In vitro Activation of heme oxygenase-2 by menadione and its analogs

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Abstract

Background: Previously, we reported that menadione activated rat, native heme oxygenase-2 (HO-2) and human recombinant heme oxygenase-2 selectively; it did not activate spleen, microsomal heme oxygenase-1. The purpose of this study was to explore some structure–activity relationships of this activation and the idea that redox properties may be an important aspect of menadione efficacy.

Methods: Heme oxygenase activity was determined in vitro using rat spleen and brain microsomes as the sources of heme oxygenase-1 and –2, respectively, as well as recombinant, human heme oxygenase-2.

Results: Menadione analogs with bulky aliphatic groups at position-3, namely vitamins K1 and K2, were not able to activate HO-2. In contrast, several compounds with similar bulky but less lipophilic moieties at position-2 (and –3) were able to activate HO-2 many fold; these compounds included polar, rigid, furan-containing naphthoquinones, furan-benzoxazine naphthoquinones, 2-(aminophenylphenyl)-3-piperidin-1-yl naphthoquinones. To explore the idea that redox properties might be involved in menadione efficacy, we tested analogs such as 1,4-dimethoxy-2-methylnaphthalene, pentafluoromenadione, monohalogenated naphthoquinones, α-tetralone and 1,4-naphthoquinone. All of these compounds were inactive except for 1,4-naphthoquinone. Menadione activated full-length recombinant human heme oxygenase-2 (FL-hHO-2) as effectively as rat brain enzyme, but it did not activate rat spleen heme oxygenase.

Conclusions: These observations are consistent with the idea that naphthoquinones such as menadione bind to a receptor in HO-2 and activate the enzyme through a mechanism that may involve redox properties.

Keywords: Heme oxygenase, Enzyme activator, Menadione, Agonist, In vitro, Redox properties

Background

Drug targets or drug receptors include a broad array of cellular entities that range from receptors for intercellular transmitters to ion channels to genetic macromolecules to enzymes. Examples of the latter include xanthine oxidase, aldehyde dehydrogenase, angiotensin converting enzyme and acetylcholinesterase for which allopurinol, disulfuram, enalaprilate and organophosphates, respectively, act as ligands. In their 2010 review, Zorn and Wells [1] highlight the observation that inhibition of enzyme function is the usual modus operandi of drug discovery programs that target specific enzymes; accordingly in all of the examples given above the ligands inhibited the enzymes. The pursuit of small molecules that activate enzymes is much less common as is the number of therapeutic agents that increase the activity of enzyme molecules.

In the realm of heme oxygenases (HO), much of the progress made toward understanding their functional roles has exploited animals that are genetically deficient in either HO-1 or HO-2, treatments that induce HO-1, or drugs that inhibit these enzymes [2,3]. Through utilization of these tools, we now appreciate that heme oxygenases and their products are involved in an interesting array of cellular activities. For example, considerable evidence revealed that HO-1 affords tissue protection in the vasculature due to the antioxidant, anti-inflammatory and anti-apoptotic properties of its products (see review by Araujo et al. [4]). Similarly,
there is substantial evidence that HO-1 protects neurons against oxidative stress [5]. The first generation of HO inhibitors comprises the metalloporphyrins, such as zinc protoporphyrin (ZnPP) and tin protoporphyrin (SnPP), which are powerful inhibitors of both HO-1 and HO-2 [6]. With respect to the cardiovascular system, Araujo et al. [4] have reviewed the evidence showing that HO-1 is protective against vascular inflammation, and cite examples such as the worsening of ischemia reperfusion injury in the presence of ZnPP [7]. Our laboratory has subsequently created a series of azole-based HO inhibitors, many of which are selective for HO-1 [8,9] and more recently a series of compounds that are selective for HO-2 [10]. As was the case for research addressing enzymes as drug targets in general, virtually all of the literature on small molecules that affect heme oxygenases was devoted to inhibitors of these enzymes. An exception to this was our recent report on the in vitro activation of HO-2 by menadione [11]. Menadione activated both the rat, native HO-2 and recombinant human HO-2 seven to thirty-fold, and did not activate HO-1. The mechanism of activation of HO-2 was through an increase in maximum velocity and not by a change in affinity for the substrate.

In this communication, we present some structure–activity relationships of menadione and its analogs as activators of HO-2, and test the hypothesis that redox properties of these compounds participate in their activation of HO-2.

Methods

Native HO-1 and HO-2 were prepared as microsomes from rat spleen and brain as described previously by Vukomanovic et al. [11]. All animals used were cared for in accordance with the principles and guidelines of the Canadian Council on Animal Care and the experimental protocols were approved by the Queen’s University Animal Care Committee. A full-length cDNA clone of full-length human HO-2 (FL-hHO-2) in pOTB7 was obtained from Open Biosystems. Following PCR amplification to engineer Ndel and EcoRI sites at the 5’ and 3’ ends, respectively, the coding region was subcloned into the pET28a vector using these restriction sites. The resultant recombinant protein contained an N-terminal histidine tag to allow purification by metal chelation using a protocol modified from previous publications [12,13]. Briefly, BL21 (DE3) cells, transformed with the FL-hHO-2/pET28a plasmid, were grown in LB supplemented with 30 μg/mL of kanamycin at 37°C until an OD_600 of ~0.8, at which point protein expression was induced with 1 mM isopropyl β-D-1-thiogalactopyranoside. Cells were grown for a further 3–4 h before harvesting and cell pellets were stored at ~80°C until ready to purify. Subsequently, thawed pellets were resuspended in lysis buffer [50 mM NaH_2PO_4 (pH 8.0), 300 mM NaCl, lysozyme (0.5 mg/mL; BioShop), DNasel (5 units/mL; Fermentas), EDTA-free protease inhibitor cocktail (1 tablet/50 mL; Roche)], incubated on ice (30 min) and lysed by sonication. Following centrifugation at 27,000 × g (15,000 rpm in a Beckman JA 25.5 rotor), the resultant pellets were solubilized in lysis buffer containing 1% n-Dodecyl β-D-Maltopyranoside (Affymetrix) overnight at 4°C. Solubilized protein was collected from the supernatant following centrifugation at 31,000 × g (16,000 rpm in a Beckman JA 25.5 rotor) and combined with the previously clarified lystate for purification by nickel affinity chromatography using Ni-NTA agarose resin. The combined supernatants were allowed to bind to the resin by rocking for 1 h at 4°C. After collecting the flowthrough, unbound protein was removed by a step-wise imidazole gradient (0–50 mM); protein was eluted by an addition step-wise gradient from 100–500 mM imidazole. Protein-containing fractions were identified by 12% SDS-PAGE and positive fractions (200–500 mM) dialyzed overnight against 20 mM potassium phosphate (pH 7.4), 0.1 mM EDTA, 10% glycerol. Following protein quantification (Bio-Rad), heme was conjugated by slow addition of hemin solution to a final molar ratio of 2:1 as described previously by Rahman et al. [13]. Further purification of the heme-bound enzyme and removal of excess heme was performed by size-exclusion chromatography over an S200 column. Protein concentration was determined by absorbance (ε_405 = 171.4 ± 1.2 mM⁻¹ cm⁻¹) [13]. Purity was assessed by measurement of the Rz ratio (A_405/A_280 > 2.1) and by SDS-PAGE analysis.

HO-1 and HO-2 activities were determined as described by Vukomanovic et al. [11]. In brief, a reaction mixture (150 μL) containing 100 mM phosphate buffer (pH 7.4), 50 μM methemalbumin and 0.5 mg mL⁻¹ brain or 1 mg mL⁻¹ spleen protein was pre-incubated with an HO activator, concentration as described below, for 10 min at 37°C. Methemalbumin was prepared by dissolving hemin in 10% (w/v) ethanolamine in water, and mixing it with a bovine serum albumin that had been dissolved in water [14,15]. The reaction was initiated by adding 1 mM NADPH and was continued for 15 min at 37°C. The reaction was stopped by instantly freezing the reaction mixture on dry ice, and the generated CO was determined by gas chromatography. For experiments involving FL-hHO-2, the concentration of the enzyme was 0.7 μM and the concentration of purified human NADPH-P450 reductase (recombiant, Becton Dickinson Canada Inc., Toronto, ON, Canada) was 0.01 μM.

Source of drugs

Drugs and compounds were obtained from the following sources: Sigma-Aldrich (Toronto, ON, Canada): menadione, vitamin K₂, NADPH, bovine serum albumin, hemin and L-ascorbic acid; Alfa Aesar- (A Johnson
Matthey, Ward Hill, MA, USA)- vitamin K1; Tokio Chemical Industry America (Portland, OR, USA)- hydroquinone, 1,4-benzoquinone, and 1,4-naphthoquinone (1,4-naphthalenedione); Acros Organics (Fair Lawn, NJ, USA)- alpha-tetralone, 1,4-cyclohexanediol; Ryan Scientific (Mt. Pleasant, SC, USA)- pentafluoromenadione; Otava Ltd. (Toronto, ON, Canada)- furan-containing naphthoquinones, furan-benzoxazine containing naphthoquinones, 2-aminophenyl naphthoquinones and 3-(piperidin-1-yl)naphtoquinones. 2-Phenylnaphthoquinones were obtained from Snieckus Innovations (Kingston, ON, Canada). 1,4-Dimethoxy-2-methylnaphthalene (methoxy analog of menadione) was a gift from Dr. D. N. Criddle, University of Liverpool, UK.

**Results**

An important biological property of menadione is its activity as a K vitamin; thus, we were interested to know

![Figure 1](http://www.medicalgasresearch.com/content/4/1/4)

**Figure 1** Lack of effect of vitamins K1 and K2 on HO-1 and HO-2 activity *in vitro*. HO-1 and HO-2 activity were measured as described in Methods using spleen (solid triangles) and brain (open circles) microsomes, respectively. The abscissa shows drug concentration and ordinate shows HO-2 activity as a percent of control in the absence of added drug. Panel a, Vitamin K1 and panel b, Vitamin K2. The structures of vitamins K1 and K2 in the panels to illustrate their differences from menadione.
if vitamin K1 or vitamin K2 also activated HO-2. In contrast to menadione, neither vitamin K1 nor vitamin K2 was able to increase the enzymatic activity of either HO-1 or HO-2 as shown in Figure 1. Thus, the HO-1 and HO-2 activity curves for vitamin K1 (panel a) were flat from 10 nM to 10 μM with a small upward trend at 100 μM for brain microsomal HO-2, and the curves for vitamin K2 (panel b) were flat throughout the full concentration range. In comparison, menadione activated HO-2 about 7-fold (Figure 2). Structurally menadione differs from vitamins K1 and K2 in that it possesses a methyl moiety at the 2-position whereas vitamins K1 and K2 each possess an additional, adjacent long unsaturated aliphatic side chain (Figure 1). Because this side chain could possibly interfere with binding of vitamin K1 or K2 to HO-2 through steric hindrance, we decided to explore the activity of other menadione analogs that possessed large non-aliphatic substituents at positions -2 and -3.

For these experiments, we tested menadione analogs that possessed various substituents at the 2- and 3-positions including naphthoquinones containing furan-carboxylic acid substituents (Table 1), naphthoquinones containing furan-benzoxazine (Table 2), naphthoquinones containing aminophenyl and piperidinyl moieties (Table 3), naphthoquinones containing 2-phenyl (Table 4) and other naphthoquinones containing various substituents at positions -2 and -3 (Table 5). Tables 1, 2, 3 and 4 show activation of rat brain HO-2 by menadione analogs possessing a wide variety of substituents at the 2- and 3-position. For example, compound 71830417 (a furan-containing naphthoquinone) activated HO-2 to 693% of control, SI-859 (a 2-phenynaphthoquinone) activated HO-2 to 691%, and 7010100237 (a 2-(aminophenyl)-3-piperidin-1-yl)naphthoquinone) activated HO-2 to 249% of control. Clearly, a variety of moieties could be present at the 2- and 3-positions of naphthoquinone, with maintenance of the ability to activate HO-2. Thus, unlike vitamins K1 and K2, the mere presence of a large substituent at the 2- and 3-positions in these compounds did not render them unable to activate HO-2.

**Mechanism of action**

According to classical receptor theory, an agonist drug such as menadione possesses affinity for its receptor on HO-2 and efficacy once it has bound. The inactivity of vitamin K1 might be explained by an inability to bind to HO-2. To explore this, we tested the effects of menadione in the presence of vitamin K1. This experiment was done in two ways- by adding vitamin K1 (10 – 100 μM) to the reaction mixture before menadione (25 μM) or after menadione. Both iterations yielded no effect on menadione activation of HO-2; vitamin K1 did not alter the menadione-induced activation of HO-2 (data not shown).

An obvious characteristic of menadione and its analogs studied herein is their naphthoquinone core, which is known to participate in redox reactions [16]. Consequently,
we obtained four menadione analogs in which redox activity was compromised to determine if there was a direct correlation between redox properties and activation of HO-2. One such analog was 1,4-dimethoxy-2-methylnaphthalene wherein the keto oxygens of menadione are replaced by methoxy groups; this analog did not activate HO-2 as shown in Figure 2. Another analog used was pentafluoromenadione, which would be compromised in redox reactions due to the strong electron-withdrawing action of the five fluorine atoms; pentafluoromenadione was without HO-2 stimulating activity (Figure 2). Table 6 shows the lack of HO-2 activation by naphthoquinones containing furan-carboxylic acid substituents.

**Table 1** HO-2 activation by naphthoquinones containing furan-carboxylic acid substituents

| Structure | HO-2 % Act. | HO-1 % Act. | Structure | HO-2 % Act. | HO-1 % Act. |
|-----------|-------------|-------------|-----------|-------------|-------------|
| ![Structure 1](image1) | 276 | 96 | ![Structure 2](image2) | 153 | 93 |
| ![Structure 3](image3) | 288 | 85 | ![Structure 4](image4) | 320 | 91 |
| ![Structure 5](image5) | 326 | 78 | ![Structure 6](image6) | 578 | 72 |
| ![Structure 7](image7) | 368 | 80 | ![Structure 8](image8) | 508 | 89 |
| ![Structure 9](image9) | 457 | 66 | ![Structure 10](image10) | 540 | 100 |
| ![Structure 11](image11) | 548 | 79 | ![Structure 12](image12) | 533 | 86 |
| ![Structure 13](image13) | 566 | 79 | ![Structure 14](image14) | 560 | 85 |
| ![Structure 15](image15) | 586 | 84 | ![Structure 16](image16) | 602 | 91 |
| ![Structure 17](image17) | 693 | 71 | ![Structure 18](image18) | 718340381 | 91 |

Rat HO-1 and HO-2 activities were determined as described in Methods. Test compounds were used at a concentration of 100 μM. Activity is reported as a percentage of activity in the absence of test compound; control = 100%. The 10-digit numbers are Otava catalog numbers.
Table 2 HO-2 activation by naphthoquinones containing furan-benzoxazine moieties

| Structure | HO-2 % Act. | HO-1 % Act. | Structure | HO-2 % Act. | HO-1 % Act. |
|-----------|-------------|-------------|-----------|-------------|-------------|
| ![Structure](image1) | 165         | 32          | ![Structure](image2) | 178         | 51          |
| ![Structure](image3) | 170         | 29          | ![Structure](image4) | 173         | 69          |
| ![Structure](image5) | 179         | 28          | ![Structure](image6) | 345         | 85          |

Rat HO-1 and HO-2 activities were determined as described in Methods. Test compounds were used at a concentration of 100 μM. Activity is reported as a percentage of activity in the absence of test compound; control = 100%. The 10-digit numbers are Otava catalog numbers.

Table 3 HO-2 activation by naphthoquinones containing aminophenyl-piperidinyl substituents

| Structure | HO-2 % Act. | HO-1 % Act. | Structure | HO-2 % Act. | HO-1 % Act. |
|-----------|-------------|-------------|-----------|-------------|-------------|
| ![Structure](image7) | 175         | 81          | ![Structure](image8) | 249         | 82          |
| ![Structure](image9) | 192         | 71          | ![Structure](image10) | 171         | 82          |

Rat HO-1 and HO-2 activities were determined as described in Methods. Test compounds were used at a concentration of 100 μM. Activity is reported as a percentage of activity in the absence of test compound; control = 100%. The 10-digit numbers are Otava catalog numbers.
toward HO-1 and HO-2. 1,4-Naphthoquinone had little effect on HO-1, but activated HO-2 five-fold.

**Menadione activation of recombinant human HO-2**

Our drug design work has relied heavily on the rat, native HOs, in part because the native enzymes were crucial in allowing us to create selective inhibitors of HO-1 [11]. Nevertheless, the ultimate application of knowledge to human health requires one to explore the novel effects of drugs in the human context. Furthermore, working with a purified recombinant protein sample provides insights at the molecular level that is often challenging when using a mixture. Accordingly, we tested the effects of menadione on recombinant human HO-2. Figure 3 shows that the FL-hHO-2, was activated ~7-fold by menadione, which compares well with menadione-activation of the rat, native HO-2 (also ~ 7-fold).

**Discussion**

The goals of this work were to explore the structure-activity relationship of menadione activation of HO-2 and to test the hypothesis that redox properties of menadione and its analogs participate in their activation of heme oxygenase-2. The main observations made in the present study were- (a) HO-2 was not activated by vitamins K1 and K2 but was activated by menadione (vitamin K3), (b) HO-2 was activated by furan carboxylic acid-containing naphthoquinones, furan-benzoxazine-containing naphthoquinones, 2-phenylnaphthoquinones, and 2-(aminophenyl)-3-(piperidin-1-yl)naphthoquinones, (c) the dimethoxy, pentafluoro and monohalogenated analogs of

### Table 4 HO-2 activation by naphthoquinones containing 2-phenyl moieties

| Structure | HO-2 | HO-1 | Structure | HO-2 | HO-1 |
|-----------|------|------|-----------|------|------|
| SI-859    | 691  | 81   | SI-2522   | 484  | 79   |
| SI-108    | 643  | 95   | SI-556    | 455  | 64   |
| SI-1606   | 596  | 84   | SI-1186   | 350  | 91   |
| SI-102    | 560  | 102  |           |      |      |

HO-1 and HO-2 activities were determined as described in Methods. Test compounds were used at a concentration of 100 μM. Activity is reported as a percentage of activity in the absence of test compound; control = 100%. MOMO is methoxymethyl ether. SI = Sniekus Innovations.
menadione did not activate HO-2, (d) 1,4-naphthoquinone strongly activated HO-2, and (e) full-length human recombinant HO-2 was activated to a similar extent as rat, native HO-2.

The absence of HO-2 activation by vitamins K1 and K2 indicates that the receptor responsible for HO-2 activation by menadione differs from the receptor for vitamin K activity. As stated above, this lack of HO-2 activation by vitamins K1 and K2 could have been due to their bulky aliphatic side chains sterically hindering binding to the enzyme or lack of inducing an appropriate activating conformational change of HO-2. The observation that addition of both menadione and vitamin K1 to the in vitro reaction mixture resulted in activation of HO-2 similar to that of menadione alone is consistent with the notion that vitamin K1 does not bind to the same site on HO-2 as does menadione. Had vitamin K1 bound to the menadione binding site, it would have antagonized the activating effect of menadione. Nevertheless, we acknowledge the fact that binding experiments with a radiolabelled ligand were not conducted.

Table 5 Lesser HO-2 activation by naphthoquinones containing various additional moieties

| Structure | HO-2 % Act. | HO-1 % Act. | Structure | HO-2 % Act. | HO-1 % Act. |
|-----------|-------------|-------------|-----------|-------------|-------------|
| ![Image](image1) | 107 | 99 | ![Image](image2) | 110 | 90 |
| ![Image](image3) | 108 | 100 | ![Image](image4) | 95 | 76 |
| ![Image](image5) | 105 | 97 | ![Image](image6) | 127 | 73 |
| ![Image](image7) | 113 | 93 | ![Image](image8) | 120 | 90 |
| ![Image](image9) | 102 | 95 | ![Image](image10) | 92 | 17 |
| ![Image](image11) | 119 | 74 | ![Image](image12) | 72071007 |

HO-1 and HO-2 activities were determined as described in Methods. Test compounds were used at a concentration of 100 μM. Activity is reported as a percentage of activity in the absence of test compound; control = 100%. The 10-digit numbers are Otava catalog numbers.
This inability of vitamin K\textsubscript{1} to bind to the menadione binding site on HO-2 does not appear to be due to steric hindrance alone because the side chains attached to the compounds listed in Tables 1, 2, 3, 4 are similar in size to the aliphatic side chains of vitamin K\textsubscript{1}. While these compounds are similar to vitamins K\textsubscript{1} and K\textsubscript{2} with respect to the presence of a significantly large moiety at carbon-2, they differ substantially in the nature of the substituent; these compounds contain hetero-atoms that would increase polarity. If there are polar sites on HO-2 in the region of the menadione binding site, they might be compatible with heteroatom-containing side chains but incompatible with non-polar moieties such as the aliphatic side chain of vitamin K\textsubscript{1}.

The hypothesis that redox properties of menadione and its analogs participate in their activation of HO-2 arose from the well-known redox properties of the quinone nucleus [16]. Our observation that the dimethoxy...
analog of menadione (1,4-dimethoxy-2-methylnaphthalene) was unable to activate HO-2 is consistent with the hypothesis. In the case of 1,4-dimethoxy-2-methylnaphthalene, the absence of the quinone nucleus would block the ability of this molecule to participate in redox reactions [17]. In the case of pentafluoromenadione, the presence of the five fluorine atoms causes a large change in the behaviour of electrons (negative inductive effect) of the naphthoquinone nucleus; accordingly, the reduction potential of pentafluoromenadione would be in the order of 400 mV compared with ~203 mV for menadione [18]. This reduction potential for the pentafluorinated analog of menadione would obviate redox function. Further support for this line of reasoning may be found in the lack of HO-2 activation by the two monohalogenated, monohydroxy naphthoquinone analogs shown in Table 6, which were found not to activate either HO-2 or HO-1. The hydroxyl groups on the latter two molecules could also contribute to this loss of HO-2 activation through resonance-based decreases in redox reactions. The HO-2 data obtained with these four compounds are consistent with the hypothesis as articulated above.

Another approach taken was to test the effects of five commercially available benzoquinone-like and naphthoquinone-like compounds on HO activation as shown in Table 7. Based on their chemical structures, we considered the hypothesis to predict that α-tetralone would be a poor activator of HO-2 because it would not be active with respect to redox activity. Similarly, 1,4-cyclohexanediione and hydroquinone were predicted to be inactive as HO-2 activators. The hypothesis predicted that 1,4-naphthoquinone would be fully active as an HO-2 activator because it possesses the complete quinone nucleus that is essential for redox reactions. While 1,4-benzoquinone shares a quinone nucleus with 1,4-naphthoquinone, it was anticipated to be inactive because it has been observed not to participate in redox reactions but rather it arylates protein thiols [19]. Thus, the data obtained from these five compounds are consistent with the hypothesis articulated above. On balance, it appears that HO-2 activating naphthoquinones require both affinity for a selective receptor and redox activity.

The observations that these naphthoquinones activated HO-2 but did not activate HO-1 are interesting from both mechanistic and application perspectives; certainly they should be useful for elucidating the roles of HO-1 and HO-2 in various physiological and metabolic functions. This observation also indicates that these HO-2 activators bind to a site on HO-2 other than the catalytic site because of the high conservation of amino acid sequence between HO-1 and HO-2 therein. We have excluded cytochrome P450 reductase as the activation site by observing that hHO-2 (1–288) activation occurred (>200%) in the absence of NADPH and cytochrome P450 reductase whereby the source of reducing capacity was ascorbic acid (data not shown). In the present experiments, the source of reducing equivalents would include NADPH. In the case of enzyme inhibitors the locus of drug binding is often at or near the active site, as would be the case for competitive inhibitors. In the case of enzyme activators, the binding site should not be the same as that of the natural substrate (i.e. active site), because binding of a non-substrate molecule there must result in competitive inhibition. It follows that an enzyme activator must bind beyond the active site, but still influence the reaction in a positive manner.

Conclusions
The activation of HO-2 by menadione and similar agonists requires affinity for a receptor site outside of the enzyme core (which it shares with HO-1) and probably also the ability to participate in redox reactions.

Abbreviations
HO: Heme oxygenase; HO-1: Heme oxygenase-1; HO-2: Heme oxygenase-2; hHO-2: Recombinant human heme oxygenase-2; ZnPP: Zinc protoporphyrin; SnPP: Tin protoporphyrin.

Competing interests
None declared for Vukomanovic, Rahman, Brien and Jia. Nakatsu and Szarek hold a patent on selective heme oxygenase-1 inhibitors. Bilokin and Golub are affiliated with Otava.

Authors’ contributions
Participated in research design: N, V. Conducted experiments: V, R. Contributed new reagents or analytic tools: B, G. Performed data analysis: N, V. Wrote or contributed to the writing of the manuscript: N, V, B, J, S. All authors read and approved the final manuscript.

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