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A rapid and sensitive method for chiroptical sensing of α -amino acids *via* click-like labeling with *o*-phthalaldehyde and *p*-toluenethiol†

Bo Li,^{†ab} Jie Zhang,^{†a} Li Li^{ID}*^a and Gong Chen^{ID}*^b

A highly practical method for comprehensive chiroptical sensing of free α amino acids with streamlined operation and high sensitivity *via* dual CD/UV measurements is developed. The assay takes advantage of an efficient and selective three-component labeling reaction of primary amines with *o*-phthalaldehyde and *p*-toluenethiol reagents to derivatize the NH₂ group of analytes into an isoindole. The covalent labeling generates sensitive UV and CD readouts, both of which show an excellent linear relationship with the concentration of analytes. The high reactivity and the novel optical reporting mechanism allow fast and accurate measurement without background interference. The sensing assay works well for a remarkably broad range of analyte concentrations, with an unprecedented lower limit of 10 micromolar concentration.

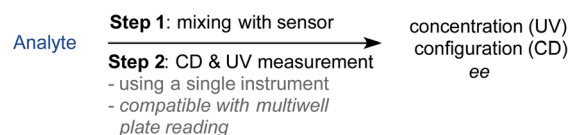
Introduction

The widespread use of α -amino acids (α AAs) and other chiral amine compounds in academic and industrial laboratories has generated substantial interest in developing sensitive and convenient methods for their chiral analysis.¹ Due to the operational limitations of chromatographic techniques, attention has been increasingly shifted to optical methods.^{2,3} Over the past decade, methods for chiroptical sensing based on circular dichroism (CD) spectroscopy with small molecule sensors have been greatly advanced by the groups of Chin, Anslyn, Wolf, Pu, Joyce, Zonta and others.⁴⁻⁷ By exploiting the mechanisms of dynamic covalent chemistry, supramolecular assembly, and metal complexation, these methods both amplify the CD signal and report the concentration to allow quantitative measurement of the enantiomeric composition of analytes with high accuracy.

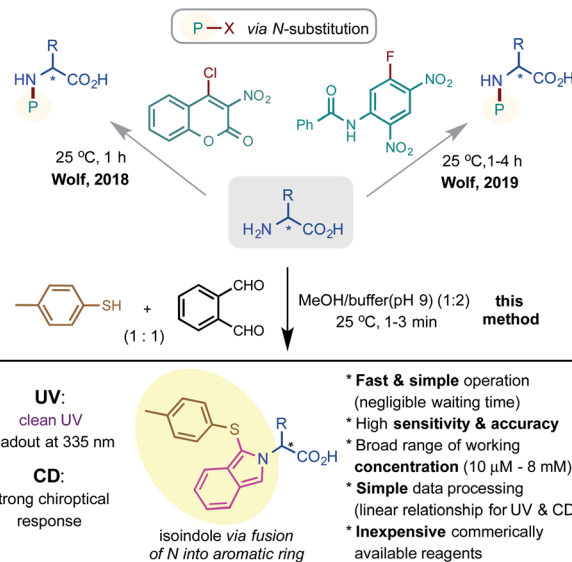
Among these chiroptical methods, the combination of CD and UV is probably the most desirable as the measurements can be conveniently performed on a single instrument, which can be readily modified with multiwell plate reading for parallel analysis (Scheme 1A). Notably, the Wolf group reported several powerful mix-and-measure protocols using organohalide

derivatizing agents (a coumarin chloride and an aryl fluoride) that selectively react with the amine analytes to generate CD and UV readouts (Scheme 1B).^{7,8} However, while labeling reactions

A) Mix-and-measure protocol using dual CD/UV measurements



B) Click-like covalent labeling of chiral amines with reporter probes



^aBeijing Key Laboratory of Active Substances Discovery and Druggability Evaluation, Institute of Materia Medica, Chinese Academy of Medical Sciences, Peking Union Medical College, Beijing 100050, China. E-mail: annaleelin@imm.ac.cn

^bState Key Laboratory and Institute of Elemento-Organic Chemistry, College of Chemistry, Nankai University, Tianjin 300071, China. E-mail: gongchen@nankai.edu.cn

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‡ These authors contributed equally to this work.

Scheme 1 CD/UV-based quantitative chiroptical sensing of α AAs with organic probes.



via N-substitution can proceed cleanly, they require relatively long reaction times (1–4 hours) at high reaction concentrations (typically > 1 mM). Herein, we report a new method for chiroptical sensing of various α -amino acids *via* a click-like three-component labeling of primary amines with *o*-phthalaldehyde

and *p*-toluenethiol reagents.⁹ The resulting N-fused aromatic isoindole moiety provides strong UV and CD responses, enabling fast, sensitive, and accurate measurement across a very broad range of analyte concentrations.

Results and discussion

The key to designing dual sensors for CD/UV-based assay is to generate strong and clear readouts for both CD and UV. While a variety of sensing mechanisms have been successfully exploited to amplify the CD signal of α AAs, methods for simultaneous UV amplification with high sensitivity are surprisingly limited. The three-component condensation reaction between the amino group of α AAs, *o*-phthalaldehyde (*o*PA) **2**, and alkyl thiol reagents such as 2-mercaptoethanol **3a** or 3-mercapto propanoic acid **3b** has long been used to derivatize α AAs for HPLC analysis (Fig. 1A).^{10–12} The *o*PA labeling reactions can proceed cleanly and quickly under mild conditions to form 1-thiolate substituted isoindole product **5**. The isoindole moiety has a strong UV absorption around 335 nm, offering a clean UV readout with little background inference from the α AA and the labeling reagents. Its emission around 450 nm has also been used for fluorescence detection. Encouraged by their favorable UV properties, we questioned whether the N-fused isoindole structure on the α AA can induce useful CD signals for chiroptical sensing. The three-component condensation reaction is believed to proceed through a cyclic hemiaminal intermediate **4**, which upon dehydration gives the heteroaromatic product.

We commenced the investigation with the reaction of a model α AA L-alanine **1** and a stoichiometric mixture of *o*PA and different thiols. To our delight, the isoindolyl derivatives of Ala indeed exhibited an excellent CD response around 335 nm with *p*-toluenethiol **3e** giving the strongest signal among the thiol reagents tested. **3e** is a commercially available solid compound and has a much weaker odor than the liquid alkyl thiols. The UV spectrum of the reaction mixture showed a distinct absorption at 335 nm (Fig. 1B).¹³ As indicated by LC-MS and UV measurements, 1.2 equiv. of **3e** and **2** reacted cleanly with 1 equiv. of **1** at 2 mM in the mixed solvents of MeOH and phosphate buffer (pH 9) (1/2) at room temperature (rt) under an air atmosphere to give product **5e** in >95% conversion in 1 min. Notably, the use of excess amounts of **2** and **3e** (*e.g.* 3 equiv.) has a negligible impact on the readout of CD and UV at 335 nm (Fig. 1B). The reaction mixture was diluted for CD and UV measurements. UV and LC-MS analyses showed that product **5e** is stable over a period of 12 hours.¹⁴ The UV and CD spectra of **5e** are not strongly affected by the solvents (see the ESI† for assaying spectra using other organic solvents). Aqueous MeOH medium is preferred for its excellent solubilizing ability for α AAs and lack of interference for UV and CD measurements.

As shown in Fig. 1C, our preliminary structural and spectroscopic analysis using time-dependent density functional theory (TDDFT) calculations showed that the tolyl group of (*S*)-**5e** could be positioned below (type (i) conformers) or above (type (ii) conformers) the isoindole plane, forming the opposite

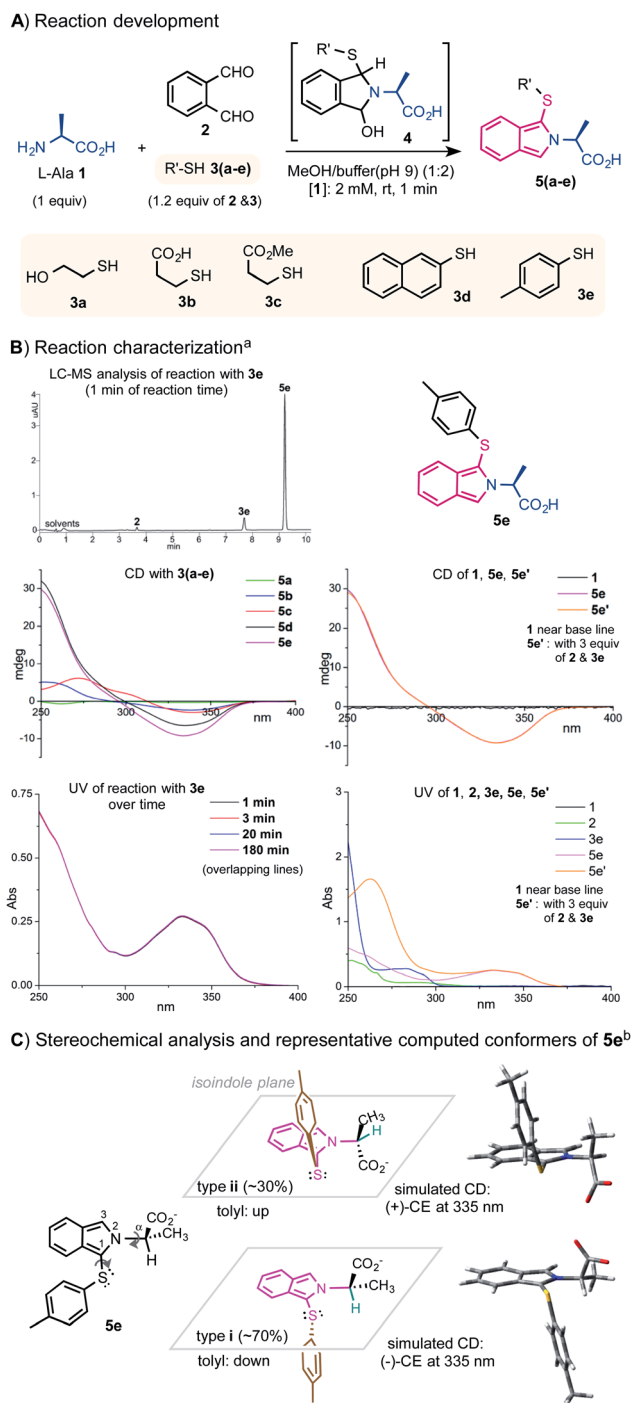


Fig. 1 Labeling of L-Ala with *o*PA and thiols for chiroptical sensing by CD/UV. (A) Reaction development; (B) Reaction characterization; a) reaction mixture was diluted 7 times and samples were measured at 0.285 mM for CD and UV. (C) Stereochemical analysis and representative computed conformers of **5e**; b) DFT calculations were performed at the M062X/6-311+G(d,p) level, see the ESI† for details.



relative stereochemical arrangement of the two aryl groups along the C₁-S bond. Both types (i) and (ii) prefer near-eclipse conformations to the C α -H bond and adopt *syn* and *anti* to the C₁-S bond respectively. Type (i) and (ii) conformers give opposite Cotton effects (CE) according to the electronic circular dichroism (ECD) simulation. Type (i) conformers are more stable than type (ii) conformers; (i) and (ii) roughly represent 70% and 30% in the conformational equilibrium mixture respectively. The overall ECD spectrum of (*S*)-**5e** matched well with its experimental data, which showed a negative CE at 335 nm (see ESI Fig. 10† for details).

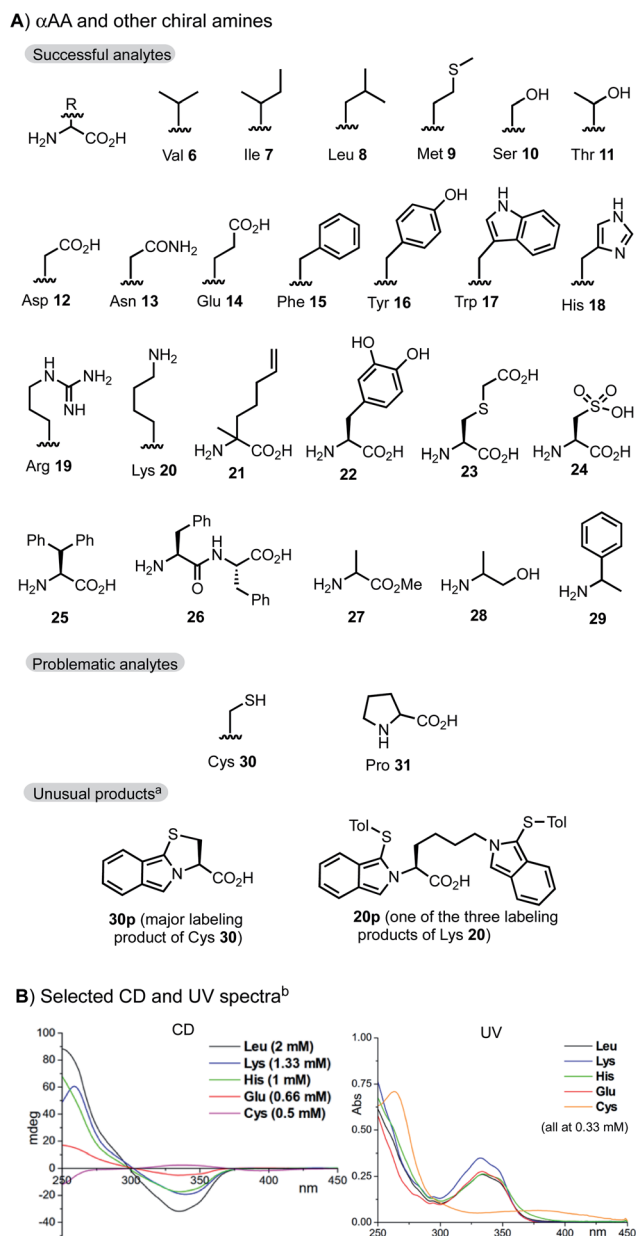


Fig. 2 Substrate scope at a normal concentration range. Standard assaying conditions (1.2 equiv. of reagents, 1 min). (A) α AA and other chiral amines; a) see the ESI† for LC-MS analysis of the reaction mixture. (B) Selected CD and UV spectra; b) samples were measured at the concentration specified.

The new protocol was next applied to sense other optically pure proteinogenic α AA and primary alkylamine samples (Fig. 2A). Typically, 1 equiv. of the analytes and 1.2 equiv. of **2** and **3e** were mixed in MeOH/buffer at 0.5–8 mM concentration for 1 min. In practice, the assaying can be performed without any deliberate aging. All α AAs except Cys and Pro showed a similar profile and sensitivity in both CD and UV spectra. The CD and UV spectra of selected α AAs are shown in Fig. 2B. Pro **31** cannot form the corresponding isoindole product due to its secondary amino group. The reaction of Cys **30** mainly formed a tricyclic isoindole product **30p** *via* the intramolecular addition of the SH side chain along with two dimeric side products (see the ESI†). The reaction of Lys **20** with 1.2 equiv. of **2** and **3e** gave a mixture of mono- and bis-labeled products (*e.g.* **20p**) at the α and ϵ amino positions (see ESI Fig. 9† for details). Side chains such as CO₂H (Glu, Asp), CONH₂ (Gln, Asn), OH (Ser, Thr, Tyr), guanidine (Arg), and imidazole (His) did not interfere with the condensation reaction. Reactions of nonproteinogenic or unnatural α AAs **21–25** and even dipeptide Phe–Phe **26** gave similar results. Ala methyl ester **27** gave an almost identical CD readout to Ala **1**. Assaying of alaninol **28** and 1-phenylethylamine **29** gave a slightly weaker CD signal (\sim 40%) in comparison with Ala **1**.

As shown in Fig. 3A, the UV absorption at 335 nm shows an excellent linear relationship with the concentration of Trp between 0.5 and 8 mM (all samples above 0.5 mM were diluted for CD and UV measurements). Moreover, the CD signals at 335 nm also showed an excellent linear relationship with the enantiomeric excess (ee) of Trp samples assayed at 2 mM using the standard protocol with 1.2 equiv. of *o*PA and **3e**. As shown in Fig. 3B, plots of the *g* factor of CD *vs.* ee% for representative α AAs Leu **8**, Glu **14**, His **18**, and Thr **11** measured at 0.33–1 mM showed an excellent linear relationship. The plot of UV absorption *vs.* concentration showed a notable deviation ($R^2 = 0.9852$) from the linear relationship for Lys probably due to the formation of mixed labeling products (see the ESI†). However, the plot of *g* factor *vs.* ee% showed excellent linear relationship for Lys.

Chiroptical sensing of analytes at low concentrations remains a difficult challenge for the existing methods due to low labeling reactivity and high background noise. We were pleased to find that our method worked well for α AA samples below 300 μ M under slightly modified conditions with excess amounts of *o*PA and **3e** (5–100 equiv.) and 3 min of mixing time. As exemplified by the test of Ala, the UV signal at 335 nm showed excellent linear relationship with the concentration between 5 and 400 μ M (assaying samples below 0.5 mM were measured without dilution). Moreover, the plot of *g* factor *vs.* ee% showed an excellent linear relationship for Ala at 10 μ M.¹⁵ Notably, excess amounts of reagents had a negligible impact on the CD and UV signals at 335 nm, due to the unique signal amplifying mechanism of this method. Sensing of the Trp sample at 10 μ M gave similar sensitivity (see the ESI†).

As shown in Table 1, our assay was tested with representative Ala and Trp samples at varied concentrations and of varied enantiomeric ratios. The simple linear relationships greatly simplified the calculation of the concentrations and ee values



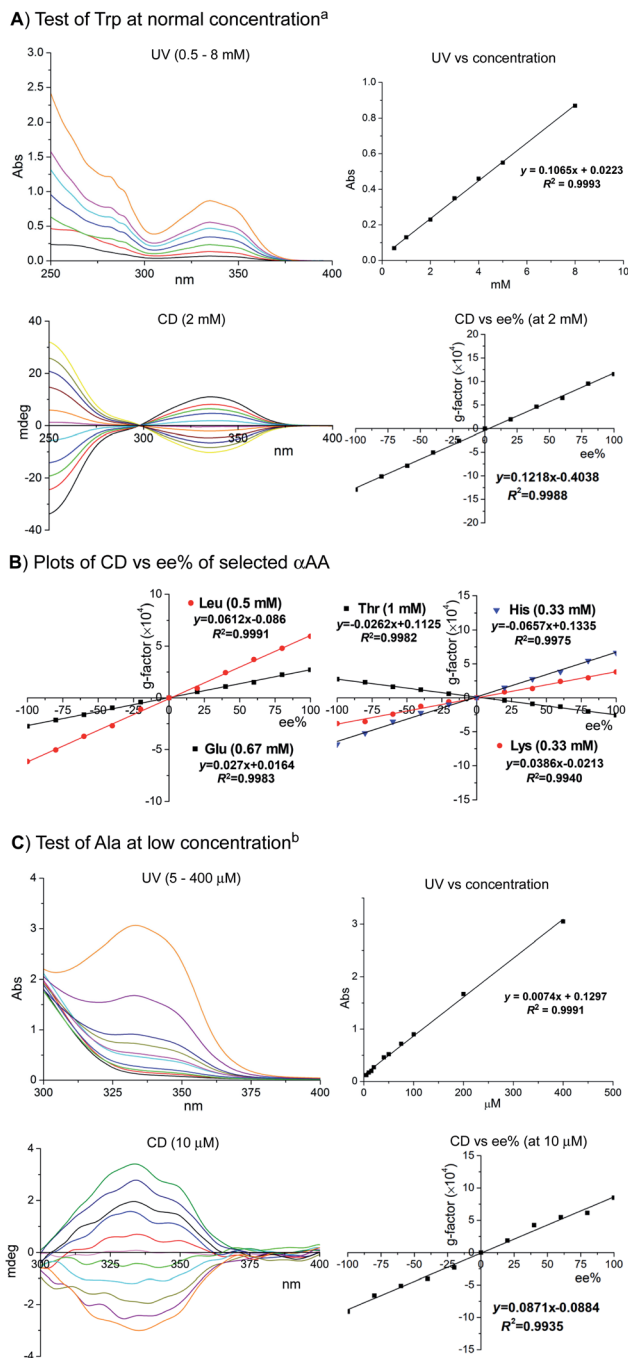


Fig. 3 Assaying α AAs at varied concentrations. Standard conditions: 1.2 equiv. of **2** and **3e**, 1 min for $[\alpha\text{AA}] > 300 \mu\text{M}$. Modified conditions: 5 equiv. and 1 min for 100–300 μM ; 20 equiv. and 3 min for 50–100 μM ; 100 equiv. and 3 min for 5–50 μM . See the ESI† for tests of Trp at 10 μM and Ala at 2 mM. (A) Test of Trp at normal concentration; a) all samples were diluted 6 times for the UV vs. concentration plot. The real measured concentration is 1/6 of the ones shown in the X axis. All samples (2 mM) were diluted 4 times for the CD vs. ee% plot, measured at 0.50 mM. (B) Plots of CD vs. ee% of selected α AA. (C) Test of Ala at low concentration; b) assaying samples were measured without dilution. The *g*-factor was the average of four measurements.

from UV and CD measurements. In all cases, the absolute configuration of the major enantiomer was correctly assigned. The measured data of concentration and ee values were mostly

Table 1 Chiroptical sensing of Ala and Trp samples at varied concentrations and ee values

Entry	Sample			Sensing		
	Conf.	Conc.	ee%	Conf.	Conc.	ee%
1	Ala-L	2.00 mM	−100.0	L	1.91 mM	−98.7
2	Ala-D	7.00 mM	37.1	D	6.87 mM	38.2
3	Trp-L	7.00 mM	−100.0	L	7.10 mM	−103.4
4	Trp-D	3.50 mM	42.8	D	3.57 mM	43.9
5	Ala-L	40.0 μM	−60.0	L	39.2 μM	−61.2
6	Ala-D	10.0 μM	100.0	D	9.5 μM	99.6
7	Trp-L	21.0 μM	−65.0	L	22.2 μM	−67.2
8	Trp-D	38.0 μM	79.0	D	35.8 μM	77.5

within 5% error of the actual value at both high (8 mM) and low (10 μM) assaying concentrations.

Conclusions

In summary, we have developed a highly practical method for comprehensive chiroptical sensing of free α amino acids which gives high sensitivity *via* dual CD/UV measurements using a single instrument. The click-like covalent labeling reaction rapidly fuses the NH_2 group of analytes into a *N*-heteroaromatic isoindole structure, which amplifies both UV and CD signals of amines with an excellent linear relationship with the concentration. The high reactivity and the novel optical reporting mechanism of the labeling reaction allow fast and accurate measurement with negligible aging time and background interference. The sensing assay works well for a remarkably broad range of analyte concentrations with an unprecedented lower limit of 10 micromolar concentration. The sensing assay is simple, robust and cheap. We expect that this method can be readily adapted for high throughput experimentation analysis using a CD instrument equipped with a multiwell plate reader.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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- 13 Compound **5e** is weakly fluorescent (emission maximum at 450 nm, Φ : 8.3% in MeOH/H₂O).
- 14 In the classic oPA labelling reactions of α AAs with alkyl thiol reagents such as **3a** and **3b**, the corresponding isoindole product **5** can slowly react with O₂ to form oxidized by-products. In comparison, **5e** with the tolyl thiolate group is stable and does not show evident decomposition in the reaction mixture within 12 hours under an air atmosphere.
- 15 The plot of *g*-factor vs. ee% showed a better linear relationship than the plot of molar ellipticity vs. ee% at 10 μ M (R^2 : 0.9935 vs. 0.9913, see the ESI† for details). Measurements of Ala at 5 μ M gave acceptable sensitivity at high ee values while the sensitivity dropped considerably at 2.5 μ M.

