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# Methods for kinetic evaluation of reversible covalent inhibitors from time-dependent IC<sub>50</sub> data†

Lavleen K. Mader and Jeffrey W. Keillor \*

Potent reversible covalent inhibitors are often slow in establishing their covalent modification equilibrium, resulting in time-dependent inhibition. While these inhibitors are commonly assessed using IC<sub>50</sub> values, there are no methods available to analyze their time-dependent IC<sub>50</sub> data to provide their inhibition ( $K_i$  and  $K_i^*$ ) and covalent modification rate ( $k_5$  and  $k_6$ ) constants, leading to difficulty in accurately ranking drug candidates. Herein, we present an implicit equation that can estimate these constants from incubation time-dependent IC<sub>50</sub> values and a numerical modelling method, EPIC-CoRe, that can fit these kinetic parameters from pre-incubation time-dependent IC<sub>50</sub> data. The application of these new methods is demonstrated by the evaluation of a known inhibitor, saxagliptin, providing results consistent with those obtained by other known methods. This work introduces two new practical methods of evaluation for time-dependent reversible covalent inhibitors, allowing for rigorous characterization to enable the fine-tuning of their binding and reactivity.

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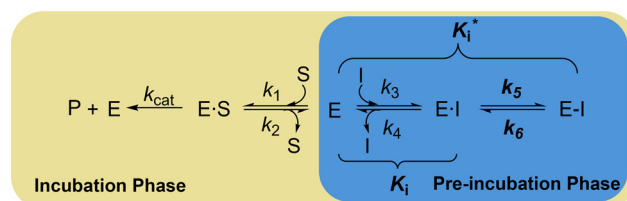
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## Introduction

Reversible covalent inhibitors are a subset of targeted covalent inhibitors (TCIs). These compounds follow a two-step inhibition mechanism (Scheme 1, blue) featuring an initial non-covalent binding event, whose equilibrium is described by the inhibition constant  $K_i$ , followed by a reversible covalent reaction between the protein target and the inhibitor's electrophilic 'warhead'.

The entire equilibrium from free enzyme E to covalently bound enzyme E-I is described by the inhibition constant  $K_i^*$ , which can be calculated from  $K_i$ ,  $k_5$ , and  $k_6$ .<sup>2-5</sup> Many warheads are known to display such reversible reactivity, including  $\alpha$ -cyanoacrylamides,  $\alpha$ -ketoamides, nitriles, and boronic acids.<sup>4,6-12</sup> While our understanding and development of reversible TCIs is still relatively new, some inhibitors of this class have shown clinical potential, and others are already on the market.<sup>13-15</sup> Reversible covalent inhibitors have been proposed to offer lower toxicity profiles compared to irreversible inhibitors, due to reversible off-target protein modification, while still providing the increased residence time and affinity afforded by a covalent interaction.<sup>6,16</sup> Interestingly, all FDA-approved drugs of this

type also display time-dependent inhibition,<sup>13,17-21</sup> where the final equilibrium between free enzyme and covalently bound enzyme is slow to establish. This time-dependent behaviour is almost always due to the breakdown of the covalent inhibitor-enzyme complex being very slow compared to covalent bond formation ( $k_6 \ll k_5$ ), often due to the intrinsic stability of the covalent bond, or steric hindrance introduced by either the scaffold itself or the surrounding protein microenvironment.<sup>22</sup> This behaviour serves to enhance on-target residence times and apparent affinity even further, and can also be tuned based on the intrinsic reactivity of the warhead or the steric environment around it (*i.e.* increase or decrease  $k_5$  and/or  $k_6$ ), as first demonstrated by Taunton *et al.*<sup>8,22</sup> Therefore, complete characterization of all relevant binding and rate constants is crucial to optimizing overall on- and off-rates for both on-target and off-target proteins, in order to ensure potency while minimizing off-target side effects. Recently, this strategy has been particularly relevant



**Scheme 1** Kinetic scheme of reversible covalent inhibition, where  $K_i = k_4/k_3$  and  $K_i^* = K_i/1 + (k_5/k_6)$ . This figure was adapted from ref. 1.

Department of Chemistry and Biomolecular Sciences, University of Ottawa, Ottawa, Ontario K1N 6N5, Canada. E-mail: jkeillor@uottawa.ca

† Electronic supplementary information (ESI) available: Derivations of all kinetic equations, table of differential equations, code for numerical modelling and kinetic data for Gly-Pro-pNA  $K_M$  and  $k_{cat}$  determination (pdf). See DOI: <https://doi.org/10.1039/d5md00050e>



in achieving selectivity among kinases, which have very similar binding sites.<sup>23,24</sup>

While the kinetic behaviour of fast-binding and fast-reacting reversible covalent inhibitors is difficult to distinguish from that of traditional non-covalent reversible inhibitors (*i.e.*, instantaneous and uniform decrease in activity, showing no time-dependence), slow-binding and/or slow-reacting covalent inhibitors display time-dependent kinetics, which allow their individual rate and inhibition constants to be measured. This time-dependence can be seen clearly in continuous assays, where product formation over time resembles a 'curve' rather than a straight line.<sup>2,3</sup> These 'curves' feature an initial linear phase representative of non-covalent binding (described by  $K_i$ ) which slowly transitions to a final linear phase once the final covalent equilibrium has also been established (described by  $K_i^*$ ). Progress curves from continuous assays can be used to evaluate time-dependent reversible inhibitors and derive all of their binding and rate constants. However, the fitting of data from this method is error prone, especially for the typically small  $k_6$  values.<sup>3</sup> Additionally, if a continuous assay is not available, attempting to mimic continuous monitoring by taking many time-based aliquots of an enzymatic reaction can be very labour- and time-intensive to perform. It has also recently been shown that these kinetic parameters can be estimated in a substrate-free manner by strategically monitoring binding kinetics at assumed pre- and post-equilibrium periods using surface plasmon-resonance, followed by complex modelling using fixed  $k_3$  and  $k_4$  (Scheme 1) values.<sup>24</sup> This is a clever approach, but this type of analysis requires not only specialized instrumentation but also advanced knowledge of mathematics and modelling.

In practice, the time-dependence of reversible covalent inhibitors is often probed by time-dependent  $IC_{50}$  assays that are performed by varying either the duration of the 'pre-incubation' phase, when the enzyme is incubated with inhibitor alone, or the 'incubation phase', when substrate is added (Scheme 1, yellow).<sup>25,26</sup> A shift to lower  $IC_{50}$  values with increasing assay times confirms the time-dependence of covalent addition, while the reversibility must be established otherwise.<sup>27,28</sup> However, no fitting methods are available to analyze this kind of time-dependent  $IC_{50}$  data and derive all relevant inhibition and rate constants, perhaps due to the mathematical difficulty the additional rate constant ( $k_6$ ) imposes. Instead, many researchers simply report a single  $IC_{50}$  value at a single time point. While this single value may be useful for screening compounds within the same library using the exact same assay, it is difficult to discern whether such an  $IC_{50}$  value is reporting on the initial linear phase (*i.e.*, pre-equilibrium), the transitional period, or the final linear phase (*i.e.*, equilibrium) of the reaction. Hence, information about the forward and reversible covalent reaction steps (related to  $k_5$  and  $k_6$ ) and the overall affinity (related to  $K_i^*$ ) may be missed, causing researchers to overlook promising candidates.<sup>2</sup>

Herein, we report, for the first time, two new methods to analyze time-dependent  $IC_{50}$  data from reversible covalent inhibitors in order to derive the inhibition constants  $K_i$  and  $K_i^*$ , and the covalent reaction rate constants  $k_5$  and  $k_6$ . The first method is an implicit equation that relates incubation time-dependent  $IC_{50}$  values (obtained without pre-incubation) to these constants. The second method is an empirical global fitting method that models pre-incubation time-dependent  $IC_{50}$  data to estimate the same constants. To demonstrate the performance of our methods and consensus with established evaluation procedures, we also present the characterization of a known, clinically relevant, time-dependent reversible covalent di-peptidyl peptidase IV (DPPIV) inhibitor, saxagliptin.<sup>14</sup> All relevant equations, derivations thereof, differential equations and embodying code relating to these methods of evaluation have been provided in the ESI,<sup>†</sup> with key fitting equations included in the Experimental section.

## Results and discussion

While the recent general resurgence in interest in covalent inhibitors has led to the development of many clever methods to evaluate irreversible binders,<sup>1,17,29,30</sup> introduction of a reversible reaction step (*i.e.*, corresponding to rate constant  $k_6$  in Scheme 1) results in much more nuance to the mathematical models that describe the kinetics of reversible covalent modulators. For a fast-binding and fast-reacting reversible covalent inhibitor there are no practical means for kinetically distinguishing it from a fast-binding (traditional) reversible non-covalent inhibitor, using activity assay-based methods. This means only its overall affinity, related to the inhibition constant  $K_i^*$ , can be measured, giving no information about the reactivity of the compound with its protein target. However, reversible covalent inhibitors that react more slowly with their targets display time-dependent kinetics, which allows their initial non-covalent binding and subsequent covalent reaction to be dissected to provide individual inhibition and rate constants.<sup>2</sup> The basic mathematical principles that form the foundation of all previous methods, and the new methods described herein for evaluation of these inhibitors, are identical to those previously derived for slow-binding reversible non-covalent inhibitors that display an 'induced fit' mechanism, as summarized by Copeland.<sup>2</sup> The only difference between the induced fit slow-binding mechanism and that of slow-reacting reversible covalent modification is that a covalent bond is formed (and broken) in the reversible step described by rate constants  $k_5$  (and  $k_6$ ) (Scheme 1). Recognition of this similarity allows for the application of eqn (S12) and (S13) in continuous, progress curve analysis for these inhibitors, as described in the ESI:<sup>†</sup>

$$k_{\text{obs}} = k_6 + \left( \frac{k_5 [I]}{[I] + K_i^{\text{app}}} \right) \quad (\text{S12})$$



$$[P]_{I(t)} = v_s t + \frac{(v_i - v_s)}{k_{obs}} (1 - e^{-k_{obs}t}) \quad (S13)$$

However, despite the growing attention given to this class of inhibitors, there is much confusion in the literature about the application and interpretation of this fitting method.<sup>22,24</sup> Continuous assays for these types of inhibitors typically require optimization for long assay times in order to observe and capture the steady state equilibrium in the fitting. Moreover, the fitting procedures for the resulting data cannot account for substrate depletion and/or enzyme degradation (*i.e.*, the uninhibited control must be perfectly linear, since no mathematical correction can be applied in the way that it can be for an irreversible inhibitor) and often lead to high errors in the fitted parameters.<sup>3</sup>

$$K_i^{app} \left( \frac{k_6}{k_6 + k_5} \right) + \left( \frac{\left( \frac{2k_6}{k_6 + k_5} \right) (K_i^{app})^2 + 2K_i^{app} IC_{50}(t)}{K_i^{app} + IC_{50}(t)} - 2K_i^{app} \left( \frac{k_6}{k_6 + k_5} \right) \right) \cdot \frac{(1 - e^{-k_{obs}t})}{k_{obs} \cdot t} = IC_{50}(t) \quad (S18)$$

Perhaps for these reasons, these inhibitors are more often evaluated using  $IC_{50}$  experiments, where time-dependency is simply demonstrated by a decrease in  $IC_{50}$  values with increasing times of either assay incubation or enzyme-inhibitor pre-incubation.<sup>13,22</sup> Prior to this work, this type of data could only be used to confirm time-dependent behaviour, before selecting a given time-point and ranking inhibitors based on their  $IC_{50}$  at that time, with no knowledge of whether the final equilibrium has been reached. This poses the obvious problem of grossly misleading structure-activity relationship data, as well as limited information about how to tune binding and reactivity, preventing researchers from harnessing the true potential of such inhibitors. Not too long ago, irreversible inhibitors were also plagued by this very issue, which has since been resolved by equations and modelling methods for irreversible binding.<sup>1,29</sup> Recognizing this disconnect and drawing on our own expertise in the methodologies for evaluating irreversible covalent inhibitors,<sup>1,31-33</sup> we have developed two new methods that allow for time-dependent  $IC_{50}$  data to be analyzed to provide values for  $K_i$ ,  $k_5$ , and  $k_6$ , and subsequently calculate  $K_i^*$ , for time-dependent reversible covalent inhibitors. For the sake of clarity, in the main body of this article we will only describe the elements of method development that relate to implementation, assay design, and drug properties, in the simplest terms possible. The detailed derivation of equations is provided in the ESI†

### Development of implicit equation for time-dependent $IC_{50}$ values

First, we took an approach similar to that of Krippendorff *et al.*<sup>28,29</sup> in recognizing the central relationship between continuous product formation (as described by eqn (S13)) and  $IC_{50}$  values measured at different incubation times, in an assay

where enzyme, inhibitor, and substrate are all combined at the same time. We will refer to this as an incubation time-dependent  $IC_{50}$  experiment (Scheme 1, yellow). Functionally, an  $IC_{50}$  value at any given time is simply the inhibitor concentration that results in 50% of the uninhibited response, where the uninhibited response (*i.e.* product formation) is described by Michaelis-Menten kinetics ( $[P] = v_0 \cdot t$ , where  $v_0$  is given by the Michaelis-Menten equation). At any given assay time, half of this response ( $\frac{1}{2}v_0 \cdot t = [P]_{IC_{50}(t)}$ ), is observed in an inhibited reaction when  $[I] = IC_{50}(t)$  (see ESI†, eqn (S16)). Understanding the derivation of eqn (S16),† and the definitions of the parameters within, allows for algebraic manipulation (shown in the ESI†) to arrive at an implicit equation (eqn (S18)) that relates a discontinuous  $IC_{50}$  value determined at time  $t$  to parameters  $K_i^{app}$ ,  $k_5$ , and  $k_6$ :

where

$$k_{obs} = k_6 + \left( \frac{k_5 IC_{50}(t)}{IC_{50}(t) + K_i^{app}} \right) \quad (S19)$$

Eqn (S18) is an implicit equation in that the dependent parameter (*i.e.*,  $IC_{50}(t)$ ) appears on both sides of the equation. However, if many discontinuous  $IC_{50}$  values are measured at different times (Fig. 1A), a curve can be generated by plotting  $IC_{50}(t)$  vs. time (Fig. 1B) which can be fitted to eqn (S18) by non-linear regression, allowing for  $K_i^{app}$ ,  $k_6$ , and  $k_5$  to be solved. Note that  $K_i$  is calculated from  $K_i^{app}$  using eqn (S4).†

It is also instructive to consider the limits of eqn (S18), at vanishingly small or infinite incubation times. As shown in the ESI†, the limit of  $IC_{50}(t)$  as time approaches zero is equal to  $K_i^{app}$  (see eqn (S22)†). Intuitively, this makes sense as it describes the initial rapid-binding equilibrium, before covalent modification has taken place. Alternatively, at infinite incubation time, the limit of  $IC_{50}(t)$  is equal to  $K_i^{*app}$  (see eqn (S24)†). This reflects the final inhibition equilibrium that is definitively established after infinite incubation time.

Eqn (S18) can also be compared to the ‘Krippendorff equation’, which was derived for two-step irreversible inhibition (see eqn (S25)†).<sup>1,29</sup> Interestingly, if  $k_6$  is set to zero, which would simplify the kinetic scheme shown in Scheme 1 to that of irreversible inhibition, eqn (S18) does indeed simplify to the Krippendorff equation (see eqn (S27)†). Likewise, setting  $k_6$  to zero in eqn (S19) results in a simplified equation (eqn (S28)†) that is equivalent to what is observed for irreversible inhibition. These comparisons serve to validate eqn (S18) and (S19), but they also demonstrate that the Krippendorff equation can be thought of as a simplified version of eqn (S18), for the special case where  $k_6$  is equal to zero. Thus, the equation derived here (*i.e.* eqn (S18)) is all-encompassing for all types of covalent inhibitors that show time-dependence.



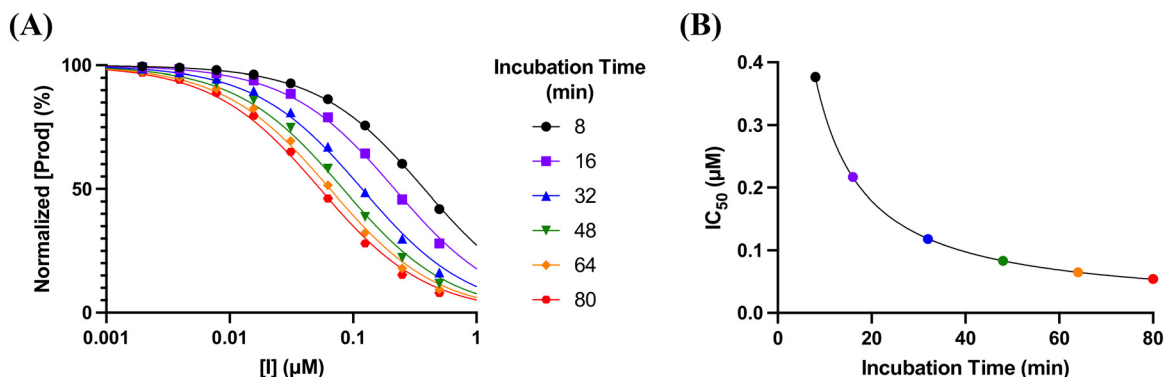


Fig. 1 Fitting of incubation time-dependent  $IC_{50}$  values to implicit eqn (S18). A) Simulated  $IC_{50}$  curves for the reaction of 1 nM enzyme ( $k_{cat} = 1000 \text{ min}^{-1}$ ) with 500  $\mu\text{M}$  substrate ( $K_M = 100 \text{ }\mu\text{M}$ ) in the presence of varying concentrations (1–500 nM) of a reversible covalent inhibitor ( $k_5 = 0.5 \text{ min}^{-1}$ ,  $k_6 = 0.005 \text{ min}^{-1}$ ,  $K_i = 0.10 \text{ }\mu\text{M}$ ), measured after incubation times ranging from 8–80 min. B) Fitting of  $IC_{50}$  values resulting from (A) to implicit eqn (S18).

Despite the apparent complexity of eqn (S18) at first glance, it allows for fitting through non-linear regression, using any standard graphing software (*e.g.*, GraphPad Prism). In the ESI† we provide a plain-text version that can be copied and pasted as a user-defined equation in any desired software and used to fit experimental data.

### Development of EPIC-CoRe

In the equally, if not more common case where a pre-incubation time-dependent  $IC_{50}$  experiment is performed, by first incubating the enzyme with inhibitor for varying times (Scheme 1, blue), before adding substrate and measuring product formation after a defined incubation period (Scheme 1, yellow), no implicit equation can be derived to describe the variation of  $IC_{50}$  with time. This is due to the biphasic nature of such an experiment, where enzyme is differentially inhibited, in the first phase rapidly by

inhibitor alone, and in the second phase more slowly due to partial protection by a competitive substrate (Fig. 2A). The  $IC_{50}$  is then measured, based on product formation at a fixed end-point time in the second (assay incubation) phase (Fig. 2B).

Similar to an incubation time-dependent  $IC_{50}$  experiment,  $IC_{50}$  values also decrease with increasing pre-incubation times (Fig. 2B); however, in this case, it is impossible to embody both phases in any one equation to fit kinetic parameters. We recently noted the absence of tools available to characterize irreversible inhibitors in this way as well – that is, to derive  $K_I$  and  $k_{inact}$  values from pre-incubation time-dependent  $IC_{50}$  data. This led us to develop a numerical modelling method, EPIC-Fit, to accomplish this important task.<sup>1</sup> Here, we describe a similar empirical approach, developed for time-dependent reversible covalent inhibitors, that we call EPIC-CoRe (Endpoint Pre-incubation  $IC_{50}$ -Covalent Reversible).

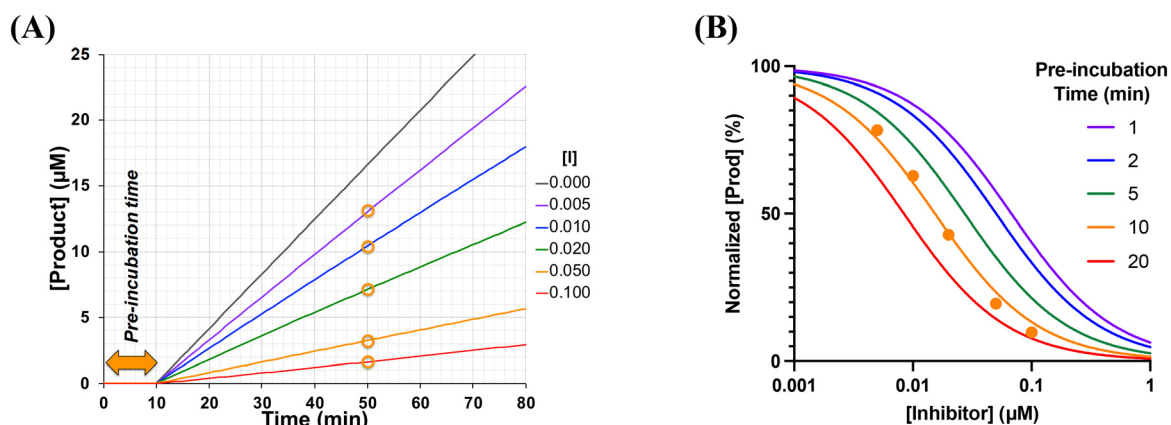


Fig. 2 Origin of pre-incubation time-dependent  $IC_{50}$  curves. A) Simulated progress curves for the reaction of 1 nM enzyme ( $k_{cat} = 1000 \text{ min}^{-1}$ ) with varying concentrations (5–100 nM) of a reversible covalent inhibitor ( $k_5 = 0.5 \text{ min}^{-1}$ ,  $k_6 = 0.005 \text{ min}^{-1}$ ,  $K_i = 0.10 \text{ }\mu\text{M}$ ) for a 10-min pre-incubation phase, prior to addition of 500  $\mu\text{M}$  substrate ( $K_M = 100 \text{ }\mu\text{M}$ ) and measurement of end-point product concentration after a 40-min incubation phase (orange circles). B) End-point product concentrations are then normalized against those of the uninhibited reaction and plotted against inhibitor concentration to generate  $IC_{50}$  curves, at various pre-incubation times (*e.g.* 1–20 min). The orange data points in (B) are those shown in (A), at a pre-incubation time of 10 min. Note how the measured  $IC_{50}$  values shift to lower values at longer pre-incubation times.



This method employs code that uses differential equations to describe instantaneous changes in enzyme, inhibitor, substrate, and product concentrations, from the beginning of the pre-incubation phase to the end of the assay incubation phase, ultimately predicting the end-point signal (e.g. product concentration) that would be observed for a given set of inhibition parameters ( $K_i$ ,  $k_5$ , and  $k_6$ ) in a pre-incubation  $IC_{50}$  experiment. This calculation is performed for every inhibitor concentration, at a given pre-incubation and assay measurement time, for a competitive inhibitor/substrate pair, to generate a predicted  $IC_{50}$  at that time. Thus, experimental parameters of the activity assay must also be known, including initial enzyme and substrate concentrations, the  $k_{cat}$  and  $K_M$  values, and the dilution of enzyme and inhibitor resulting from addition of substrate to initiate the incubation phase. When experimental  $IC_{50}$  datasets (comprising product signal values measured at different inhibitor concentrations) are entered at many different pre-incubation times, and initial estimates for  $k_5$ ,  $k_6$  and  $K_i$  are provided (see below for practical considerations), the difference between each observed endpoint value recorded in the global dataset and the endpoint concentration predicted according to initial estimates are reported as “residuals”. Least squares regression can then be performed to arrive at  $k_5$ ,  $k_6$  and  $K_i$  values that provide a global minimum to the sum of the squares of these residuals (i.e., where the predicted signal values match the experimental signal values as closely as possible) for all data points.

This method can be implemented using any spreadsheet or modelling software that allows user code to be inputted. We have chosen to use Microsoft Excel, as it offers readily accessible spreadsheet calculations, user-defined code input (in visual basic for applications (VBA)) for rapid iterative calculations, and least-squares fitting using the Solver add-in, to create a functional embodiment of this method. Our spreadsheet contains cells to input values of the experimental parameters of the assay, as well as predicted and experimental  $IC_{50}$  plots from all the pre-incubation time-dependent datasets that are updated in real-time as data is entered. It also presents a column of predicted endpoint values, the calculated residual of each data point, and the residual sum of squares (RSS). The VBA code used for the numerical modelling calculations is also presented in the Supporting Information, showing how differential equations are used iteratively to arrive at the predicted end-point concentrations. The Solver add-in included in Excel has been set up such that the cells containing the values of  $k_5$ ,  $k_6$  and  $K_i$  contain initial estimates and are subsequently optimized by least square regression, using Generalized Reduced Gradient (GRG2) code.<sup>34</sup> We have also included a cell that calculates  $K_i^*$ , based on fitted parameters. A fully functional EPIC-CoRe spreadsheet that includes all aforementioned features, as well as instructions and formatting to guide user data input, is freely available upon request.

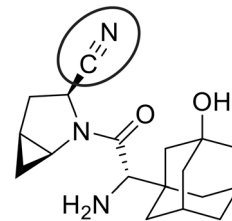


Fig. 3 Structure of reversible covalent DPPPIV inhibitor saxagliptin, featuring a nitrile warhead.

### Evaluation of saxagliptin

To demonstrate the application of the new methodologies reported herein, we evaluated the well-known DPPPIV inhibitor, saxagliptin, which bears a reversibly reactive nitrile warhead (Fig. 3).<sup>14</sup> We chose to evaluate saxagliptin because it is one of the very few inhibitors that has been previously shown to exhibit time-dependent, reversible covalent inhibition and for which a chromogenic continuous activity assay has already been developed, in the form of a commercially available kit. This makes saxagliptin nearly uniquely amenable to independent evaluation according to all of the methods discussed above, allowing us to validate our two new methods against the existing one. The fitted kinetic parameters determined by each method are summarized in Table 1. The enzymatic assay follows the chromogenic DPPPIV-mediated hydrolysis of substrate glycine-proline-*para*-nitroanilide (Gly-Pro-*p*NA), which releases the coloured product *para*-nitroaniline (Scheme 2).<sup>14,35,36</sup> We used this assay in both continuous (real-time monitoring) and discontinuous (end-point) mode, depending on the method of evaluation, for demonstration purposes. Since each of the methods described here employs a substrate that is competitive with inhibitor, a  $K_M$  value is required for each method, and a  $k_{cat}$  value is required for analysis by EPIC-CoRe, as discussed above. Using standard Michaelis–Menten kinetic analysis (Fig. S3†) we obtained values of  $K_M = 70.7 \pm 6.0 \mu\text{M}$  and  $k_{cat} = 17.6 \pm 0.5 \text{ s}^{-1}$  (or roughly  $1056 \text{ min}^{-1}$ ), which are consistent with literature values.<sup>14</sup> These values were used for subsequent inhibitor characterization.

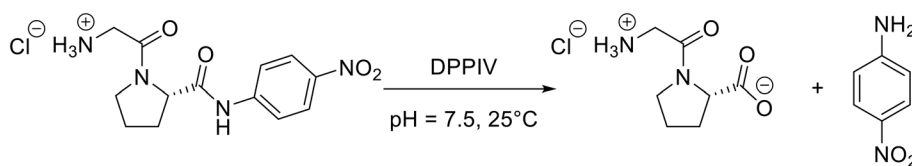
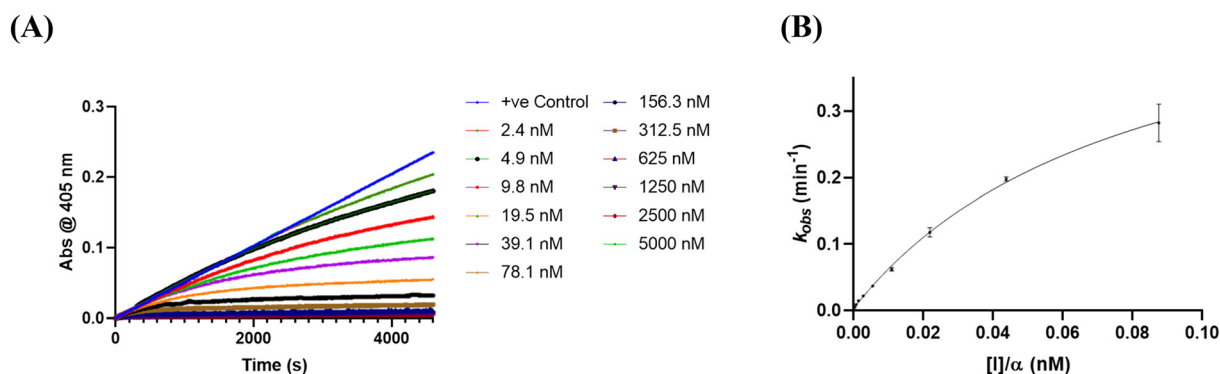
Firstly, we evaluated saxagliptin in a continuous assay fashion (Scheme 1, yellow) to obtain progress curves at different inhibitor concentrations. This type of reaction, initiated in the presence of both inhibitor and substrate, can be described by the explicit equation for time-dependent product formation, eqn (S13). Incubation of 12 different concentrations of saxagliptin with DPPPIV in the presence of Gly-Pro-*p*NA, in an experiment where enzyme is added to initiate the reaction, resulted in biphasic ‘curves’ of absorbance vs. time, representing time-dependent inhibition (Fig. 4A). These curves were monitored over 80 minutes to allow sufficient time for steady state slopes to be reached, and were fitted to eqn (S13) to give  $k_{obs}$  values for the transition phase at each concentration.<sup>14</sup> These rate



**Table 1** Summary of kinetic parameters for the inhibition of DPPiV by saxagliptin obtained from different methods of evaluation

Assay	Fitting method	$k_5$ (min <sup>-1</sup> )	$k_6$ (min <sup>-1</sup> )	$K_i$ (nM)	$K_i^{*d}$ (nM)
Continuous assay	Saturation <sup>a</sup>	$0.54 \pm 0.03$	$0.0033 \pm 0.0031$	$81 \pm 7$	$0.49 \pm 0.41$
	IC <sub>50</sub> from initial slopes	—	—	$89 \pm 6$	—
	IC <sub>50</sub> from final slopes	—	—	—	$0.74 \pm 0.18$
Incubation time-dependent IC <sub>50</sub>	Implicit equation <sup>b</sup>	$0.85 \pm 0.15$	$0.0042 \pm 0.0015$	$120 \pm 15$	$0.52 \pm 0.20$
	EPIC-CoRe <sup>c</sup>	$0.63 \pm 0.14$	$0.0028 \pm 0.0009$	$83 \pm 17$	$0.35 \pm 0.12$
Pre-incubation time-dependent IC <sub>50</sub>	EPIC-CoRe	$0.64 \pm 0.05$	$0.0045 \pm 0.0015$	$141 \pm 8$	$0.99 \pm 0.35$

<sup>a</sup> Errors in parameters shown for this method are from the fitting of saturation kinetic data ( $k_{\text{obs}}$  vs.  $[I]$ ). <sup>b</sup> Errors in parameters shown for this method are from the fitting of IC<sub>50</sub> values vs. incubation time, according to implicit equation eqn (S18). <sup>c</sup> Errors in parameters shown for this method are from the standard deviation of duplicate values determined from the global fitting of pre-incubation time-dependent IC<sub>50</sub> data. <sup>d</sup>  $K_i^*$  was calculated using eqn (S11).

**Scheme 2** Hydrolysis of Gly-Pro-pNA by DPPiV to form *para*-nitroaniline, which absorbs at 405 nm.

**Fig. 4** Continuous activity assay with saxagliptin, DPPiV, and Gly-Pro-pNA. (A)  $k_{\text{obs}}$  values were fitted from reaction progress curves of absorbance vs. time at different concentrations of saxagliptin using eqn (S12). (B) Saturation fitting of these  $k_{\text{obs}}$  values according to eqn (4) provided values for  $k_5$ ,  $k_6$ , and  $K_i$ .

constants were then plotted against inhibitor concentrations (corrected for competition with substrate), and then fitted to hyperbolic eqn (S12), representative of saturation kinetics, taking care not to constrain the  $y$ -intercept to zero (Fig. 4B). This analysis provided values of  $k_5 = 0.54 \pm 0.03$  min<sup>-1</sup>,  $k_6 = 0.0033 \pm 0.0031$  min<sup>-1</sup>,  $K_i = 81 \pm 9$  nM, and  $K_i^* = 0.49 \pm 0.41$  nM. Note that the rather large relative error in  $k_6$  and consequently in  $K_i^*$  are typical for this method of analysis.<sup>3</sup>

The fitting of these reaction progress curves to eqn (S12) also provides initial and steady state slopes of inhibition,  $v_i$  and  $v_s$ , respectively. As previously shown,<sup>3</sup> these slopes can be normalized as a percentage of the uninhibited slope and plotted against the inhibitor concentration to provide an IC<sub>50</sub> plot. The IC<sub>50</sub> values obtained from analysis of  $v_i$  and  $v_s$  are equivalent to  $K_i$  and  $K_i^*$ , respectively, when competition from substrate is accounted for.<sup>2,3</sup> However, this analysis cannot

provide individual rate constants  $k_5$  and/or  $k_6$ . For saxagliptin, these IC<sub>50</sub> plots provided values of  $K_i = 89 \pm 6$  nM (Fig. 5A) and  $K_i^* = 0.74 \pm 0.18$  nM (Fig. 5B). Interestingly, although this type of IC<sub>50</sub> analysis cannot provide individual reaction rate constants, it is the recommended method of analysis of progress curves due the lower error it typically provides in  $K_i^*$  values.<sup>3</sup> Moreover, it is still possible to analyze the ratio of the rate constants as  $k_5/k_6 = (K_i/K_i^*) - 1$  ( $\approx 120$  in the case of saxagliptin analyzed in this way), which may provide useful insight and distinction into how much the forward reaction is favoured compared to the reverse reaction, within a library of inhibitors.

Next, we carried out an incubation time-dependent IC<sub>50</sub> experiment, which is performed essentially in the same way as the continuous assay method, except that one absorbance value (related to product concentration) is measured at a final end-point at a pre-determined time for each inhibitor



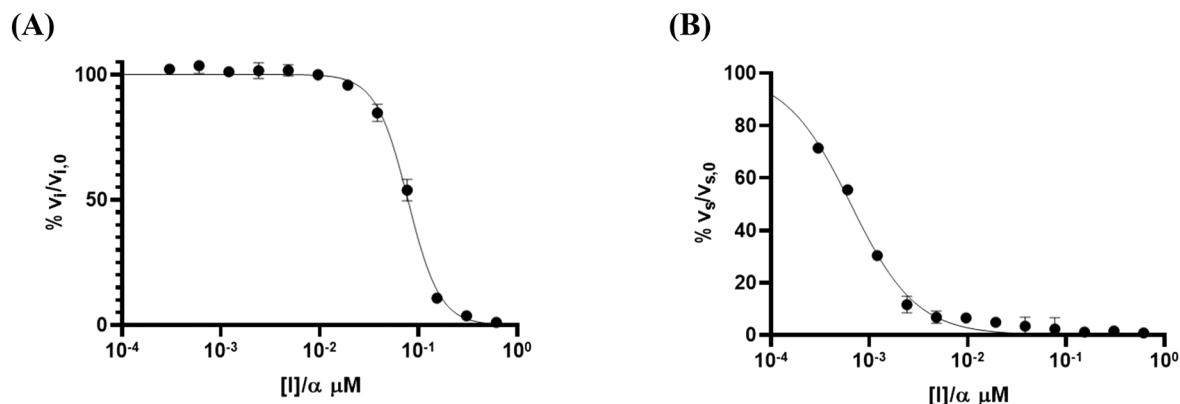


Fig. 5  $IC_{50}$  analysis of initial and final slopes obtained from progress curve fitting. (A) Plotting normalized initial slopes against corrected inhibitor concentrations gives an  $IC_{50} = K_i$ . (B) Plotting normalized final slopes against corrected inhibitor concentrations gives an  $IC_{50} = K_i^*$ .

concentration, from which an  $IC_{50}$  plot is generated. This is then repeated at different assay incubation times to generate many time-dependent  $IC_{50}$  curves (Fig. 6A).  $IC_{50}$  values decrease with increasing incubation time (*i.e.*, apparent potency increases) as the inhibitor is allowed more time to establish the final equilibrium with the covalently bound enzyme. These  $IC_{50}$  values change dramatically at short incubation times, but very little at long incubation times, as the steady state-equilibrium is approached.  $IC_{50}$  values for saxagliptin were determined at 10 different time points up to 80 minutes, then were plotted against assay incubation time and fitted to implicit eqn (S18) (Fig. 6B) to obtain the kinetic parameter values  $k_5 = 0.85 \pm 0.15 \text{ min}^{-1}$ ,  $k_6 = 0.0042 \pm 0.0015 \text{ min}^{-1}$ ,  $K_i = 120 \pm 15 \text{ nM}$ , and  $K_i^* = 0.52 \pm 0.20 \text{ nM}$ , showing excellent agreement with the values determined by progress curve analysis (above).

Interestingly, the  $IC_{50}$  curves obtained from this incubation time-dependent  $IC_{50}$  experiment can also be analyzed using our EPIC-CoRe spreadsheet (Fig. S4<sup>†</sup>), by setting the dilution factor to 1 and a pre-incubation time to zero. Global fitting of the same data by this method provided values of  $k_5 = 0.63 \pm 0.14 \text{ min}^{-1}$ ,  $k_6 = 0.0028 \pm$

$0.0009 \text{ min}^{-1}$ ,  $K_i = 83 \pm 17 \text{ nM}$ , and  $K_i^* = 0.35 \pm 0.12 \text{ nM}$ , with an average correlation of  $R^2 = 0.99$ , and RMSE = 1.2, very similar to those obtained by fitting to implicit eqn (S18). This suggests that either fitting procedure would be suitable for analysis of incubation time-dependent  $IC_{50}$  datasets.

Finally, a pre-incubation experiment was performed by incubating 10 different concentrations of inhibitor with enzyme alone, for six different pre-incubation times up to 60 minutes, before initiating the assay by addition of substrate. An end-point absorbance was then measured after 20 minutes of running the activity assay. The endpoint data were used to generate  $IC_{50}$  curves at each pre-incubation time. These  $IC_{50}$  values decrease with increasing pre-incubation time (Fig. 7), phenomenologically similar to the incubation experiment above. However, since no equation exists to fit these pre-incubation  $IC_{50}$  datasets, they can only be analyzed by global fitting using EPIC-CoRe (Fig. 8). This fitting provided values of  $k_5 = 0.64 \pm 0.05 \text{ min}^{-1}$ ,  $k_6 = 0.0045 \pm 0.0015 \text{ min}^{-1}$ ,  $K_i = 141 \pm 8 \text{ nM}$ , and  $K_i^* = 0.99 \pm 0.35 \text{ nM}$ , with an average correlation of  $R^2 = 0.98$ , and RMSE = 4.4.

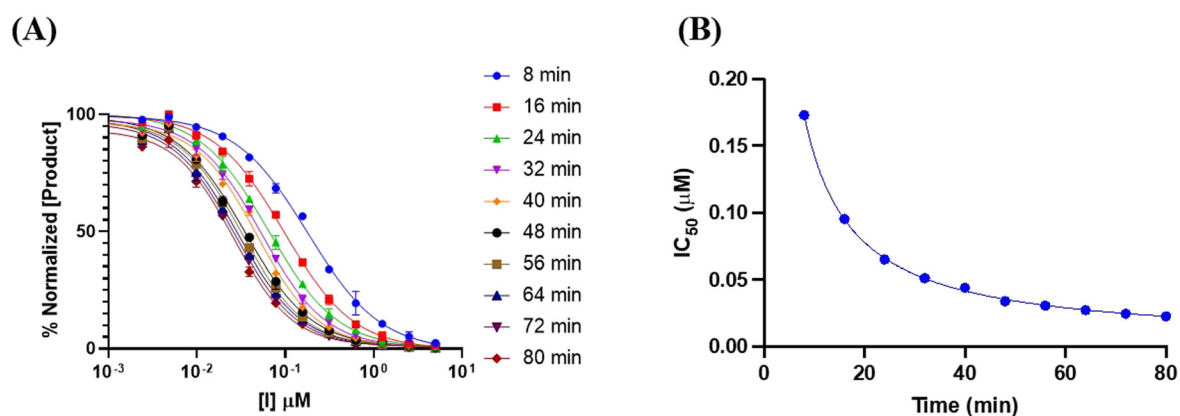


Fig. 6 Incubation time-dependent  $IC_{50}$  experiment with saxagliptin, DPPiV, and Gly-Pro-pNA. (A)  $IC_{50}$  value decreases with increasing incubation time. (B) Fitting of time-dependent  $IC_{50}$  values to implicit eqn (S18) provides  $k_5$ ,  $k_6$ , and  $K_i^{app}$ .



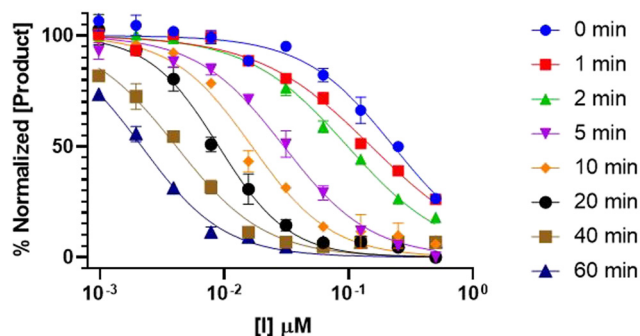


Fig. 7 Time-dependent  $IC_{50}$  curves for the inhibition of DPPiV with saxagliptin, showing decreasing  $IC_{50}$  values with increasing pre-incubation times.

Evaluation of the reaction rate constants and inhibition constants describing the time-dependent reversible covalent inhibition of DPPiV by saxagliptin using the known progress curve analysis method, as well as the new time-dependent  $IC_{50}$  analysis methods presented herein, allows for direct comparison of the precision and

congruency of each method. We have confirmed that saturation fitting from progress curves produces the largest relative error in  $k_6$  and  $K_i^*$ , with errors essentially as large as the fitted values, providing little confidence in the fitting, even though the values do match reasonably with the other methods. All methods provided a  $k_5$  value of around  $0.5\text{--}1.0\text{ min}^{-1}$ , which is consistent with typical forward covalent reaction rate constants for a nitrile warhead (*i.e.*, analogous to  $k_{\text{inact}}$  for an irreversible inhibitor).<sup>7,37,38</sup> Values for  $k_6$  ranged from roughly  $0.003\text{--}0.005\text{ min}^{-1}$ , around 200-fold less than  $k_5$ , which is consistent with the manifestation of time-dependent inhibition that almost resembles irreversibility due to the reverse reaction being very slow (Fig. 4A). This value is also consistent with the observed off-rate constant ( $k_{\text{off}}$ ) that approximates  $k_6$ , as painstakingly determined by Kim *et al.* in an independent jump-dilution experiment.<sup>14</sup> In their experiment, the recovery of enzymatic activity was monitored, to measure  $k_{\text{off}} \approx k_6 \approx 0.0030 \pm 0.0002\text{ min}^{-1}$ , as the slowest, rate-determining step of inhibitor dissociation.<sup>14,39</sup> Our  $K_i$  values ranged from  $\sim 80\text{--}140\text{ nM}$  and were an order of magnitude higher than our

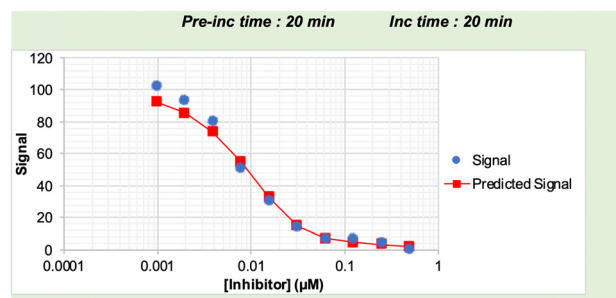
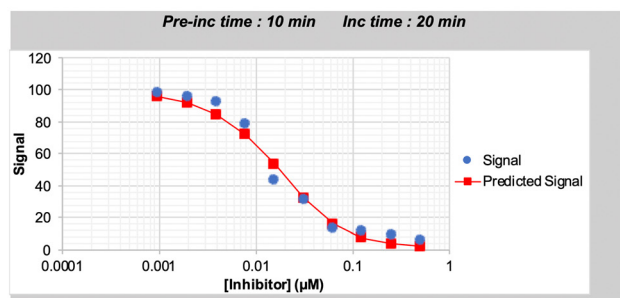
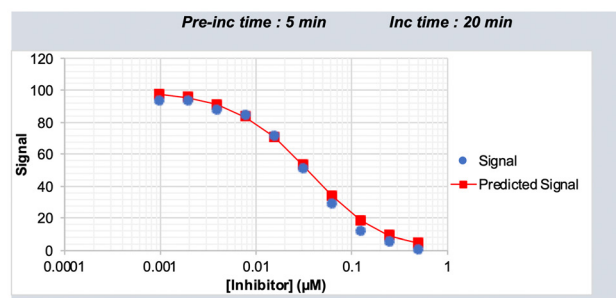
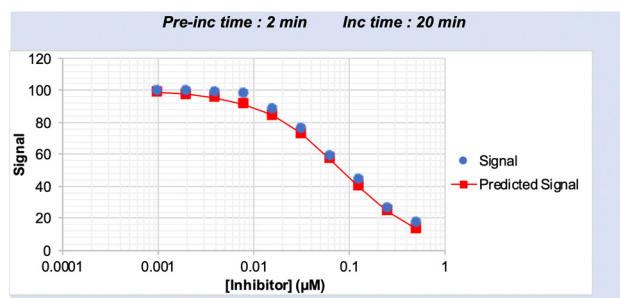
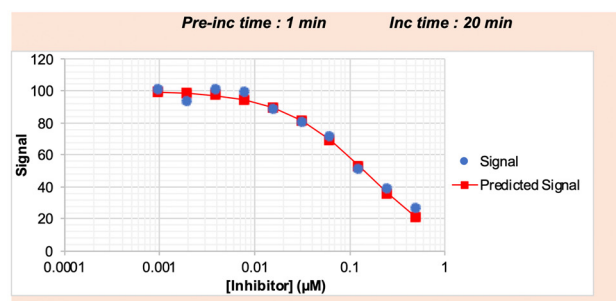
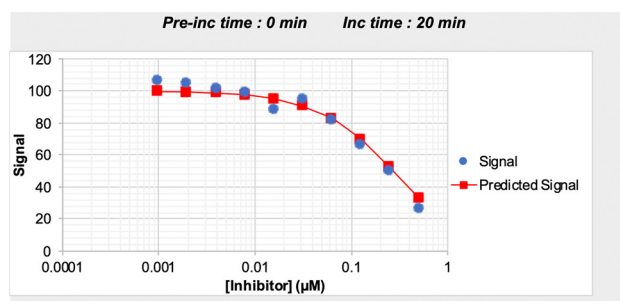


Fig. 8 Global fitting of pre-incubation time dependent  $IC_{50}$  datasets obtained for saxagliptin, using EPIC-CoRe.  $IC_{50}$  curves were obtained for pre-incubation times ranging from 0–20 min and incubation times fixed at 20 min.



calculated  $K_i^*$  values ( $\sim 0.4\text{--}1$  nM), highlighting the importance of taking time-dependency into account to evaluate true affinity. Again, this is consistent with the value of  $K_i^* \approx 0.35$  nM determined by Kim *et al.*, using continuous progress curve fitting and subsequent calculations.<sup>14</sup> Overall, all methods produced reasonably consistent results with each other and literature sources,<sup>14</sup> generally within 1–2 standard deviations. This analysis provides confidence in the use of our new implicit equation, as well as our EPIC-CoRe method, for the robust evaluation of independent values of  $k_5$ ,  $k_6$ ,  $K_i$ , and  $K_i^*$  for time-dependent reversible covalent inhibition from commonly acquired time-dependent  $IC_{50}$  data.

### Practical considerations

From an assay design perspective, continuous methods are generally the most powerful for kinetic characterization. They require relatively little material and are easy to implement, while providing nearly instantaneous information. For irreversible inhibitors, continuous assay methods are indeed the most robust option not only for assay implementation but also subsequent kinetic analysis and fitting.<sup>33</sup> However, in the case of a time-dependent reversible inhibitor, these types of assays can be particularly difficult to optimize and analyze, if they are available at all. Generally, long incubation times on the course of hours are required to fully observe steady state equilibria, and over this period the uninhibited control must be strictly linear, as no algebraic correction can be applied to account for substrate depletion and/or enzyme degradation.<sup>3</sup> This requires considerable optimization of the concentrations of enzyme, substrate, and inhibitor. It also requires that the enzyme itself must be stable over this extended time period. Even when these requirements are met, progress curve fitting may result in high relative error in  $k_6$ , as this value is typically very small and sensitive to any non-linearity in the uninhibited control. This then leads to high errors in the calculated  $K_i^*$  value. In a model system, such as the one demonstrated by our evaluation of saxagliptin, this fitting method worked well and provided reasonable values of kinetic parameters. However, for researchers studying new inhibitors and/or new targets it can be quite cumbersome to perform,<sup>14</sup> especially to extract  $k_5$  and  $k_6$  values. It may be easier to use  $v_i$  and  $v_s$  values to arrive at a ratio of  $k_5/k_6$ , as shown above, but dissection and subsequent tuning of the forward and reverse reactions is not possible.

For the same amount of effort and material, an incubation time-dependent  $IC_{50}$  experiment can be performed, with the added benefit that substrate depletion or degradation of enzyme can be accounted for by normalizing against the uninhibited control reaction. The major consideration for this method is selecting time-points at which to measure  $IC_{50}$  values. Time points that encompass the initial  $v_i$  phase, the transitional phase,

and the final  $v_s$  phase provide the most robust fitting of all parameters, of which the final phase is the most crucial for fitting of  $k_6$ . The times at which these phases occur will depend on the specific inhibitor/substrate/target combination. Generally, it is recommended to run the experiment out to  $>5$  half-lives of the slowest transition phase, in order to allow the final equilibrium to be observed completely. The slowest transition, at the lowest concentration of inhibitor, may be approximated roughly by a  $k_{obs}$  of  $\sim 0.1 \times k_5$  (see eqn (S19)), corresponding to a recommended experiment duration of  $\sim 35/k_5$ .

While fitting to the implicit equation with the saxagliptin model system worked well, we noted that 10 data points and long incubation times were still required to provide precise fitting. Interestingly, we have found that for this type of experiment using EPIC-CoRe requires fewer time points overall to arrive at similar solutions to the implicit equation. For the sake of demonstration, the results shown above include 6 time-points up to 80 minutes; however, similar results were obtained by monitoring for up to only 40 minutes with 4 data sets (see ESI,† Fig. S5), cutting the assay time in half. This is likely due to the robustness provided by global fitting of entire datasets rather than fitting the variance of  $IC_{50}$  values. Although this was not our original intended application, we note that EPIC-CoRe may be better suited for estimation of kinetic parameters from incubation time-dependent  $IC_{50}$  datasets that contain more error or noise.

While a pre-incubation time-dependent  $IC_{50}$  experiment does require more material (particularly more enzyme) to perform, it is commonly employed in industry as it lends itself well to automated assay procedures where end-point observations are taken at a pre-defined time into the activity assay. Similar to its utility in incubation time-dependent  $IC_{50}$  experiments, we have found that fewer time-points (*i.e.*, at least 4) and shorter assay times are required for robust fitting (see ESI,† Fig. S6). With a pre-incubation phase included, the final steady state equilibrium is reached much faster than in a single-phase incubation experiment, allowing shorter subsequent (incubation phase) assay times to provide sufficient information for fitting. We generally recommend using a broad range of pre-incubation times, such as zero,  $1/k_5$ ,  $2/k_5$ , and  $5/k_5$ , which would normally ensure broad variation of  $IC_{50}$  dataset values and robust fitting.<sup>1</sup>

For both methods, successful fitting requires datasets that show broad variance of end-point signal with respect to both inhibitor concentration and time, such that the dynamic range and inflection point of each  $IC_{50}$  curve is well-defined and time-dependence is clear between the different data sets. These methods do not account for more complex inhibition behaviour that result in binding isotherms with atypical Hill slopes (*i.e.*,  $n_H \neq 1$ ). In our experience, EPIC-CoRe provides robust fitting from initial estimates that may be orders of magnitude different from optimized values, whereas the implicit equation requires



more accurate initial estimates for  $k_5$  and  $k_6$ . However, given that time-dependent reversible inhibition generally results when  $k_6$  is at least an order of magnitude less than  $k_5$ , and  $k_5$  is analogous to well-catalogued  $k_{\text{inact}}$  values for a broad range of irreversible warheads,<sup>1</sup> initial estimates of 0.1–1.0 min<sup>-1</sup> for  $k_5$  and 0.001–0.010 min<sup>-1</sup> for  $k_6$  are appropriate for a broad range of inhibitors. We recommend constraining  $k_6 < k_5$  during fitting, as this must be true for time-dependent inhibition to be observed (where  $\nu_s < \nu_i$ ). Initial estimates for  $K_i$  require less accuracy, and a value of 0.1–1.0  $\mu\text{M}$  is generally suitable. Further guidance for an initial estimate for  $K_i$  can be obtained from the IC<sub>50</sub> value measured at the shortest incubation or pre-incubation time, as the  $K_i$  value will be higher than this.<sup>1</sup> For this reason, we also recommend including a pre-incubation time of zero among the pre-incubation data sets. It is also important to note that inhibitors must show appreciable reversibility to allow  $k_6$  to be fitted. As  $k_6$  gets smaller, inhibition becomes effectively irreversible, and the fitting methods discussed in this work will result in a fitted value of  $k_6$  approaching zero. For these inhibitors it may be more expedient to treat them as irreversible and to fit the data sets as described previously.<sup>1</sup>

If IC<sub>50</sub> values or global datasets do not demonstrate sufficient predictable variance of signal as a function of time, fitting by implicit equation or EPIC-CoRe may not converge to a unique solution. In this case, it may be necessary to include additional datasets covering a larger time frame, as recommended above. It is also possible to fix one parameter (*i.e.*, fix  $K_i$  to a value larger than the one obtained from the shortest incubation/pre-incubation time, or fix  $k_5$  based on known values for similar warheads), fit the rest of the parameters, and then fix those parameters and re-fit the first. This will also allow one to confirm convergence to a global minimum over a local minimum. In the spreadsheet we have developed for EPIC-CoRe, real-time visual comparison of the experimental data and predicted values allow the user to gauge how well their initial estimates of fitted parameters describe the data sets. It is therefore possible to manually adjust  $K_i$ ,  $k_5$ , and  $k_6$  values to arrive at reasonable estimates prior to Solver-powered fitting. Adjustment of  $K_i$  will largely shift the inflection points of all of the predicted IC<sub>50</sub> curves along the  $x$ -axis, while the adjustment of  $k_5$  and  $k_6$  will more prominently alter the degree of their time-dependence. This provides a user-friendly and intuitive way to guide the fitting procedure.

Drug design optimized for strong on-target interaction and weak off-target reactivity/binding is generally best guided by tuning overall  $k_{\text{off}}$  values (for drug release) rather than  $k_{\text{on}}$  values (for drug binding).<sup>39</sup> This is because the *in vivo* rate of association and actual concentration of drug reaching the target is heavily influenced by pharmacokinetics. Additionally, diffusion, desolvation, and orientation effects are difficult to predict.

However, the rate of dissociation is almost entirely dependent on specific drug-target interactions alone, and can be tuned systematically by structure-based design.<sup>16,39</sup> Time-dependent reversible covalent inhibitors provide a unique avenue to do this, as the overall  $k_{\text{off}}$  value and therefore drug-target residence time will be most influenced by  $k_6$  (note that  $k_{\text{off}}$  and residence time are actually a collection of rate constants).<sup>39</sup> Knowledge of on- and off-target protein microenvironments and the mechanism of the reverse covalent reaction allows for fine-tuning of warhead structure to alter  $k_6$  values to lower values for the target protein and to higher values for off-targets, providing additional potency and selectivity.<sup>22,24</sup> From a practical standpoint, reversibility is only relevant *in vivo* if the drug dissociates from a protein faster than its rate of degradation. In some cases, residence times of reversible covalent inhibitors can be so long that the inhibitor does not dissociate before the protein is degraded; effectively, this is equivalent to irreversible modification.<sup>22</sup> However, a reversible inhibitor may be able to recapture newly synthesized protein, once the original to which it was bound has been degraded. If the residence time with off-targets is much lower, this provides the advantage of fewer side effects while maintaining prolonged desired target engagement.<sup>23</sup> For these reasons, it is highly recommended to fully characterize these inhibitors with  $k_6$  values for rational drug design, which is now possible from IC<sub>50</sub> experiments, using the methods reported in this work.

Finally, it is relevant to discuss the limitations of the scope of these new methods, which illustrates possible directions for future work. First, it is important to reiterate that all of the methods discussed herein are only applicable to assays that employ competitive inhibitor-substrate pairs. While this is currently the most common application for covalent inhibition, we acknowledge that it would be possible to design covalent inhibitors that operate at an allosteric site and are non-competitive in modality.<sup>40</sup> Neither our EPIC-CoRe spreadsheet nor implicit eqn (S18) account for this mode of inhibition. Furthermore, if the E-S-I complex from a non-competitive inhibition scheme can lead to (slower) product formation, it would be possible to observe a non-zero lower plateau in experimental IC<sub>50</sub> datasets. Our numerical modelling does not account for this pathway for product formation, and both eqn (S18) and Krippendorff's equation were derived on the assumption that saturation with inhibitor leads to zero product formation (see eqn (S15)†).<sup>29</sup> It is also important to clarify that our work focuses on truly reversible covalent inhibitors, and does not account for covalent inhibitors that may be slowly consumed by the enzyme as a pseudo-substrate.<sup>41,42</sup> Finally, apart from these very unusual cases where a covalent inhibitor may be non-competitive or undergo enzyme-mediated degradation, it is possible that an optimised covalent inhibitor may be bound very tightly, such that its  $K_i^{\text{app}}$  value approaches the concentration of



enzyme used in the assay. Eqn (S18), like Krippendorff's equation, was derived using the free ligand approximation (see eqn (S4)†) and does not account for tight binding. Experimentally, the use of low enzyme concentrations and high competing substrate (such that  $K_i^{\text{APP}} \geq [\text{Enz}]$ ) would extend the range of applicability of these implicit equations. However, the numerical modelling embodied in EPIC-CoRe uses Morrison's quadratic equation to calculate the fraction of E-I,<sup>43</sup> which means it is uniquely appropriate to be applied to the evaluation of tight-binding reversible covalent inhibitors, as well.

## Conclusions

In recent years there has been an emerging interest in reversible covalent inhibitors that provide enhanced covalent affinity without permanent off-target modifications. Time-dependent inhibitors of this type that dissociate slowly from their target, due to the reverse covalent reaction rate constant ( $k_6$ ) being small, are particularly notable, as they allow for fine-tuning of residence time based on warhead structure and protein microenvironment. The goal is to modulate  $k_5$  and  $k_6$  to allow for a moderately fast on-target addition reaction and slow dissociation from the desired target, but relatively rapid dissociation from any off-target reactive proteins. This strategy has proven successful for many kinases and growth receptors. Best practice to achieve this sort of fine-tuning requires individual characterization of all relevant inhibition and rate constants ( $k_5$ ,  $k_6$ ,  $K_i$ , and  $K_i^*$ ) for this complex kinetic scheme, by taking time-dependency into account. Previously, the only practical way to achieve this was through progress curve analysis, which requires the availability of a convenient continuous assay and often results in high errors. Recognizing that these inhibitors are often assessed more practically using discontinuous (end-point)  $\text{IC}_{50}$  experiments, we developed two methods that allow for time-dependent  $\text{IC}_{50}$  data from time-dependent reversible covalent inhibitors to be analyzed, to provide all aforementioned constants. For incubation  $\text{IC}_{50}$  experiments, we present an implicit equation that can be used to fit  $\text{IC}_{50}$  value vs. time data to arrive at these constants. For pre-incubation  $\text{IC}_{50}$  experiments, we have developed EPIC-CoRe, a numerical modelling method that relates the global fitting of  $\text{IC}_{50}$  data sets to all relevant parameters. EPIC-CoRe features versatility that allows it to be used for incubation  $\text{IC}_{50}$  data sets (*i.e.*, obtained with no pre-incubation phase) as well, providing an alternative to the complex implicit equation. These methods can be implemented quite easily by any practicing medicinal chemist as they only require common fitting (*e.g.*, GraphPad Prism) or spreadsheet (*e.g.*, Microsoft Excel) software. Both methods were shown to provide values that are consistent with other known methods of evaluation, as demonstrated here for the DPPiV inhibitor, saxagliptin, confirming their robustness and utility. To the best of our

knowledge, this is the first demonstration of approaches that allow discontinuous  $\text{IC}_{50}$  data to be used to separately derive inhibition and rate constants of a time-dependent reversible covalent inhibitor. This is critical to drug design and optimization, allowing chemists to discern and improve overall on-target affinity ( $K_i^*$ ) and reactivity ( $k_5$  and  $k_6$ ), while maintaining intrinsic and off-target stability. Our methods serve to bridge the gap between the most commonly used assay methods for reversible covalent inhibitors, namely discontinuous  $\text{IC}_{50}$  experiments, and the most powerful kinetic characterization required for the fine-tuning of effective inhibitors of this type.

## Experimental section

### General

No unexpected, new, or significant hazards/risks were encountered in any of the experimental procedures. All commercially available chemicals, enzymes, and solvents were used without further purification. Saxagliptin hydrochloride (item no. 23697) and Gly-Pro-*p*NA hydrochloride (item no. 21244) were purchased from Cayman Chemical with a reported purity of >95% as determined by HPLC analysis. Active recombinant, and untagged, DPPiV was purchased from Enzo Life Sciences Inc. as part of a commercial activity assay kit (BML-AK499-0001). The provided stock enzyme concentration was  $0.071 \text{ mg mL}^{-1}$  with an activity of  $0.867 \text{ U mL}^{-1}$ . Biochemical assays were conducted directly in standard sterile 96-well plates and absorbance readings were taken in either continuous or discontinuous fashion using a BioTek Synergy 4 plate reader at 405 nm at 25 °C. Raw and/or processed readout data were analyzed by either GraphPad Prism or our EPIC-CoRe (Microsoft Excel) spreadsheet. All experiments were performed in at least duplicate. Derivations of all equations are provided in the Supporting Information, as well as commentary on how these relate to the differential equations used to develop EPIC-CoRe.

### Determination of $K_M$ and $k_{\text{cat}}$ of Gly-Pro-*p*NA with DPPiV

Buffered solutions of 50 mM TRIS (pH 7.5) and various concentrations of Gly-Pro-*p*NA (from 10–500  $\mu\text{M}$  in the final assay solution, representing roughly  $0.1\text{--}5 \times K_M$ ) were prepared in 96-well plates. To initiate the enzymatic reactions, a solution of DPPiV was added to the wells, ensuring that the final concentration was 2.5 nM ( $2.6 \text{ mU mL}^{-1}$ ), and additional buffer was added to the blank. The final well volume was 100  $\mu\text{L}$ . Stock solutions of Gly-Pro-*p*NA were prepared by diluting a 10 mM DMSO stock in assay buffer, ensuring that the final concentration of DMSO did not exceed 2.5% v/v. The enzyme stock solution was prepared by dilution in assay buffer. The formation of the hydrolysis product, *para*-nitroaniline (*p*NA) was monitored



at 405 nm for 80 min. No background hydrolysis was observed over this time period. Initial rates were calculated over 10% of substrate conversion (Fig. S1†). A standard curve was generated (Fig. S2†) for product *p*NA by taking an absorbance reading of solutions of concentrations varying from 10–500 μM, in assay buffer. Stock solutions of *p*NA were prepared using a 10 mM DMSO stock and diluting in assay buffer. A blank absorbance reading with assay buffer alone was subtracted from the readings. Absorbance readings were plotted against [*p*NA] and the slope of this line (representing the experimental extinction coefficient) was used to convert the initial rates from Abs/s to μM s<sup>-1</sup>. These initial rates were then plotted against [Gly-Pro-*p*NA] to generate a saturation plot (Fig. S3†) that was fitted by non-linear regression to the Michaelis–Menten equation (eqn (1), cf. eqn (S1)†) to obtain  $K_M$  and  $V_{max}$ . The value of  $k_{cat}$  was then calculated using eqn (2).

$$v_0 = \frac{V_{max}[S]}{K_M + [S]} \quad (1)$$

$$k_{cat} = \frac{V_{max}}{[E]} \quad (2)$$

### Continuous activity assay

Enzymatic assays were run according to a modified procedure reported by Kim *et al.*,<sup>14</sup> in the presence of final concentrations of 500 μM ( $\approx 7 \times K_M$ ) Gly-Pro-*p*NA and 0.64 nM (0.65 mU mL<sup>-1</sup>) DPPiV. These concentrations were chosen to ensure a significant and linear signal in the uninhibited positive control over the 80 minute assay. Buffered solutions of 50 mM TRIS (pH 7.5), 500 μM Gly-Pro-*p*NA, and various concentrations of saxagliptin (2.4–5000 nM in the final assay well, 12 2-fold serial dilutions, representing roughly 0.01–10 ×  $K_i^{app}$ ) were prepared in a 96-well plate. Saxagliptin stocks were prepared by dilution of a 10 mM stock in assay buffer, ensuring that the concentration of DMSO did not exceed 2.5% v/v in the final assay well. To initiate the enzymatic reactions, DPPiV was added to a final concentration of 0.64 nM (0.65 mU mL<sup>-1</sup>) and additional buffer was added to the blank. The final well volume was 100 μL. Formation of *p*NA was followed at 405 nm for 80 minutes. No background reaction was observed. Observed pseudo-first order rate constants of inactivation ( $k_{obs}$ ), initial inhibited rates  $v_i$ , and final inhibited rates  $v_s$ , were obtained by fitting the resulting progress curves to eqn (3) (cf. eqn (S13)).

$$Abs_t = v_s t + \frac{(v_i - v_s)}{k_{obs}} (1 - e^{-k_{obs}t}) \quad (3)$$

Fitted  $k_{obs}$  values were plotted against inhibitor concentrations corrected for the presence of substrate ( $[I]/\alpha$ ) by dividing inhibitor concentrations by ( $\alpha = 1 + [S]/K_M = 8.14$ ). The resulting hyperbolic curve was fitted to eqn (4) (cf.

eqn (S12)) to obtain values for  $k_5$ ,  $k_6$ , and  $K_i$ . A value for  $K_i^*$  was subsequently calculated using eqn (S11).

$$k_{obs} = k_6 + \left( \frac{k_5 \frac{[I]}{\alpha}}{\frac{[I]}{\alpha} + K_i} \right) \quad (4)$$

$$K_i^* = \frac{K_i}{1 + \frac{k_5}{k_6}} \quad (S11)$$

Alternatively, the fitted  $v_i$  and  $v_s$  values were normalized with respect to the uninhibited control by dividing by the slope of the control (note the slope of the control should be the same throughout the assay, *i.e.*,  $v_{i,0} = v_{s,0}$ ) and multiplying by 100. The resulting % inhibition values were plotted against  $[I]/\alpha$  on a logarithmic scale and IC<sub>50</sub> values were determined using four-parameter fitting according to eqn (5) (the upper plateau (top) can be fixed at 100, the lower plateau (bottom) can be fixed at 0, and a Hill slope can be fixed at 1 if needed). The IC<sub>50</sub> value obtained from analysis of initial slopes is equal to  $K_i$  and the IC<sub>50</sub> value obtained from analysis of steady state slopes is equal to  $K_i^*$ .

$$Y = \frac{\text{Top} - \text{Bottom}}{1 + \left( \frac{IC_{50}}{[I]/\alpha} \right)^{\text{HillSlope}}} + \text{Bottom} \quad (5)$$

### Incubation time-dependent IC<sub>50</sub> assay

Enzymatic assays were run under the same conditions as the continuous activity assay described above. An absorbance reading at 405 nm was taken from each reaction with different saxagliptin concentrations at 8, 16, 24, 32, 40, 48, 56, 64, 72, and 80 min observation times. No background reaction was observed. The absorbance reading of each inhibited reaction was divided by the absorbance reading taken from the uninhibited control at the same time to give a percent relative absorbance according to the following equation (eqn (6)):

$$\% \text{ Relative Abs}_t = \frac{Abs_{inhib}}{Abs_{uninhib}} \times 100\% \quad (6)$$

These percent relative absorbance values for each time point were plotted against the inhibitor concentration on a logarithmic scale to generate IC<sub>50</sub> curves. IC<sub>50</sub> values at each time point were determined using four-parameter fitting as described above (eqn (5)). These IC<sub>50</sub> values were then plotted against their corresponding observation times to generate a curve that was fitted to user-defined implicit eqn (S18) using initial estimates of  $k_5 = 0.1 \text{ min}^{-1}$ ,  $k_6 = 0.01 \text{ min}^{-1}$ , and  $K_i = 0.1 \text{ μM}$  to obtain  $k_5$ ,  $k_6$ , and  $K_i^{app}$ ; then  $K_i$  was calculated from the Cheng–Prusoff equation (eqn (S4)†) and  $K_i^*$  was calculated using eqn (S11).



$$IC_{50(t)} = K_i^{app} \left( \frac{k_6}{k_6 + k_5} \right) + \left( \frac{\left( \frac{2k_6}{k_6 + k_5} \right) (K_i^{app})^2 + 2K_i^{app} IC_{50}(t)}{K_i^{app} + IC_{50}(t)} - 2K_i^{app} \left( \frac{k_6}{k_6 + k_5} \right) \right) \cdot \frac{(1 - e^{-k_{obs}t})}{k_{obs}t} \quad (S18)$$

where

$$k_{obs} = k_6 + \left( \frac{k_5 IC_{50}(t)}{IC_{50}(t) + K_i^{app}} \right) \quad (S19)$$

Alternatively, the percent relative absorbance values were entered into our EPIC-CoRe Excel sheet, using pre-incubation times of zero and incubation times of 8, 24, 40, 56, 72, and 80 min for the corresponding datasets. The response coefficient for each data set (which corresponds to the signal sensitivity and normalisation) was adjusted to give a value of approximately 100% in the presence of 'zero' inhibitor. The substrate and enzyme concentrations were 500  $\mu$ M and 0.64 nM, respectively. The  $k_{cat}$  value used was 1056  $\text{min}^{-1}$  and  $K_M$  was 70.7  $\mu$ M, as measured herein (see above). The pre-incubation and incubation volumes were both set as 100  $\mu$ L, representing a dilution factor of 1 (*i.e.* no dilution). Initial estimates of  $k_5 = 0.1 \text{ min}^{-1}$ ,  $k_6 = 0.01 \text{ min}^{-1}$ , and  $K_i = 0.1 \mu\text{M}$  were entered before fitting these values using Solver. The value of  $K_i^*$  was automatically calculated within the spreadsheet according to eqn (S11). This fitting procedure was repeated on each individual dataset, as well as on a dataset of averaged endpoint values, to generate a standard deviation reflective of experimental error, as well as  $R^2$  and RMSE values to assess goodness of fit.

### Pre-incubation time-dependent $IC_{50}$ assay

A 96-well microplate was set up with 12 different reactions, in duplicate, representing an uninhibited control, 10 different saxagliptin concentrations (2-fold serial dilutions starting from 1  $\mu$ M), and a blank, each to an initial volume of 35  $\mu$ L in assay buffer (50 mM TRIS, pH 7.5). Then, 15  $\mu$ L of a 0.0043  $\mu$ M (4.3 mU  $\text{mL}^{-1}$ ) solution of DPPiV in assay buffer was added to each well using a multi-channel pipette. This gave a final concentration of 0.0013  $\mu$ M (1.3 mU  $\text{mL}^{-1}$ ) of DPPiV and 2-fold dilutions of saxagliptin starting from 0.5  $\mu$ M in the pre-incubation phase. After pre-incubation, 50  $\mu$ L of a 1 mM stock solution of Gly-Pro-pNA in assay buffer (prepared from a 20 mM DMSO stock) was added to each well using a multi-channel pipette to initiate the activity assay, giving a final concentration of 500  $\mu$ M Gly-Pro-pNA in the assay and approximately 2.5% DMSO v/v fixed in each well, with the final well volume being 100  $\mu$ L. After a 20 min assay incubation period, an absorbance reading at 405 nm was taken from each well. This was repeated, in duplicate, for pre-incubation times of 0, 1, 2, 5, 10, and 20 min. No background reaction was observed.

The absorbance reading of each inhibited reaction was divided by the absorbance reading taken from the uninhibited control to give a percent relative absorbance

(eqn (6)). The percent relative absorbance values were entered into our EPIC-CoRe Excel sheet with the pre-incubation times set to 0, 1, 2, 5, 10, and 20 min and an incubation time of 20 min for the corresponding datasets. The response coefficient for each data set was adjusted to give a value of approximately 100% in the presence of 'zero' inhibitor. The substrate and enzyme concentrations were entered as 500 and 0.0013  $\mu$ M, respectively. The  $k_{cat}$  was entered as 1056  $\text{min}^{-1}$  and  $K_M$  as 70.7  $\mu$ M, as determined herein (see above). The pre-incubation volume was set to 50  $\mu$ L and incubation volume was set to 100  $\mu$ L, representing a dilution factor of 0.5. Initial estimates of  $k_5 = 0.1 \text{ min}^{-1}$ ,  $k_6 = 0.01 \text{ min}^{-1}$ , and  $K_i = 0.1 \mu\text{M}$  were entered before fitting these values using Solver. The value of  $K_i^*$  was automatically calculated within the spreadsheet according to eqn (S11). This fitting procedure was repeated on each individual dataset, as well as an averaged data set to generate a standard deviation reflective of experimental error, as well as  $R^2$  and RMSE values to assess goodness of fit.

## List of abbreviations

TCI	Targeted covalent inhibitor
EPIC-CoRe	Endpoint Pre-Incubation $IC_{50}$ fitting-Covalent Reversible
Gly-Pro-pNA	Glycine-proline- <i>para</i> -nitroanilide
DMSO	Dimethyl sulfoxide
DPPiV	Dipeptidyl peptidase 4
RMSE	Root mean squared error
VBA	Visual basic for applications
RSS	Residual sum of squares
TRIS	Tris(hydroxymethyl)aminomethane

## Data availability

The data supporting this article have been included as part of the ESI.† The code for the fitting spreadsheet can be found at <https://github.com/JeffreyKeillor/JeffreyKeillor>. The version of the code employed for this study is version 1.1.

## Author contributions

Conceptualization: J. W. K. and L. K. M.; funding acquisition: J. W. K.; supervision: J. W. K.; investigation and methodology: J. W. K. (method development), L. K. M. (characterization and enzymatic assays), writing – original draft: L. K. M.; reviewing and editing: L. K. M. and J. W. K.

## Conflicts of interest

There are no conflicts of interest to declare.



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