

Elektrochemiese en adsorpsiestudies van die metaalorganiese netwerk, NH₂-MIL-53(Al)

**Authors:**

F.H. Peens¹
E.H.G. Langner¹

Affiliations:

¹Department of Chemistry,
University of the Free State,
South Africa

Corresponding author:

F. Peens,
peensfh.ufs@gmail.com

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Electrochemical and Adsorption Studies of the Metal Organic Framework, NH₂-MIL-53(Al).

NH₂-MIL-53(Al) is an amino-functionalised derivative of the metal organic framework MIL-53(Al). When successfully synthesised, it has the ability to be functionalised with metal complexes. A new solvo-intrusion method was used to study the amidation with butyric acid and ferrocenecarboxylic acid.

NH₂-MIL-53(Al) is 'n amiengefunksionaliseerde derivaat van die mikroporeuse metaalorganiese netwerk, MIL-53(Al) (Ahnfeldt *et al.* 2009). 'n Netwerk van 1-dimensionele diamantvormige kanale (~10 Å in deursnee) word gevorm deur aluminium- (III) oktahedrale hokstene deur middel van 2-aminotereftaalzuurligande onderling te verbind. Aluminium (III), afwisselend gekoördineer met suurstof, vorm liniëre kettings op die hoeke van die diamantvormige kanale. Die 2-aminotereftaalzuurligande vorm dus die vier sye van die kanale. Vanweë sy organiese verbindingsligande is hierdie netwerkstruktuur besonder buigsaam afhangend van die temperatuur en die aard van die geadsorbeerde molekules. Ná 'n solvotermiese hoëdruk-skoonmaakproses is die gevormde NH₂-MIL-53(Al) termies hoogs stabiel (~400 °C) en ook omkeerbaar higroskopies.

NH₂-MIL-53(Al) is sonder enige geadsorbeerde spesies 'n goeie heterogene katalis vir die Knoevenagel-kondensasiereaksie (Gascon *et al.* 2009). Palladiumnanodeeltjies in NH₂-MIL-53(Al) het tydens die Suzuki-Miyaura-kruiskoppelingreaksie goeie katalytiese aktiwiteit getoon (Huang *et al.* 2011). Ongefunksionaliseerde MIL-53(Al) kan gesublimerende ferroseen adsorbeer (Meilikhov, Yusenko & Fischer 2009a) en is ook al gefunksionaliseer met 1-1'ferroseendiildimetilsilaan om as redokskatalis vir benseenoksidasie op te tree (Meilikhov, Yusenko & Fischer 2009b). Ons studie is egter daarop gemik om die chemiese adsorpsie van karbokselsure en die proses daarvan in NH₂-MIL-53(Al) te verbeter.

In hierdie studie is 'n nuwe solvo-intrusiemetode ontwerp om die adsorpsie van bottersuur en ferroseenkarbokselsuur in NH₂-MIL-53(Al) te bestudeer. Dit is 'n baie belangrike konsep omdat die netwerkkanale slegs molekule kleiner as 13.04 Å kan deurlaat (Horcajada *et al.* 2008). 'n Vergelykende tydstudie is gedoen om die minimum tyd te bepaal wat die hoogste lading van beide die bogenoemde molekules gee. Die hoeveelheid lading is bepaal deur 'n termogravimetriese analise (TGA) en kernmagnetiese resonansspektroskopie (KMR). Postsintetiese, direkte amidasie van NH₂-MIL-53(Al) is met bottersuur en ferroseenkarbokselsuur bereik. Hierna is die elektriese geleidingseienskappe van ferroseenkarbokselsuur in NH₂-MIL-53(Al) bepaal deur 'n nuut ontwerpte vastetoestand-sikliese voltametriemetode.

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