.0.87	116	0.136	136	0.158
D5 - 01	15.7	1.68	228	0.267	136	0.159
D6 - 01	6.7	2.54	347	0.40 6	137	0.160
D7 - 01	3.75	3.19	435	0.509	136	0.160

Core No. 2:

Fissile solution $C_U = 217.9 \text{ g.}1^{-1}$ $\text{H}^{+\text{t}} = 2.84 \text{ M}$ d = 1.3562

core bottom thickness = 36 mm

critical height $H_c = 23.27$ cm

Effects of one millimeter $\Delta \rho$

Experiment	<i>—</i>	A11	Δρ		<u>Др</u> Дн				
NO.	12 S	ΔH mm	р.с.т.	\$	pcm/mm	\$/mm			
D1 - 115	31.1	0.97	151	0.177	156	0.182			
D2 - 195	9.3	2.0	298	0.349	149	0.175			
D3 - 195	3.9	2.91	430	0.504	148	0.173			
D1 - 200	39.7	0.85	127	0.149	149	0.175			
D2 - 200	13.9	1.67	243	0.285	146	0.171			
D3 - 200	6.5	2.41	350	0.410	145	0.170			
D4 - 200	3.36	3.09	454	0.532	147	0.172			

5. Results of kinetic experiments

These results are recorded in Table 1 and illustrated on graphs 3 to 35.

The notations used are the following:

 U_t : Total uranium concentration (in g.1⁻¹) (92.7 % ²³⁵U enriched uranium)

H_c : Delayed critical height (in cm)

V_c : Critical volume (in liters)

- M_c : Critical mass in U_t (in kg)
- H_f : Final fissile solution height (in cm)
- V_f : Final volume (in liters)

ΔH : Excess solution introduced above H_c , i.e. $H_f - H_c$ (in nm) : Total number of fissions Nf ∆kp : Total potential reactivity inserted in the core ^{∆k}eff₁ : Effective reactivity present at the first peak : Mean rise in temperature of the fissile solution at the end of the experiment (in $^{\circ}C$), θ_i designating the initial temperature and θ_{f} the temperature reached at the equilibrium state t. T₂ : Doubling time of power rise (in s) Τ. : Period of divergence (in s) : Reciprocal of period (in s⁻¹) ω Ε : Maximum power at the top of the first peak (in fissions/s) : Time corresponding to the first peak t₁ : Dynamic pressure wave on the core tank bottom during the first Δp peak (relative value) : Energy integrated up to the bottom of the first peak (in E_{p1} fissions) E_{p 2} : Total energy integrated during the first peak (including residual power) : Time required for the system to reach temperature equilibrium t., E.q : Energy integrated at time t_{eq} (in fissions).

4

Calibration of reactor energy is obtained by gamma spectrometric analysis of the fissile solution. This radiation-chemical data was obtained in experiments D1-12, D4-195, and LE1.201.

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EXPERIMENTS RESULTS AT 220g.1-1

							1 st PEAK							TEMP	PERATURE OYENNE				[
NUMBER	C _H tg/I	H _c cm	∆H _{mm}	H _{fcm}	v _{fl}	<i>T</i> 2 ₅	^w s ⁻¹	E fissions	^E P ₁	E _{p1}	∆ P bar	$\Delta k_{1_{PCW}}$	^{∆t} 1s	θ,		∆k P _{pcm}	∆k _{ps}	DURATION	N _{f fissions}	NEUTRO SOURC
D4-195	218	23.25	2.4	23.59	22.88	6.5	0.107	5.5 1014	2.3 1016			352	0.413	18.8	12.8	352	0.413	700	4.2 10 16	OUI
S1-196	218	23.28	5.6	23.84	23.13	0.30	2.31	8.2 1015	2.0 1016	2.5 1016		762	0.893	18.8	24.8	762	0.893	260	8.1 1016	NON
S2-196	218	23.28	7.7	24.05	23.33	0.0043	161	1.0 1018	3.1 1016	3.4 1016		1069	1.25	18.8	33.0	1069	1.25	160	1.1 1017	NON
S1-197	218	23.28	8.4	24.12	23.40	0.0024	289	2.9 1018	4.1 1016	4.6 1016	0.16	1238	1.45	18.8	33.2	1238	1.45	160	1.2 1017	NON
S1-198	218	23.28	9.5	24.23	23.51	0.00171	405	6.6 1018	5.4 1016	5.6 1016	1.1	1391	1.63	18.9	36.1	1391	1.63	160	1.2 1017	NON
LE 1-199	218	23.25	15.0	24.75	24.01	0.018	38.5	3.5 1017	2.8 1016	3.1 1016		902	1.06	17.9	60.3	2025	2.37	310	2.2 1017	OUI
LE 1-201	218	23.25	19.1	25.16	24.40	0.010	69.3	3.0 1017	2.6 1016	3.0 1016		937	1.10	19.1	69.3	2540	2.98	280	2.5 1017	OUI
LE 1-207	218	23.13	26.8	25.81	25.04									19.1	101	3484	4.08	640	3.8 1017	OUI
LE 2-207	221	23.13	27.2	25.85	25.07	0.010	69.3	2.7 1017	3.0 1016	3.2 1016		946	1.11	19.0	101	3510	4.11	550	3.5 1017	OUI
LE 1-208	221	23.20	27.0	25.90	25.12	0.010	69.3	3.0 1017	3.2 1016	3.5 1016		946	1.11	19.0	101	3510	4.11	550	3.5 1017	011



Variation du keff en fonction de la hauteur de solution

Résultats MORET



et la reactivite Δk

Fig: 3 I Expérience du type "Divergence" Evolution de la puissance SILENE



御養 業産業の まちぬい

Fig: 4 _ Experience du type pive gence Evolution de la température au sein du réacteur -Expérience du type SILENE "Divergence"



Fig: S Débit d'exposition Expérience du de l'axe SIL du réacteur type **ENE** \sim ۵, a "Divergence" 1m et **4** m



















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pression au sein du reacteur





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RESULTATS DES APPROCHES SOUS-CRITIQUES SANS BARRE

Numéro	Concentration g/1 Ur	Concentration S/1 U275	H_ (cm)	N _c (litres)	M (s) ² u _r	(g) ^C U ₂₇₅
/c 6-01	220	205	- 25,02	24,21	5326	4963
/c 1-02 }	146,3	1963	25.79	24,95	1652	3402
/o 1-03 - 1	101,4	94. *	29,61	26.67	2907÷	2706
/c 1-04 _ ,	80,9	J .75.4	, st.re	. 264	* 264 2	2463
/c 1-06	70,2		38,8 7		2644	2463
/6 1-05	60,9	- 1563	46.20		A 19729	2541
10 1-07	7.6	63 C	36 <i>6</i> 9		2612.	2475
/c 1-08	70,7	es.	. ورياد		2616	2475
/c 1-11	° 71. 2		38,68		2630	2450
/c 1-12			. sat		2019	2438
/c 1-14 '	79.8		18.02	36.83	2611."	2431
/c 1-15	7927	65,8	38,45	7.5	2634	2451
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SILENE

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			SILENE RESULTS OF THE CALLERATION ON THE PAIRT CORE														
NUMBER C. H.				Teis	1ªPEÁK				TEMPERATURE MOYENNE								
	C.,	#	∆ <i>ll mm</i>	H _{fen}	B ,	72.	or of	}	e		EP3 provide	ΔΡ	e Bi e	ET IC2	DURATION	N f _{fissions}	NEUTRON Source
D7-01	228	3.01	- 32 -	25.34	24.52	200 T (1)			1.3 1016	A THE COLLECT			21	13.7	300	4.6 1016	OUI
D3-03	1014	29.61	4.7	30.08	29.13	-1.			2.0 1014		A STATES	Nag 75	22	12.6	240	5.3 1016	OUL
D3-86	70.5	MIT	1 72	79.39	38.36				2 3.6 1010				22	10.8	240	5.7 1016	OUI
D1-12	70.7	38.24	8.2	3411	37.00			1 1.4 101	21 1014				22	13.9	240	7.4 1016	OUI
D4-23	70.3	38.33	43	38.76	37.56	11.6	NO.	2.1 101	*1.4 10 ¹⁴			din din	21.5	8.3	540	4.3 1016	OUI
S1-06	70.2	39.12	14	40.52	39.28	\$ 0.42	1.65	1.8.6 1017	1.8.1019	3.0 1018	3.7 1016		22	17.7	180	9.6 1016	NON
S1-08	70.7	38.15	17	39.85	38.61	4.028	24.8		1.0.10	3.6 1016	4.0 1016		22	21.7	180	1.2 1017	NON
52-08	70.7	38.14	20	.40.14	38.90	0.010	69,3	7.7 1017	2.5 194	5.5 1016	5.7 1016		21	25,6	180	1.4 1017	NON
\$3-08	70.2	38.14	25	40.64	39.38	0.0049	1.14	3.4 1014	4.7.102	8.2 10 ¹⁸	8.7 1016	0.5	21	21,3	<60	8.7 1016	NON
S4-08	0 70.7	39.14	31	41.24	39.96	0.0040		3.5 101	1.1.1	· 9.3 1018	9.8 1014		21	20,6	<60	9.8 1016	NON
S5-08	76.2	39.14	35	41.64	49.35	8.0023		Cal Singe	IN SUCCESSION	1.3-1017	1.4 1017	2.95	.22	41.7	180	2.5 1017	NON
51-11	71.3	37.87	313	41.60	39.72	8.0031	22	- 1.3 1145	14 A.S 1914	1.1 10 ¹²	1.4 1017	1.75	20	37.8	180	2.1 1017	NON
\$1-12	* 10.2	48.24	49.6	2.3	41.92		1 999 A		A CANOLY	Es ters	17 1017	4.8	- 20	293	<60	1.7 1017	NON
\$2-12	78.7	31.24	A1.6	32.40	41.09	0.00200	1 341	Estion.	12 1010	1.5 101	1.5 1017	4	21	41.2	120	2.7 1017	NON
S1-16	70 F.	38.28	41.7	3243		A ACOLL	1 14	STR. P. D. A.		AP 1.4 101	1.6 1017	4.3	18.6	26.8	<60	1.6 1017	NON
\$2-16	70.7	38.26	43.6	12.62	4140	0.00160	413	2.4 10	9.3 (pto	12,1017	1.7 1017	4.95	18.6	29	<60	1.7 1017	NON
S1-22	37126	17.63	£ 30	40.63	39.31-	0.0031	224	7,8 104	6.7 1014	1.0 1017	1.1 1017	1.55	20	26.4	<60	1.1 1017	NON
\$1-23	1423	87.63	5 31	. 01.03 .	46.12	-11.0023	18 391	13 10 %	14.1 B	1210	, 13 10 ¹⁷ 55	2.95	20.5	24.9	<60	1.3 1017	NON
\$3-12	124	17.63	42.1		10.54	1.00172	1403	22 1014	1.99.0 161	1.6 1017	14 1017	4.95			<60	1.6 1017	NON
\$4-22		31.4	24		39.18	0.0019	177	4.9 1018	35 101	8.7 1010	9.2 1016	0.9	20.5	17.6	<60	9.2 1016	NON
SS-22	- 213	37.85	115.5	39.18	37,96	0.090	7.7	2.2 1010	13 1010	3.4 1010	3.8 1016		20.5	9.3	<60	3.8 1016	NON
\$6-22	712	57.51	4	42.03	40.73	0.00177	392	2.1 1019	9.1 1010	1.5 1017	1.6 1017	4.65			<60	1.5 1017	NON
\$1-23	70.3	38.51	44	42.91	41.58	0.0037	187	6.7 1018	6.4 1016	9.9 1016	1.0 1017	1.3	21	. 19	<60	1.0 1017	NON
S1-23	70.3	38.51	44	42.91	41.58	0 0037	187	6.8 1018	6.4 1016	1.0 1017	1.0 1017	1.3	21	19.1	<60	1.1 1017	NON
LE1-12	70.7	38.24	60.1	44.25	42.88	0.019	36.5	3.1 1018	1.6 1016	5.0 1016	5.4 1016		21	64.3	360	4.0 1017	000

56-22 Find / top just polan = 242.3 5/0 Total final pulse 3.9515 5/0

1 - A table of results from all supercritical tests.

2 - Critical characteristics of the core (result table and graph).

3 - The variation of the effect produced on reactivity by one millimeter of solution above the critical height (result table and graph).

4 - The variation of the feedback term as a function of concentration (graph).

5 - The neutron constants determined by calculation and used in evaluating reactivity.

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APPENDIX 2

RESULTS OF THE SILENE REACTOR CALIBRATION CAMPAIGN

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ST STATES

In order to select the optimal characteristics for routine operation of the Silene reactor and meet safety requirements, a reactor calibration campaign was conducted in 1974. This campaign made it possible to determine, through computation or measurement, the principal parameters and neutron constants necessary for operation and far interpretation of kinetic experiments.

The results of these tests are presented separately because they were obtained from the first reactor core tank mode characteristics differ slightly from those of the tanks used previously. The only differences were the thickness of the stainlass steel for the cover (20 mm) and for the tank bottom (16 mm). The critical conditions thus varied only slightly from current measurements. Hometheless, where these tests covered a wine contentration range, from 68 to 270 gfd, and included kinetic parameter measurements (affect of some slillingter and feedback effects). It appeared uneful to add this information to the present summary of blasme experiments.

Detailed remains are found in technical note BSE/SEVENC No. 124.

This summarized presentation consists of:



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Evolution de la température au sein du réacteur Fig:37-Expérience du type SILENE



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Concestination 325Ug/1

Concentration ³²⁹Ugn 300 SILENE Variation du terme de freinage en fonctien de la concentration en ²³⁵U (Shuldown Coefficient)




APPENDIX III

RAPPORT SRSC 93. 220 - Décembre 1993

A REVIEW OF THE SILENE CRITICALITY EXCURSIONS EXPERIMENTS

Francis Y. BARBRY

Jan 199

CEA INSTITUT DE PROTECTION ET DE SURETE NUCLEAIRE DE PARTEMENT DE RECHERCHES EN SECURIT SERVICE DE RECHERCHES EN SURETE ET CRITICIT Centre d'Etudés de VALDUC - SRSC - 21120 Issur-Tille - France **2** 80 23 40 de

A REVIEW OF THE SILENE

CRITICALITY EXCURSIONS EXPERIMENTS

Francis Y. Barbry Institut de Protection et de Sûreté Nucléaire CEA Valduc, 21120 Is-sur-Tille France 80 23 50 20

ABSTRACT

More than one thousand experiments reproducing criticality excursions in aqueous solutions of uranium highed enriched in 235 U have been conducted to date at VALDUC in the SILENE facility by the French Institute for Nuclear Safety and Protection (IPSN). This document summarizes results of selected experiments with reactivity insertions ranging from 4 cents up to 7 \$. Valuable results relating to the first peak characteristics and integrated fissions yields provide reference data for evaluating and improving analytical models describing criticality excursions. In addition acquired knowledge is given concerning pressure increase, radiolytic gas formation and the general behaviour expected in the accidental situations which may occur in fuel cycle installations. The lessons drawn from those experiments concerning criticality accidents studies are given in this paper.

I. INTRODUCTION

It is generally assumed that criticality accidents are most likely to occur during the processing of fissile solutions. So as to be able to cope with such an eventuality, the CEA has been engaged since 1967 in an important programme of criticality accident studies¹ using the CRAC facility² to begin with and at present since 1974 the SILENE reactor³. The general objective of the work undertaken at VALDUC is to study the phenomenology as well as the radiological consequences of an accidental criticality excursion so as to acquire, on the one hand, indispensable knowledge for criticality asfety assessment and, on the other hand, to be in a position to define a protection and intervention policy.

II. DESCRIPTION OF THE SILENE FACILITY

SILENE is a homogeneous experimental reactor using a fissile solution of uranyl nitrate (U enriched in 235 U at 93 %) as fuel. The core is in the form of a small annular tank (36 cm diameter) located in the middle of a large concrete room referred to as the "cell". The fissile solution required for reactor operation of the reactor is prepared in a laboratory located in the cell basement³

The general operating principle of the reactor is as follows :

The fissile solution, previously adjusted to SILENE operation concentration in a special large capacity tank, is pumped into the core up to a predetermined supercritical level. During this phase, a control rod is present in the core so as to avoid divergence.

The "divergence" power excursion is then produced by withdrawing the rod from the core according to a procedure that depends on the selected operating mode for each experiment.

When the experiment has been completed, the fissile solution containing the radioactive fission products is dumped into a tank located in a shielded room so as to allow quick access to the cell.



A ventilation circuit blows continuously through the upper part of the core so as to dilute the radiolysis gases that are formed. After the period required for their radioactive decay, these gases are removed through ad hoc filtration systems.

SILENE is designed to operate in three different modes depending on core reactivity, the control rod withdrawal rate and the presence or absence of an auxiliary neutron source.

- . "PULSE" operation is obtained by the rapid removal of the control rod (at a rate of 0.2 or 2 m.sec⁻¹) with or without an additional source in order to obtain a very high power peak in a very short time (up to 1000 megawatts in a few milliseconds). With fast transients of this kind the reactivity is limited to 3.0 \$.
- "FREE EVOLUTION" is achieved by slowly removing the control rod at speeds lower than 2 cm.sec⁻¹ with a neutron source. Reactivity cannot exceed 4 \$ in normal operation but may reach 7 \$ for solution "boiling" experiments. This mode simulates an accidental criticality excursion allowed to evolve freely.
- "STEADY-STATE" operation involves automatic control of the rod position, with very slow displacement rates (approx 2 mm.sec⁻¹) in order to maintain the reactor at a predetermined stable power level.

III. RESULTS

A. General behaviour of a criticality excursion in a solution

In general terms the development of an accidental criticality excursion in a solution is governed by the following principal parameters:

- . the reactivity addition
- , the initial neutrons source
- . the effects due to the temperature, i.e.
 - the nuclear temperature effect

the expansion effect which brings about density variation of the medium as well as a "buckling" variation.

 the effects due to void; this void arises in the first instance from the formation of bubbles of radiolytic gas, but void also occurs if boiling of the solution develops.

Once delayed critical is exceeded, a divergent chain reaction may be established leading to an exponential evolution of the power. This power excursion, whose kinetics is dependent on the reactivity input, is accompanied by the release of energy which is manifested mainly in thermal form. The heating of the fissile material results, on the neutronic side, in the appearance of feedback mechanisms. These feedback reactions overcome the current reactivity. The result is the appearance of a first power peak; Figure 1 is a sketch of the result of a typical transient.

The first feedbacks to occur are due to the thermal expansion of the system and to the neutronic temperature effect. To these initial mechanisms arresting the chain reaction is added a second phenomena specific to solutions, the formation of radiolytic gas due to the decomposition of water along the length of the trajectories of the fission fragments. This second mechanism, due to voiding, supplements the first feedback reactivities which reduces the power level considerably. However, the radiolytic gas bubbles migrate towards the surface of the liquid, the accompanying reverse reactivity disappears and the power excursion starts again. It is this process of the appearance of gas, then the release out of the solution which is the origin of the power oscillations observed in the evolution of power. The essential difference which should be emphasized between the feedback mechanisms lies in the effect of the void formed by the radiolytic gas bubbles being a transitory effect, whilst the effect of the temperature, consequent on heating of the fissile solution, is permanent on the time scales being considered.

In certain cases the energy generated is sufficient to either bring the system to boiling, or produce a large pressure pulse resulting in significant equipment damage or solution dispersal.



Figure 1 - Typical Criticality excursion

B. Results obtained at the 71 g.1^{-1 c}oncentration

The most representative results of the SILENE experiments in the unshielded configuration are reviewed in table I. The experiments have been classified into different categories :

- . category 1 : reactivity steps below prompt criticality ($\rho < \beta$)
- . category 2 : reactivity steps near prompt criticality ($\rho \# \beta$)
- . category 3 : reactor "pulse" operation with large reactivity input ($\rho >> \beta$)
- . category 4 : the same as category 3 but with presence of an external additional neutron source
- . category 5 : reactivity ramps below solution boiling
- . category 6 : large reactivity ramps ($\rho >> 4$ \$) leading the solution to boiling
- . category 7 : series of experiments performed with the same total potential reactivity but in various initial conditions

The effective reactivity at the first peak Δk_1 is calculated by the inhour equation and appropriate values of neutron lifetime and β_{eff} . The total potential reactivity Δk_p is calculated by the Monte Carlo MORET code. The relation used between the K value and the solution height for the SILENE 71 g.l⁻¹ solution concentration is

Keff = $0.68276 + 1.194 \ 10^{-2} \text{ x H} - 9.905 \ 10^{-5} \text{ x H}^2$

. Neutronics data and symbols used

Lifetime of prompt neutrons at 71 g/l $l = 36\mu s$

GROUP	1	2	3	4	5	6
εί	1.28	1.24	1.26	1.23	1.26	1.20
βi	0.021	0.139	0.126	0.252	0.074	0.026
λi	0.0124	0.0305	0.111	0.301	1.13	3.0

$$\beta_{eff} = \sum_{i=1}^{5} \varepsilon_i \times \beta_i = 794 \, pcm \qquad (pcm = 10^{-5})$$

Characteristics					Results								
Cu _t = 71.g ⁺¹ Hc ≆ 37 cm					1 st peak						Final		
Experiment A*	v _f	Akp s	Neutron Source	Duration s	T ₂	∆kį S	fissions.s ⁻¹	Ê fissions	Ep ₁ fusions	Ap bars	^{∆0} •C	N _F fissions	Category
LE3-229	36.1	0.035	yes	14400	235	0.035	1.4 10 ¹²	1.1 1015		•	ı	3.1 1015	1
LE1-214	36.9	0.14	ym	8000	44	0.14	4.5 10 ¹³	1.0 1016	-		4	2.4 10 ^{16*}	1
LE2-229	36.6	0.42	7 94	14400	6.2	0.42	7.1 10 ¹⁴	1.9 1016	•	•	11	7.5 10 ^{16•}	1
\$1-300	37.0	0.51	yes	1100	3.8	0.51	1.3 1015	2.2 10 ¹⁶		•	14	6.5 10 ¹⁶	1
52-300	37.7	0.97	no	260	0.13	0.97	1.7 10 ¹⁶	1.3 10 ¹⁶	4.2 1016	•	22	1.1 1017	2
\$2-258	38.7	1.32	RO	006	0.010	1.32	7.1 1017	1.8 1016	4.5 1016	•	28	1.5 1017	3
53-258	39.3	1.84	90	420	0.0038	1.84	4.9 1018	5.4 1016	\$.1 10 ¹⁶	0,95	36	1.9 1017	3
\$4-258	39.7	2.14	no	420	0.0028	2.14	9.1 1018	7.0 1016	1.0 10 ¹⁷	1.9	40	2.3 1017	3
\$2-259	40.1	2.60	80	420	0.00198	2.60	1.7 1019	8.2 10 ¹⁶	1.3 10 ¹⁷	3.7	47	2.7 1017	3
\$3-259	40.5	2.86	90	420	0.00171	2.86	2.1 10 ¹⁹	7.9 1016	1.4 1017	4,8	51	2.9 1017	3
\$4-346	40.2	3.0	30	180	0.00162	3.0	2.4 10 ¹⁹	9.8 10 ¹⁶	1.5 10 ¹⁷	5.6	\$3	2.9 1017	3.7
\$1-362	40.9	3.0	788	210	0.0039 (zzatabie)	1.79 1.89 34.5 s ⁻¹	5.3 10 ¹⁸	5.6 10 ¹⁶	8.5 10 ¹⁶	1.1	50	2.8 10 ¹⁷	4.7
LE1-362	40.9	3.0	78	700	0.136 (unstable)	0,95	2.0 1016	9.6 10 ¹³	4.1 1016	•	52	3.1 10 ¹⁷	5.7
LE2-362	40.9	3.0	yes	210	0.025 (unstable)	1.12	1.8 1017	1.2 10 ¹⁶	4.5 1016	•	50	2.8 10 ¹⁷	5.7
LEJ-273	42.4	3.6	yes	1300	0.025 (unstable)	1.12 map 8.10 \$ s ⁻¹	1.7 1017	1.1 10 ¹⁶	4,4 1016	•	63	4.6 1017*	5
LE2-176	50.5	6.0	' yes	900	0.01\$ (unstable)	1.17 rump 4.16 \$. s ⁻¹	4.1 1017	1.7 10 ¹⁶	3.7 1016	•	boiling	7.4 10 ¹⁷	6
LE1-281	52.8	7.2	yes	600	0.017 (magable)	1.18 mme.16.2.s ⁻¹	4.2 1037	1.7 1016	3.8 10 ¹⁶	•	boiling	8.6 10 ³⁷	6

* Experiments lasting up to power rustarting (post-accident phase)

Table I - SILENE data

- ϵ_i represents the efficiency of each group of delayed neutrons
- C_{Ut} Total uranium concentration (in g.l⁻¹) (93 % 235U enriched uranium)
- H_c Delayed critical height (in cm)
- V_f Final volume (in liters)
- N_f Total number of fissions
- Δkp Total potential reactivity addition in the core
- Δk_{eff_1} Effective reactivity present at the first peak
- $\Delta \theta$ Mean rise in temperature of the fissile solution at the end of the experiment (in °C), θ_i designating the initial temperature and θ_f the temperature reached at the equilibrium state te
- T₂ Doubling time of power rise (in s)
- ω Reciprocal of period (in s⁻¹)
- *E* Maximum power at the top of the first peak (in fissions/s)
- Δp Dynamic pressure wave on the core tank bottom during the first peak (relative value in bars)

- E_{p_1} Energy integrated up to the botton of the first peak (in fissions)
- \vec{E} Total energy integrated up to the maximum of the peak power

The figures 2 to 5 illustrate the evolution of the power, energy and temperature during the experiments. Several typical excursions have been reproduced, namely:

- . Figure 2 A fast transient (S4-346) produced by a 3 \$ reactivity step and two intermediate transients (LE2-362 and LE1-362) resulting of reactivity ramps (0.17 and 0.035 \$. s⁻¹), all three experiments performed with exactly the same total reactivity excess.
- . Figure 3 A very slow transient (LE2-229 produced by a small reactivity addition (0.42 \$) but showing that, even in this case, a restarting must be expected.
- . Figure 4 An experiment (LE1-281) with a very large excess of reactivity ($\Delta k \equiv 7.2$ \$) leading the solution to boiling.
- . Figure 5 Fast Transient 1st peak



Figure 2 - Experiments Performed at the Same 3 \$ Total Reactivity



Figure 3 - Slow Kinetics Excursion with Power Restarting





IV. KNOWLEDGE ACQUIRED

The lesson we have learnt here takes into consideration the overall results acquired during the CRAC and SILENE experiments, namely the 72 experiments conducted in the CRAC facility with a reactivity insertion obtained through continuous pumping of the solution into 30 and 80 cm diameter cylindrical tanks and several hundred excursions performed with SILENE.

The range of the parameter and result variations were as follows :

- . uranium concentrations between 20 and 340 g x l⁻¹ (homogeneous configurations only)
- volume of fissile solution up to 250 liters
- . total potential reactivity input ranging up to 10 \$
- . reactivity ramps and "steps" reaching 2 \$ x s⁻¹ and 3 \$ respectively
- . presence or not of an additional neutron start-up source
- . heating of the solution up to boiling

Our observations were :

- maximum power observed $\leq 10^{20}$ fissions x s⁻¹
- . maximum total number of fissions $\leq 5 \times 10^{18}$ fissions
- . doubling time ranging from a few hundred µs to a few minutes

We can also mention other valuable information :

. Relation between N_f , V, Δk_p , t

For one configuration, namely a certain diameter ϕ of the vessel, it is possible to approximately relate the total specific fission yield during the chain of pulses to the potential reactivity Δk_p (except if boiling). The total number of fissions in the excursion normalized to the total volume of solution (hence to its heat capacity) fits an expression :

$$\frac{N_f(yield)}{total \ volume \ V} \cong k . \Delta k_p$$

k depends on the diameter of the vessel with $k # 3.4 \ 10^{12}$ fissions.(p.c.m)⁻¹. I⁻¹ for SILENE

. Relation between
$$\frac{N_f}{V}$$
 and t (ref.4)

From another standpoint, and this may useful for the accident estimation, it is possible using the maximum energies measured during CRAC and SILENE excursions up to 7 S reactivity insertion, to fit an empirical correlation N

between the specific total number of fissions $\frac{N_{f}}{V}$ (t) and

the time as follows :

$$\frac{N_f}{V} = \frac{t}{3.55 \, 10^{-15} + 6.38 \, 10^{-17} \times t}$$
with t in seconds and V volume in liters

. Relation between the first peak power \vec{E} and ω (fig 6)

For fast transients ($\rho > \beta$) the maximum specific peakpower \dot{E}/V is varying with the reciprocal period ω as the following relation

$$\frac{\dot{E}}{V} = c^{\text{te}} \times \omega^{1,8}$$

. Radiolytic gas formation

The volume of radiolysis gas formed is proportional to the number of fissions⁵ reaching approximately 1.1 x 10⁻¹³ cm³/fission (i.e. 110 liters of gas for 10¹⁸ fissions). The threshold for the formation of radiolysis bubbles is estimated at 1.5 x 10⁻¹⁵ fissions per liter of fissile solution.

. Pressure increase during the 1st peak

No significant overpressure was observed for doubling times T_2 greater than 10 ms but for very fast transients the energy generated and gas formed within the liquid may cause material ejection and some mechanical damage to the vessel (i.e. CRAC 44).

Dosimetry

A list of dosimetric measurements have been established⁶. It should be kept in mind that for different configurations the number of fissions is not proportional to the dose. The maximum value observed was 5.8×10^2 Gy at 1 m from the CRAC source for 10^{18} fissions.

. Solution boiling

The fissile solution is brought to its boiling point for a released energy level of about 0.33 MJ per liter ($\equiv 1.1$. 10¹⁶ fissions/l) and the boiling pseudo-plateau level depends on the excess of reactivity.

. Startup neutron source influence

The presence of an initial neutron source (external or intrinsic) strongly influence the behaviour of the excursion by the tendency to significantly delay the initiation of the burst when the source decreases. It results in a larger burst height because of a strong reactivity step in place of moderated reactivity ramps. It could be to recommanded to provide a neutron background in some processing operations when the internal source is weak to limit, in the case of an excursion, the size of the first peak and to maintain the resulting pressure below the level that would othervise result in significant equipment damage or solution dispersal.

Post accident situation

Experiments carried out without solution dumping (See table I and Fig 3) show that in most cases of accidental situations the power excursion is likely to restart after a delay depending on heat transfers with the environment, unless some means of intervention has been prepared so as to stop the accident process.

Criticality accident detection systems, if designed for that purpose and this is the case for the French system, may be helpful in this kind of situation for providing information on the evalution of the post-accident phase and contribute to decision making, for example in the event of intervention.

V. CONCLUSION

The CRAC and SILENE criticality accident study programmes have provided a wealth of information on all the aspects of criticality excursions in aqueous fissile media : physics, dosimetry, detection, post-accident situation.

The practical lessons thus drawn will be helpful for evaluating and coping with the consequences of such an event. The accurate SILENE results will allow improvement of excursion modeling and new calculation code development ^{7,8}

The overall work will undoubtedly contribute to a better knowledge of the phenomena that may be encountered in accident situations and to the elaboration of a nuclear installation safety policy.



Figure 6 - Relation Beetwen the Specific Power and the Reciprocal Period

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