Differential sensing of sugars by colorimetric arrays
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While the complexes between boronic acids and diols have been studied for decades, researchers continue to design new and interesting methods to use these interactions to produce saccharide sensors that are more sensitive and selective. Herein we discuss how the use of pattern-based colorimetric arrays from a collection of crossreactive sensors have been developed as new differential sensing platforms for sugars and related saccharides.

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Introduction
Since the first synthesis of phenylboronic acid in the late 19th century, scientists have been interested in the unique properties associated with boron chemistry. Not until the mid-20th century, however, were the interactions between boronic acids and diol-containing compounds discovered [1]. While the reaction was debated for several years, it soon became apparent that the formation of cyclic boronate esters dominated. In a review published by Shinkai and coworkers in 1996 [2], numerous synthetic receptors for the optical detection of saccharides based on boronic acid receptors gained notoriety and helped launch a new field of chemical sensors. In 2002, Wang et al. discussed work on boronic-acid-based sensors [3], leading the group to later publish several articles related to the pH and pKₐ dependence of boronic acid–diol interactions. It should be noted that while fluorescence-based saccharide detection is not within the scope of this review, significant work has also been reported with that approach [4–9].

Pattern-based sensing, similar to what we now know of the human olfactory and gustatory systems, relies on a selection of crossreactive semiselective indicators or receptors, the responses of which are analyzed as a whole in order to gain information on a diverse chemical library. While the concept has been well documented, only recently has pattern-based, differential sensing made its way into the forefront of sensing platforms and has found useful applications to a wide range of analytes, including peptides [10–12], proteins [13], volatile organic compounds [14–16], toxic industrial gases [17,18**], organic compounds in water [19], and many others. Herein we discuss the use of differential sensing for the colorimetric detection of sugars and sugar substitutes.

Interactions between boronic acids and diols
Boronic acids bind strongly with diol-containing compounds with high affinities via boronate ester formation (Figure 1). The reaction, discovered in 1954 by Kuivila et al., has been the basis for many synthetic molecular receptors, but considerable recent efforts have been made to improve the strength of binding, thus paving the way for more sensitive sensing methods for diol-containing targets [20,21]. Shinkai and James have spent the better part of a decade designing new and interesting sensing platforms based on the complexation of the two species [22–24]. The selectivity relies in part on the differences in association constants of different boronic acids with diols resulting in changes in equilibrium between the boronic acid and the cyclic boronate ester.

It has been shown that changing the parameters of the experiment (e.g. solution pH, boronic acid pKₐ, etc.) can result in drastically different binding constants between boronic acids and diols. As a result, derivatization techniques have been employed to produce a plethora of molecular sensors for saccharides based on phenylboronic acid. In many cases, phenylboronic acids are coupled with chromophores to give a color response to binding of the boronic acid moiety. In this manner, several color-changing motifs have been utilized including, indicator displacement [26**,27,28*], boronic acid appended chromophores [24,29–31,32*], and pH induction [33,34*], and these have been extended to produce differential sensing platforms optimized to human body temperature and physiological pH [23,35–37].

Colorimetric sensors and sensing methods
Indicator displacement assays
The use of indicator displacement assays (IDAs) for colorimetric sensing has been pioneered most recently by Anslyn and coworkers [26**,38,39]. The basic principle
relies on the affinity for two competing guests to one host. For sensing, the analyte displaces the indicator from the host or receptor (or vice versa), causing the indicator to change its spectroscopic properties (c.f. Figure 2). For saccharide detection, color-changing indicators containing cis-diols were optimized to create an IDA sugar sensor. Among the most common of these indicators is alizarin red and bromopyrogallol red. IDA inherently represents an equilibrium competition between the indicator and the analyte of interest; this has a potential downside of limiting sensitivity, particularly for lower-affinity analytes.

Indicator displacement has recently gained new ground via a sensing motif based on dye displacement using boronate hydrogels [28*]. Making use of a new development, boronate affinity in saccharide electrophoresis, or BASE, that relies on the modulation of saccharide mobility within a flexible porous hydrogel, the group went on to quantitatively detect a wide range of fructose concentrations at physiological pH; the indicator displacement scheme used can be seen in Figure 3. Recently, the synthesis of a similar sensing platform has been reported based on phenylboronic acid derivatization of solid silica supports [27].
**Boronic-acid appended chromophores**

**Chemosensors**

Recently, various colorimetric sensors based on boronic acid appended chromophores have been designed and studied for detection of a host of analyte classes including amines, aldehydes, alcohols, and in particular, *cis*-diols. Mohr has published numerous articles focused on the use of reversible covalent-bonding sensing motifs combining a chromogenic center with a reactive terminus capable of color-change upon reaction with the desired analyte [40–43]. Several chromogenic receptors designed by Shinkai *et al.* are based on photo-induced electronic transfer (PET) created by complexation between the receptors’ boronic acid functionality and saccharides. This strengthens the Lewis acid–Lewis base interaction within the receptor, resulting in PET from the amino nitrogen to the chromophoric center, thereby causing a change in color (Figure 4a) [2,31]. Since a shift in the absorbance of most azo-dyes is directly associated with changes in the environment of one of the bridging nitrogens, some sugar sensors have been designed that make use of a spectral change based on the formation and cleavage of the B–N bond between the boronic acid and the azo linkage (Figure 4b) [30,32]. In other examples, the boronic acid group is directly attached to the chromophore, whose color changes upon chelation of the boronic acid group to a monosaccharide (see Figure 4c) [29]. In this manner, facile sugar detection has been possible, particularly involving the differentiation of glucose from fructose.

**Macrocyclic ring opening and oxidation**

A new class of color-changing saccharide sensors was developed earlier this decade by Strongin and coworkers.
This sensing motif takes advantage of phenylboronic acid appended macrocycles undergoing significant color changes in the presence of carbohydrates as well as glucose phosphates, carboxylic acid, and amino sugars. It was found that solutions containing the tetra-arylboronic acid resorcinarene macrocycle (Figure 4d), undergo a condensation reaction in situ to form a xanthene chromophore. The authors argue that ring opening of macrocycles to acyclic oligomers could be a prerequisite for xanthene formation from resorcinarenes. In this way, complexation of a sugar with the newly formed acyclic chromophores was shown to semiselectively discriminate among a family of closely related saccharides.

**pH-induced color changes**

Oddly, one of the least exploited characteristics regarding the reactions of boronic acids with saccharides had been the production of protons resulting from the formation of a cyclic boronate ester. The pH depression associated with the complexation of boronic acids with saccharides is, of course, buffer dependent, and in most of the detection methods mentioned thus far, systems have been strongly buffered. That being said, several groups have developed sugar-sensing systems based mostly, if not solely, on induced pH depression. Among these, Chang and coworkers have presented a wonderful piece of work wherein pH indicators were employed to monitor and report the change in solution pH upon the addition of sugars [34]. More recently, pH reporter dyes have been incorporated with one of several phenylboronic acid candidates (c.f. Figure 4e) [33]. This new sensor technique can selectively and quantitatively differentiate between fructose and glucose at millimolar concentrations.

**Array-based (pattern-based) sensing**

The detection and identification of chemicals is, fundamentally, supramolecular chemistry and intrinsically relies on the interactions between molecules [46]. The classification and relative strength of intermolecular interactions is of course very well established and includes Lewis acid–base interactions (the limiting case of which is bond formation and metal ion coordination), Brønsted acid–base interactions, hydrogen-bonding, charge-transfer and π–π molecular complexation, dipolar and multipolar interactions, and van der Waals interaction and physical adsorption. The use of sensors in arrays that probe this full range of intermolecular interactions is essential to the further development of array technology.

Array-based sensing has emerged as a potentially powerful approach toward the detection of chemically diverse analytes. On the basis of crossresponsive sensor elements, rather than analyte-specific receptors for individual analytes, sensor arrays produce composite responses and it is the composite response that is unique to an odorant or taste, just as in the mammalian olfactory and gustatory systems. In this design architecture, one receptor responds to many analytes and many receptors respond to any given analyte. It is the distinct pattern of responses produced by the array that provides differential sensing of multiple analytes.

For nearly 50 years, the concept of pattern-based discrimination of a broad range of analytes, also known as ‘differential sensing’, has emerged as a powerful sensing tool. Recently, Anslyn and coworkers have successfully used arrays of crossreactive sensors in combination with IDA and kinetic measurements to differentiate among a wide selection of aqueous analytes including, peptides [12], nucleotide phosphates (e.g. ATP, AMP, and GTP) [47], proteins [48], and recently, saccharides. Using boronic-acid-based peptidic receptors; chemosensors were developed capable of discrimination among a number of monosaccharides, disaccharides, and saccharide derivatives such as sucralose (the sweetening agent used in Splenda brand sweetener) and maltitol (a sugar alcohol derived by the reduction of maltose) [26]. Multiple liquid wells were assembled using a randomly chosen set of receptors from a combinatorial library of pentapeptidic boronic acids optimized to work at
physiological pH. A three-dimensional plot relating analytes, receptors, and average rate constants for indicator adduct displacement is shown in Figure 5. The plot shows how each semiselective receptor responds to each analyte via rate constants. Indicator displacement assays were carried out, generating a set of response data that, when coupled with dimension-reducing chemometric tools such as linear discriminant analysis (LDA), can classify both structurally similar and chemically diverse analytes.

Another example of differential sensing is described by Chang and coworkers, who have recently published their work on a pH change-induced carbohydrate sensing ensemble that allowed for the differential recognition of 23 monosaccharides, disaccharides, and trisaccharides at millimolar concentrations [34]. Twelve dyes were employed in an array of aqueous solutions in well plates, covering a large pH range along with two boronic acids: boric acid and phenylboronic acid. Experiments for each system were carried within a pH range that optimizes the production of the cyclic tetrahedral boronate ester and thus depress the solution pH. A principal component analysis (PCA) scatter plot showing the discrete clusters of saccharides can be seen in Figure 6.

The Suslick research group has been developing colorimetric sensor arrays to detect a myriad of analytes for over a decade. This work started as an application of metalloporphyrins as sensor elements for array-based vaporsensing [49]. The arrays were broadened to include a wide range of chemically diverse sensors, adding pH indicators (which actually indicate much more than just pH), solvatochromic dyes, and shape-selective receptors to detect and differentiate between chemically diverse
analytes [50–52]. We have successfully designed colorimetric sensor arrays capable of differential sensing of volatile organic compounds [14], organics in water [19], amines [53], toxic industrial chemicals (TICs) [17, 18**], and even coffees [54], beers [55], and soft drinks [56]. Our previous printing formulations had been based on plasticized hydrophobic colorants that showed resistance to humidity while successfully detecting gaseous analytes and hydrophobic organic analytes in aqueous solutions.

The sensitivity of our earlier hydrophobic sensing platform [19], however, proved to be problematic for the sensing of hydrophilic analytes, including carbohydrates. We determined that rapid and sensitive detection of these classes of analytes requires an immobilization method whereby all chromogenic centers are accessible to analytes while the dyes themselves must remain impervious to leaching or blooming upon exposure to aqueous solutions. After several techniques were tested, we developed a printable sensor array based on nanoporous organically modified polysiloxanes (ormosils) to encapsulate a diverse set of chemically responsive dyes. These nanoporous pigments were printed onto hydrophilic, porous membranes, which permits the user to avoid tedious handling of multiple aqueous solutions in well plates. One simply exposes the analyte solution once to a single printed array and images on an ordinary flat bed scanner. We successfully used this approach to detect and identify 14 structurally similar monosaccharides and disaccharides as well as reduced sugars and sugar substitutes at low millimolar concentrations [25**]. Most of this discrimination was based on the induced solution pH changes from 3-nitrophenylboronic acid–diol complexation an initially neutral pH. Color difference maps are useful in showing how colorimetric sensor arrays can provide ‘molecular fingerprints’ of the analytes tested. As visualized qualitatively in difference maps, discrimination can be easily achieved, as shown in Figure 7a.

For display of the discrimination among the sweeteners and sugars, we use a standard chemometric approach, hierarchical cluster analysis (HCA), which is based on the grouping of the analyte vectors according to their spatial distances in their full vector space [57]. HCA has the advantages of being model-free (unlike, e.g. LDA or neural nets) and of using the full dimensionality of the data. As shown in Figure 7b, HCA generates dendrograms based on clustering of the array response data. Using HCA, we showed that our array was capable of discrimination of the analytes with discreet clustering and no misclassifications among 14 different analytes in 75 trials. As an extension of this work, we were also able to test our array’s performance in real-world situations using common table-top sweetener packets readily found in any restaurant or cafe, both in aqueous solution and in sweetened iced-tea. Differential sensing among these analytes was observed without error. Color difference maps representing the arrays response to one serving of several representative sweeteners and a control are shown in Figure 8.
Conclusion and future endeavors

In recent years, we have seen substantial progress in the design, optimization, and application of semiselective, crossresponsive indicators for the differential sensing of a host of analytes, including sugars, the focus of this overview. By manipulating the old and well-understood reaction of boronic acids with diols, scientists have developed arrays of chemoresponsive chromophores for the detection and identification of sugars and sugar analogs. Using biomimetic pattern-based sensing similar to the mammalian olfactory and gustatory systems, the development of electronic noses and electronic tongues based on colorimetric sensor arrays are providing advancements in sensitivity and pattern-based selectivity above and beyond that which nature provides.

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References and recommended reading

Papers of particular interest, published within the period of review, have been highlighted as:

- of special interest
- of outstanding interest

13. Miranda OR, Chen H-T, You C-C, Mortenson DE, Yang X-C, Burz UHF, Rotello VM: Enzyme-amplified array sensing of...
Differential sensing of sugars

Musto and Suslick 765


19. A colorimetric sensor array made from crossreactive nanoporous pigments was developed for rapid and sensitive detection and identification of 20 different toxic industrial chemicals (TICs).


27. Colorimetric sensor solid-state arrays made from nanoporous pigments were used to differentiate 14 sugars, sugar alcohols, and artificial sweeteners at low millimolar concentrations, imaged with an ordinary flat bed scanner. Real-world applications of the array are demonstrated using tea sweetened with coffee-house sweeteners.


29. This comprehensive work describes, in detail, the workings of indicator displacement assays as they relate to the differential sensing of sugars and sugar substitutes.


32. An interesting semisolid-state indicator displacement assay. Easily synthesized hydrogel spheres incorporate boronate units that function as sugar sensors.


37. A culmination of several studies performed by the group. In this latest publication, multilayer films based on boronic acid appended azo-dyes were synthesized and employed to differentiate glucose from fructose.


40. A pattern-based sugar-sensing technique relying on the pH depression at nonphysiological pH associated with cyclic boronate ester formation when diols interact with boronic acids.


53. A compelling review of recent advances in analytical chemistry using synthetic receptors meant to mimic natural receptors. Several techniques were discussed including indicator displacement assays (IDA), and photo-induced electron transfer (PET), and their employment as differential receptors.


