

Characterisation of natural organic matter (NOM) and its removal using cyclodextrin polyurethanes

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Abstract

Natural organic matter (NOM) occurs in all natural water sources when animal and plant material breaks down. NOM in water may react with chlorine and other disinfectants to produce disinfection by-products (DBPs), many of which are either carcinogenic or mutagenic. In this study the NOM content of the raw water from the Vaalkop Water Treatment Plant (which uses both chlorination and ozonation as treatment protocols) was characterised after fractionation on ion-exchange resins. Fractionation at different pH values resulted in the isolation of a neutral, a basic and an acidic component of either predominantly hydrophobic or hydrophilic NOM. In addition, NOM results from 3 open water bodies in Johannesburg were evaluated in the same manner. As expected, NOM from all water sources was predominantly hydrophobic (~60%). Each of the 6 isolated NOM fractions was percolated through synthetic cyclodextrin (CD) polyurethanes to determine the extent to which the CD polymers can remove NOM from water. The hydrophobic basic fraction and the hydrophilic acid fraction were most efficiently removed (24% and 10%, respectively). The remaining fractions were not much affected by the polymer treatment.

Keywords: cyclodextrin polyurethanes, disinfection by-products, fractionation, natural organic matter (NOM)

Nomenclature

β -CD	= beta cyclodextrin
CD	= cyclodextrin
BSH	= Braamfontein Dam
CDK	= Coronation Dam
DBP	= disinfection by-product
FLR	= Florida Lake
GAC	= granular activated carbon
HAA	= haloacetic acid
HMDI- β -CD	= polyurethane made by co-polymerising β -CD and hexamethylene diisocyanate
HpoA	= hydrophobic acid fraction
HpoB	= hydrophobic base fraction
HpoN	= hydrophobic neutral fraction
HpiA	= hydrophilic acid fraction
HpiB	= hydrophilic base fraction
HpiN	= hydrophilic neutral fraction
HS	= humic substances (a generally hydrophobic fraction of NOM)
KHP	= potassium hydrogen phthalate
DST/	= Nanotechnology Innovation Centre
Mintek NIC	= Water Platform
NOM	= natural organic matter
oligomers	= short polymers (usually <10 monomers)
ppb	= parts-per-billion (weight/weight i.e. mg/kg)
SUVA	= specific ultraviolet absorbance
THM	= trihalomethane

Introduction

Natural organic matter (NOM) present in most water bodies may react with chlorine or other disinfectants to produce disinfection by-products (DBPs), many of which are either carcinogenic or mutagenic (Xie 2003). For example, haloacetic acids (HAAs), a DBP component, are considered harmful to human health and have diverse toxicological effects, including reproductive and developmental effects, in laboratory animals (Kanokkantapong, 2006). In addition NOM can cause problems of membrane fouling, aesthetically displeasing flavour and malodour, and NOM has even been implicated in the corrosion of turbines and engineering systems (McDonald, 2004). An understanding of NOM in water and its removal is therefore important for human health and in industrial processes.

NOM occurs in all water sources when animal and plant material breaks down. Since this material is different worldwide, so too is the NOM present in each body of water. NOM can be broadly categorised into humic substances (HS), microbial by-products and colloidal natural organic matter, depending on their source and structure.

Humic substances are generally the hydrophobic fraction of NOM and exhibit relatively high specific ultraviolet absorbance (SUVA) values as they usually contain relatively large proportions of aromatic moieties (Chen et al., 2002). Secondly, microbial by-products are composed of acids, with relatively high charge density, polysaccharides, amino sugars and proteins (Vanboon et al., 2005). Finally, colloidal natural organic matter contains relatively polar amino sugars and may exhibit high membrane fouling potential due to its neutrality.

Granular activated carbon (GAC) is widely used to remove organic matter from water, but it often fails to remove these to levels as low as parts-per-billion (ppb) (McDonald et al., 2004). Ozonation, often used in combination with other treatment processes, should lead to the degradation of NOM and formation of low molecular weight compounds (Karnik et al., 2005). These small fragments should be better adsorbed by GAC; however,

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when not well adsorbed on GAC they tend to be more difficult to remove due to their mobility and generally increased polarity (De Wuilloud et al., 2003).

We have previously reported on the use of cyclodextrin polyurethanes for the generation of ultra-pure water in the power industry (Mamba et al., 2008) mainly through the removal of some of these small organic fragments. It is therefore anticipated that some of the NOM and the ozonation fragments, being similar to heat-generated fragments, will be removed efficiently by the use of water-insoluble cyclodextrin polyurethanes.

To better understand the mechanism of NOM removal from water, it is necessary to understand the chemistry of NOM. In addition, this information may allow us to tailor the β -CD polyurethane adsorbents, based on our comprehension of the fundamental interactions of these molecules. Ultimately this should lead to better tailor-made solutions for the treatment of contaminated water. NOM, which is usually found in drinking water at concentration levels between 2 and 15 mg/l (Hepplewhite et al., 2004), can be fractionated or separated into 3 hydrophobic and 3 hydrophilic fractions using suitable ion-exchange resins. The hydrophobic fractions contain mostly humic and fulvic acids whereas the hydrophilic fractions comprise low molecular weight carbohydrates, proteins and amino acids. This fractionation will allow for the evaluation of the extent with which each organic fraction present in the water sample can be effectively removed by the CD polymers, and hence allow for optimisation of adsorbents.

The synthesis of cyclodextrin polyurethanes has previously been reported (Mhlanga et al., 2007). These polymers demonstrate a good capacity to remove a wide range of organic contaminants present in water at concentration levels of ng/l (Mhlanga et al., 2007).

The aim of this work is to determine the different organic fractions of NOM in 'raw water' samples from a water treatment plant (Vaalkop) and 3 open water bodies in Johannesburg, and then to apply water-insoluble cyclodextrin polyurethanes in the removal of these NOM fractions from water.

Materials and methods

Sample collection and preservation

Water samples from the Vaalkop Water Treatment Plant and from 3 dams (Coronation, Braamfontein and Florida Lake) in Johannesburg were collected. At the Vaalkop Water Treatment Plant samples were collected before and after ozonation for the evaluation of the effect of ozone on the NOM present in water. All samples were filtered through a 0.45 μ m cellulose filter paper and stored in the refrigerator at 4°C for not more than 48 h before analysis. Two litres of each sample were then fractionated over ion-exchange resins according to the method by Marhaba et al., (2003).

To evaluate the removal of NOM by the cyclodextrin polymers, each fraction was treated with hexamethylene diisocyanate co-polymerised with *beta*-cyclodextrin (HMDI- β -CD) polymer.

Fractionation procedure

Fractionation of the NOM was performed using 3 types of ion-exchange resins, namely XAD-7HP, Diaion-WA-10, and Dowex © 88, following the procedure described by Marhaba et al., (2003). The result is that the organic matter was divided into 6 fractions, namely; hydrophobic acid (HpoA), hydrophobic base

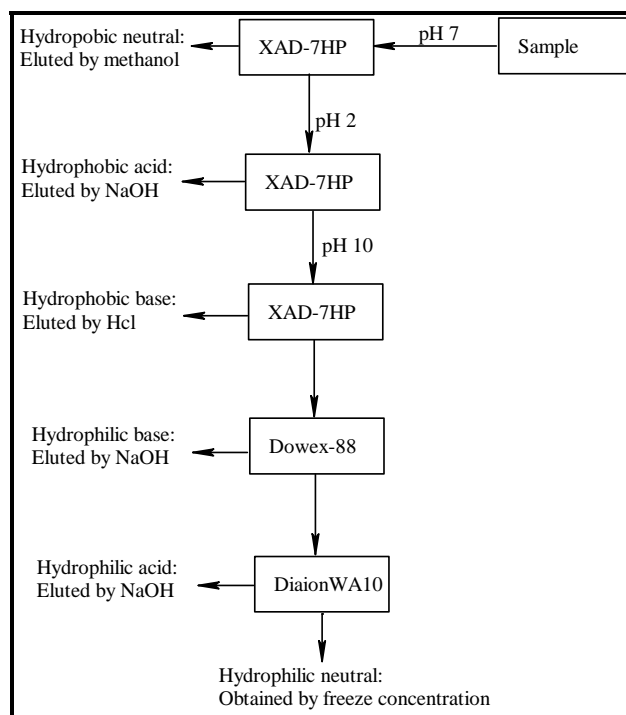


Figure 1
Fractionation method

(HpoB), hydrophobic neutral (HpoN), hydrophilic acid (HpiA), hydrophilic base (HpiB) and hydrophilic neutral (HpiN). The fractionation procedure is illustrated in Fig. 1.

Organic carbon analysis

For the purposes of this study, the dissolved organic carbon (DOC) was used as an indicator for the organic content in the water samples. DOC is that fraction of organic matter passing through a 0.45 μ m filter paper (i.e. it excludes suspended organic matter). The DOC of each NOM fraction was measured with a total organic carbon (TOC) analyzer (Tekmar Dormann Apollo 9000). Standards of 0, 1, 2, 5, and 10 mg/l of total carbon were prepared with potassium hydrogen phthalate (KHP), using ultra-pure water for all dilutions. At least 3 replicates of each measurement were carried out and more replicates performed where the variation between each measurement exceeded 5%.

Ultraviolet (UV) visible spectroscopic analysis

The water samples were also analyzed on a UV-Vis spectrophotometer at a wavelength of 254 nm, as the absorption at this wavelength has been reported to represent the aromatic character of the organic species. UV-254 was measured with a Cary-50 UV spectrophotometer with a 1 cm quartz cell.

Passage of each NOM fraction through polymer

The synthesised polymer was first cleaned to remove un-reacted cyclodextrin and synthesis solvents that could interfere with TOC measurements. This was done by first heating the polymer for an hour in an oil bath at 150°C under vacuum. It was then cooled to room temperature, loaded into empty solid phase extraction (SPE) cartridges, and washed with de-ionised water until the TOC reading of the filtrate was less than 0.5 mg/l of

Sample name	TOC (mg/ℓ)	SUVA (ℓ/mg·m)	Turbidity (NTU)	pH
BSH	4.38	4.22	2.28	7.76
CDK	4.91	5.12	5.93	7.72
FLR	6.46	2.98	2.97	7.60

Note: BSH = Braamfontein Dam, CDK = Coronation Dam, and FLR = Florida Lake

Sample	Raw water (mg/ℓ)	HpoB (mg/ℓ)	HpoA (mg/ℓ)	HpoN (mg/ℓ)	HpiB (mg/ℓ)	HpiA (mg/ℓ)	HN (mg/ℓ)	% Surplus
CDK	4.91	0.12	0.74	0.54	1.12	1.45	1.10	+3.3
BSH	4.38	0.23	1.89	0.57	1.05	0.47	1.02	+19.4
FLR	6.46	0.53	2.12	0.64	1.49	0.72	1.24	+4.3

Raw water (mg/ℓ)	HpoB (mg/ℓ)	HpoA (mg/ℓ)	HpoN (mg/ℓ)	HpiB (mg/ℓ)	HpiA (mg/ℓ)	HpiN (mg/ℓ)	% surplus	SUVA [ℓ/(mg·m)]	pH
12.6	2.91	2.66	2.18	1.06	0.78	4.23	+12.7	2.96	7.8

total carbon. The isolated NOM fractions were then treated with the polymer by passing 30 cm³ of each organic fraction through the polymer at a filtration rate of 10 cm³/min. Filtration was enhanced by the use of a vacuum pump. The polymer-treated water was then analyzed again for its carbon content using the TOC analyzer to determine the per cent carbon absorbed by the polymer.

Results and discussion

Fractionation of organic species

The dissolved organic carbon (DOC) was used as an indicator for the organic content in the water samples. Details of the average mass balance and some chemical parameters of the raw water from the reservoirs around Johannesburg are summarised in Table 1 and Table 2 while those from the Vaalkop Water Treatment Plant are summarised in Table 3.

The specific UV absorbance (SUVA) was used as an indication of the amount of humic substances versus non-humic substances in the water samples. The SUVA characterisation, results from Table 1, indicates that sample CDK and BSH are composed mainly of humic substances while FLR consists mainly of non-humic substances. SUVA results as shown in Table 3, on the other hand, imply that the raw water from Vaalkop is composed mainly of humic substances. The infrared spectra in Fig. 1 display the different functional groups associated with the NOM in raw water.

The mass balance summarised in Table 2 and Table 3 indicates a percentage surplus which meant that the quantity of the 6 organic fractions exceeded the total amount of the initial TOC. This was due to organics released from the resin during elution and a smaller percentage due to organics in the eluants used. It was found that organics released by the resin were at times as high as 0.5 mg/ℓ. This resulted in a weight surplus in the resulting organic fractions.

These data (summarised in Table 2) indicate that all the water samples were composed of mostly hydrophilic fractions, namely

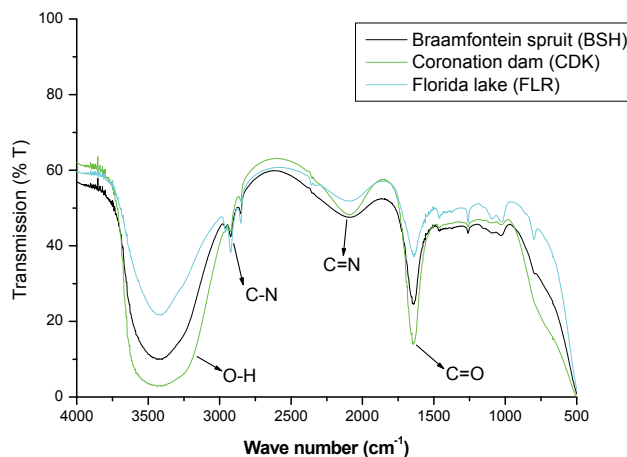


Figure 2
An IR spectrum displaying the functional groups mainly associated with the NOM in raw water

71.8%, 54.6% and 51.9% for CDK, BSH and FLR, respectively. The relatively high percentage of the hydrophilic substances as compared to the hydrophobic substance suggests a contamination of the water source.

Similar to the open water sources around Johannesburg, data from the Vaalkop treatment plant (summarised in Table 3) indicates that the hydrophobic fractions are more abundant than the hydrophilic organic fractions; 56.1% and 43.9%, respectively.

These findings are in agreement with accepted knowledge that hydrophobic substances would constitute a larger fraction than the hydrophilic fractions in natural water sources.

Passage through β -HMDI CD polymer

Only the samples from the Vaalkop Water Treatment Plant were passed through the polymer for this part of the study. Also, although samples were taken both before and after ozone treat-

NOM fraction	TOC before passing through polymer (mg/ℓ)	TOC after passing through polymer (mg/ℓ)	% absorbed
HpoB	2.91	2.21	24.1
HpoA	2.66	2.52	5.26
HpoN	2.18	2.18	0
HpiB	1.06	1.05	0.9
HpiA	0.78	0.70	10.3
HpiN	4.23	4.20	0.71

ment at the plant, it was later discovered that the ozone generator at the plant was malfunctioning, and there was no change in the relative proportions of the 6 fractions.

The data presented in Table 4 show that the percentage NOM absorbed by the polymer was not very high. The hydrophobic fractions were generally better adsorbed on the polymer than the hydrophilic fractions, with almost a quarter of the hydrophobic basic fraction (HpoB) being removed. This was followed by the hydrophilic acidic (HpiA) fraction with 10.3% removal. The hydrophobic neutral was the least absorbed fraction with 0% absorbance on the polymer.

Conclusion

NOM present in 3 natural water bodies and 1 treatment plant in South Africa was characterised by fractionation on ion-exchange resins. As expected in an urban environment, the hydrophobic fractions from both the reservoirs around Johannesburg and the Vaalkop Water Treatment Plant was composed mainly of humic substances, apart from the water from Florida Lake (FLR), which is composed mainly of non-humic substances. The extent to which each of the 6 NOM organic fractions could be removed by a synthetic CD polymer was evaluated to obtain a base-line for future research. Although the polymer showed a relatively low potential for removing the NOM fractions from the source water, the hydrophobic basic fraction showed an encouraging 24% removal. Future work involves pre-treating the samples with different ozone dosages in order to break down the NOM. This post-ozone treated water will then be passed through the cyclodextrin polymers to evaluate the efficiency of this combined removal process.

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