

# EFFECT OF CREVICE CONFIGURATION ON CORROSION BEHAVIOUR OF TENSILE WIRES OF FLEXIBLE PIPES UNDER DIFFERENT CONDITIONS

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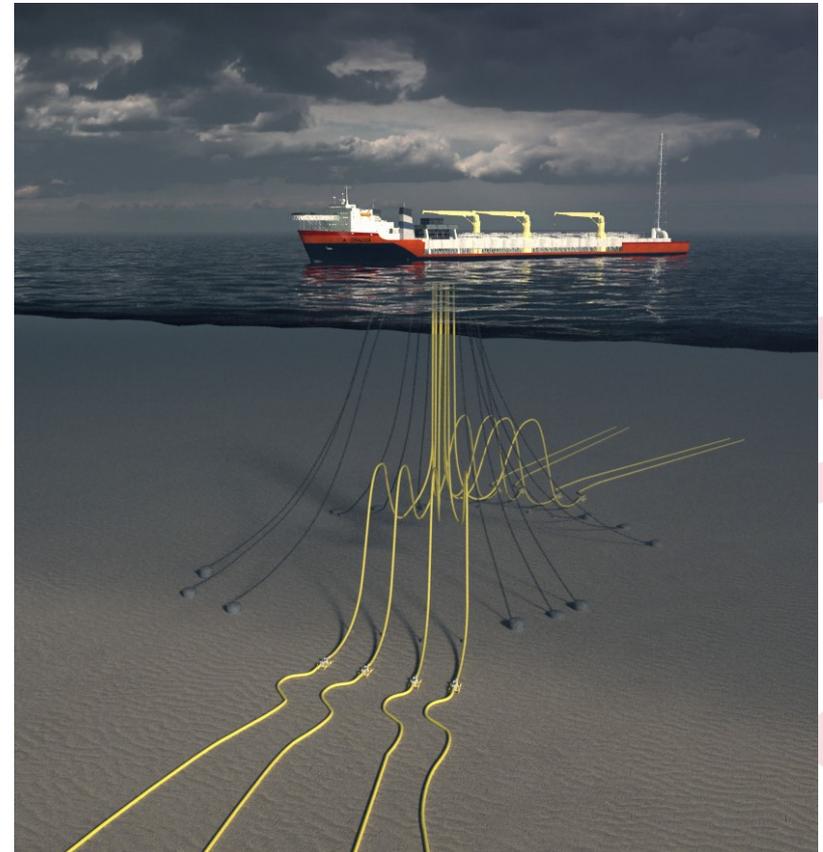
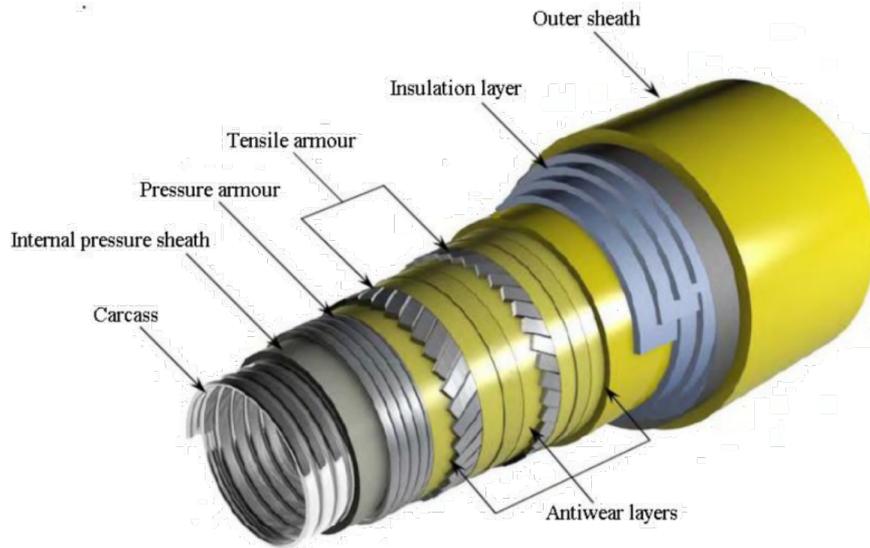


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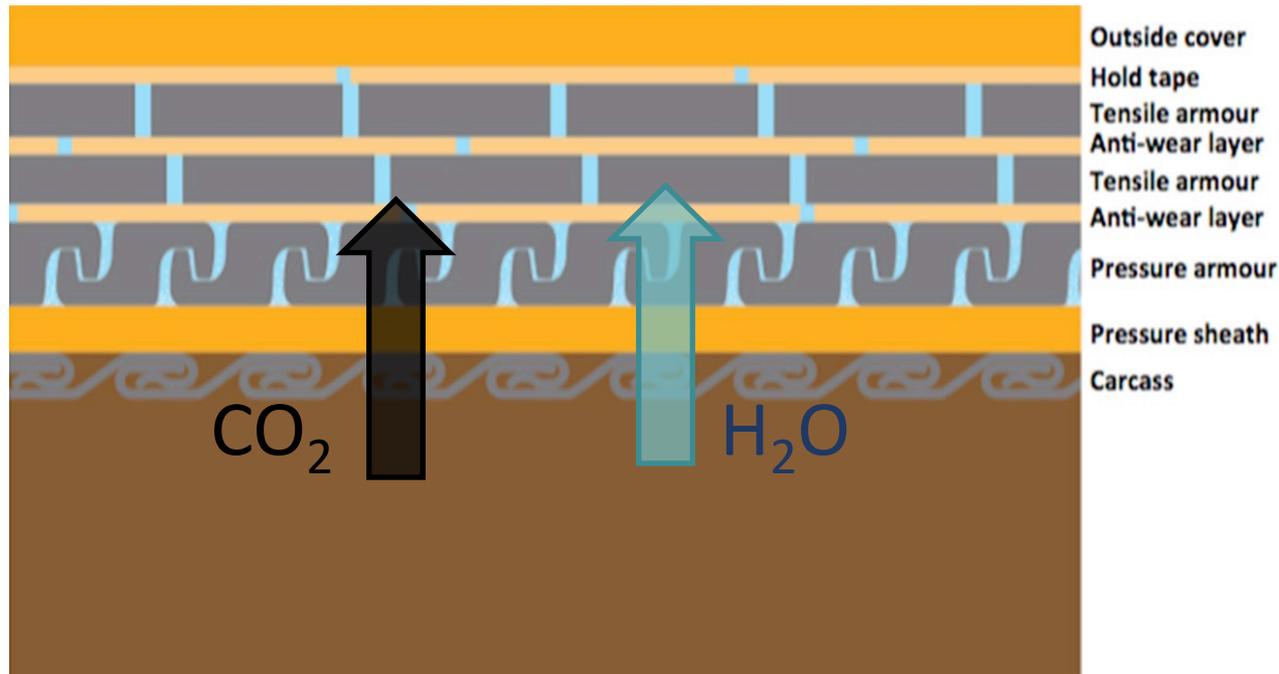
# Background

- ❑ Schematic indicating the multi-layer construction of flexible pipe designs



# Background

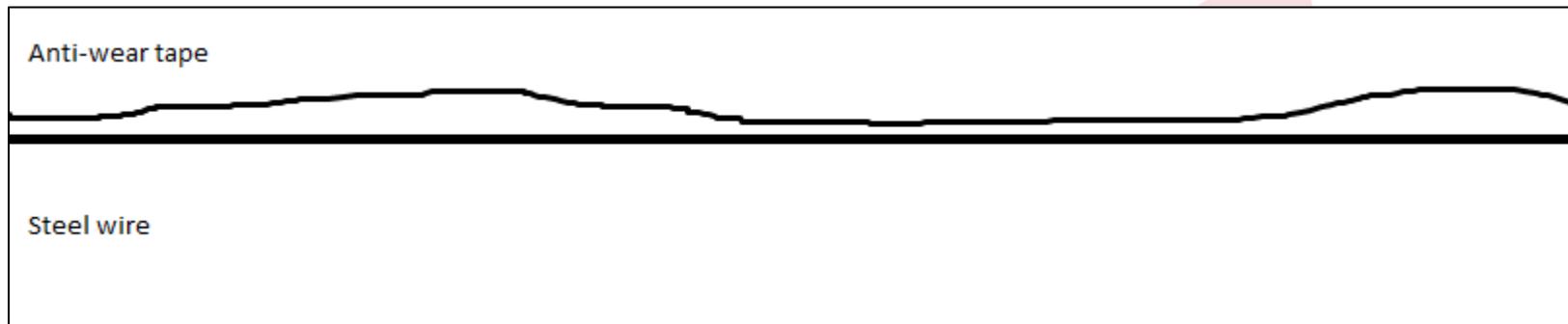
- The high strength steel wires are confined in the annulus and in contact with a corrosive environment formed by gases and water molecules diffusing from the bore through the inner polymer layers.



# Background

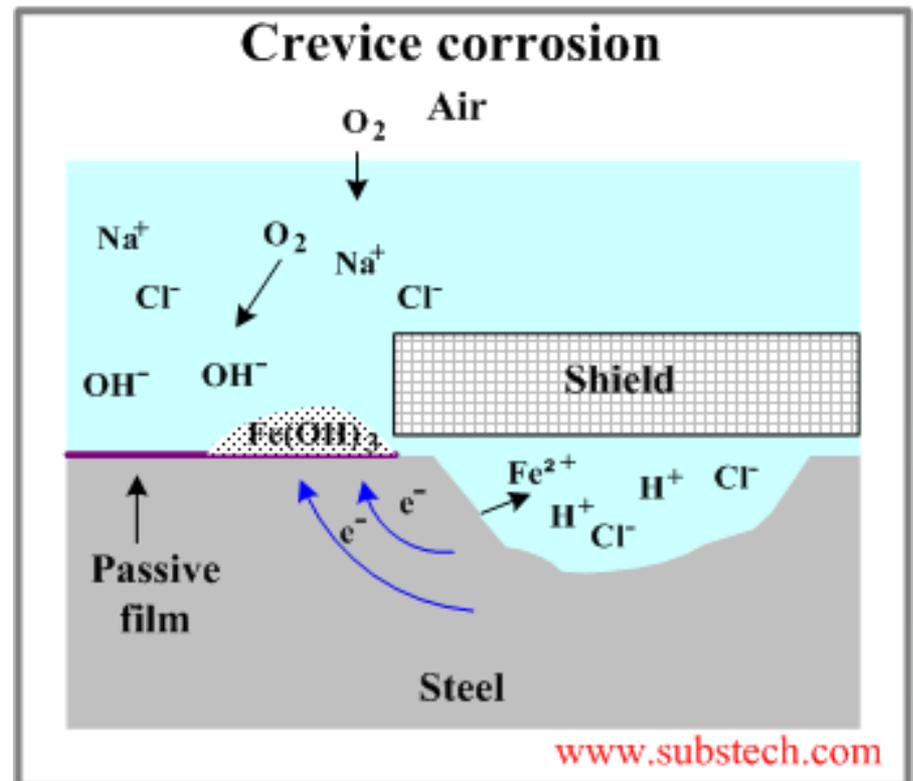
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- ❑ The tensile wires are usually covered by antiwear tapes and this configuration can also interfere in their corrosion behaviour.
- ❑ The position of the tape relative to the wire is not uniform due to many variations in the manufacturing process of the pipe.
- ❑ **Some regions** → almost **absolute intimate contact**.  
**Other region** → there is a **gap** between the polymer and the steel → **allowing water and gases** to reach the **surface of the wire**.



# Background

- ❑ If this gap is wide enough to guarantee sufficient mass transport of chemical species, the composition of the fluid in the annulus in this region will be relatively homogeneous.
- ❑ If this gap is so narrow that locally the chemistry of the corrosive environment develops differently than in the bulk, localised corrosion may occur.



# Two questions

1) Does the polymer prevent the liquid from reaching the metal surface completely?

R: YES → corrosion does not occur under the polymer.

R: NO → **corrosion occurs** under the polymer.

2) But what is the morphology of the attacks, **localised** or **uniform**?

Uniform corrosion → where the **opening is large** enough to **allow mass transport** and **prevent the development of a different local environments**

Localised corrosion → where the **opening is so narrow** that the environment **develops differently in these regions.**



# What conditions would we like to simulate in this work?

## General investigation of the effect of oxygen on the CO<sub>2</sub> corrosion

### Step 1: Annulus initially flooded: ↑[Fe<sup>2+</sup>] and deaerated

- CO<sub>2</sub> corrosion
- Iron carbonate precipitation

### Step 2: Outer layer damaged: ↑O<sub>2</sub>

- O<sub>2</sub> and CO<sub>2</sub> corrosion
- Iron carbonate film formation is affected
- Iron oxides formation

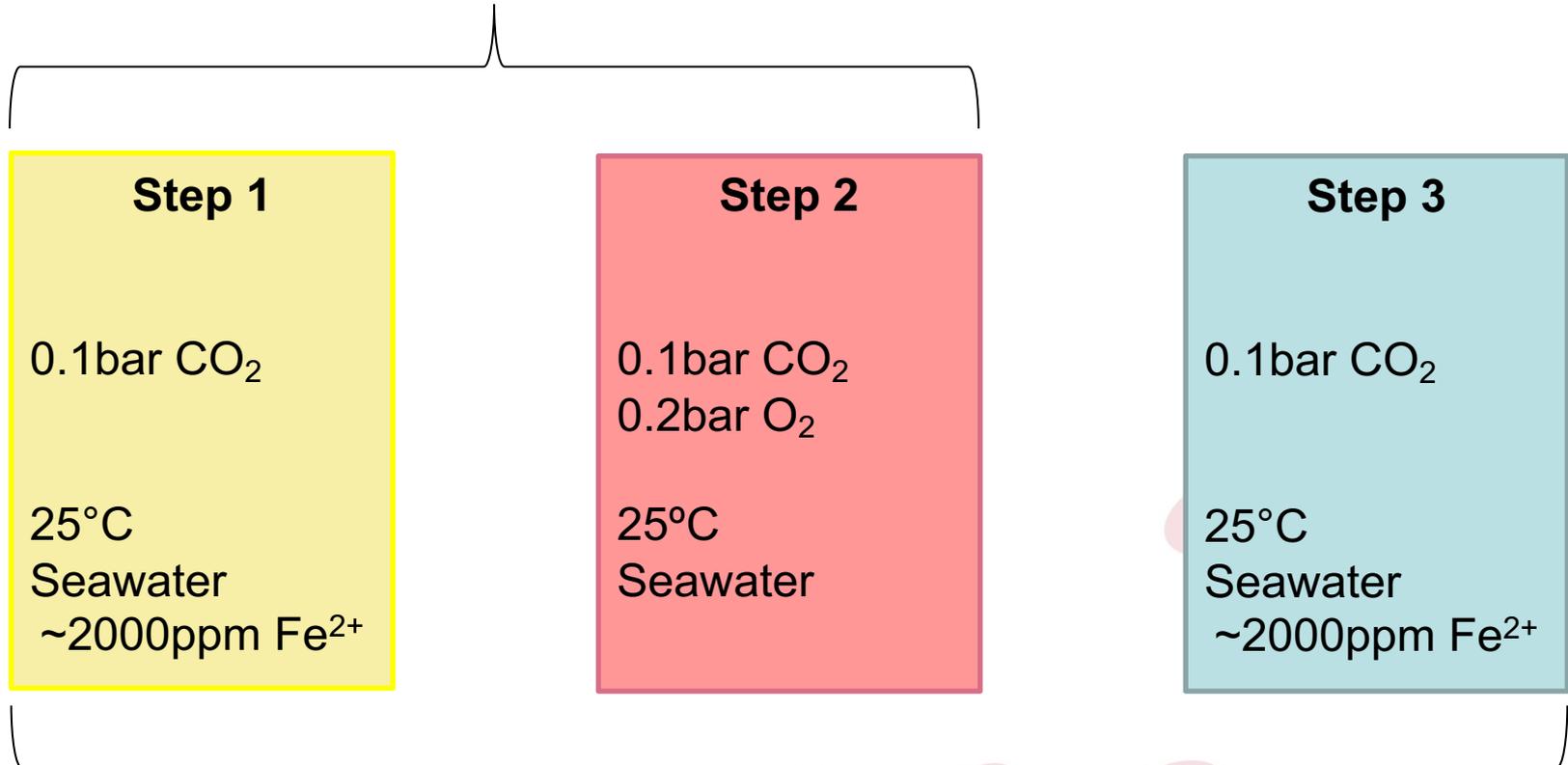
### Step 3: Outer layer damage is fixed: O<sub>2</sub> ~ 0

- CO<sub>2</sub> corrosion
- Iron carbonate precipitates again.

Arne Dugstad's previous works on history effect were used as theoretical background for this setup.

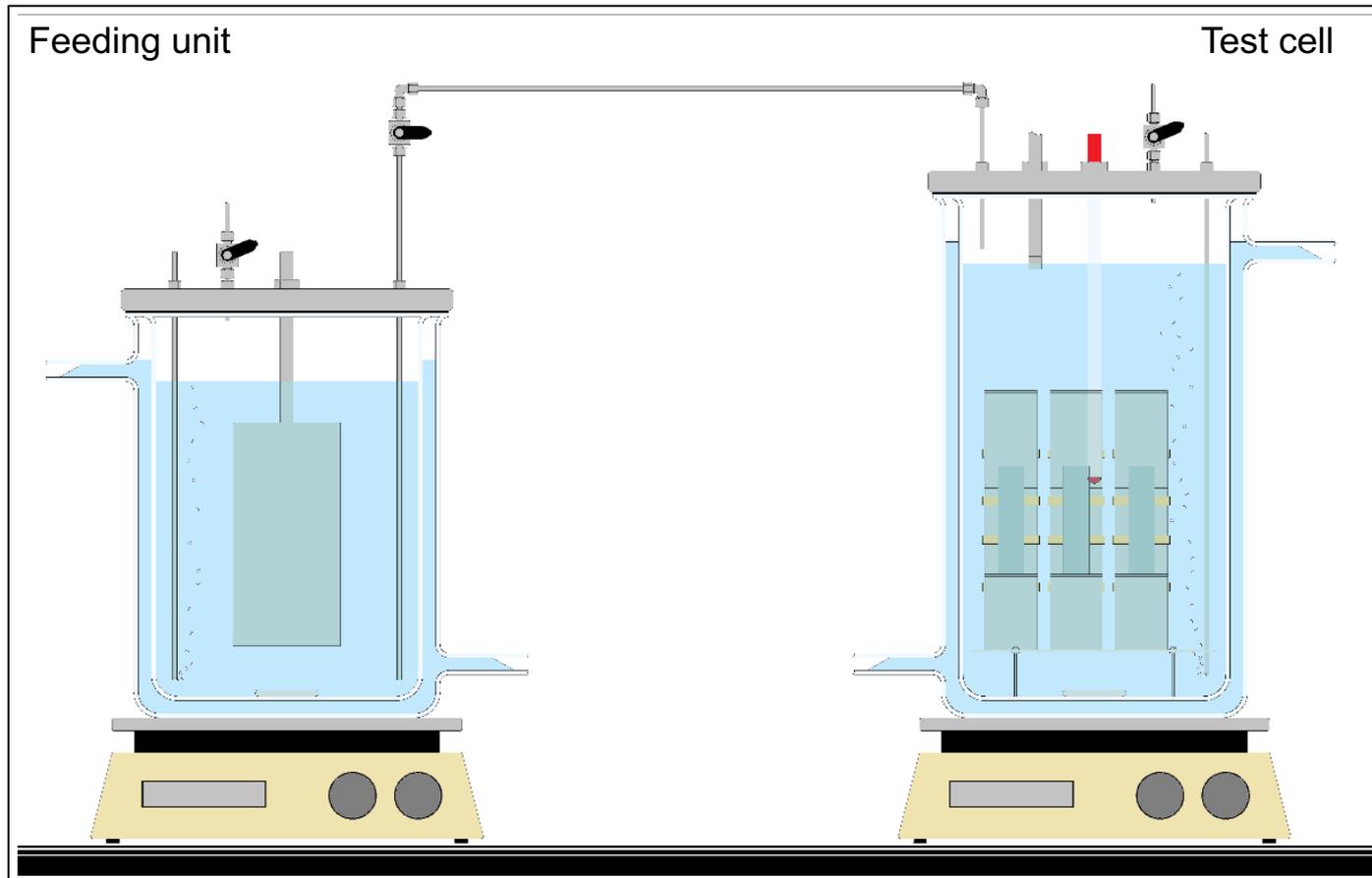
# Test Parameters

Test 1: 30 days – Finished



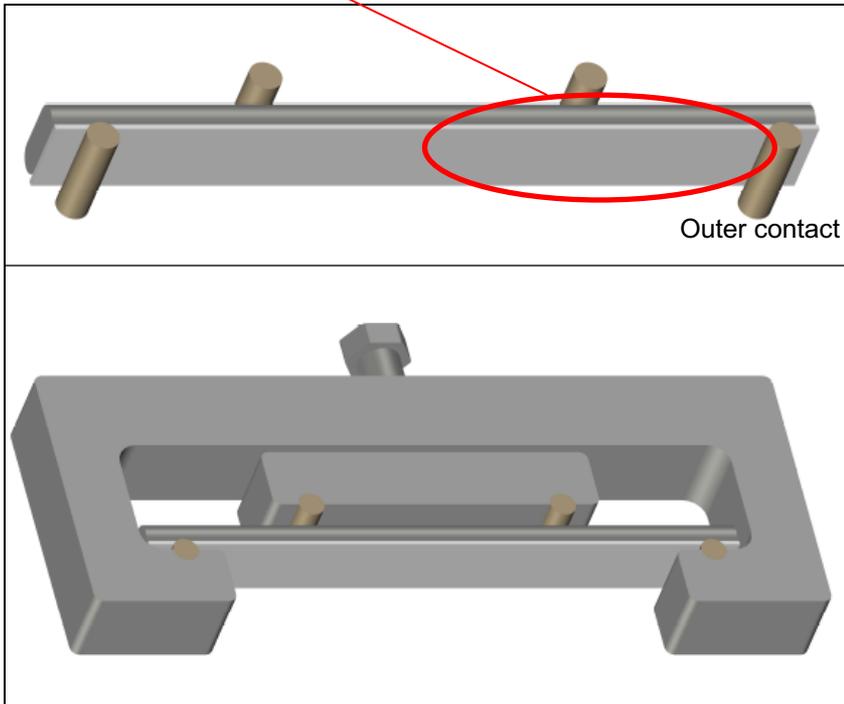
Test 2 and 3: 90 and 180 days – On going

# Experimental Setup

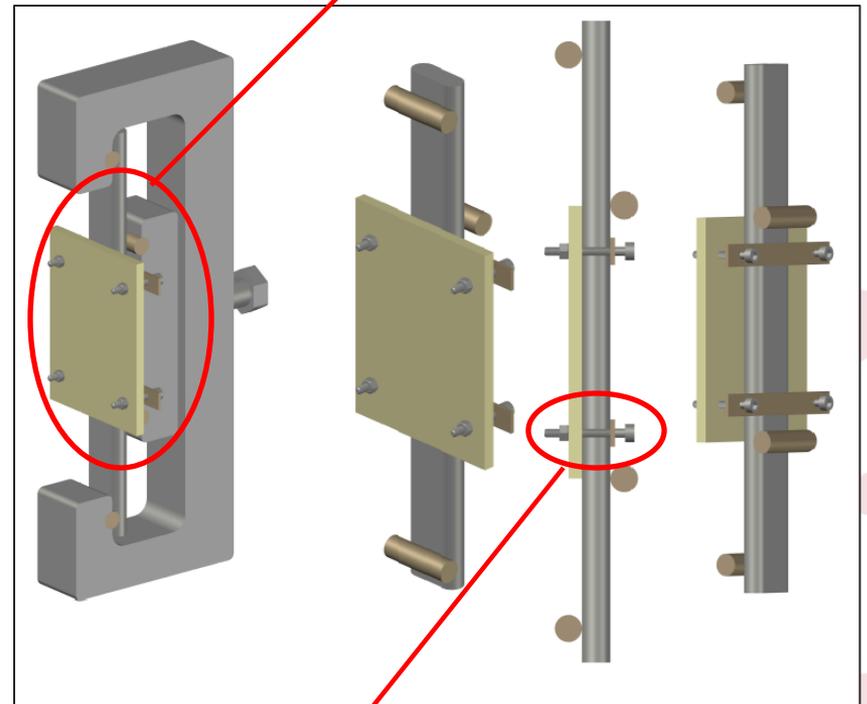


# Configuration of creviced specimens

TEFLON: Low friction coefficient →  
 sliding during the deformation  
 process → difficult a uniform contact



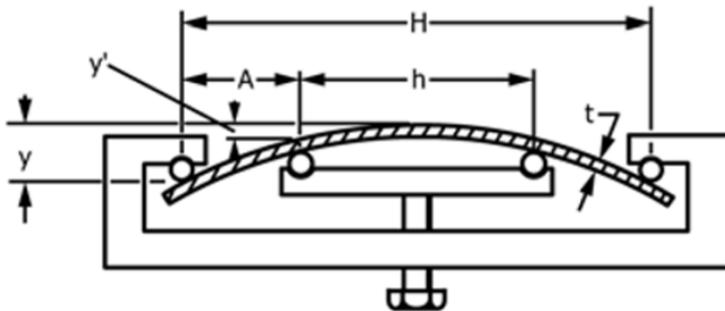
PA11: Higher friction  
 coefficient. Covering only the  
 region A (central portion of



Nuts and bolts to facilitate the adjust the gap  
 between polymer and specimen before loaded

# The four-point bend tests

Test specimens were loaded to 90% YS.  
Creviced specimens had their deflection measured already with the polymer tape.



Dial indicator



$$\sigma = \frac{12 \cdot E \cdot t \cdot y}{(3 \cdot H^2 - 4 \cdot A^2)} \quad \therefore \quad y = \frac{\sigma \cdot (3 \cdot H^2 - 4 \cdot A^2)}{12 \cdot E \cdot t}$$

y	deflection
$\sigma$	stress on the outer fiber (convex surface)
H	distance between outer supports
A	distance between inner support and closest outer support
E	Young's modulus (usually E=210GPa for steels)
t	specimen thickness

INPUT			OUTPUT	
%Y.S.	Y.S.		$\sigma$	
(MPa)	(MPa)		(MPa)	
90	1085		976.5	
INPUT			OUTPUT	
H	A	E	t	y
(mm)	(mm)	(GPa)	(mm)	(mm)
100	23	210	5.88	1.84

# Test 1. TEFLON polymer layer

(1 month)

Test cell aspects according to the variations of the test environment.



**Step 1**



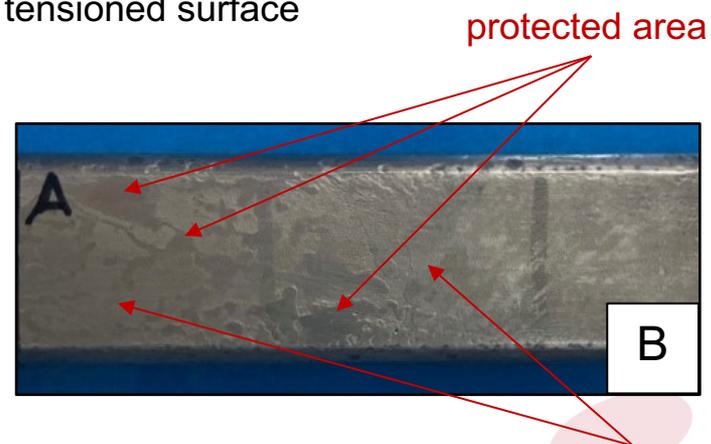
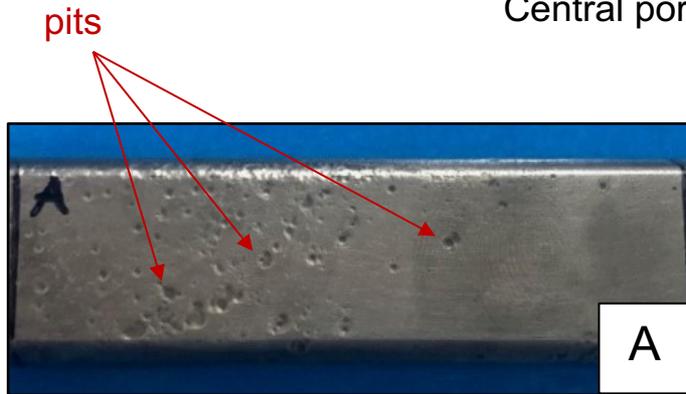
**Step 2**

# Test 1. Results

(1 month)

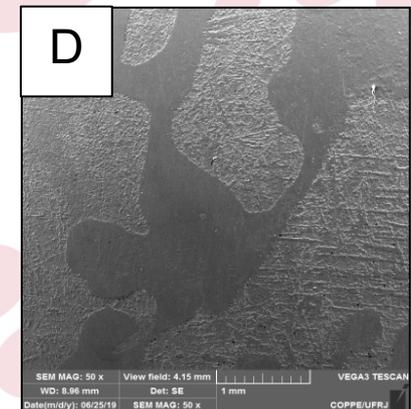
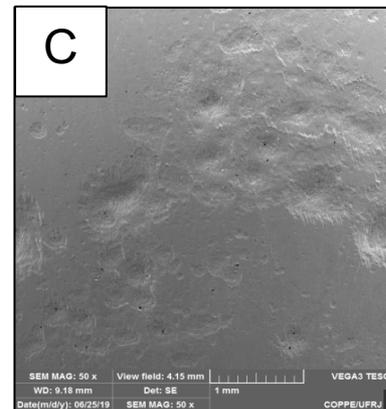
## Surface analysis – Macroscopic and SEM

Central portion of the tensioned surface



Macroscopic analysis: (A) Bare specimen  
(B) creviced specimen

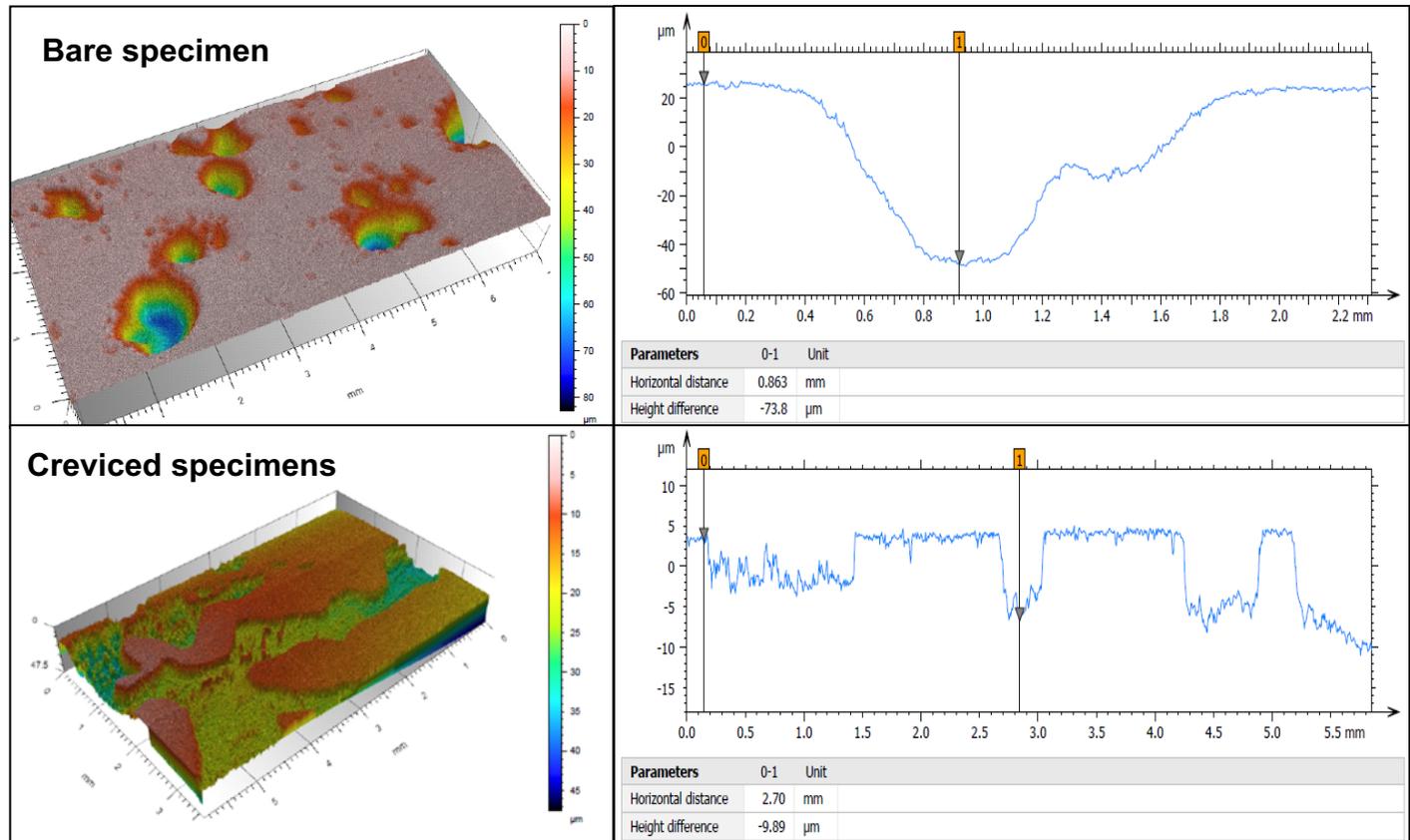
SEM analysis: (C) Pitted area: Bare specimen  
(D) Protected regions and uniform corrosion: creviced specimen



# Test 1. Results

(1 month)

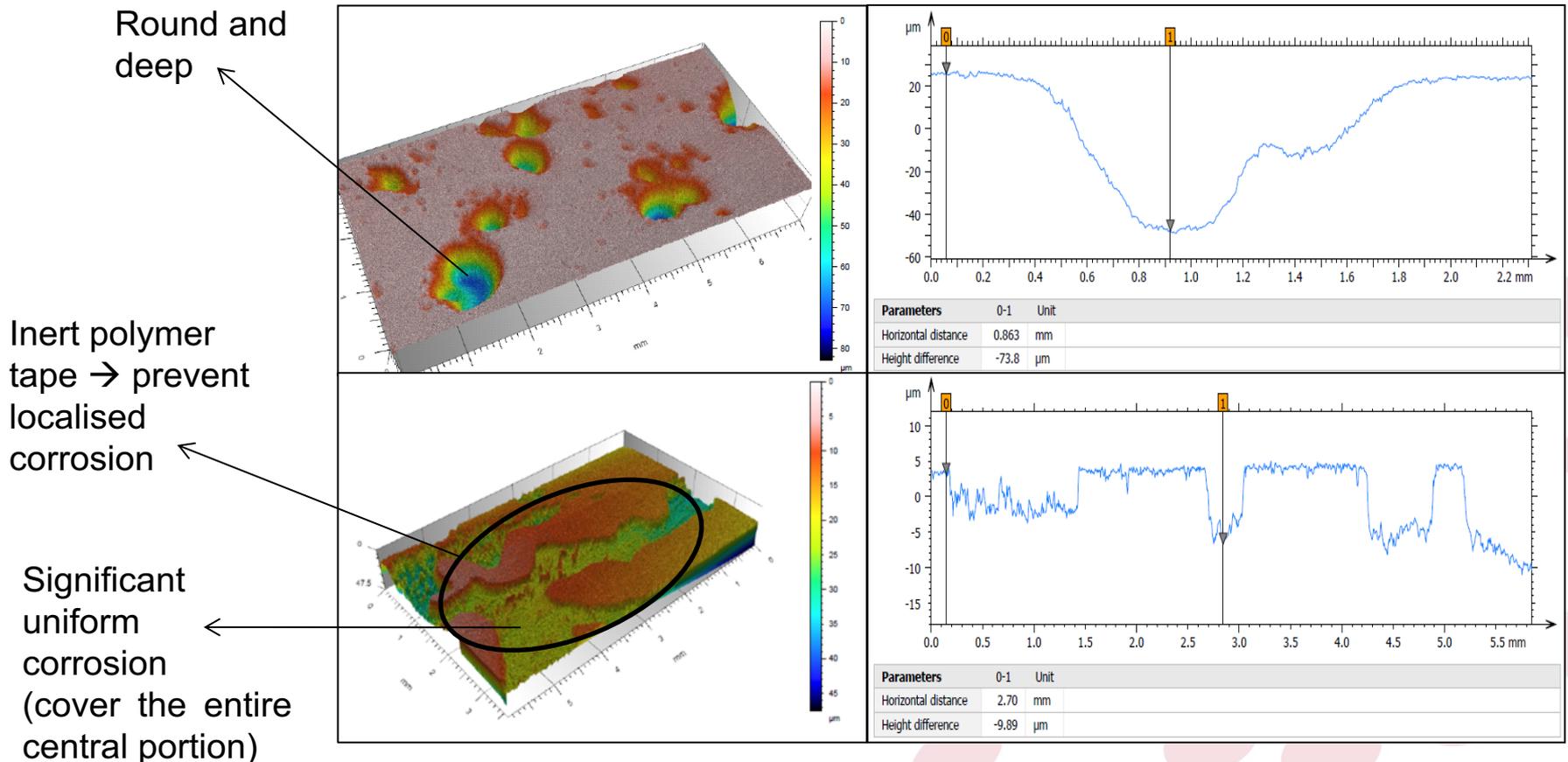
## Surface analysis – Confocal microscopy



# Test 1. Results

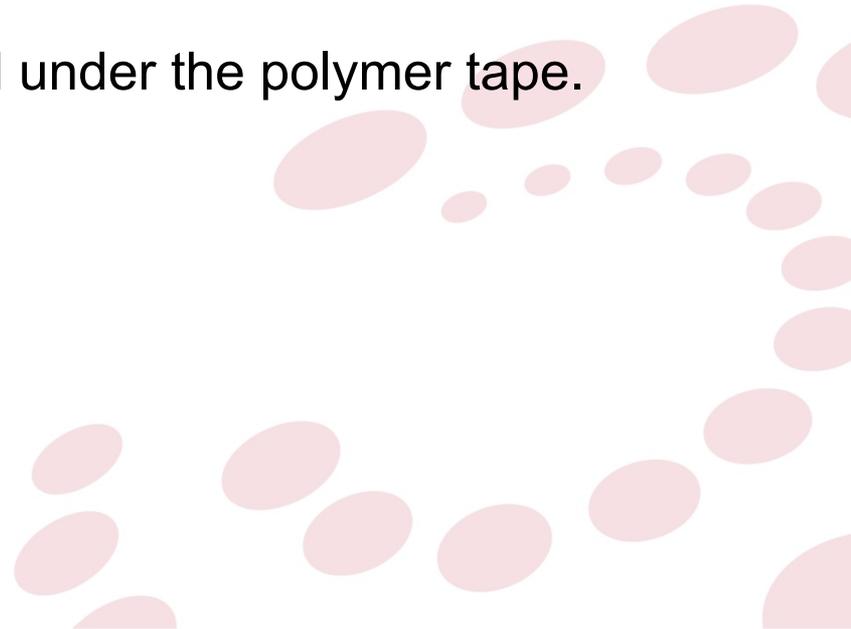
(1 month)

## Surface analysis – Confocal microscopy



# Main conclusions of Test 1

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- No cracks were observed.
  - Pitting was observed in the specimen tested without crevice.
  - The polymer tape on the surface of the steel prevented pitting.
  - Severe uniform corrosion was observed under the polymer tape.
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# Test 1.2. Setup at IFE (Norway)

Experiment	01.1
Status	Finished
Polymer	Teflon
Duration	1 month
Cracks	No

**Before cleaning**



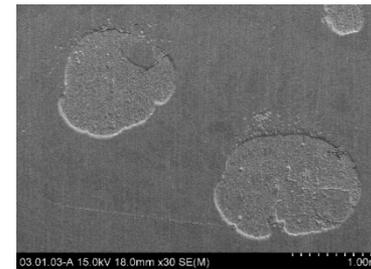
Bare specimen



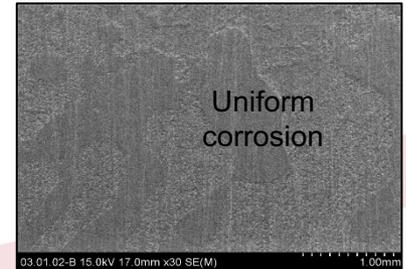
Creviced specimen

**No pitting – iron carbonate film**

Localised attacks resulted from CO<sub>2</sub> bubbles retained at the surface of the bare specimen.

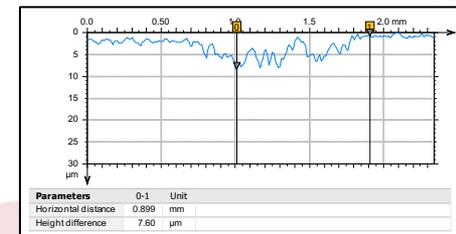
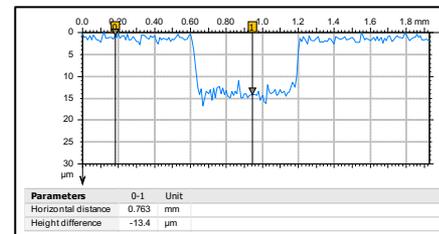


Bare specimen



Uniform corrosion

Creviced specimen



# Main conclusions of Test 1.2

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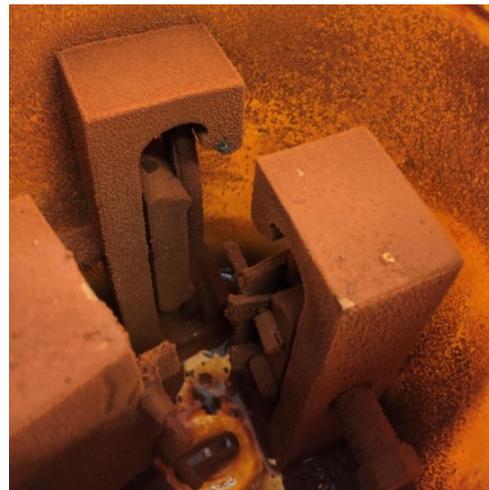
- No cracks were observed.
- No pitting was observed in the specimen tested without crevice.
- Localised attacks resulted from CO<sub>2</sub> bubbles retained at the surface of the bare specimen. As they were large and shallow, did not characterise pitting.
- The bare specimens were relatively protected by the iron carbonate film and almost did not corrode.
- Uniform corrosion was observed under the polymer tape, although not as severe as in the specimens exposed to oxygen.

# Tests 2 and 3. PA11 polymer layer



# Tests 2 and 3. PA11 polymer layer

Test cell aspects according to the variations of the test environment.



A: first day of experiment, **with only CO<sub>2</sub> and N<sub>2</sub> in the gas mixture. Before heavy precipitation of iron carbonate.**

B: gas mixture is changed and **oxygen reacts to the dissolved iron and disrupts the iron carbonate films.**

C: test cell after the test solution was drained at the **end of step 2, and before step 3 started.**

D: test cell **during step 3.**

# References

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**Thanks for your  
attention!**

