

Metal Dusting Prevention by Two Protection Systems in One Coating, Oxide Barrier and Catalytic Inhibition

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Metal Dusting

Metal Dusting destroys material's integrity due to immense **carbon precipitation** along **grain boundaries**.

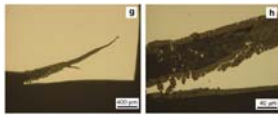
Affected materials are **iron, nickel, cobalt**, and their alloys.

MD occurs in high carbon containing atmospheres (**carbon activity >1**) such as in petro chemistry plants or steam reformers.

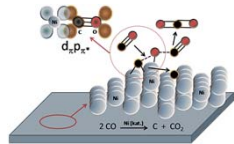
Critical temperature range: **450-900°C**



Coupon sample of alloy 600 after exposure in H₂-24%CO-2%H₂O at 620°C



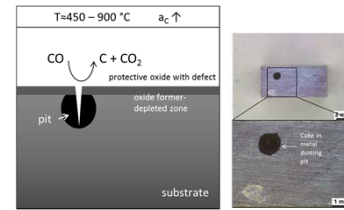
Internal graphite deposition in nickel after 100 h at $a_c=19$
J. Zhang, D.J. Young (2007)



Certain metals catalyze the dissociation of carbon monoxide. The emerging carbon diffuses into the metal.

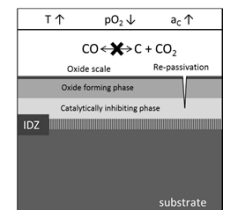
Coating concept

Conventional **metal dusting protection** by **high alloyed materials** always leads to an **oxide former-depleted zone** beneath the oxide scale. In the case of **local scale damage**, the corrosive environment gets access to the depleted material, which initiates **pit formation** and **metal dusting**.



Conventional metal dusting protection by high alloyed materials (left). Metal dusting pit in alloy 602CA (right).

The **combined coating concept** consists of two systems: At first **oxide forming elements** build dense oxide scales. Secondly, in the case of scale damage, an underlying **catalytically inhibiting phase** suppresses carbon precipitation until **re-passivation** of the scale.



Schematic illustration of the combined coating concept.

Ni-Sn-Al system

Ni₃Sn₂ coatings inhibit coking and metal dusting.

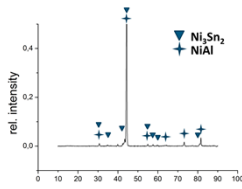
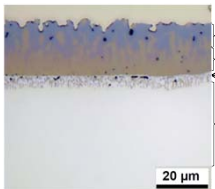
Coating process via **powder pack cementation (CVD)**

(Sn, Al, NH₄Cl, Al₂O₃, 800°C, 6 h)

→ **co-deposition** of **Sn** and **Al** into Alloy 600 (72 wt.% Ni)

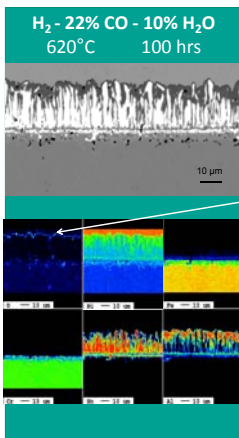


Uncoated (l) and Ni₃Sn₂-coated (r.) low alloyed steel after metal dusting exposure

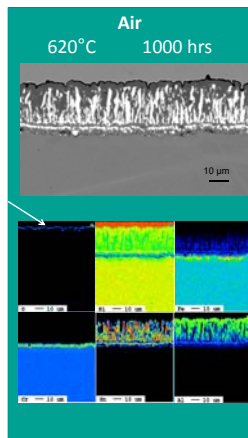


Cross section image of Ni-Sn-Al-coated sample.

Phase analysis by XRD (surface)



Formation of protective Al₂O₃ scale



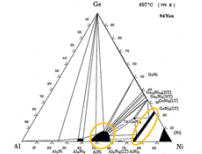
- Al₂O₃ formation on the surface in **carbonaceous** atmosphere with 10% **steam** content as well as **oxidizing** environments.
- No change in **microstructure** due to phases at **equilibrium** state
- Suitable for **Metal Dusting protection** for applications with temporarily oxidizing conditions (start-up or shut-down processes).

Ni-Ge-Al system

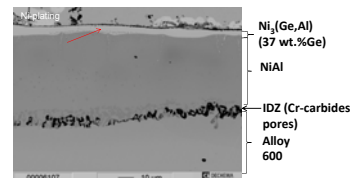
Coating process via **powder pack cementation (CVD) (Al and Ge)**

(Al, NH₄Cl, Al₂O₃, 1000°C, 4 h; Ge, NH₄Cl, Al₂O₃, 1050°C, 2 h)

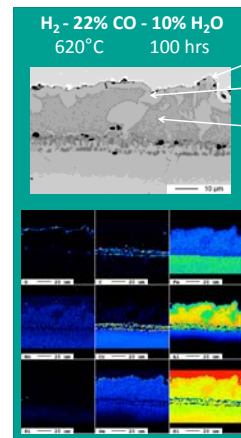
→ **Subsequent deposition** of **Al** and **Ge** into Ni-plated Alloy 600



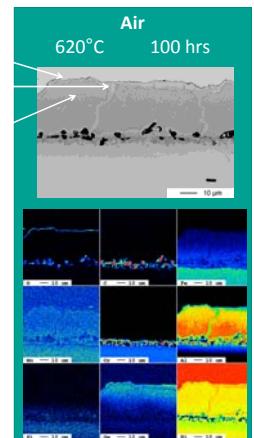
Ternary phase diagram of Al, Ge, and Ni at 497 °C
P. Villars et. al (1997)



Cross section image (l.) and elemental distribution maps (r.) of Ni-Ge-Al-coated sample.



Al₂O₃ scale
Ni₃(Ge,Al) (6 wt.%Ge)
Ni(Al,Ge) (up to 5 wt.% Ge)



- Al₂O₃ formation on the surface in **carbonaceous** atmosphere and air
- **No coking or metal dusting observed**
- Initial **Ge content** at the surface: **37 wt.%**. Ge dissolves in NiAl by formation of Ni(Al,Ge)
- Ni₃(Ge,Al) as well as Ni(Al,Ge) are suitable compounds for metal dusting-protective coatings.