

RESEARCH ARTICLE

EXPONENTIAL-TIME DIFFERENCING METHOD FOR SOLVING REACTOR POINT KINETICS EQUATIONS WITH TEMPERATURE AND POWER FEEDBACK AND ITS VALIDATION BY ANALYSINGTHERMAL REACTOR BENCHMARKS

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Abstract

..... Exponential Time Differencing method with Taylor Series Expansion (ETD-TS) is used for solving nonlinear systems of point kinetics equations (PKEs). Integral forms of PKE are expanded using Taylor Series Expansion method. Time step value and Taylor series expansion order (n) is chosen to minimize the truncation error.PKEs are highly stiff and nonlinear in nature in presence of thermal feedback. So, advanced method (ETD-TS) is used in the present work to estimate the power transients. Thermal Reactors benchmark with stepandcompensated ramp reactivity insertions are considered for estimation of power transients with feedback. Comparison of results with other accurate numerical methods likeCATS (Convergence Accelerated Taylor Series), SCS (Shifted Chebyshev Series), ITS2 (Integral Taylor Series)etc confirms the validity and accuracy of this method. Time step of $10^{-2} - 10^{-4}$ sec is considered for estimation of power and reactivity transients.

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Introduction:-

The solution of linear Boltzmann transport equation is coupled with precursor concentration equationsthat are in different time scales. To estimate the accurate value of neutron densities and precursor concentrations, temperature and power feedbacks needs to be applied. Due to the presence of feedback, nonlinearity is introduced in PKEs.PKEs are derived from the time dependent neutron transport equations. After, considering weight based approximations, these integro-differential equations are converted to ODEs. For reactor safety, it is important to estimate the power values at different time intervals by solving point kinetics equations. These equations provide useful information on the dynamics of the reactor core during transients that involve considerable power change in the nuclear power plant. Temperature feedback effect is considered in PKEs for improving the accuracy of results. Different numerical schemes are available in literature to solve PKEs with temperature and power feedback. Recent examples are, methods based on Reactivity Piecewise Constant Approximations(Kinardand Allen,2004), ET technique, based on Backward Euler (ET-BE) and Crank Nicholson (ET-CN) approximations (Nahla,2011), The Converged Accelerated Taylor Series Method (CATS) (Ganapol and Picca et al, 2012), The Enhanced Piecewise Constant Algorithm, EPCA (Picca etal,2013), Integral Taylor Series, ITS-2 (Serigo.Q et al, 2016) and Shifted Chebyshev Series,SCS (Hamda,2018.Each of these methods are having different levels of accuracy, computational time and complexity in implementing the algorithm, The numerical scheme used in this paper is based on Exponential Time Differencing

(2)

with Taylor series expansion (ETD-TS), which involves the exact integration of PKEs followed by multiplying by integrating factor. The integrand is expanded with Taylor series expansion method. Recently, (M.M.A.Razak etal., 2015) used ETD method with Taylor series for solving the PKEs with step and ramp reactivity insertions in the absence of feedback, however in this paper we had used ETD-TS method for estimation of power transients by solving PKEs in the presence of temperature and power feedback. The coefficient matrix 'A' used here is 8×8 matrix, here the last column of the coefficient matrix is corresponds to the function U(t) which is dependent on temperature of the reactor core, which is different from (M.M.A.Razak etal., 2015) ETD-TS algorithm.

Point kinetics equations and ETD-TS method

Point kinetics equations are coupled equations and has to be solved simultaneously. ETD – TS method has four different steps:

Multiplying the PKE's with exponential Integrating factor

- 1. Integrating the PKEs within the time limits ' t_n ' to ' t_{n+1} '.
- 2. Expanding the neutron density and precursor concentrations using the Taylor series expansion
- 3. Estimation of neutron density and precursor concentrations at n+1th time step.

$$\frac{\mathrm{d}P(t)}{\mathrm{d}t} = \frac{(\rho(t)-\beta)}{\Lambda}P(t) + \sum_{i=1}^{6}\lambda_i * C_i(t) \dots (1)$$
(1)

 $\frac{dC_{i}(t)}{dt} = \frac{\beta_{i}}{\Lambda} P(t) - \lambda_{i} * C_{i}(t)$ $P(t) = \text{power in watts}; C_{i}(t) = i^{\text{th}} \text{ group precursors concentration (atoms/cm}^{3});$

P(t) = power in watts; C_i(t) = 1th group precursors concentration (atoms/cm²) β = effective value of beta; Λ = mean neutron generation time; λi = decay constant of ith group precursor; βi = ith group precursor beta value;

11.11

For simplicity P(t) which is the neutron number density can be assumed as power of the reactor by assuming praportinal

From Eq.(1) and Eq.(2) we can write PKE with feedback in the matrix form as shown below:

$\begin{pmatrix} dP(t)/dt \\ dC1(t)/dt \\ dC2(t)/dt \\ dC3(t)/dt \\ dC4(t)/dt \\ dC5(t)/dt \\ dC6(t)/dt \end{pmatrix} =$	$\begin{pmatrix} (\beta(t) - \beta) \\ \Lambda \\ \beta_1/\lambda 1 \\ \beta_2/\lambda 2 \\ \beta_3/\lambda 3 \\ \beta_4/\lambda 4 \\ \beta_5/\lambda 5 \end{pmatrix}$	λ1 - $λ1$ 0.0 0.0 0.0 0.0 0.0	$\lambda 2$ 0.0 $-\lambda 2$ 0.0 0.0 0.0 0.0	$\lambda 3$ 0.0 0.0 $-\lambda 3$ 0.0 0.0 0.0	$\lambda 4$ 0.0 0.0 $-\lambda 4$ 0.0	$\lambda 5$ 0.0 0.0 0.0 0.0 $-\lambda 5$	λ6 0 0.0 0.0 0.0 0.0 0.0	0.0 0.0 0.0 0.0 0.0 0.0	$ \begin{pmatrix} P(t) \\ C1(t) \\ C2(t) \\ C3(t) \\ C4(t) \\ C5(t) \\ C6(t) \end{pmatrix} $
dC6(t)/dt	β ₅ /λ5	0.0	0.0	0.0	0.0	$-\lambda 5$	0.0	0.0	C6(t)
\dTemp(t)/dt/	$\beta_6/\lambda 6$ U(t)	0.0	0.0	0.0	0.0	0.0	-x6 0.0	0.0	Temp(t)

Right hand side of above 8×8 square matrix is the coefficient matrix 'A(t)'.

The general form of first order stiff ODE (Cox,2002)is written as:

$$\frac{du}{dt} = cu + F(u, t)$$

(3)

Here 'c' is stiffness coefficient. The value of $u(t_{n+1})$ can be written as follows after applying the procedure of exponential differencing method(Cox,2002): $u(t_{n+1})=u(t_n)e^{ch}+e^{ch}\int_0^h e^{-ct}F(u(t_n+t_n)t_n+t_n)dt(4)$

$$P(t_n + 1) = P(t_n)e^{\frac{-h}{\Lambda}} + \frac{1.0}{\Lambda}\int_0^h P(t_n + \varepsilon)e^{\frac{\varepsilon - h}{\Lambda}} (1.0 - \beta + \rho(t_n + \varepsilon))d\varepsilon$$

$$+ \int_0^h e^{\frac{\varepsilon - h}{\Lambda}} \sum_{i=1}^6 \lambda_i C_i(t_n + \varepsilon) d\varepsilon(5) C_i(t_n + 1) = C_i(t_n) e^{\frac{-h}{\Lambda}} + \left(\frac{\beta_i}{\Lambda}\right) \int_0^h e^{\frac{\varepsilon - h}{\Lambda}} P(t_n + \varepsilon) d\varepsilon + \left(\frac{1.0}{\Lambda} - \lambda_i\right) \int e^{\frac{\varepsilon - h}{\Lambda}} C_i(t_n + \varepsilon) d\varepsilon$$
(6)

Eqns.(5) and (6) can be obtained after integration of (1) and (2) and multiplying both sides of equation (1) and (2) with integrating factor $et/\Lambda(M.M.A.Razak et al,2015,Cox et al,2002)$: With Taylor series expansion power at time $t_n + t$ can be written as shown below: $P(t_n + r) = \sum_{i=0}^{n} \frac{\partial^{i} P(t_n)}{\partial r^{i}} \frac{r^{i}}{i!} (7)$

DifferentialCoefficients of power procurer concentrations, Reactivity and Temperature functions are obtained using the above matrix equations as follows:

$$\frac{\mathrm{d}\mathbf{F}(\mathbf{t}_{\mathrm{n}})}{\mathrm{d}\mathbf{t}} = \mathbf{A}(\mathbf{t}_{\mathrm{n}})\mathbf{F}(\mathbf{t}_{\mathrm{n}})$$

While estimating the higher order derivatives as we are choosing the smaller time steps, derivativitives of $A(t_n)$ can be neglected. matrix equation can be written as follows:

$$\frac{d^{2}F(t_{n})}{dt^{2}} = A^{2}(t_{n})F(t_{n})\dots\dots\dots\frac{d^{n}F(t_{n})}{dt^{2}} = A^{n}(t_{n})F(t_{n})$$

These Higher order derivatives are used in the taylor series expansion of $P(t_n + t_i)$, $C_i(tn + t_i)$, $\rho(tn + t_i)$ functions estimations in Eq(5) and Eq(6).

TemperatureFeedback and it'sNumerical Solution

Step change in Reactivity with thermal feedback (adiabatic model), which is mentioned in (Hetrick, 1971; Stacey, 2001) is shown below:

$$\rho(tn) = \rho_0 - \alpha [\text{Temp}(t) - \text{Temp}_0] - \dots \qquad (8)$$

$$\frac{d\text{Temp}(t)}{dt} = \text{Hn}(t) \dots \dots \dots \qquad (9)$$

Here, Temp(t) is the reactor temperature at time 't' and Temp₀ is the initial temperature of the reactor ' α ' is the Temperature coefficient of reactivity, H=AKc, whose units are K/MW.s is the reciprocal of the thermal capacity of the reactor and units of A' are MW.cm3. Differtinating Eq. (8) and substituting in Eq. (9), we obtain:

$$\frac{d\rho(t)}{dt} = -\alpha Hn(t) \dots \dots \dots \dots (10)$$

After applying the procedure of (COX et al,2002) Eq. (9) & (10) can be written in the following integral form:

$$\rho(t_{n+1}) = \rho(t_n)e^{-h/\Lambda} + e^{-\frac{n}{\Lambda}}(-\alpha H \int_0^h n(t_n + \varepsilon)e^{\frac{\varepsilon}{\Lambda}}d\varepsilon + \frac{1.0}{\Lambda}\int_0^h \rho(t_n + \varepsilon)e^{\frac{\varepsilon}{\Lambda}}d\varepsilon) \dots (11)$$

$$\text{Temp}(t_{n+1}) = \text{Temp}(t_n)e^{-h/\Lambda} + e^{-\frac{h}{\Lambda}}(H\int_0^h n(t_n+\tau)e^{\frac{t}{\Lambda}}d\tau + \frac{1.0}{\Lambda}\int_0^h \text{Temp}(t_n+\tau)e^{\frac{t}{\Lambda}}d\tau)). (12)$$

Equations (11)&(12) are solved using Taylor series expansion method, These equations can be used for estimation of temperature and reactivity values at 'tn+1' sec.

Temperature Feedback and it's Numerical Solution Power Transients of thermal reactor caused by step reactivity with Temperature Feedback

ETD-TS method has been applied to solve the point kinetics equations for thermal reactors, in presence of Newtonian temperature feedback. Six groups of delayed neutrons are considered in this problem. The decay constants of neutron precursors, delayed neutron fractions and feedback parameters for thermal reactor are taken as $\lambda 1 = 0.0124s-1$, $\lambda 2 = 0.0305s-1$, $\lambda 3 = 0.111$ sec-1, $\lambda 4 = 0.301$ sec-1, $\lambda 5 = 1.13$ sec-1, $\lambda 6 = 3.00$ sec-1, $\beta 1 = 0.00021$, $\beta 2 = 0.00141$, $\beta 3 = 0.00127$, $\beta 4 = 0.00255$, $\beta 5 = 0.00074$, $\beta 6 = 0.00027$, $\beta eff = 0.00645$, $\Lambda = 0.00005$ (sec), $\alpha = 0.00005$ (K-1) and Kc = 0.05 (K/MWs).

Transient power, reactivity and temperature values initiated by step reactivity addition of 1.0 \$, 1.5\$ and 2.0 \$ with temperature feedback was studied using ETD-TS method. The results are summarized in Table 1, 2 and 3 respectively for 1.0 \$, 1.5\$ and 2.0 \$. Results are compared with other numerical methods ET-BE, ET-CN, ITS2 and CATS method values. It is observed that even with higher step reactivity addition of 2 \$, ETD-TS method is more accurate than ET-BE and ET-CN methods while estimating the transient power and reactivity values. Estimated values using ETD-TS method are giving identical results when compared with highly accurate i.e, ITS2 method values. Integrands in the Equations (4)-(6) & (8) expanded up to five Taylor series terms for better accuracy of ETD-TS method.

In the Table.1 ET-BE, ET-CN and NAM methods are using fixed time step of h=0.001 sec, which is much larger than the time step sizes h=0.01 sec used in the ETD-TS method. computational time for ETD-TS method is 26 sec

for 1.0, 1.5 and 2.0\$ step reactivity insertion case, with Intel \mathbb{R} Core TM i9 computer and processer speed 2.8 GHz, which is less when compared with ITS2 method i.e,28.0 45.0, 52 sec.ITS2 method takes more computational time when compared with ETD-TS method. But, whereas ETD-TS method uses constant computational time for all the three cases.

Table 1:- Comparison of transient values of power and reactivity values initiated due to step reactivity 1.0 \$. * Numbers in the parenthesis denotes powers of 10.

Tim	Power (Wat	tt)			ρ (\$)				Temp
e		-	-			-	-		ETD-
(sec)	ET-BE	ET-CN	ITS-2	ETD-TS	ET-BE	ET-	ITS2	ETD-	TS
	Method	Method	Method	Method(h=0.	Metho	CN	Metho	TS	Method
	Ref.	Ref.	Ref.	01 sec)n=5	d	Metho	d	Metho	([°] K)
					Ref.	d	(Ref.)	d	
						Ref.			
0.0	1.00000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	1.0000	293.000
					0	0	0	0	0
10.0	1.3210E+0	1.3206E+0	1.3203E+0	1.3203E+02	-	-	-	-	446.032
	2	2	2		0.1857	0.1860	0.1862	0.1862	7
					9	4	9	9	
20.0	5.1727E+0	5.1716E+0	5.1699E+0	5.1699E+01	-	-	-	-	487.303
	1	1	1		0.5058	0.5059	0.5062	0.5062	8
					5	5	2	3	
30.0	2.8189E+0	2.8187E+0	2.8174E+0	2.8174E+01	-	-	-	-	506.316
	1	1	1		0.6533	0.6532	0.6536	0.6536	3
10.0					2	1	1	1	
40.0	1.8154E+0	1.8155E+0	1.8146E+0	1.8146E+01	-	-	-	-	517.602
	1	1	1		0.7408	0.7407	0.7411	0.7411	2
50.0	1.07005+0	1.070(E+0	1.0770E+0	1.27705 .01	/	0	0	0	505 012
50.0	1.2/88E+0	1.2/86E+0	1.2779E+0	1.2779E+01	-	-	-	-	525.213
	1	1	1		0.7998	0.8001	0.8001	0.8001	4
60.0	0.4929E+0	0.4909E+0	0.4740E+0	0.4740E+00	2	0	0	0	520 719
00.0	9.4626E+0	9.4696E+0	9.4/49E+0	9.4749E+00	-	-	-	-	350.718
	0	0	0		0.0424	0.6412	0.0427	0.8427	0
70.0	7 2485E±0	7 2324E±0	$7.2444E\pm0$	7.2444E+00	1	2	/	/	53/ 86/
70.0	7.2405E+0	7.2324E+0	7.2444D+0	7.2444L+00	0.8746	0.8781	0.8740	- 0.8740	7
	0	0	0		3	8	1	1	/
80.0	5 6467E+0	5 6534E+0	5 6462E+0	5 6462E+00	-	-	-	-	538.066
00.0	0	0	0	5.01021100	0 8999	0 8977	0 8997	0 8997	7
	Ũ	Ŭ	Ŭ		7	1	4	4	,
90.0	4.4588E+0	4.4684E+0	4.4568E+0	4.4568E+00	-	-	-	-	540.578
	0	0	0		0.9189	0.9161	0.9192	0.9192	6
	-	-	-		2	9	1	1	-
100.	3.5507E+0	3.5531E+0	3.5501E+0	3.5501E+00	-	-	-	-	542.570
0	0	0	0		0.9347	0.9346	0.9346	0.9346	4
					4	8	5	5	

|--|

	ET-BE Method	ET-CN Method	ITS2 Method	ETD-TS Method(h=0.	ET- BE	ET- CN	ITS2 Metho	ETD- TS	
	Ref.	Ref.	Ref.	01 sec)n=5	Metho d Ref.	Metho d Ref.	d (Ref.)	Metho d	
0.0	1.00000	1.0000	1.0000	1.0000	1.5000 0	1.5000 0	1.5000 0	1.5000 0	293.000 0
10.0	1.08546E+ 02	1.08231E+ 02	1.07911E+ 02	1.07911E+02	- 0.8200 8	- 0.8256 2	- 0.8315 0	- 0.8315 2	593.764 7
20.0	4.18325E+ 01	4.17205E+ 01	4.16042E+ 01	4.16042E+01	- 1.0789 9	- 1.0838 0	- 1.0889 5	- 1.0889 7	626.975 6
30.0	2.34123E+ 01	2.33593E+ 01	2.32989E+ 01	2.32989E+01	- 1.1996 1	- 1.2038 1	- 1.2090 2	- 1.2090 3	642.464 4
40.0	1.53686E+ 01	1.53360E+ 01	1.53034E+ 01	1.53034E+01	- 1.2733 1	- 1.2773 2	- 1.2821 0	- 1.2821 1	651.890 7
50.0	1.09320E+ 01	1.09096E+ 01	1.08901E+ 01	1.08901E+01	- 1.3238 4	- 1.3277 3	- 1.3321 3	- 1.3321 4	658.345 4
60.0	8.13017E+ 00	8.11509E+ 00	8.10103E+ 00	8.10104E+00	- 1.3606 8	- 1.3646 9	- 1.3685 7	- 1.3685 8	663.046 0
70.0	6.20461E+ 00	6.19337E+ 00	6.18269E+ 00	6.18269E+00	- 1.3884 9	- 1.3917 7	- 1.3960 4	- 1.3960 5	666.589 5
80.0	4.81225E+ 00	4.81004E+ 00	4.79331E+ 00	4.79331E+00	- 1.4090 4	- 1.4102 5	- 1.4171 7	- 1.4171 8	669.316 0
90.0	3.76957E+ 00	3.76577E+ 00	3.75561E+ 00	3.75561E+00	- 1.4260 6	- 1.4287 4	- 1.4336 5	- 1.4336 6	671.441 1
100. 0	2.97798E+ 00	2.96556E+ 00	2.96607E+ 00	2.96607E+00	- 1.4386 9	- 1.4472 2	- 1.4466 1	- 1.4466 2	673.112 7



Fig.1:- Comparison of Transient Power (Watt) with initial stepreactivity (1.0\$).



Fig.2:- Comparison of Transient Reactivity values with initial step reactivity (1.0\$).









Tim	Power (Wat	tt)			ρ (\$)				Temp
е	ET-BE	ET-CN	ITS2	ETD-TS	ET-BE	ET-	ITS2	ETD-	ETD-
(sec)	Method	Method	Method	Method(h=0.0	Metho	CN	Metho	TS	TS
· ·	Ref.	Ref.	Ref.	1 sec)n=5	d	Metho	d	Metho	Method
				, í	Ref.	d	(Ref.)	d	(⁰ K)
						Ref.	l` í		
0.0	1.00000	1.0000	1.0000	1.0000	2.0000	2.0000	2.0000	2.0000	293.000
					0	0	0	0	0
10.0	1.05111E+	1.04263E+	1.03381E+	1.03381E+	-	-	-	-	727.684
	02	02	02	02	1.3174	1.3420	1.3696	1.3696	9
					0	1	3	5	
20.0	3.97746E+	3.94644E+	3.91388E+	3.91389E+	-	-	-	-	759.122
	01	01	01	01	1.5650	1.5877	1.6133	1.6133	5
					2	6	3	5	
30.0	2.23353E+	2.21750E+	2.20038E+	2.20038E+	-	-	-	-	773.716
	01	01	01	01	1.6801	1.7018	1.7264	1.7264	2
					2	2	7	8	
40.0	1.46961E+	1.45965E+	1.44937E+	1.44937E+	-	-	-	-	782.632
	01	01	01	01	1.7504	1.7710	1.7955	1.7955	0
					6	1	8	9	
50.0	1.04532E+	1.03871E+	1.03186E+	1.03186E+	-	-	-	-	788.747
	01	01	01	01	1.7986	1.8184	1.8429	1.8429	6
					6	1	9	9	
60.0	7.76180E+	7.70673	7.66332E+	7.66336E+	-	-	-	-	793.198
	00	E+00	00	00	1.8332	1.8553	1.8774	1.8775	7
	5 00 (00 T)		5 000 (0 7)	5 000 (0 T)	8	7	9	0	
70.0	5.90482E+	5.86898	5.82940E+	5.82943E+	-	-	-	-	796.545
	00	E+00	00	00	1.8593	1.8/84	1.9034	1.9034	9
	4.55000T	4.52522	4 400 40 7	4.400.4675.	2	2	4	2	700.111
80.0	4.55980E+	4.53523	4.49943E+	4.49945E+	-	-	-	-	799.111
	00	E+00	00	00	1.8/89	1.8969	1.9233	1.9233	2
00.0	2.5555E	0.50051	0.5074071	0.50744EL	0	0	3	4	001 101
90.0	5.000050世+	5.53251 E+00	3.50/42E+	3.50/44E+	-	-	-	-	801.101
	00	E+00	00	00	1.8949	1.9153	1.938/	1.938/	5
100	0.7055AE	0.76655	0.7551071	0.7551472	ð	δ	0	0	000.650
100.	2.79004E+	2.70000 E+00	2.75512E+	2.70014E+	-	-	-	-	802.008
0	00	ETU0	00	00	1.9003	1.9338	1.9508	1.9508	4
					С	0	5	4	

Table 3:- Comparison of transient values of power and reactivity values initiated due to step reactivity (2.0 \$).



Fig.5:- Comparison of Transient Power (Watt) with initial step reactivity (2.0 \$).



Fig.6:- Comparison of Transient Reactivity values with initial step reactivity (2.0 \$).

Table 4. shows transient power values corresponding to different 'h' values using ETD-TS methods and results are compared with ITS2 method values for the transient power estimated up to 100.0 sec. Results are in excellent agreement with reference values.

'able 4:- Comparison of transient values of power and reactivity initiated due to step reactivity (2.0\$) with differe	nt
n' values.	

Time (sec)	Power (Watt)			
	ETD-TS Method (ITS2–Method (Reference)		
	(h=0.1 sec) n=5	(h=0.05 sec) n=5	(h=0.01 sec) n=5	
0.0	1.0000	1.0000	1.0000	1.0000
20.0	38.8049	39.0941	39.1389	39.1389
50.0	10.2598	10.3090	10.3186	10.3186
80.0	4.4771	4.4951	4.4994	4.4994
100.0	2.7415	2.7524	2.7551	2.7551

Similarly, Table 5 shows the comparison of ETD-TS results with NAM (New Analytical Method) (Nahla, 2011), PCA (Piecewise Continuous Approximation method) (Kinard, 2004) and SCS methods. Comparison shows that transient power values estimated with ETD-TS method is quite accurate when compared with NAM and PCA method values. Also, ETD-TS method uses the time step size of (h=0.01 sec), which is larger than the time steps used in NAM and PCA methods (h=0.001 sec). This indicates that present method is efficient and accurate even with finer time steps and the values are in good agreement with most accurate ITS2 method values.

	Power (Watt)		
Time (sec)	NAM Method h=0.001 sec	PCAMethod h=0.001 sec	ETD-TS Method (Observed) (h=0.01 sec) n=5
0.0	1.0000	1.0000	1.0000
20.0	39.1793	39.0022	39.1389
50.0	10.3277	10.2872	10.3186
80.0	4.4855	4.4994	4.4994
100.0	2.7575	2.7463	2.7551

Table 5:- Comparison of ETD – TS power values with differentnumerical methods (with step reactivity of 2.0 \$).

Table.6, Shows comparison of Peak transient power values and corresponding time to reach peak power at the initial step reactivity values i.e,1.0, 1.5 and 2.0 \$ with other numerical methods. ETD-TS Method estimated values are in good agreement with ITS2 method values. From the comparison, it is observed that ETD-TS method is quite accurate and similar values when compared with highly accurate method ITS2 method.

Table 6:- Comparison of ETD – TS power peak values with different numerical methods with different initial reactivity values.

Initial Decetivity	Time to Peak (se	c)	Peak Neutron Density		
$\rho_0(\$)$	ITS2	ETD-TS	ITS2	ETD-TS	
1.0	0.953478	0.953476	8.078681E+02	8.078687E+02	

1.5	0.168288	0.168289	4.302461E+04	4.302461E+04
2.0	0.098390	0.098399	1.678457E+05	1.678456E+05

* Numbers in the parenthesis denotes powers of 10.

Compensated ramp reactivity Insertion with neutron density feedback and its numerical solution

Compensated ramp functions which are defined previously, as in (Aboanber and Hamada, 2002). The reactivity at time 't' can be described by the following equation.

 $\rho(t) = a(t - t_0) - b \int_{t_0}^t n(t') dt \dots (13)$ After applying the procedure in (2002,Cox) the above equation can be rewritten as shown below:

$$\rho(t_{n+1}) = \rho(t_n)e^{-\frac{n}{\Lambda}} + e^{-\frac{n}{\Lambda}} \int_0^{t} (a - b.n(t_n + t) + \frac{1.0}{\Lambda}) dt \dots \dots \dots (14)$$

After expanding $n(t_n+r)$ using Taylor series expansion, we can estimate the reactivity values at different time steps t_{n+1} , using Eq.(14).

Power Transients of thermal reactor caused by compensated ramp reactivity with Power Feedback

We next consider response to insertion of ramp reactivity as shown in Eq (13)in the thermal reactor. The power transients caused by compensated ramp reactivity is considered. Six group kinetic parameters are $\lambda_1 = 0.0124 \text{sec}^{-1}$, λ_2 $= 0.0305 \text{sec}^{-1}, \ \lambda_3 = 0.111 \text{ sec}^{-1}, \ \lambda_4 = 0.301 \text{ sec}^{-1}, \ \lambda_5 = 1.13 \text{ sec}^{-1}, \ \lambda_6 = 3.00 \text{ sec}^{-1}, \ \beta_1 = 0.00021, \ \beta_2 = 0.00141, \ \beta_3 = 0.00127, \ \beta_4 = 0.00255, \ \beta_5 = 0.00074, \ \beta_6 = 0.00027. \ \beta_{\text{eff}} = 0.0065, \ \Lambda = 0.00005 \text{ (sec)}. \ \text{Here, time step chosen for all the}$ calculations using ETD-TS method is h = 0.000025 sec.Neutron Density values are estimated up to 10.0 seconds, with fixed shutdown coefficient $b = 10^{-11}$ cm³ /sec and 'a' values are changed from 0.003-0.1 sec⁻¹. Transient Neutron Density values are in excellent agreement with ITS2method values at all the time steps and results are shown in Table 7-8. Transient Neutron density and reactivity values are plotted against ITS2 and ETD-TS values corresponding to fixed shutdown coefficient ($b=10^{-11}$ cm³/sec) up to 10.0 sec with a=0.003sec⁻¹ and up to 1.0 sec with $a=0.1 \sec^{-1}as$ shown in Fig.6-9. In Most of the cases, Neutron density and Reactivity values are well agreement with most accurate ITS2 method values up to 4decimal places.

Table 7:- Comparison of Compensated response to a ramp change of reactivity with $a = 0.003 \text{ sec}^{-1}$ and $b = 10^{-11} \text{ cm}^3$ /sec.

Time (sec)	Neutron Density	(/cm ³)	Reactivity (\$)	
	ITS-2	ETD-TS	ITS-2	ETD-TS
		n=3 ;h=0.000125 s		n=3 ;h=0.000125 s
0	1.0000	1.000	0.0	0.0
0.5	1.3247E+00	1.3247E+00	2.3256E-01	2.3255E-01
1.0	2.0532E+00	2.0532E+00	4.6512E-01	4.6511E-01
1.5	4.3472E+00	4.3472E+00	6.9767E-01	6.9767E-01
2.0	2.3921E+01	2.3922E+01	9.3023E-01	9.3024E-01
2.5	1.4390E+04	1.4397E+04	1.1628E+00	1.1628E+00
3.0	8.0571E+08	8.0525E+08	6.4673E-01	6.4669E-01
4.0	3.4133E+08	3.4134E+08	5.0440E-01	5.0439E-01
5.0	3.2933E+08	3.2934E+08	4.5283E-01	4.5283E-01
10.0	3.1459E+08	3.1460E+08	2.7934E-01	2.7934E-01

* Numbers in the parenthesis denotes powers of 10.



Fig. 6:- Comparison plot of Reactivity (\$) in compensated ramp with $a = 0.003 \text{ sec}^{-1}$ and $b = 10^{-11} \text{ cm}^3/\text{sec}$.



Fig. 7:- Comparison plot of Neutron density in compensated rampwith $a = 0.003 \text{ sec}^{-1}$ and $b = 10^{-11} \text{ cm}^{3}/\text{sec}$.

Table 8:	 Comparison o 	f Compensated re	sponse to a ram	p change of reactivit	$y a = 0.1 \text{ sec}^-$	and $b = 10^{-1}$	¹ cm ³ /sec.
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Time (sec)	Neutron Density (/cm ³)		Reactivity (\$)	
	ITS-2	ETD-TS	ITS-2	ETD-TS
		n=3 ;h=0.000125 s		n=3 ;h=0.000125 s
0.00	1.00	1.00	0.00	0.00
0.10	2.4734E+01	2.4730E+01	1.5504E+00	1.5504E+00
0.20	7.2379E+08	7.2371E+08	3.0965E+00	3.0964E+00
0.30	8.1620E+08	8.1619E+08	-3.8677E-01	-3.8682E-01
0.40	3.7509E+09	3.7504E+09	9.0487E-01	9.0481E-01
0.50	9.9500E+09	9.9503E+09	7.2456E-01	7.2456E-01
0.60	1.1466E+10	1.1466E+10	8.7409E-01	8.7411E-01
0.70	9.7644E+09	9.7643E+09	8.2555E-01	8.2555E-01
0.80	1.0086E+10	1.0086E+10	7.9324E-01	7.9324E-01
0.90	1.0153E+10	1.0153E+10	7.8014E-01	7.8014E-01
1.00	1.0104E+10	1.0104E+10	7.6207E-01	7.6207E-01

* Numbers in the parenthesis denotes powers of 10.







Fig. 9:- Comparison plot of Neutron density in compensated rampwith $a = 0.1 \text{ sec}^{-1}$ and $b = 10^{-11} \text{ cm}^3/\text{sec}$.

Similarly, Neutron Density values are estimated up to 10.0 seconds, with fixed shutdown coefficient b = 10-13 cm³ /sec and for three different 'a' values 0.003,0.01 and 0.1 sec⁻¹. Transient Neutron Density values are in excellent agreement with other accurate methods likeEPCA and CATS at all the time steps and results are shown in Table 9. Transient Neutron density and reactivity values are plotted against CATS and ETD-TS values corresponding to fixed shutdown coefficient (10^{-13} cm³ /sec) up to 10.0 sec as shown in Fig.10. In Most of the cases, Neutron density and Reactivity values are well agreement with most accurate CATS method.

Time (sec)	EPCA	ITS-2	ETD-TS			
	(Reference)	(Reference)	(Observed) n=3 (h=0.000125 sec)			
Case 1. $a = 0.1 (sec^{-1})$						
0.1	2.4734E+01	2.4734E+01	2.4730E+01			
0.5	1.5434E+12	1.5434E+12	1.5434E+12			
5.0	1.0030E+12	1.0030E+12	1.0030E+12			
7.5	1.0018E+12	1.0018E+12	1.0018E+12			
10.0	1.0012E+12	1.0012E+12	1.0012E+12			
Case 2. $a = 0.01 (sec^{-1})$						
0.1	1.1672E+00	1.1672E+00	1.1672E+00			
0.5	4.2699E+00	4.2700E+00	4.2697E+00			
5.0	1.0339E+11	1.0339E+11	1.0339E+11			
7.5	1.0195E+11	1.0195E+11	1.0195E+11			
10.0	1.0124E+11	1.0124E+11	1.0124E+11			
Case 3. a =0.003 (sec-1)						

Table 9:- Comparison of Neutron densities in compensated ramp changes of reactivity with $b = 10^{-13}$ cm³/secfor different 'a' values.

0.1	1.0453E+00	1.0453E+00	1.0453E+00
0.5	1.3246E+00	1.3246E+00	1.3246E+00
5.0	3.2156E+10	3.2156E+10	3.2156E+10
7.5	3.2102E+10	3.2102E+10	3.2102E+10
10.0	3.1456E+10	3.1456E+10	3.1456E+10



Fig. 10:- Comparison plot of Neutron density in compensated ramp change with $b = 10^{-11}$ cm3/sec and different a (sec-1) values.

Conclusion:-

ETD-TS method is used for solving point kinetics equations with multi group of delayed neutrons in the presence of temperature and power feedback. Thermal reactor benchmark problems with stepand compensated ramp reactivity insertions are considered for validation of this method. In ETD-TS method integral form of PKEs are solved using Taylor Series expansion method with temperature and power feedbacks. Time constant '-1/ Λ ' is chosen as stiffness constant. PKEs are integrated using exponential factor. Smaller time step ~ 10⁻²sec is sufficient for estimation of power transients with step reactivity insertion. But, for transients with compensated ramp reactivity insertion, time step size ~10⁻⁴sec was used in ETD-TS method to estimate the power transients. Computational time is found to be relatively less without any loss of accuracy when compared with ITS2 method. It is observed that ETD-TS method estimates the power transients for longer duration of time with good accuracy. The method was found to give accurate results similar to PCA and NAM methods even with larger time steps in the case of compensated ramp reactivity insertion case. ETD-TS method can be further used for estimation of power transients in 3D space-time calculations.

References:-

- 1. Kinard, M., Allen, E.J., 2004. Efficient Numerical Solution of Point Kinetics Equations in NuclearReactor Dynamics, Annals of nuclear energy, vol. 31, page 1039-1051.
- 2. Nahla, A.A., 2011. An efficient technique for point reactor kinetics equations with Newtonian temperature feedback effects, Annals of nuclear energy, vol.38,2810.
- 3. Ganapol, B., Picca, P., Previti, A., Mostacci, D., 2012. The solution of the point kinetics equations via converged accelerated Taylor series (CATS). In: Advances in Reactor PhysicseLinking Research, Industry and Education (PHYSOR 2012), Knoxville, Tennessee, USA, April 15-20, 2012.
- 4. Picca, P., Furfaro, R., Ganapol, B.D., 2013. A highly accurate technique for the solution of the non-linear point kinetics equations. Ann. Nucl. Energy 58, 43.
- 5. Serigo Q.B.L., Marco, T. D. V., 2016. Solution of the point reactor kinetics equations with temperature feedback by the ITS2 method, Progress in Nuclear Energy, Vol .91, pages 240-249.
- 6. Hamada, Y.M., 2018. A New Accurate Numerical Method Based on Shifted Chebyshev Series for Nuclear Reactor Dynamical Systems. Sci. Technol. Nucl. Install. 2018.
- 7. Cox, S., M., Mathews, P., C., 2002. Exponential time differencing method for stiff systems, J. Comput. Phys., vol. 176, pages 430-455.
- Mohideen Abdul Razak, M., Devan, K., Shathiyasheela, T., 2015. The modified exponential time differencing (ETD) method for solving the reactor point kinetics equations, Annals of nuclear energy, vol. 76, pages 193-199.
- 9. Hetrick, D.L., 1971. Dynamics of Nuclear Reactors. University of Chicago Press, Chicago and London.
- 10. Stacey, W.M., 2001. Nuclear Reactor Physics. John Wiley and Sons Inc., USA
- 11. Aboanber, A.E., Hamada, Y.M., 2002. PWS: an efficient code system for solving space-independent nuclear reactor dynamics. Ann. Nucl. Energy 29, 2159.