



Coatings for phase change heat exchange

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Phase change heat exchange

Why phase change heat exchange?

Phase change is essential for energy applications, because latent heat is much larger than sensible heat.

Example:

For water, and for 1 g:

Latent heat to boil = 2260 J

Heat required to increase the temperature by 5 K= 21 J

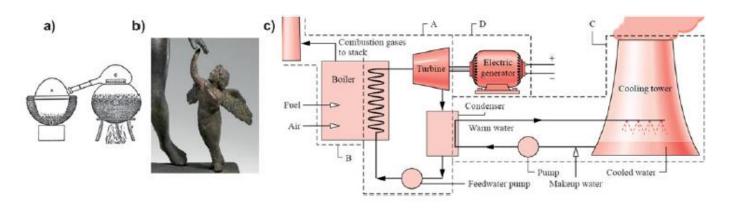
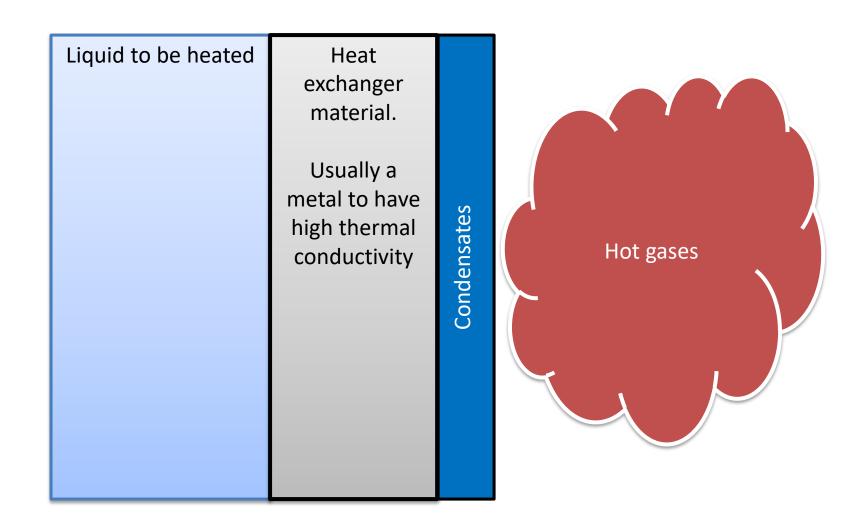


Figure 1. Human beings have relied on phase change heat transfer from Antiquity to now, to satisfy needs such as (a) alcohol production by distillation¹ (ca. 600 BCE, India), (b) metal casting for military or artistic purposes (Eros with Aphrodite, bronze, ca. 100 BCE), and (c) boiling and condensation for mechanical power, since the industrial revolution of the 19th century.⁴ Figures (a, b, c) reprinted with respective permissions of ASME, by courtesy of the Getty's Open Content Program, and with the permission of the Taylor and Francis Group.



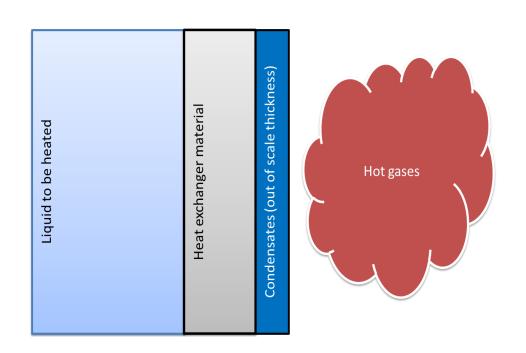
_Phase change heat exchange



(Relative dimensions are not in same scale)



Phase change heat exchange-Need for coatings

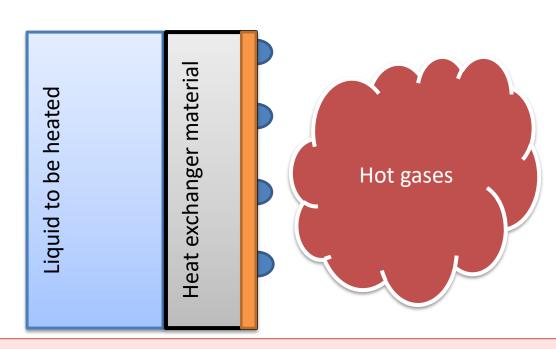


What is the problem here?

- 1. Condensates might be corrosive, if hot gases contain SO_x, Cl, etc.
- 2. Filmwise condensates introduce additional thermal resistance.
- 3. Condensates that are not being easily self-removed from the surface prohibit additional condensates to perform thermal exchange



Phase change heat exchange-Ultimate goal



Why do we need a coating?

- 1. Corrosion protection: Condensates might be corrosive if hot gases contain SO_x, Cl, etc.
- 2. <u>To enhance dropwise condensation over filmwise condensation</u>: Filmwise condensates introduce additional thermal resistance.
- 3. <u>Ease removal of condensates</u>: Condensates shall be easily self-removed from the surface to allow for additional thermal exchange.

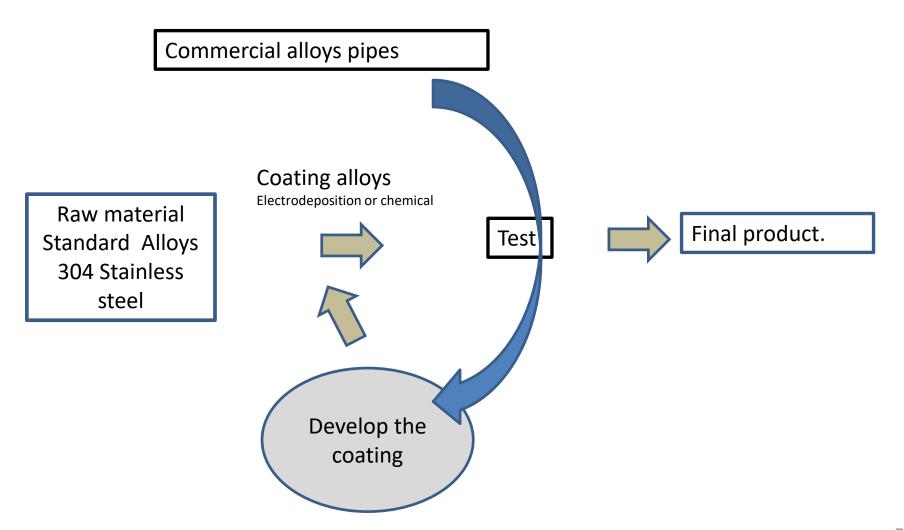
The coating shall also exhibit high thermal conductivity



Two approaches were followed and are presented hereafter

- Coatings with formulations close to conventional alloys with desirable properties. Here we will present the Alloy 59coating, which provides exceptional corrosion protection against H₂SO₄ and HCl
- Coatings with robust hydrophobic properties with very good corrosion protection against H₂SO₄ and functional wetting phenomena







Testing









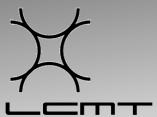


Figure 1. Experimental Setup for fast quality assessment of coatings



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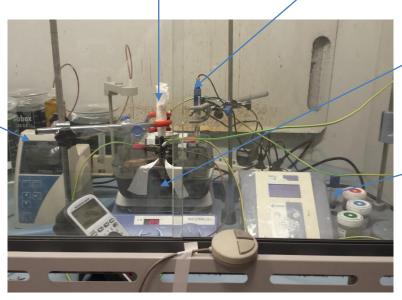


lab-. ELECTRODEPSITION

Sample

Counterelectrode.

pH controler



Ni- anode

pH-meter

- Stainless steel 304
- Stainless steel 316
- Carbon steel





ELECTRODEPSITION – pilot plant.

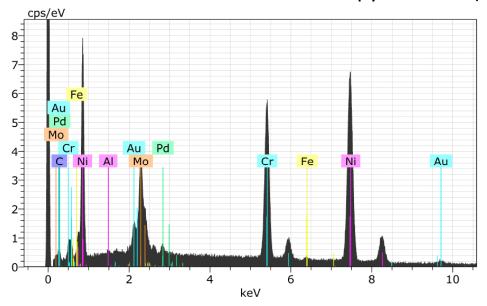
50-70 liters.



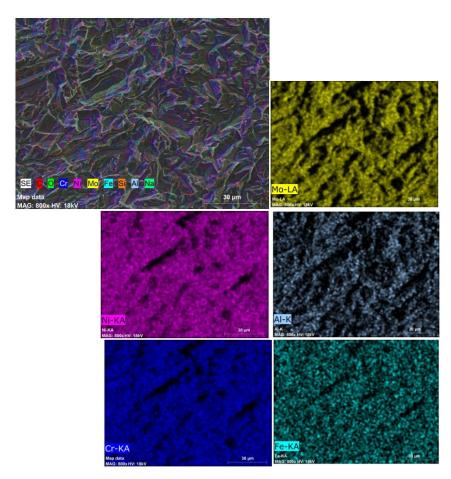




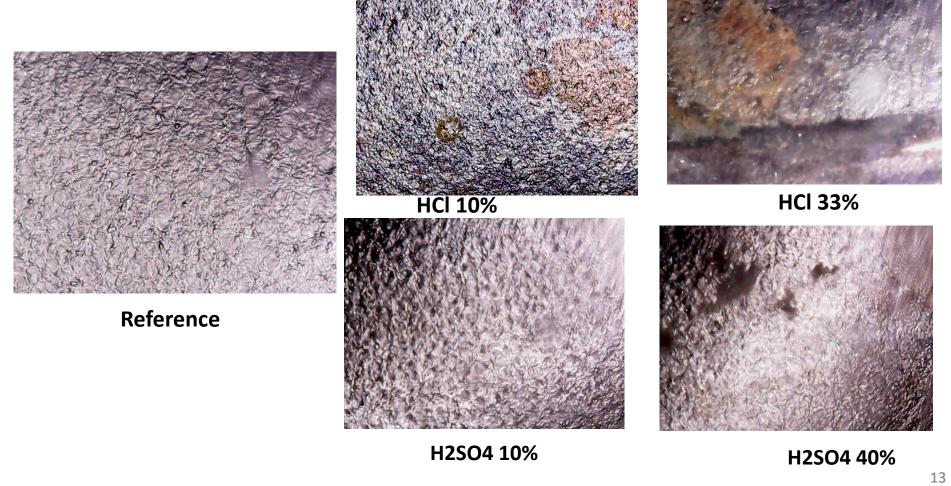
The formation of the Alloy59-like phase has been confirmed with Scanning Electron Microscopy and EDX (element mapping)



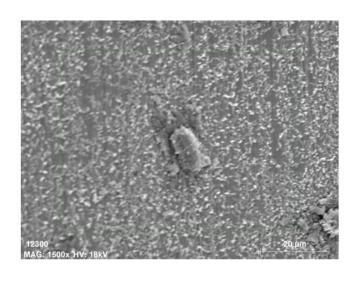
Element	AN	Series			Atom. C [at.%]		Sigma) [wt.%]
Carbon	_				0.00		0.00
Chromium	24	K-series	19.43	20.62	24.79		1.71
Nickel	28	K-series	55.80	59.22	63.08		4.78
Molybdenum	42	L-series	13.18	13.99	9.12		1.52
Palladium	46	L-series	1.14	1.21	0.71		0.22
Gold	79	L-series	4.09	4.34	1.38		0.72
Iron	26	K-series	0.42	0.44	0.50		0.15
Aluminium	13	K-series	0.17	0.18	0.43		0.12
		Total:	94.23	100.00	100.00	 	



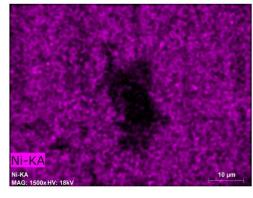


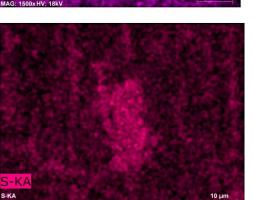


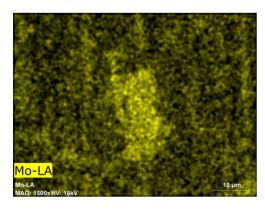




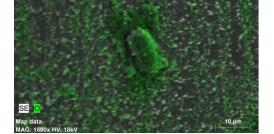
A59 40% H2SO4 48 h



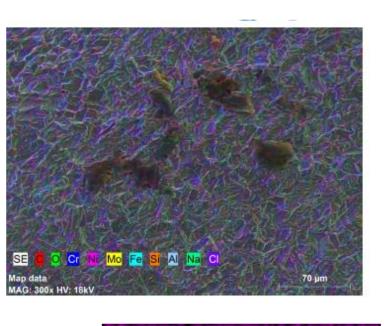




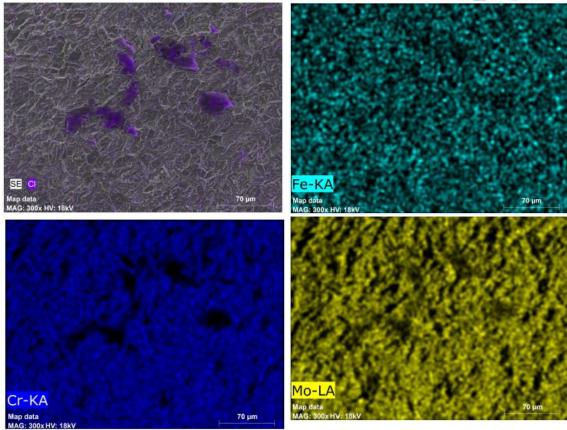








A59 33% HCl 48 h





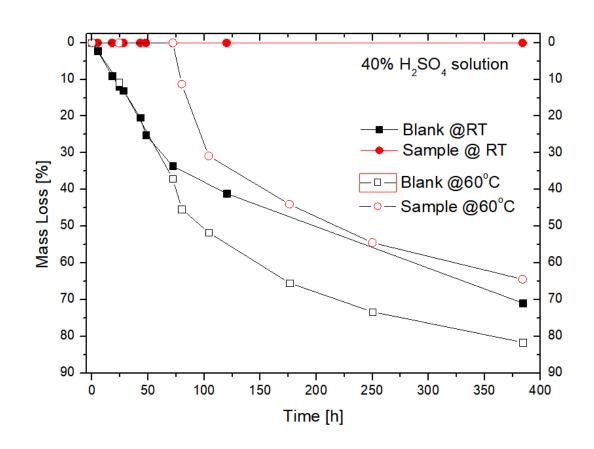
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- All dry process
- ✓ Hydrophobic coating
- ✓ High thermal conductivity

- ✓ Easy to implement ✓ No problem with adhesion
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Fully immersed sample at H₂SO₄ @ RT



Acknowledgements



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_Thank you

Thank you for your attention