Brodifacoum pharmacokinetics in acute human poisoning: implications for estimating duration of vitamin K therapy

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ABSTRACT
Standard of care follow-up therapy for patients poisoned by long-acting anticoagulant rodenticides (LAARs) is daily high-dose (up to 100 mg per day) oral vitamin K1 (VK1) for weeks to months to over a year. The availability of CLIA-certified quantitative testing for plasma LAAR concentrations can now assist health care providers in determining when to safely discontinue VK1 therapy. We present estimates of treatment duration required to reach safe concentrations (< 10 ng/ml) using serial measurements of plasma brodifacoum (BDF, a potent LAAR) concentrations obtained from patients poisoned after inhaling synthetic cannabinoids containing BDF. We fit the data to zero-order (linear) and first-order (exponential) curves, the latter to account for enterohepatic circulation of BDF. The results show that estimates of therapy duration are significantly longer when exponential clearance is assumed. Accordingly, we recommend that plasma BDF concentrations be monitored simultaneously with international normalization ratio (INR) during follow-up of poisoned patients, and that concentrations be determined after VK1 therapy is discontinued to document persistence of safe concentrations.

KEYWORDS
Brodifacoum; synthetic cannabinoids; rodenticide; vitamin K1

Introduction
A recent outbreak of long-acting anticoagulant rodenticide (LAAR)-associated coagulopathy and life-threatening bleeding in Illinois and several other States [1–3] focused attention on long-term treatment of this condition with vitamin K1 (VK1). Current standard of care treatment of patients poisoned by the LAAR brodifacoum (BDF) is VK1 given intravenously or orally, thereafter requiring supplementation for durations of weeks to months. Extended treatment is due to the long biological half-life of BDF, resulting from a combination of insignificant liver metabolism, lack of renal excretion, accumulation in lipophilic tissues, and enterohepatic circulation [4]. Since VK1 does not increase BDF clearance, treatment is required until gradual elimination leads to a safe serum level of <10 ng/ml [5]. Premature discontinuation of VK1 before this threshold is reached may lead to recurrence of coagulopathy and bleeding. Hence, both international normalized ratio (INR) and plasma BDF concentrations must be monitored during follow-up. One means of estimating the duration of VK1 treatment is to carry out serial measurements of plasma BDF concentrations, then fitting the data to pharmacokinetic models of BDF clearance which can be described as either 1- or 2-compartment models, the latter to account for biphasic elimination and enterohepatic circulation [6]. The purpose of this study was to estimate the time required for VK1 treatment after acute BDF poisoning by fitting quantitative data of plasma BDF concentrations obtained from patients acutely poisoned in Peoria, Illinois.

Methods
We previously quantified plasma BDF concentrations using a high pressure liquid chromatography-mass spectrophotometric method in blood samples obtained from patients poisoned due to inhalation of synthetic cannabinoids contaminated with BDF [7]. For this study, ‘initial BDF concentration’ refers to the concentration measured in the first blood sample collected after a patient was admitted to the...
emergency department; however, the time between BDF exposure to admission is not known. CurveExpert Professional v2.7.3 (Hyams Development, Chattanooga, TN) was used to directly fit plasma BDF values to zero-order (linear) or first-order (exponential) decay without linearization of the data, and we used the resulting equations to calculate the time needed to reach 10 ng/ml. Zero versus first-order derived times were compared by paired t-test, and associations by Pearson correlation analysis.

**Results**

Of the 33 patients whose samples were available, there were 19 for whom 3 or more serial blood samples were collected. When we fit the data directly to zero-order (linear) decline, the average time to reach 10 ng/ml was 12.2 ± 8.3 days (Table 1). In contrast, in all cases the time to reach 10 ng/ml was longer when the data were fitted by exponential regression to a first-order decay (43.4 ± 35.4 days; \( p < 0.0001 \) versus zero-order, paired t-test). Although R-squared values for zero and first-order regressions were similar (0.622 versus 0.645, Table 1), they were significantly higher (\( p < 0.01 \)) when first-order regression was used. We did not detect any significant associations between initial BDF concentrations measured and either the zero or first-order calculated times to 10 ng/ml. However, the values determined by zero-order regression were more similar to first-order values at lower compared to higher initial BDF concentrations, and the ratio (first-order: zero-order) was positively correlated to the initial BDF concentrations (\( p = 0.020 \)) (Figure 1).

**Discussion**

Clearance of BDF is complex due to distribution into multiple compartments, minimal metabolism, and enterohepatic circulation. While most studies have used first-order, exponential modeling to describe its pharmacokinetics [7–9], several have also used linear modeling [5, 10]. Measurements of plasma BDF concentrations made at early times relative to the initial exposure, as done here, will reflect a combination of partitioning into multiple compartments, enterohepatic circulation, and elimination. In contrast measurements done at later times when the distribution phase is complete will primarily reflect the elimination phase and ongoing enterohepatic circulation. In this study, the times between initial exposure to BDF and the first blood draw (the initial measurement) were not known, therefore samples with low initial BDF concentrations could reflect longer duration since exposure; or exposure to lower amounts. Our finding that zero-order fitting more closely paralleled first-order fitting at lower initial BDF concentrations suggest that the initial distribution phase was already completed in those cases, although it remains possible that the distribution phase at low BDF concentrations may, in some patients, follow linear kinetics.

**Table 1.** Calculation of time to reach safe plasma BDF concentrations.

| Case # | First BDF [ng/ml] | \( T_{1/2} \) (days) | Zero Order R^2 | First Order R^2 |
|--------|-------------------|----------------------|----------------|----------------|
| 29     | 130               | 7.6                  | 14.1           | 31.1           |
| 20     | 207               | 8.9                  | 13.0           | 38.8           |
| 27     | 211               | 4.4                  | 7.1            | 17.0           |
| 24     | 270               | 2.6                  | 4.0            | 11.8           |
| 23     | 294               | 4.6                  | 8.7            | 20.8           |
| 4      | 355               | 2.0                  | 3.5            | 9.1            |
| 28     | 355               | 17.8                 | 26.7           | 100.0          |
| 19     | 431               | 2.3                  | 2.9            | 6.8            |
| 21     | 460               | 12.4                 | 17.0           | 54.8           |
| 31     | 463               | 3.9                  | 7.4            | 21.6           |
| 22     | 563               | 7.1                  | 10.8           | 39.8           |
| 30     | 666               | 18.6                 | 28.0           | 112.8          |
| 33     | 678               | 6.6                  | 9.9            | 35.4           |
| 10     | 739               | 8.7                  | 15.7           | 60.2           |
| 32     | 822               | 17.6                 | 26.6           | 104.8          |
| 14     | 1443              | 14.0                 | 21.0           | 100.4          |
| 18     | 1451              | 3.1                  | 5.6            | 19.4           |
| 15     | 1458              | 3.4                  | 5.8            | 24.1           |
| 26     | 1862              | 2.5                  | 4.2            | 16.0           |
| average|                  | 7.8                  | 12.2           | 43.4 ^a        |

^a\( p < 0.0001 \) versus zero-order, paired t-test.

^b\( p < 0.01 \) versus zero-order, paired t-test. First BDF, BDF level measured in the first blood sample collected after patient was admitted to the emergency department. \( T_{1/2} \), half-life determined for BDF. Data adapted from [7].
Determining when to discontinue VK1 therapy for BDF poisoned patients is critical to avoid recurrence of life-threatening bleeding. Although INR values can indicate return to normal clotting function, that may not reflect safe plasma BDF concentrations. In our review of 21 case reports of LAAR poisoning [11], about half the patients had plasma BDF concentrations of <10 ng/ml at discharge from hospital, presumably INRs were within normal range at discharge. More recently, in our analysis of poisonings due to inhaling synthetic cannabinoids laced with BDF [7], we found that in most cases (24 of 27 cases) although INRs had reached normal values (<1.7) at discharge, the average plasma BDF concentration at discharge was 454 ng/ml, more than 40 times the considered safe level. In Wisconsin [12], a cohort of 19 patients also poisoned by using contaminated synthetic cannabinoids was followed for up to 102 days; and of those BDF rebound was observed in 5 cases. Whether due to patient non-adherence to VK1 treatment, repeated use of BDF-tainted cannabinoids, or if VK1 therapy was discontinued prematurely when INRs were normalized but BDF was still present leading to recurrence of anti-coagulation is not known.

Conclusions

We strongly endorse recommendations to perform serial measurements of plasma BDF to guide the post-exposure duration of oral VK1 therapy in poisoned patients [5]. Adequate modeling requires multiple (at least 3) sampling of plasma concentrations be done; and based on the reported half life for BDF (7.5 ± 1.3 days, mean ± se, [7]), we recommend those be collected over a period of 1 week. Using first-order modeling to describe its pharmacokinetics can provide an estimate of the required duration of continuous VK1 treatment. Alternatively, assuming zero-order decline can provide an estimate when the current phase of clearance will be complete. We further recommend that at least one additional measurement of plasma BDF be carried out at the end of treatment to ensure that concentrations of 10 ng/ml or less are reached, and that a second measurement be made within 2 weeks after cessation of VK1 to ascertain maintenance of safe concentrations. The availability of certified laboratories in the USA to quantify plasma BDF concentration now enables such testing to be ordered by healthcare providers at point of care thereby improving patient care [5].

Disclosure of interest

The authors report no conflict of interests.

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