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Glossary of Words [165-167]

Activation energy (Q): The energy required to initiate a reaction, such as diffusion.

Anion: A negatively charged, nonmetallic ion

Anode: The electrode in an electrochemical cell or galvanic couple that experiences oxidation, or gives up electrons.

Capacitance (C): The charge-storing ability of a capacitor, defined as the magnitude of charge stored on either plate divided by the applied voltage

Carburizing. The process by which the surface carbon concentration of a ferrous alloy is increased by diffusion from the surrounding environment.

Cathode. The electrode in an electrochemical cell or galvanic couple at which a reduction reaction occurs.

Cation. A positively charged metallic ion

Corrosion. Deteriorative loss of a metal as a result of dissolution environmental reactions

Corrosion rate (CR). Thickness loss of material per unit of time as a result of corrosion; usually expressed in terms of mils per year or millimeters per year.

Diffraction (**x-ray**). Constructive interference of x-ray beams that are scattered by atoms of a crystal.

Diffusion. Mass transport by atomic motion.

Diffusion coefficient (D). The constant of proportionality between the diffusion flux and the concentration gradient in Fick's first law. Its magnitude is indicative of the rate of atomic diffusion.

Dislocation. A linear crystalline defect around which there is atomic misalignment. Plastic deformation corresponds to the motion of dislocations in response to an applied shear stress.

Ductile-to-brittle transition. The transition from ductile to brittle behavior with a decrease in temperature exhibited by some low-strength steel (BCC) alloys; the temperature range over which the transition occurs is determined by Charpy and Izod impact tests

Ductility. A measure of a material's ability to undergo appreciable plastic deformation before fracture; it may be expressed as percent elongation (%EL) or percent reduction in area (%RA) from a tensile test.

Erosion–corrosion. A form of corrosion that arises from the combined action of chemical attack and mechanical wear.

Ferrite (iron). Body-centered cubic iron; also, iron and steel alloys that have the BCC crystal structure.

Grain. An individual crystal in a polycrystalline metal or ceramic.

Grain boundary. The interface separating two adjoining grains having different crystallographic orientations

Grain size. The average grain diameter as determined from a random cross section

Hardness. The measure of a material's resistance to deformation by surface indentation or by abrasion

Hexagonal close-packed (HCP). A crystal structure found for some metals. The HCP unit cell is of hexagonal geometry and is generated by the stacking of close-packed planes of atoms.

Intergranular corrosion. Preferential corrosion along grain-boundary regions of polycrystalline materials

Isothermal transformation (T-T-T) **diagram.** A plot of temperature versus the logarithm of time for a steel alloy of definite composition. Used to determine when transformations begin and end for an isothermal (constant-temperature) heat treatment of a previously austenitized alloy.

Kinetics. The study of reaction rates and the factors that affect them.

Lattice. The regular geometrical arrangement of points in crystal space.

Metallic bond. A primary interatomic bond involving the nondirectional sharing of nonlocalized valence electrons ("sea of electrons") that are mutually shared by all the atoms in the metallic solid.

Microstructure. The structural features of an alloy (e.g., grain and phase structure) that are subject to observation under a microscope.

Octahedral position. The void space among close-packed, hardsphere atoms or ions for which there are six nearest neighbors. An octahedron (double pyramid) is circumscribed by lines constructed from centers of adjacent spheres.

Oxidation. The removal of one or more electrons from an atom, ion, or molecule.

Passivity. The loss of chemical reactivity, under particular environmental conditions, by some active metals and alloys, often due to the formation of a protective film

Phase. A homogeneous portion of a system that has uniform physical and chemical characteristics

Pitting. A form of very localized corrosion wherein small pits or holes form, usually in a vertical direction.

Plastic deformation. Deformation that is permanent or nonrecoverable after release of the applied load. It is accompanied by permanent atomic displacements

Polarization (corrosion). The displacement of an electrode potential from its equilibrium value as a result of current flow

Reduction. The addition of one or more electrons to an atom, ion, or molecule

Slip. Plastic deformation as the result of dislocation motion; also, the shear displacement of two adjacent planes of atoms.

Solid-solution strengthening. Hardening and strengthening of metals that result from alloying in which a solid solution is formed. The presence of impurity atoms restricts dislocation mobility

Strain hardening. The increase in hardness and strength of a ductile metal as it is plastically deformed below its recrystallization temperature.

Tetrahedral position. The void space among close-packed, hardsphere atoms or ions for which there are four nearest neighbors.

Thermal stress. A residual stress introduced within a body resulting from a change in temperature

Toughness. The ability of a material to absorb energy and plastically deform before fracturing;

Yield strength (*y*). The stress required to produce a very slight yet specified amount of plastic strain; a strain offset of 0.002 is commonly used.

Appendices

Appendix A:	A brief literature survey on high temperature oxidation of austenitic
	stainless steels.

S.No.	Alloy	Temperature	Environment	Findings	References
1.	17.8Mn,	700-1000 °C	Air Oxidation	Three layered	Douglas et al.
	9.5 Cr, 1.0		1500 min	structures.	[34]
	Ni, 0.27 C,			No internal oxidation	
	and 0.03 N			up to 700°C	
2.	Mn-	800-1000°C	O ₂ , 100 h	MnCr ₂ O ₄ at the	FH Scott et
	containing			interface	al.[100]
	Fe-28% Cr			Depletion of Mn at	
	alloys			surface.	
3.	Fe-18Cr-	525, 625, and	Air Oxidation	Interstitials (N) does	James
	18Mn	725°C	500 h	not affect the	Rawers.
	stainless			oxidation.	[35]
	steels			Two layered	
				structure of oxide	
				layer.	
4.	304L steel	500-800°C	Humid air	Cr evaporation in	H Asteman et
			0.02–13 cm/sec	500-800°C	al.
			flow rate 168 h	High flow rate early	[168]
				breakdown (800°C).	[]
5.	Fe-Cr-Mn-	800°C	Ageing/oxidati	Precipitation of Cr2N	B Kartik et al
	N (Ni-free)		on	σ phase above 850°C	[13]
			200 h	Mn, Fe and Cr oxide	
				layer	
				Mn-depleted ferrite	
				zone	
6.	Low Nickel	600-700°С	Dry air	Oxide layer shows	FJ Preez et al.
	austenitic		500 cycles of 1	good adhesion to	[97]
	stainless		h	surface.	
	steel			FCC to BCC	
				transition at the	
				surface reduces the	
				thermal stresses	
7.	Fe-Cr18-	950-1150°С	Static air for 10	Parabolic rate law	Xianli Liu et
	Mn16-		h	was followed and	al.[157]
	N0.83-			Mn ₃ O ₄ oxide layer	
	Nb0.15			was formed	
8.	Nitronic 32	800-1000°C	Static air for 24	Presence of Mn	F.Gesmundo
			h	deteriorates the	et al.[169]
				oxidation resistance.	
9.	Fe-9Cr, Fe-	700°C	500 Cycle of 1	Cr vaporization was	N.K. Othman
	17Cr and		hr Ar-20O ₂ , Ar-	observed with	et al.
	Fe-25Cr		$20O_2$ -5H ₂ O and	increasing Cr content	[170]
			Ar-5O ₂ -20H ₂ O	in steels. Water vapor	
				presence increases	
				the formation of	
				Chromium oxide.	
10.	Cr_2O_3 and	850-1050°C	50-1000 h	Cr ₂ O ₃ shows higher	Alexander
	MnCr ₂ O ₄		Synthetic air	Cr volatilization	Stenzel et al.
			with 10% H ₂ O	compared to	[171]
			(5 l/h)	MnCr ₂ O ₄ in humid	
				air environments.	

S.No.	Alloy	Temperature	Environment	Findings	References
1.	304L	650°C	25%CO+	Carbide's formation	P. Szakalos
			$3\%H_2O + H_2$	followed by oxidation.	et al.
			1000 h		[49]
2	Ea and Ni	550 650°C	U.O.(0.21	Dependencial offect of the U.O.	A Douoir
۷.	hased allows	550-650 C	$H_2 \cup (0-21)$ wt%) CO (31-	for low allowed steels	A. KOUAIX- Vande Put
	based anoys		$43 \text{ wt\%}) \text{ H}_2$	Do not prevent pitting	et al $[62]$
			(31-59wt%),	Thick oxide layer was	
			$CO_2(2-6 \text{ wt\%})$	formed.	
			500 h		
3.	Ni based	550-750°C	$20 \text{ vol}\%\text{H}_2$	Experimental Setup affects	F. Di
	alloys		and 80 vol%	initiation and damage	Gabriele et
			100-1000 h	sample noider affects the	a1.[1/2]
			100-1000 II	process	
4.	Fe-Ni-Cr-	650°C	50%CO-	Thermal and Mechanical	Jianqiang
	Al based		$49\% H_{2}$	stress causes spallation and	Zhang
	alloys		1%H ₂ O	initiation of metal dusting.	et al.[173]
			325 cycles of		
			1 h		
5.	Fe–Cr–C	680°C	68% CO–26%	Spallation of Cr ₂ O ₃ due to	C. H. Toh
	and Fe-Ni-		H2-6% H2O	thermal cycle. Internal	et al.[59]
	Cr–C		200 h	Carburization (Fe	
	systems	105000		containing).	
6.	$MnCr_2O_4$	1050°C	2%CH4 +	Better performance of	Hao Li
	sninel		98% HZ at a flow rate of	$\operatorname{America}_{2O4}$ than Cr_{2O3} due to	et al.
	spiner		55.4 mL/min	manganese	[107]
			100 h	manganese	
7.	Inconel 601	650°C	50CO:50H2	Defects in coating in	C.M. Chun
	(substrate)		gas 160 h and	NiCrAlY leads to	et al.
	NiCrAlY.		1000 h	carburization and metal	[174]
8	800HT	570 ∘C and	47.2500-	uusung. Initial carburization and	Aurélien
0.	000111	atmospheric	47.25H ₂ -	subsequent oxidation were	Fabas et
		pressure	$5.5H_2O$ gas	the primary mechanism of	al.[83]
			mixture for	metal dusting	
			4000 h.	-	
9	Fe and Ni	350°C to	H ₂ :CO ratio	Highest metal dusting rate	C.M. Chun
	foils of	1050°C.	ranging from	with 1:1 ratio. Metal	et al.[175]
	99.99%		2:98 to 90:10	disintegration through	
			for 500 h	graphite tubes.	
10.	304	700°C	CO/H ₂ /H ₂ O	Internal carbide precipitation	Jianqiang
	Stainless		for 300 h.	and volume expansion was	Zhang et al.
	steels			primary mechanism	[55]

Appendix B: Metal dusting of behavior of various austenitic stainless steels.

Appendices

				1	
11.	Ni based	550 °C	10% CO in Ar	The higher carbon formation	P.V.D.S.
	Inconel 601		$(a_{\rm C} >> 1)$ gas	appears to be associated with	Gunawarda
			mixture for 20	inclusion of Ni and/or Fe	na et
			h	species in the surface oxide	al[176]
				layer.	
12.	Austenitic	593°C	53.4H2-	Ni based alloys performs	Z. Zeng
	stainless		5.7CO ₂ -	better than Fe base alloys.	[177]
	and nickel-		18.4CO-	High humidity can reduce	
	based		$22.5H_2O$ for	the susceptibility of the	
	superalloys.		1000 h	alloys to metal dusting	
				attack.	
13.	Pure iron	700°C	24.81–94.81	Heavy deposition of coke	J
	sample.		vol% H ₂ , 5–75	with iron carbide particle.	Zhang[178]
			vol% CO and	Increasing CO increases the	
			0.19 vol%	intensity of attack.	
			H_2O		
			for 4 h		
14.	Alloy 800,	600 °C	24vol %CO,	Surface finish influences the	HJ
	18Cr 8Ni		74vol %H ₂ ,	metal dusting attack. Easy Cr	Grabke[110
	steel,		2vol %H ₂ O	diffusion in ferritic stainless-]
	10CrMo 9			steel leads to Cr_2O_3	
	10, P 91			formation.	
15.	304L, 321,	650°C	CO-18.9%	Ferritic material (430&441)	GA
	316, 316L		H ₂ -79.1%,	performed better than that of	Slabbert et
	type 430		$H_2O-2\%$ for	austenitic material.	al.
	and type		30 days		[179]
	441				
16	Austanitia	70090	A = 750/ CO	Deterioretion of orida lawar	Num 1
10.	Austennuc SS 204	700 C	AI-75% CO_2	ot higher temperature and it	nurui
	35 304		75% CO:	at inglier temperature and it	chariff of
			12% H ₂ O for	of water vapor	
			10 20 30 40	of water vapor.	ai.[100]
			and 50 h		
17	Fe20%Cr32	600 °C	24%CO-	Addition of W Nh and Mo	S Strauß*
17.	% Ni	000 C	24%CO- 74%H ₂₋	initially retard the metal	et al [181]
	balanced		2% H ₂ O for	dusting and after	
	balancea.		1500 h	precipitation of carbides	
			1500 II	metal dusting increases	
18	High	600°C	N2-CO-CO2-	Internal carburization of the	L Niewolak
10.	chromium	000 0	H ₂ O	steel results in formation of	et al [182]
	ferritic and		for 300 h	(chromium)	
	austenitic		101 200 11	carbide precipitates.	
	steels				
19.	MnCr ₂ O ₄	1050°C	2%CH4 +	MnCr ₂ O ₄ spinel was found to	Hao Li et
	and Cr ₂ O ₃		98%H ₂ for	be better than Cr_2O_3 .	al.[109]
			100 h		· · · · · · · · j
20	Cr/Cr	800 °C	CH. U. for	Improvement of motol	I Mala
20.	$C1/C1_2O_3$	000 C	20 h	dusting resistance due to	L. IVICIO- Mávimo of
	deposited		20 11	Cr_2O_2 costing	al [107]
	on 304I			Cr2O3 Coaung.	a.[10/]
1		1			1

S.No.	Alloy	Temperature	Erodent & Parameter	Findings	References
1.	13/4 martensitic and 21-4-N nitronic steels	Room Temperature	SiC, 120 m/s, 30° and 90°, 120 min	12-4-N steel performed better than martensitic grade due to superior mechanical properties.	AK Chauhan et al. 2008[77]
2.	AISI 310S, AISI 316, AISI 1020	Room Temperature	Al ₂ O ₃ , 15°-90° with impact velocity of 30- 100m/s	Erosion rate increases with decrease in hardness of material. Ductile mechanism of erosion.	J Malik et al. [74]
3.	AISI 316 and 304L	25,350,650°C	SiO ₂ , 15-45° with impact velocity of 25 m/s	Erosion oxidation was prominent. Quick formation of thin oxide layer.	M Antovo et al.[130]
4.	Cast 13/4, Cast 21-4-N and hot rolled 21-4-N	Room Temperature	SiC, 30 and 90° with impact velocity of 120m/s for 120 min	Erosion resistance of cast 13/4 was better while 21-4-N in hot rolled condition performed better.	A.K Chauhan et al. 2009[79]
5.	SUS 304	Room Temperature	Al ₂ O ₃ , 0-90 ^o with impact velocity of 200m/s	Micro-cutting and ploughing were the primary mechanism of erosion.	QB Nguyen et al.[183]
6.	316L and coated with WC-Cr ₃ C ₂ – Ni	500-650°C	Al ₂ O ₃ at 30 and 90° impact angle with impact velocity of 30 m/s for 10 min.	Uncoated samples performed better than coated samples owing to lower hardness of uncoated samples.	DG Bhosale et al. 2020[184]
7.	High Cr cast iron	900°C	Alumina at 30, 60 and 90° with discharge rate of 26 g/sec.	Type of heat treatment influences the erosion rate. Better performance of as quenched samples.	Kazumichi Shimizu et al. [185]
8.	Borided X12CrNiMoV12- 3 steel	Room temperature and 400°C	$\begin{array}{c} \text{SiO}_2 \text{ at } 30, 60 \\ \text{and } 90^\circ \text{ with} \\ 30 \text{ m/sec for} \\ 10 \text{ min} \end{array}$	Coating of FeB and FeB $_2$ on steel increases the erosion resistance.	A. Ruiz-Rio et al.[186]
9.	AISI 304, 316 and 420 stainless steels	Room temperature	SiC at 30-90° at impact velocity of 24 m/sec for 10 min.	AISI 420 shows best resistance to erosion and it shows ductile Behavior for erosion.	J.R. Laguna- Camacho et al.[187]
10.	SUS 304	Room temperature	Al ₂ O ₃ at impact angle of 0-90 ^o at	Plastic deformation induced indentation, ploughing and cutting	QB Nguyen et al. 2019 [188]

			velocity of 30m/sec	was primary mechanism of erosion.	
11.	316L (Cr3C225 (Ni20Cr) coating)	Room temperature, 200, 400 and 600°C	Alumina at impact angle of 30, 60 and 90° with velocity of 100m/sec	For both coated and uncoated specimen ductile Behavior of erosion. Increase in temperature increases erosion rate.	Hemant Nautiyal et al.[138]
12.	SS 304	Room Temperature	Alumina at impact angle of 30 and 60° at velocity of 40m/sec	Highest metal removal was at 30° impingement angle and alloy exhibited ductile Behavior.	Mayank Patel et al.[189]
13.	304, 316 and 410 stainless steels	Room temperature	SiC with impact angle of 30, 60 and 90° at velocity of 100 m/sec	410 SS shows better erosion resistance. Soft zone formation beneath the eroded layer in 410 SS compared to 304 and 316.	Trilok Singh et al.[190]
14.	Martensitic and nitrogen alloyed austenitic stainless steels	Room temperature	Alumina at impact angle of 30, 45, 60 and 90° with impact velocity of 41m/sec	Mechanical properties affect the erosion rate. At oblique and normal angle of impingement lower erosion rate for nitrogen alloyed stainless steel.	Ashish Selokar et al.[191]
15.	AISI 444, AIS 439 and AISI 304	Room temperature	Silica sand with impact velocity of 40, 66 and 85 m/sec at impact angle of 15, 30, 45 and 90°	Peak erosion at 30° impingement angle. Hutching's normal model of erosion was followed.	A Aazad Hussian et al.[192]
16.	Nitrogen alloyed austenitic stainless steel and 316L	Room temperature	Alumina at 30, 60 and 90° impingement angle with impact velocity of 41m/sec	Nitrogen alloyed austenitic stainless steel performed better than 316L due its superior mechanical properties.	Ashish Shelokar et al. [139]
17.	AISI 446	550,650 and 750°C	Al ₂ O ₃ with angle of 60, 75 and 90° at velocity of 100 m/sec	Erosion rate increases with increase in temperature.	A Mishra et al. [125]
18.	API X120	Room temperature	Al ₂ O ₃ at 30-90 with velocity of 43-167 m/sec	With increase in impingement angle and velocity erosion rate increases.	Paul C. Okonkwo et al.[193]

S.No.	Alloy	Ageing	Environment	Findings	References
1.	Fe-17Cr- 21Mn-0.59N	700, 800 and 900°C for 14 h	0.5 M NaCl solution, Cyclic potentiodynamic Test	Precipitation of Cr ₂ N. Disc shape precipitates increase the pitting resistance.	K.Krishna Kumar et al.[80]
2.	Type 316L and Ni free high Mn austenitic stainless steel	650 C for 2 h	0.5% NaCl and acidic solution of 0.5 M NaCl + 0.5 M H_2SO_4	Solution treated Ni free austenitic stainless steel show high corrosion resistance compared to 316L in both the solutions.	Xinqiang Wu et al. [15]
3.	High nitrogen austenitic stainless steel	Solution treated condition	0.05 M H ₂ SO ₄ + 0.5 M NaCl and 0.05 M H ₂ SO ₄ + 0.5 M Na ₂ SO ₄	Alloys exhibit self- passive Behavior in both solutions. Cl diffusion leads to passive film breakage.	Y.X. Qiao et al.[194]
4.	High manganese nickel free stainless steel with N variation	Room temperature	3.5% NaCl and 0.5 M H ₂ SO + 0.5 M NaCl solutions	Increase in N content enhances the dissolution resistance in both the solutions.	Yao Fu et al.[195]
5.	High nitrogen stainless steel Effect of cold work and Sensitization	650 °C for 2 h	0.5 M H ₂ SO ₄ + 0.5 M NaCl, 3.5% NaCl and 0.5 M NaOH + 0.5 M NaCl	Increasing cold work increases the corrosion rate. Precipitation of Chi phase during the sensitization.	Yao Fu et al. 2009 [196]
6.	Fe-18.4Cr- 15.8Mn- 2.2Mo- 0.66N-0.04C	Room temperature	3.5 % NaCl	Ultrafine grain size increases the ability of oxide film to re- passivate easily.	H. Zhang et al.[197]
7.	High nitrogen CrMn stainless steel.	875°C for 2 h	3.5 % NaCl solution at 30°C	Increasing amount of low angle grain boundary will increase in corrosion resistance.	Jianjun Qi et al.[198]
8.	High nitrogen nickel free austenitic stainless steel and 316L	Room temperature	0.9%NaCl solution (saline), phosphate- buffered saline, Hanks' solution	High nitrogen nickel free stainless steel shows lower corrosion rates and has advantage for re- passivation in all solutions	Daisuke Kuroda et al.[199]
9.	High nitrogen nickel free	Room temperature	3.5 % NaCl	Increase in applied potential leads to deterioration in passive film	H. Shi et al.[200]

Appendix D: Aqueous corrosion Behav	or of high nitrogen austenitic stainless steels.
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	austenitic stainless steel				
10.	Fe-16Cr-Mn- Mo-N (Varying Mo and N)	Room Temperature	3.5 % NaCl	N content increases the passive Behavior of film while Mo addition enhances the pitting resistance.	KL Chao et al.[81]
11.	19Cr-18Mn- 0.69N, Type 316L SS and 14Cr-8-Mn SS	Room Temperature	3.5% NaCl	Mn sulfide inclusions reduces the combined efficiency of Mo and N towards pitting corrosion.	P. Saravanan et al.[201]
12.	Fe-19% Cr- 5% Ni-5% Mn-3% Mo- 0.024% C- 0.69% N	Ageing at 400-1000°C for 0.1-1000 h of samples.	19.5 N H ₂ SO ₄ + 0.01 M KSCN	Corrosion resistance decreases with ageing time and temperature owing to Cr ₂ N precipitation.	B.S. Covino et al.[202]
13.	Nickel free manganese bearing stainless steel	Room temperature	0.1 M NaOH solutions with 1–5 M NaCl	Passive film contains chromium oxy- hydroxides. Depletion of Mn below the passive film.	B. Elsener et al.[203]
14.	High nitrogen nickel free stainless steel.	Room temperature	0.1 M NaCl, 0.5 M NaCl, and 0.05 M H 2SO ₄ + 0.5 M NaCl	High nitrogen nickel free stainless steel possesses excellent corrosion resistance in acidic chloride environment	Yanxin Qiao et al.[23]
15.	1Cr18Ni9Ti austenitic stainless steel	Room temperature	3% NaCl test aqueous solution	Two layered passive layer structure formed first with iron hydroxides followed by chromium oxides/hydroxides.	M. K. Lei et al.[204]
16.	Cr23N1.2 and 304 austenitic stainless steels.	20, 40, 60 and 80°C of electrolyte	3.5 % NaCl solution	Increasing temperature of solution leads to increase in corrosion rate. No re- passivation occurred at all temperature.	B. R. Tzaneva[205]
17.	High nitrogen nickel free and conventional austenitic stainless steel	Room temperature	3.5 wt.% NaCl solutions	NH ₃ molecularly adsorbed on the Cr ₂ O ₃ passive film improves resistance of film and initiation of pitting corrosion on high nitrogen steel.	Shicheng Sun et al.[151]
18.	Mn–Cu–C-N austenitic	Koom temperature	0.5M H ₂ SO ₄	Cu was shown to	M. Milititsky et al.[206]

Appendices

	stainless steel and AISI 304		and 0.5M H ₂ SO ₄ plus 0.4M NaCl solutions	be beneficial in decreasing the dissolution current values. Nitrogen enhances the re- passivation ability.	
19.	Fe-22Cr- 1.9Ni- 2.3Mo-0.2N- xMn	Ageing at 800°C from 30-930 min	3.5 wt.% NaCl solutions	Precipitation of σ phase and increase in Mn content leads to deterioration of corrosion resistance.	Zaiqiang Feng et al.[207]

List of Publications

- Sharvan Kumar and G. S. Mahobia. "Cyclic oxidation of Fe–18Cr–21Mn–0.65 N austenitic stainless steel at 400–700° C." *Transactions of the Indian Institute of Metals* 73.10 (2020): 2457-2470.
- Sharvan Kumar and G. S. Mahobia. "The features of metal dusting process in the extremely low nickel austenitic stainless steel (18Cr-21Mn-0.65 N-Fe)." *Corrosion Science* 176 (2020): 108926.
- Sharvan Kumar, Ankitendran Mishra, Sunil Mohan and G.S. Mahobia. "The solid particle erosion of pre oxidized high manganese nitrogen stabilized austenitic stainless steel (18Cr-21Mn-0.65 N-Fe) at 400 to 700° C." *Surface Topography: Metrology and Properties* 9.3 (2021): 035002.
- Sharvan Kumar, Dheeraj Jaiswal, C.K. Behera and G.S. Mahobia Potentiodyanmic Corrosion of Fe-18Cr-21Mn-0.65N austenitic stainless steel at 400-700°C. (Communicated)