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Influence of Cr₂AlC on the thermal shock and corrosion resistance of low carbon Al₂O₃-C refractory

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1 Background

2 Experiment

3 Results

4 Conclusions





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1. Background

Steel products are now widely used in various fields, and steel metallurgy technology provides important support for the development of modern industry. At present, continuous casting technology is widely used in the steel industry. The steel smelting and continuous casting system is generally composed of carbon composite functional refractories.









1. Background



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Chemical attack (corrosion)

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- Physical/mechanical wear (such as erosion/abrasion)
 - **Oxidation** behavior
 - **Thermal shocks**



Formation of micro-cracks inside the carbon composite refractory, which may fail suddenly in extreme cases.





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1. Background





Addition of nano-BN



20.0 kV 3.0 20000x SE 10.0 33945 AS verkekt 1000*C

Addition of alumina sheets

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These nanoscale carbon sources and additions played a variety of roles.

- The promotion of whisker growth
- Larger surface area and higher reactivity

The improving the performance of Al₂O₃-C refractory
The reduction carbon content

V. Roungos, C.G. Aneziris, Improved thermal shock performance of Al₂O₃-C refractories due to nanoscaled additives, Ceram. Int. 38 (2) (2012) 919–927.

Y.X. Luo, X. Wang, Z.L. Liu, et al. Enhanced oxidation resistance and thermal shock resistance of low-carbon Al₂O₃-C refractories with nano-BN: a synergistic of mullitization behavior, J. Alloys Compd. 25 (2024) 172937.



1. Background

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MAX phase is a new type of ternary layered compound with highly ordered structure.







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3. Results Mechanical properties

		Linear change (%)	Bulk density (g/cm ³)	Apparent porosity (%)	Modulus of Elasticity (GPa)	Rupture strength (MPa)
AC-9	220°C×24h	00	2.90±0.04	15.59±0.06	23.78±3.33	10.03±2.41
	1100°C×3h	0.07±0.08	2.88±0.05	16.86±0.99	16.58±3.20	5.72±1.52
	1400°C×3h	0.09±0.05	2.89±0.09	17.38±2.95	24.02±1.89	9.35±0.33
	1600°C×3h	0.06±0.05	2.85±0.04	19.78±0.14	24.43±0.23	10.27±0.61
ACr-2.5	220°C×24h	÷	2.97±0.01	14.62±0.36	28.70±6.00	10.21±0.91
	1100°C×3h	0.05±0.02	2.97±0.01	16.47±0.59	15.89±1.53	4.96±0.71
	1400°C×3h	0.05±0.05	2.95±0.05	17.26±1.43	31.20±1.38	9.22±0.57
	1600°C×3h	0.28±0.06	2.91±0.04	19.88±0.89	24.43±1.47	6.38±0.72
ACr-5	220°C×24h	(. .)	3.00±0.01	14.66±0.31	31.95±1.78	9.45±1.27
	1100°C×3h	0.03±0.02	3.00±0.01	17.46±0.13	20.47±1.99	5.18±0.48
	1400°C×3h	0.09±0.01	3.07±0.01	15.89±0.32	36.28±0.57	8.88±0.70
	1600°C×3h	0.56±0.15	2.89±0.07	20.85±1.93	26.94±2.94	6.37±0.88

CMOR decreased at 1600°C

The higher heat treatment temperature led to volume instability of materials structure which resulted in the decrease in the CMOR of samples

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3. Results Microstructures



- Al element firstly migrated from the crystal site into the grain boundaries of Cr₂AlC grains, and then spread along grain boundaries to the outer layer, finally forming Al₂O₃ layer on the outer layer of Cr₂AlC particles.
- At 1600°C, the out-diffused and then oxidized aluminum was filled between the external Cr₂AlC particles, and little aluminum could be detected inside the particles.

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3. Results High temperature mechanical properties



- The excellent thermal conductivity of Cr₂AlC.
 - A dense coating with Al₂O₃ formed on the Cr₂AlC grain surface contributes to the bond strength of matrix.



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3. Results Corrosion resistance





Cross-section of the samples with different Cr₂AlC contents after the corrosion test at 1600°C

The corroded microstructure consisted of slag layer with macro-pore structure, reaction layer, penetration layer and original layer.

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3. Results Corrosion resistance



(1) $\operatorname{Cr}_{2}\operatorname{AlC}(s)+(3x/2)\operatorname{CO}(g) \rightarrow \operatorname{Cr}_{2}\operatorname{Al}_{1-x}\operatorname{C}(s)+(x/2)\operatorname{Al}_{2}\operatorname{O}_{3}(s)+(3x/2)\operatorname{C}(s)$ (2) $\operatorname{Cr}_{x}\operatorname{C}_{y}(s)+(3x/4+y/2)\operatorname{O}_{2}(g) \rightarrow (x/2)\operatorname{Cr}_{2}\operatorname{O}_{3}(s)+y\operatorname{CO}(g)$ (3) $\operatorname{Cr}_{2}\operatorname{O}_{3}(s)+\operatorname{Al}_{2}\operatorname{O}_{3}(s) \rightarrow (\operatorname{Al},\operatorname{Cr})_{2}\operatorname{O}_{3}(ss)$

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4. Conclusions

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- > The HMOR of the Al₂O₃-C sample with 5wt% Cr₂AlC-adding after heat treatment at 1400°C increased to 5.15 MPa from 3.14 MPa. Furthermore, the residual CMOR ratio did **not significantly decrease** when carbon content was reduced from 9 to 4wt%.
- \succ The addition of Cr₂AlC in Al₂O₃-C refractories led to the **improvement in corrosion resistance**. The Al₂O₃ and Cr₃C₂ grains were first generated in the original Cr₂AlC grain boundaries at a lower temperature. Then the Al_2O_3 layer of Cr_2AlC surface suffered chemical attack from the penetrated slag, and reacted with Cr₂O₃ generated by further slag corrosion to form $(Cr, Al)_2O_3$ solid solution. The $(Cr, Al)_2O_3$ solid solution dissolved in the slag, **increasing the viscosity** of the slag and suppressing the further penetration of slag. The original Cr₂AlC gradually decomposed and finally dissolved into slag.



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Thank you for your attention!

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