

Pore radius distribution and fractal dimension derived from spectral induced polarization

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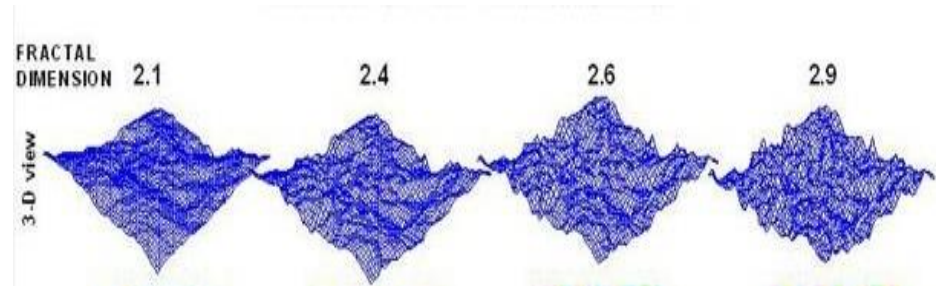
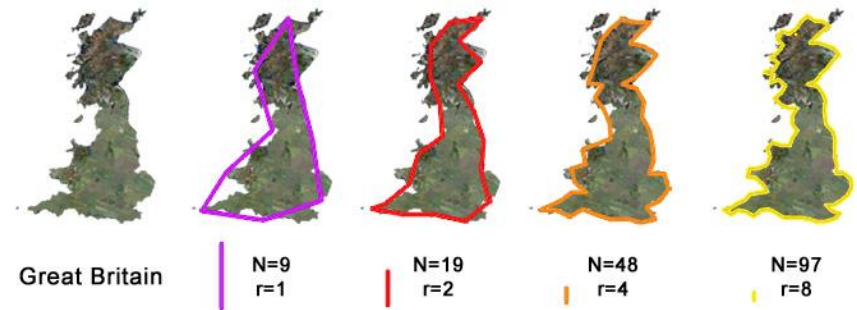
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Introduction

- Fractal theory is applied to describe the structure of geometric objects.
- At molecular-size and microscopic range, surfaces of most materials including those of natural rocks show self-similar features that can be quantified by the parameter of fractal dimension D .

Introduction

- For a straight line, $D=1$;
for a winding line, $D>1$
- For smooth surface, $D=2$;
for a rough surface, $D>2$
- For a not fully occupied volume, $D<3$



Introduction

- Lesmes & Morgan (2001) tried to relate SIP to fractal dimension.
- Here we demonstrate the use of SIP to determine the fractal dimension of the pore volume.
- We compare the fractal dimension and pore size distribution with the results of nuclear magnetic resonance (NMR) and mercury intrusion capillary pressure method (MICP).

Methods

- We transformed the relaxation time distribution of complex conductivity spectra determined by Debye decomposition (Nordsiek and Weller, 2008) into a curve showing the cumulative intensity as a function of increasing pore radius r calculated from relaxation time τ .
- It is assumed that the **distribution of relaxation times** is controlled by the **distribution of the pore sizes**.

Methods

- We adopt an equation proposed by Schwarz (1962) and applied by Revil et al. (2012) for the Stern layer polarization model:

$$r = \sqrt{2 \tau D_{(+)}}$$

- r originally refers to radius of spherical particles
- diffusion coefficient of the counterions in the Stern layer. We assume a constant $D_{(+)} = 3.8 \times 10^{-12} \text{ m}^2/\text{s}$ as proposed for clayey material (Revil, 2013).

Samples

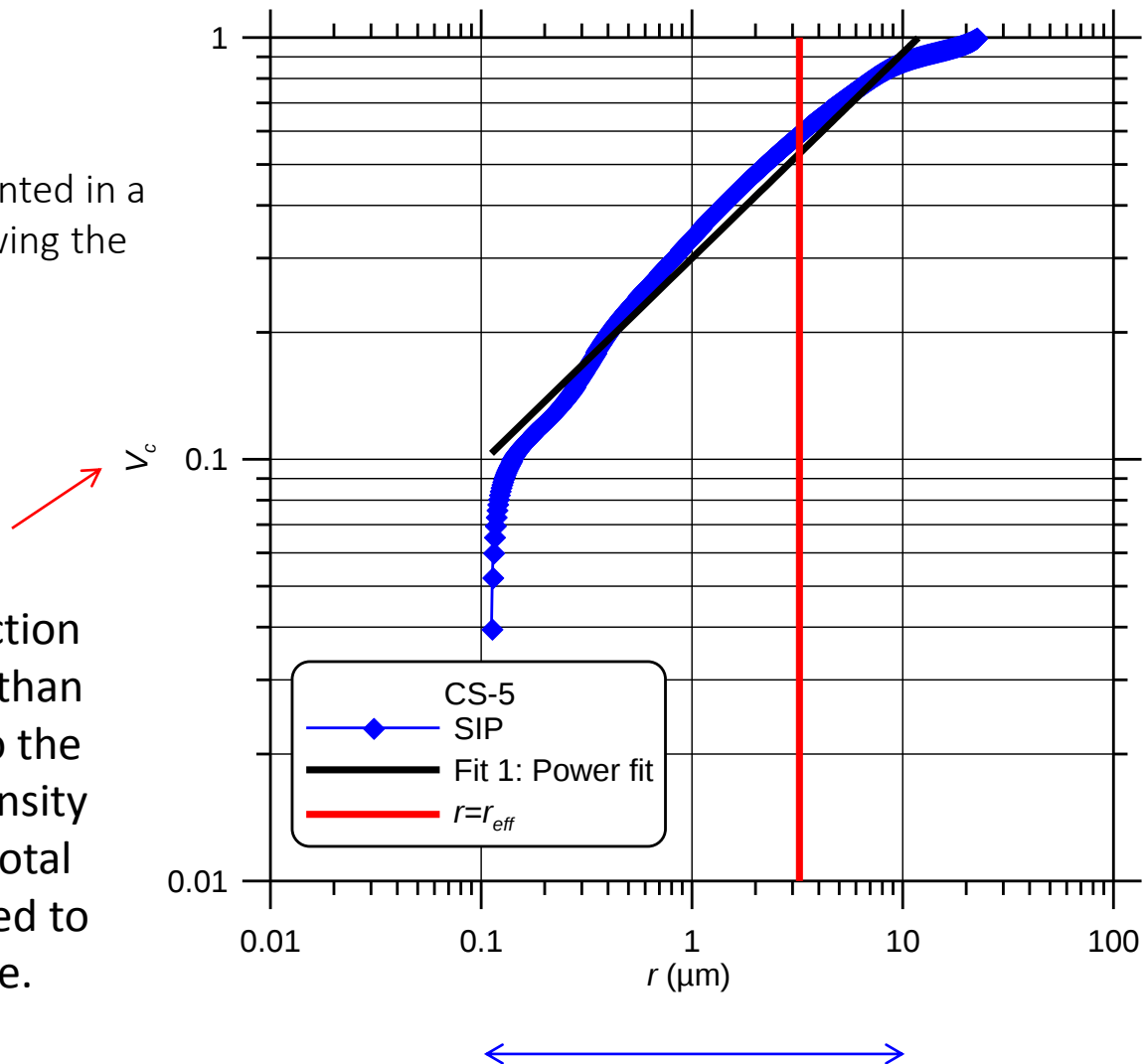
- This approach was applied to 24 rock samples from an Eocene sandstone formation in China (Zhang and Weller, 2014).



Results

The cumulative curve is presented in a double logarithmic plot showing the relation $\log(V_c) \sim \log(r)$.

$V_c = V(<r)/V$ is the cumulative volume fraction of pores with radii less than r , which corresponds to the ratio of cumulative intensity to total intensity. The total chargeability is attributed to the total pore volume.



Results

- In the case of fractal behaviour of the pore volume distribution, the slope s of the fitting line is used to get the fractal dimension $D_{SIP} = 3 - s$. The resulting fractal dimension D_{SIP} varies for the investigated samples in a range between 2.507 to 2.761.
- The red line marks the effective hydraulic radius, which has been determined from permeability k and formation factor F :

$$r_{eff} = \sqrt{8 F k}$$

Results

permeability range of the samples:
0.02 to 60 mD

High permeability
(H-samples)



low clay and carbonate content
and a good sorting

Medium permeability
(M-samples)

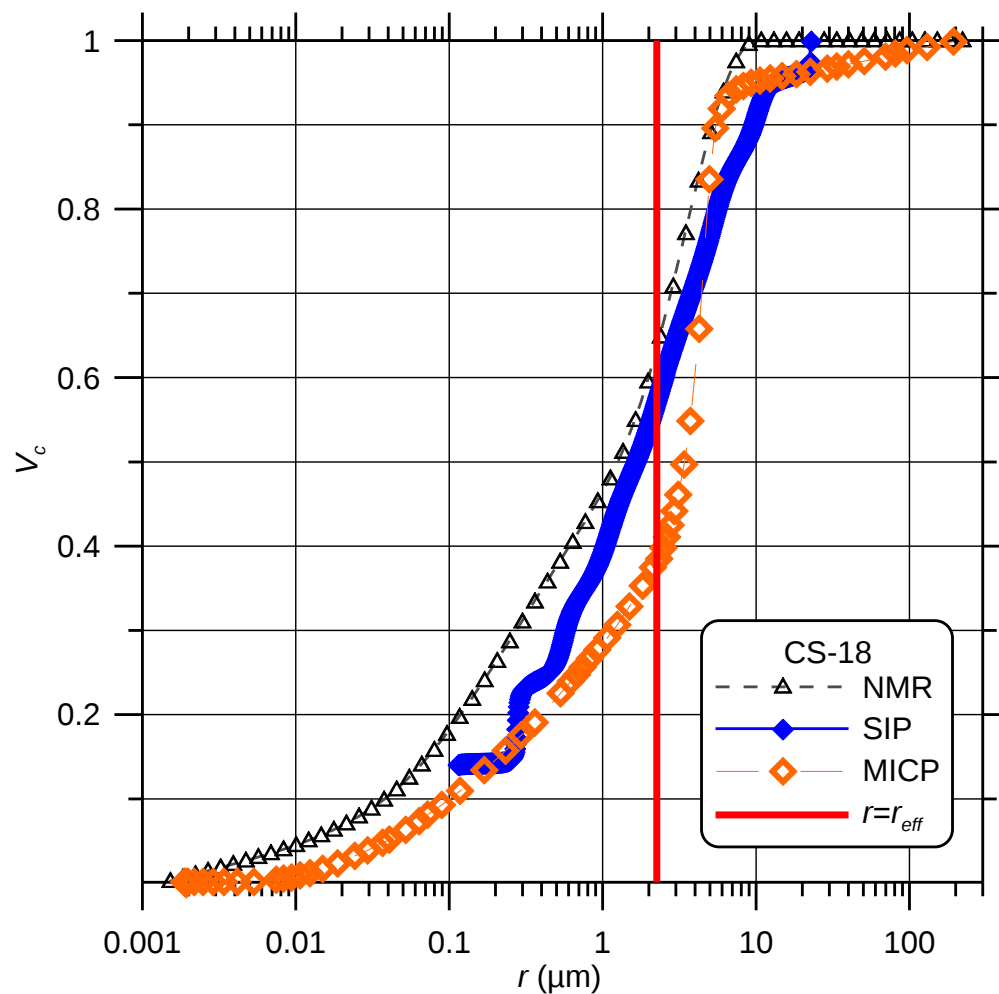
Low permeability
(L-samples)



either a strong compaction or a rich
sedimentation of carbonates in the pore
space that results in low porosity and small
pore radii

Results

H-Samples:
the V_c - r curves of the
methods NMR, MICP and
SIP show similar
characteristics

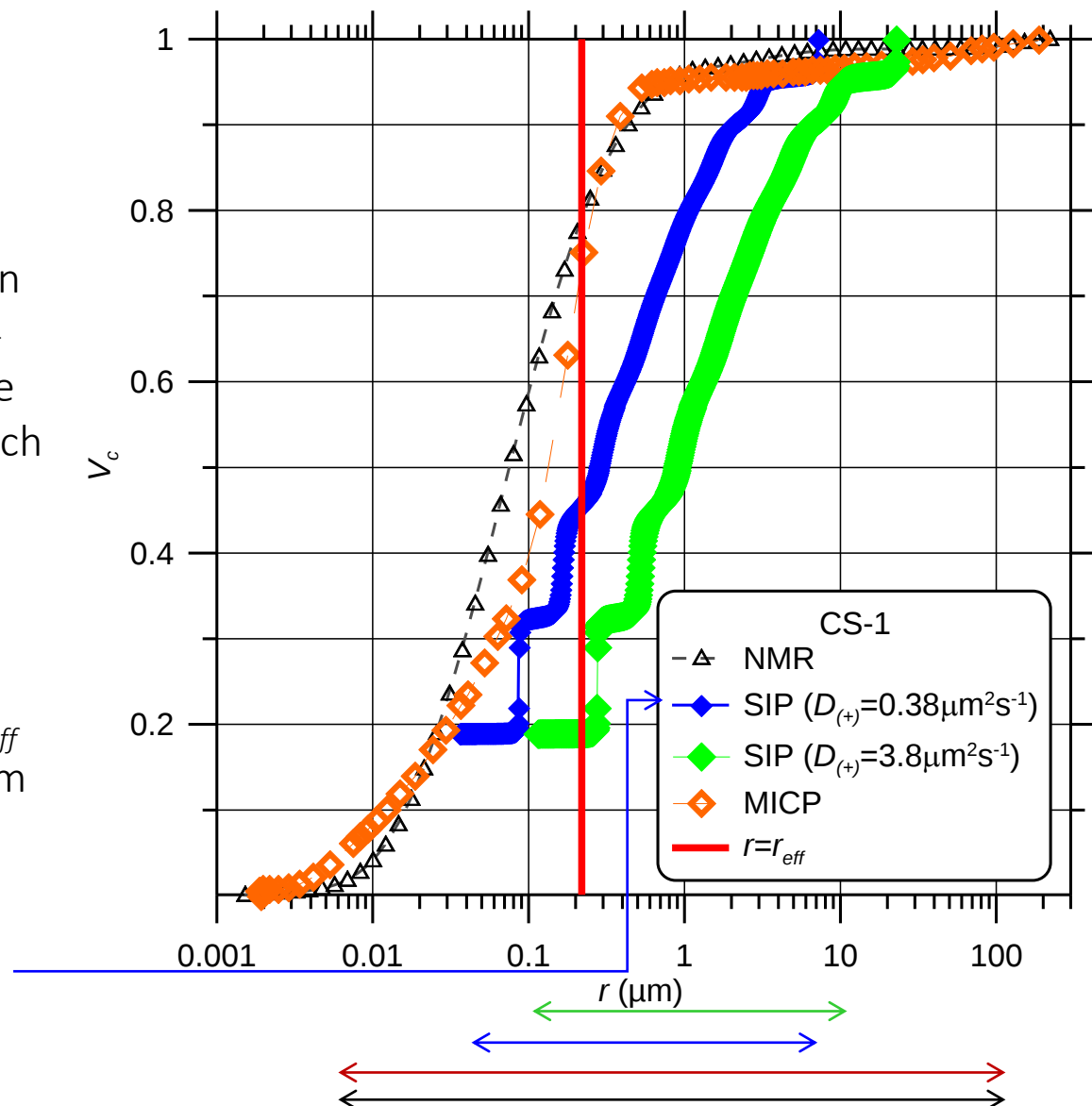


Results

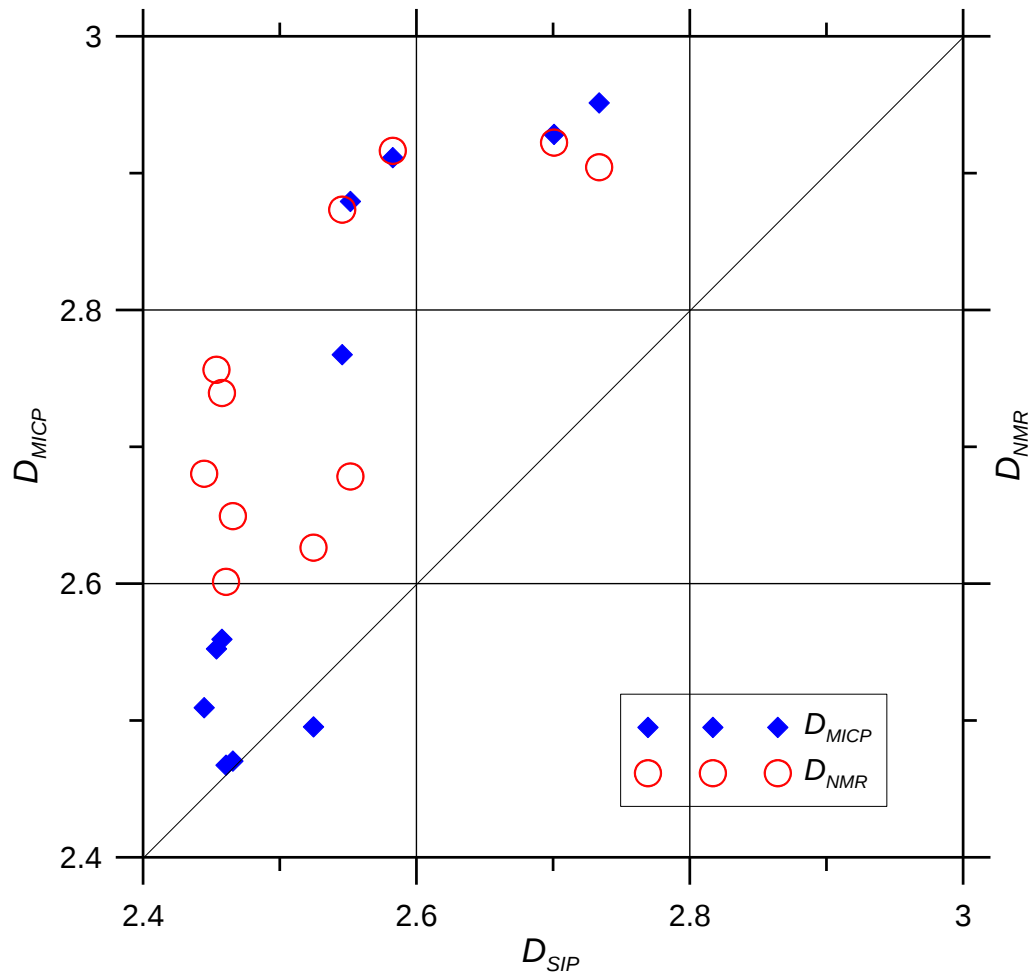
L-Samples:
good agreement between
MICP and NMR with r_{eff}
close to r_{50} . However, the
SIP-curves appear at a much
higher r -range.

$D_{(+)}$ determined using r_{eff}
and τ_{peak} that results from
the frequency of the
maximum of the
imaginary part of
conductivity $\sigma''(f)$
(Weller et al. 2016)

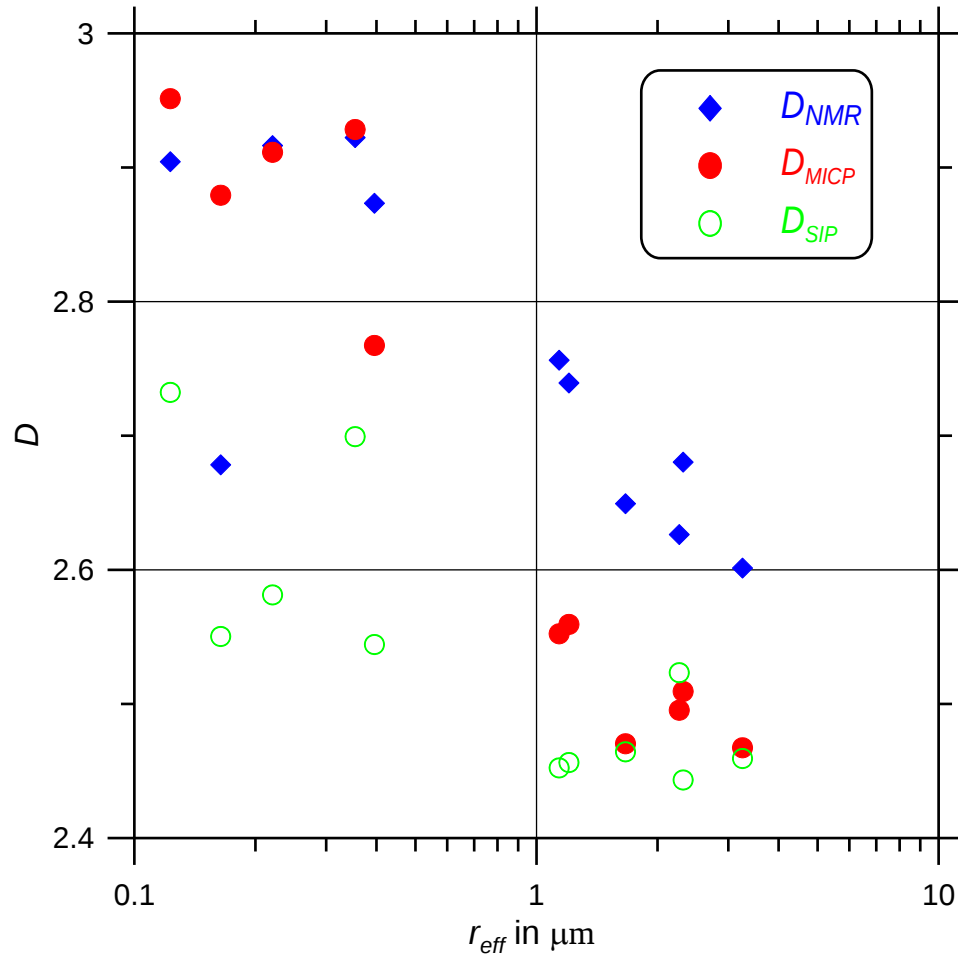
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Results



Results



Conclusions

- A fractal dimension can be determined on the basis of an IP relaxation time distribution.
- If we use a constant diffusion coefficient a good agreement can be observed with the V_c-r curves of MICP and NMR for samples with $r_{eff} > 1 \mu\text{m}$.
- Samples with smaller pore radii indicate larger deviations in the resulting V_c-r curves. This problem can be overcome if instead of a constant diffusion coefficient the apparent diffusion coefficient is used.
- Further investigations will demonstrate the integration of fractal dimension in models of permeability prediction.

Thanks for your attention!

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References

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