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## COMMUNICATION

A 2<sup>nd</sup>-Generation Scalable Synthesis of the HIV-1 Entry Inhibitor CJF-III-288 Enabled by Photoredox CatalysisReceived 00th January 20xx,  
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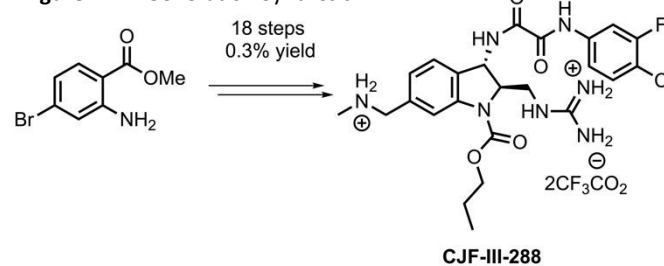
**The development and optimization of a 2<sup>nd</sup>-Generation process synthesis for the human immunodeficiency virus-1 entry inhibitor CJF-III-288 bis-trifluoroacetate salt **1** is reported. The route eliminates the use of noble metals, decreases the step count nearly by half, and increases the overall yield tenfold.**

As of 2023, 39.9 million people worldwide were living with the human immunodeficiency virus (HIV), with 1.3 million new infections that year alone.<sup>1</sup> HIV infection without treatment leads to acquired immunodeficiency syndrome (AIDS), a range of conditions that weaken the immune system thereby increasing the risk of acquiring other common and/or opportunistic infections, greatly reducing life-expectancy.<sup>2</sup> The Highly Active Antiretroviral Therapy (HAART) regimen results in almost total restoration of normal immune function and a normal lifespan, but does not eliminate viral reservoirs, necessitating lifelong adherence to treatment.<sup>3</sup> Thus, in order to end the HIV epidemic, curative treatment is needed.

Targeting the viral envelope glycoprotein (Env), the only virus-specific protein on the surface of virus particles and infected cells, comprises a selective tactic to block the viral entry process and target the viral reservoir.<sup>4</sup> CD4-mimetic compounds (CD4mcs) developed in the Smith Group bind to Env and expose gp120 epitopes recognized by otherwise “non-neutralizing” antibodies, blocking viral entry and allowing the neutralization of infectious HIV-1 virus particles. Our lead CD4mc CJF-III-288 displays promising low micromolar activity against many primary strains of HIV-1 and enhances binding of HIV+ plasma to HIV infected cells and increases HIV+ plasma mediated antibody-dependent cellular cytotoxicity (ADCC) compared to previous generations of CD4mc.<sup>4</sup> These results justified the development of a more efficient synthesis of CJF-III-288 to access larger quantities of the compound to facilitate animal studies. The established route to synthesize CJF-III-288, developed initially to enable structure-activity relationship (SAR) studies of the indoline

CD4mcs, is a laborious and time-consuming 18 steps long, with 11 flash column purifications and an overall yield of 0.3% (Fig. 1).<sup>4</sup> The use of noble and toxic metals, such as osmium, palladium, and tin, is also undesirable from a cost and safety standpoint, and triphosgene's use poses a significant safety hazard in the lab when used on scale. The length derives from protecting group and functional group manipulations, and the reliance on two-electron transformations for installing the amine functionality. The need to build up negative charge at the transmetalating carbon of the alpha-amino coupling partners may lead to poor reactivity; the same is true for the corresponding Michael addition donors.<sup>5</sup> Additionally, these reactions often require harsh nucleophiles and bases that pose issues for functional group sensitivity. Alternatively, alpha-amino radicals have seen extensive use in photoredox catalysis as their high stability and nucleophilicity makes them useful reagents in a variety of reactions.<sup>6</sup> Radical chemistry is also generally more functional group tolerant and avoids many of the substrate sensitivity issues seen in two electron chemistry.

The methylamino methyl side chain and methyl guanidine substituent were identified as areas where the implementation of photoredox chemistry would lead to significant improvements in the efficiency of the route. The 2<sup>nd</sup>-generation retrosynthesis of CJF-III-288 begins with late-stage introduction of the aryl oxalamide moiety via amidation of amine **2**, which in turn could be prepared via Curtius rearrangement from carboxylic acid **3** (Scheme 1). Installation of the methylamino methyl side chain was envisioned using Nickel/photoredox dual catalysis with aryl bromide **4** as the electrophile. Diastereoselective Giese addition into Michael acceptor **5** with judicious choice of chiral auxiliary (X) was proposed to set the

Figure 1. 1<sup>st</sup>-Generation Synthesis

- Toxic and Expensive Metals (Ru, Sn, Pd, Os)
- 11 column purifications
- Toxic triphosgene
- 2-e<sup>-</sup> transformations to install nitrogen functional groups → long step count

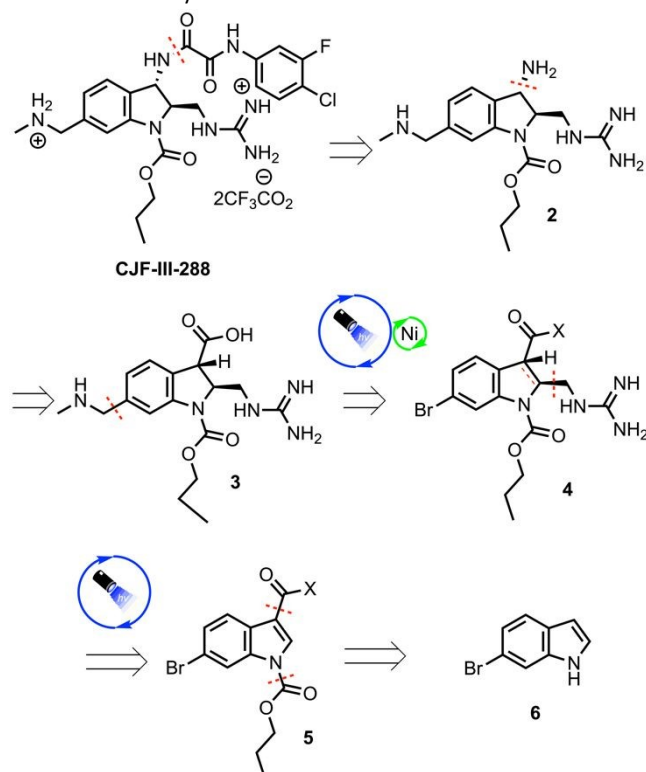
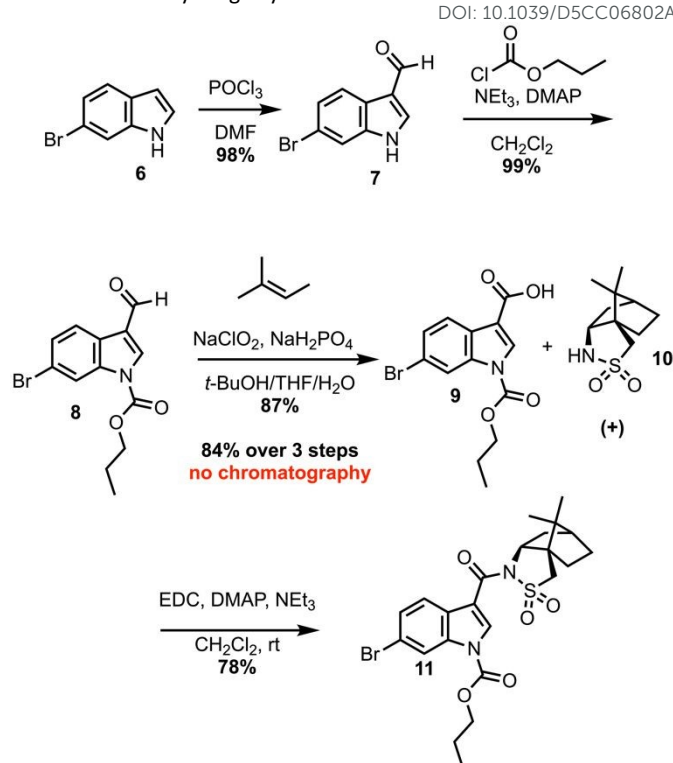
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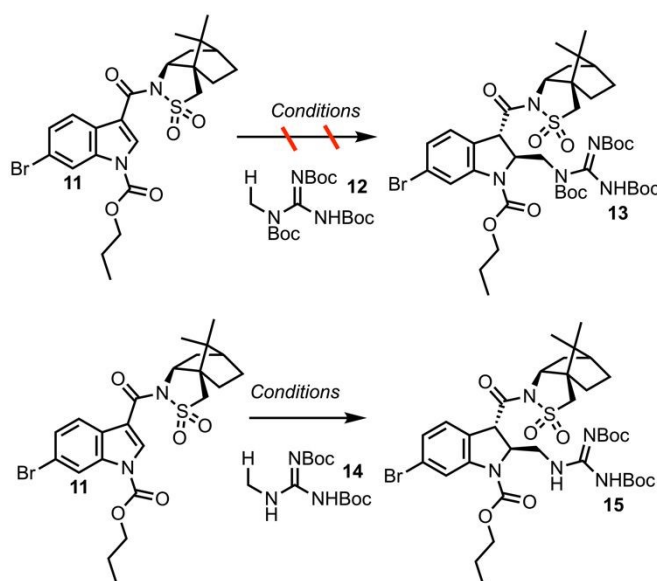
**Scheme 1.** Retrosynthesis of CJF-III-288**Scheme 2.** Early-Stage Synthesis of CJF-III-288View Article Online  
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stereogenicity of the 2 and 3 positions of the indoline core and install the guanidine motif in one step. Early stage carbamylation would obviate the need to swap out the carbamate group at a later stage, as in the 1<sup>st</sup>-Generation synthesis. Finally, an early stage Vilsmeier-Haack formylation would be implemented to install the carbonyl group at the 3 position of the indoline core, beginning from commercially available 6-bromoindole **6**.

Investigation of a 2<sup>nd</sup>-generation synthesis of CJF-III-288 begins with Vilsmeier-Haack formylation of **6** (Scheme 2), which proceeds in near quantitative yield on 105-gram scale.<sup>7</sup> An earlier attempt to convert **6** directly to the carboxylic acid was unsuccessful, as the resulting acid lacked enough solubility in common organic solvents to make it useful for further chemistry. Carbamylation of **7** with propyl chloroformate then furnishes propyl carbamate **8**, again in excellent yield on nearly an 80-gram scale. A Pinnick oxidation of aldehyde **8** next leads to carboxylic acid **9** in an excellent 87% yield on a 91-gram scale.<sup>8</sup> These first three steps proceed without silica-gel chromatography or recrystallization in an overall 84% yield. Acid **9** serves as the coupling partner with the commercially available, inexpensive (+)-camphorsultam **10** to yield **11** in 78% yield after silica-gel chromatography. The resulting amide **11** was envisioned as a Giese acceptor in a photoredox Giese addition using a guanidine-based radical donor.

The dearomatization of indoles bearing a camphorsultam moiety as a chiral auxiliary is established chemistry, with high diastereoselectivities observed in previous investigations.<sup>9</sup> Hydrogen atom transfer (HAT) catalysis was selected for its good atom economy, operational simplicity, and the ease by which the C-H radical precursors can be accessed (Scheme 3). Investigations began with globally protected precursor **12** to minimize potentially reactive guanidine nitrogen sites. HAT catalysts are chemical species that

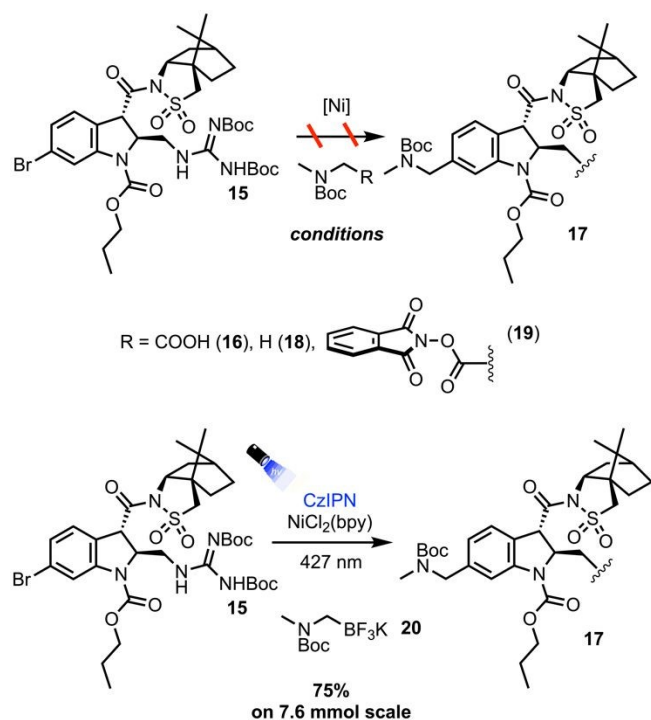
abstract homolytically labile hydrogen atoms from molecules. Under numerous protocols employing HAT catalysts and catalytic systems, such as Eosin Y, the “push-pull” benzophenone catalyst 4-Methoxy-4'-trifluoromethylbenzophenone and copper cocatalyst, and a dual catalytic system employing iridium and quinuclidine catalysts, no formation of the adduct **13** was observed.<sup>10,11,12</sup> A prohibitively large C-H bond dissociation energy (BDE) of **12**, or diminished nucleophilicity (due to either enhanced stability or steric hindrance) of the corresponding radical, was hypothesized as potential reasons for the lack of reactivity. Consequently, precursor **14**, with no *tert*-Butoxycarbonyl (Boc) group at the nitrogen alpha to the reactive

**Scheme 3.** Investigation of the Giese Addition

center, was employed for further investigations. Utilizing tetrabutylammonium decatungstate (TBADT) as the HAT catalyst, the desired Giese adduct **15** was observed by liquid chromatography – mass spectrometry (LC-MS) analysis, in low conversion. After additional optimization, the Giese adduct **15** was isolated in a good yield of 59% with 10.4:1 trans:cis diastereoselectivity, and the absolute stereochemistry was confirmed by X-ray crystallography (see Supporting Information). The reaction can be performed in up to 60.3 mmol scale in batch, and the TBADT catalyst can be synthesized in house on multi-gram scale for less than a dollar per gram. Discussion of the optimization of the Giese addition may be found in the Supplementary Information.

Numerous protocols exist for the dual photoredox/Ni catalyzed cross coupling of alpha-amino bearing coupling partners and aryl halides. The protocols first investigated were those employing the most easily synthetically accessible alpha-amino radical precursors (Scheme 4). With **15** as the aryl halide coupling partner, Dual Ir/Ni catalysis with Boc sarcosine **16** afforded only decomposition of the starting materials and none of the desired product **17**.<sup>13</sup> A HAT protocol employing 4-Methoxy-4'-trifluoromethylbenzophenone and Boc dimethylamine **18** afforded the desired cross-coupling adduct **17** as observed by LC-MS analysis, albeit in low conversion, likely due to the poor photo physics of the benzophenone triplet state sensitizers.<sup>14</sup> To remedy this, protocols and radical precursors that would lead to more efficient generation of the desired alpha amino radical were sought out. Employing redox-active ester **19** in a net-reductive cross coupling protocol with stoichiometric photoreductant Hantzsch ester affords **17** in 13% yield on 2 mmol scale.<sup>15</sup> While an appreciable amount of the desired product was obtained, the yield was low due to a largely inefficient reaction producing significant protodehalogenation side product and leaving a significant amount of unreacted starting material. The net-reductive reaction conditions and rapid fragmentation of **19** likely leads to fast generation of the corresponding alpha amino radical, but in the absence of an appreciable amount of oxidative addition

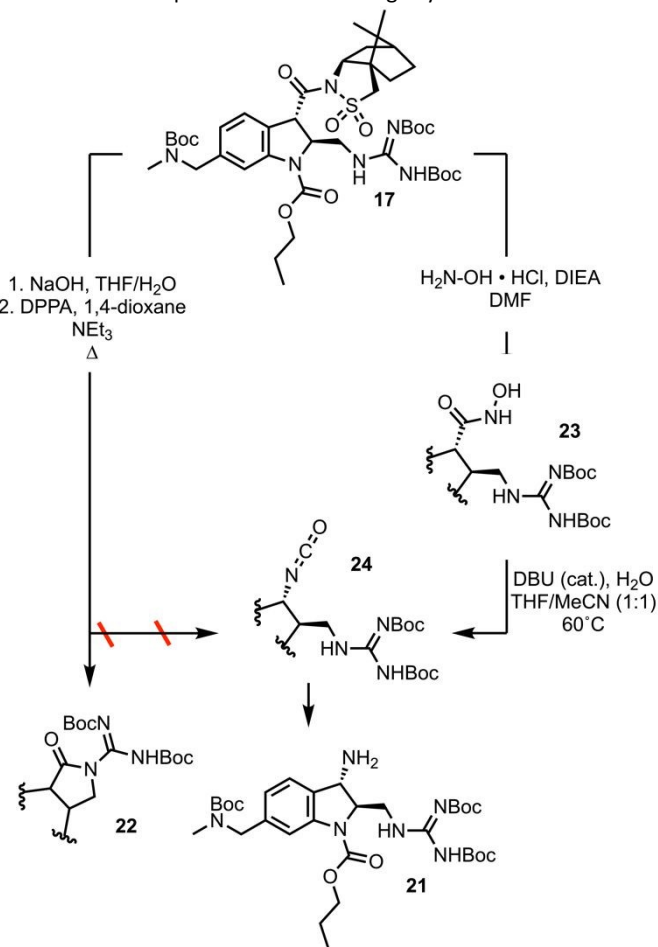
**Scheme 4.** Investigation of the Cross-Coupling



complex, non-productive processes such as radical quenching and ultimately protodehalogenation of **15** predominate. The solution was to employ a net-neutral process where the build-up of reactive intermediates could be minimized. Thus, employing CZIPN as the organic photocatalyst and (2,2'-Bipyridine)nickel dichloride ( $\text{NiCl}_2(\text{bpy})$ ), along with trifluoroborate precursor **20**, the desired adduct **17** was obtained in 75% yield after flash column chromatography on 7.6 mmol scale.<sup>16</sup> Discussion of the optimization of the cross-coupling may be found in the Supplementary Information.

The final synthetic challenge left to be overcome in the 2<sup>nd</sup>-generation synthesis of CJF-III-288 was the conversion of the carbonyl group at position 3 of the indoline to the primary amine **21** (Scheme 5). Applying Curtius conditions to **17** with treatment with diphenyl phosphorazide (DPPA) under refluxing conditions, instead of resulting in intermolecular capture of the resulting phosphate ester with azide to afford acyl azide, results in intramolecular capture by the guanidine moiety to give **22**. The solution was to employ a reaction mechanistically analogous to the Curtius rearrangement but proceeding through less electrophilic intermediates, the lesser known Lössen rearrangement.<sup>17</sup> Aminolysis of **17** with hydroxylamine furnishes hydroxamic acid **23**. Compound **23** is used in a self-propagative, "pseudo-catalytic" Lössen rearrangement, wherein condensation with acetonitrile first occurs to activate the substrate *in-situ*, which undergoes rearrangement to isocyanate **24**. Compound **24** is either hydrolyzed, or enters the self-propagative catalytic cycle with another equivalent of **23** to afford primary amine

**Scheme 5.** Development of the Late-Stage Synthesis



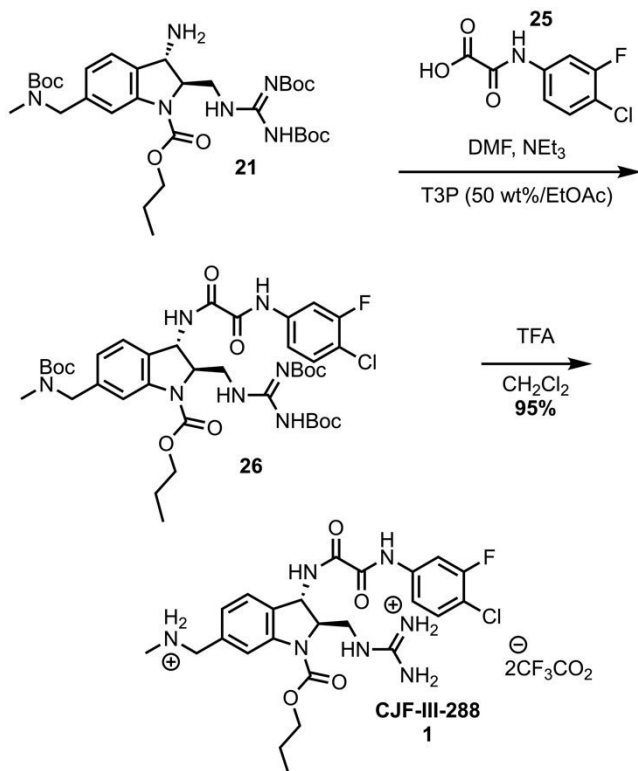


**21.** Data detailing the optimization of these two transformations is found in the Supplementary Information.

The synthesis was completed by coupling of the *meta*-fluoro-*para*-chloro acid **25** to amine **21** using propanephosphonic acid anhydride (T3P), effective for sterically encumbered substrates on large scale (Scheme 6).<sup>18,19</sup> Global Boc deprotection of **26** with TFA yields **CJF-III-288** in nearly quantitative yield. The synthesis was further validated by scaling up the cross coupling of **15** and **20**, which can be performed on a 33.16 mmol scale in a single batch. The sequence from intermediate **15** to **26** was performed on a 79 mmol scale, affording 5.66 grams of **26** after a single flash column purification. The global Boc deprotection yielded 4.90 grams of **1**.

In summary, the 2<sup>nd</sup>-generation synthesis of CJF-III-288, beginning on over a hundred-gram scale, proceeds in 10 steps to yield **1** in an overall yield of 3%. This reduces the step count almost by half compared to the original of 18 steps, and the overall yield has increased tenfold compared to the original yield of 0.3%. This was achieved by reducing the step count of the installation of the guanidine group and stereochemistry from 5 steps to 1 step, and of the methylamino methyl group from 3 steps to 1 step, compared to the first-generation synthesis. The route has been used to synthesize 5.7 grams of **1**, which are currently being used in *in-vivo* biological studies to determine the efficacy of CJF-III-288 to prevent and cure HIV infection in monkey models. Additionally, the photochemistry developed for the synthesis holds promise for use in further analogue synthesis. The modular nature of both reactions allows incorporation of diverse alkyl, aryl, methylamino, and methyloxy functionality at both the 2 and 6 positions of the indoline core and further optimization of the potency and physicochemical properties of the indole CD4mcs.

**Scheme 6.** Endgame Synthesis



## Conflicts of interest

There are no conflicts to declare.

## Data availability

The data underlying this study, including experimental procedures, characterization data, <sup>1</sup>H-NMR/<sup>13</sup>C-NMR spectra, and crystal structures are available in the published article and in its Supporting Information. Raw FID data is openly available in Dryad Databank at DOI: 10.5061/dryad.4b8gttsf. Crystallographic data for compound **15** has been deposited at the Cambridge Crystallographic Data Center under deposition number 2484437.

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