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An LC-MS/MS analysis of opiate residues on Thomas Chatterton's (1752–1770) memorandum book – Did he die from a laudanum overdose?†

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250 years after his death, Thomas Chatterton continues to create discussion amongst the literati and enchant the people of Bristol. The controversy of his life was entwined with his writings where he passed his work off as that of a fictional medieval poet – Thomas Rowley. His premature death at the age of 17 in 1770 is also shrouded in controversy – did he commit suicide from arsenic poisoning (as stated at the original inquest into his death), or did he accidentally overdose on laudanum (as suggested by the 1947 forensic analysis)? The purpose of this study is to use state-of-art analytical methods (namely ultra-high-performance liquid chromatography tandem mass spectrometry using an Orbitrap mass spectrometer) to investigate the brown stain found on his memorandum book. The conclusion of this study is that that stain is made up of, amongst other things, 18 opiate and 1 opioid degradation product – 9 of which are previously unpublished, and that the spillage was indeed due to laudanum.

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Introduction

The life, work and death of Thomas Chatterton 1752–1770 (a life span of only 17 years 277 days) has always been shrouded in myth and mystery. Born to a widow in Redcliffe, Bristol, Chatterton was expelled from school at the age of six for being a dullard, incapable of improvement.^{1,2} As a result, he was essentially self-taught. He wrote mock medieval poems in the guise of a 15th Century Bristol priest called Thomas Rowley that confounded the literati of the day. A furore raged over their authenticity for many years after his death, with some still claiming that they were really written by Rowley as late as the 1850s.^{1,2} Chatterton left Bristol in April 1770 and after residing at Shoreditch in London he moved to Holborn where he tragically died on 24th August 1770.

There are conflicting stories surrounding the events leading up to Chatterton's death. A neighbour stated that Chatterton's landlady told her that on 24th August she knew that he had not eaten anything for two or three days. She had begged that he would “take some dinner with her”; but he was offended at her expressions, “which seemed to hint he was in want” and assured her that he was not hungry. Another account was that of a baker, from whom Chatterton obtained his bread, had

refused to trust him with an additional loaf, so he immediately went home and committed suicide.^{1,2}

The Coroner had no minutes of the Chatterton inquest, and could not recall the circumstances, however did produce a “memorandum”, the finding of which stated that Thomas Chatterton “swallowed arsenick in water, on the 24th August 1770; and died, in consequence thereof, the next day”.^{3,4}

Amongst Chatterton's belongings was a small memorandum book (see Fig. 1) in which he kept his accounts. This book had a brownish stain running through eighteen or so pages. In February 1947, the South-Western Forensic Science Laboratory, Bristol, studied the stained paper and concluded: “It gives a positive reaction for opium alkaloids (i.e. morphine, etc.). The stain was doubtless, therefore, produced by tincture of opium (laudanum)”.⁵

It is now considered likely that Thomas Chatterton did not commit suicide but perhaps died of an accidental overdose of his medicinal treatment. Chatterton showed no signs, in the final four months of his life in London, of the introspective melancholy that often accompanies suicide. Further, contrary to myth, he was financially secure at the time of his death. He may have been taking laudanum (opium tincture) as a sedative. Historic laudanum is a “whole opium” alcoholic extract containing all the main opiates with morphine being at the highest concentration, followed by codeine and thebaine (see Table 1).

Herein, due to the continued interest in the life and work of Thomas Chatterton (it also being the 250th anniversary of his death) and as a result of the considerable advancements in

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Fig. 1 A photograph of Chatterton's memorandum book clearly showing the brown stain running through several pages. The paper sample analysed in this study was from a blank (unprinted) page with the same stain.

Table 1 The structures, formulae and molecular weights (M_w) of the opiate alkaloids: morphine, codeine and thebaine found in laudanum (opium tincture)

Opiate	R_1	R_2	Formula	M_w
Morphine	OH	OH	$C_{17}H_{19}NO_3$	285
Codeine	OMe	OH	$C_{18}H_{21}NO_3$	299
Thebaine	See right hand side		$C_{19}H_{21}NO_3$	311

analytical methodologies and technologies since the original forensic investigation of 1947, is contained an up-to-date study of the stain in Chatterton's memorandum book. This study makes use of Ultra performance liquid chromatography (UPLC) coupled to ultra-high-resolution tandem mass spectrometry (MS/MS).

Experimental

A 2 mm² section of unprinted paper from Chatterton's memorandum book, having the same brown stain as seen in Fig. 1 was obtained in person from Bristol Central Library. This section of paper was cut up with a scalpel and soaked in approximately 1 mL of HPLC grade water. This was left

untouched for about 10 minutes. The solution was then pipetted into an LC vial for storage and analysis. A sample of pure codeine was obtained *via* cold water extraction⁶ from a pharmaceutical grade co-codamol tablet obtained from a local pharmacy.

Direct infusion screening mass spectra were recorded by nanospray on a Synapt G2S (Waters, Manchester, UK) fitted with a Triversa Nanomate chip-based nanospray system (Advion Biosciences, Norwich, UK). Samples were diluted in methanol:water (50:50) and 5 μ L aspirated through a nanomate 400 chip by the nanospray robot at 1.5 kV nanospray voltage, and 1.5 psi gas pressure. The Synapt's cone temperature was set at 25 °C. Nanospray-MS/MS spectra were recorded by collision induced dissociation (CID) at 30 eV on isolated precursor ions at a resolution of approximately 30 000. Reverse phase UPLC-MS/MS analyses were performed on an Orbitrap Elite mass spectrometer equipped with a Dionex Ultimate 3000 UPLC system (Thermo Scientific, Hemel Hempstead, UK). The column was a Waters Acuity UPLC C18 (2.10 \times 100 mm, 1.7 μ m ID) obtained from Phenomenex (Macclesfield, UK). Solvent A was water + formic acid (1%), solvent B was acetonitrile + formic acid (1%). The 30 minute gradient was 5% B to 95% B over 26 minutes, followed by 2 minutes hold and then return to initial conditions over 2 minutes. Mass spectra were recorded using a heated electrospray ionisation source (HESI) at a resolution of 120 000. A source voltage of 3 kV, source temperature of 350 °C and a capillary temperature of 275 °C were used. High purity (oxygen free) N₂ was used for both the drying and CID gas. Tandem mass spectra (MS/MS) were recorded during the UPLC analysis on isolated precursor ions (1 Da window) using CID in the Ion Tap of the Orbitrap instrument at 30 eV normalised collision energy. Centred MS/MS spectra were recorded for every ion in the expected mass range for opium degradation products. Direct infusion MS/MS was also performed on the Orbitrap using the higher energy collisional dissociation (HCD) mode to enable the generation of an increased number of sub fragmentations. All Orbitrap analyses were recorded at a resolution of 120 000. For all MS analysis, the solvents used were HPLC gradient grade obtained from Fisher Scientific (Hemel Hempstead, UK).

Results and discussion

After leaving the stained paper sample in water for 10 minutes it was seen that a slight brown discolouration occurred, suggesting that something had gone into solution. Fig. 2 shows the mass spectrum resulting from a direct infusion analysis of a 10% diluted sample in methanol:water (50%). It is clear from this initial analysis that there are many compounds extracted from the paper into the water – this is totally to be expected from a sample of this age. The peaks with the masses labelled in green either match to known opiate degradation products identified in separate studies by Small and Rapaport^{7,8} or to proposed new degradation products. The two ions labelled in red (m/z 301 and 317) are due to the $[M + Na]^+$



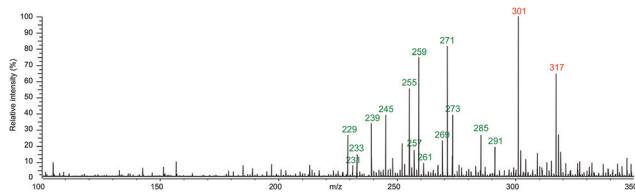


Fig. 2 The direct infusion Nanospray-MS analysis of the paper extract. Peaks labelled in green correspond to the masses believed to be due to opiate degradation products, those in red are the $[M + Na]^+$ and $[M + K]^+$ of the dibutyl phthalate impurity.

and $[M + K]^+$ of dibutyl phthalate, respectively. The presence of the phthalate contaminant peaks is not unexpected as the book was stored in an air-tight plastic bag and both the book and paper samples were handled with plastic gloves. These peaks will be ignored in all further analysis and were excluded from the UPLC-MS/MS analysis. This spectrum indicates that a wide range of possible degraded opiates were present in the paper extract and to obtain full structural information on them, a systematic analysis by UPLC-MS/MS was performed.

A 10% diluted sample of the paper extract was then subjected to a systematic UPLC-HESI-MS/MS analysis with every peak from m/z 220 to 300 fragmented by CID-MS/MS. Opiates have very characteristic fragmentation spectra with groups of peaks clustered by 2 m/z .^{9–13} Fig. 3 shows MS/MS spectra of a pure sample of codeine recorded under a range of conditions. Spectrum (a) is the Nanospray-CID-MS/MS obtained on the Synapt instrument, spectrum (b) is the HESI-CID-MS/MS and spectrum (c) is the HESI-HCD-MS/MS both obtained on the Orbitrap instrument. The pattern of peaks is due to the multitude of possible competing fragment routes which result in product ions that are different in mass by 2 Da. This effect is also apparent in the direct infusion spectrum in Fig. 2, where

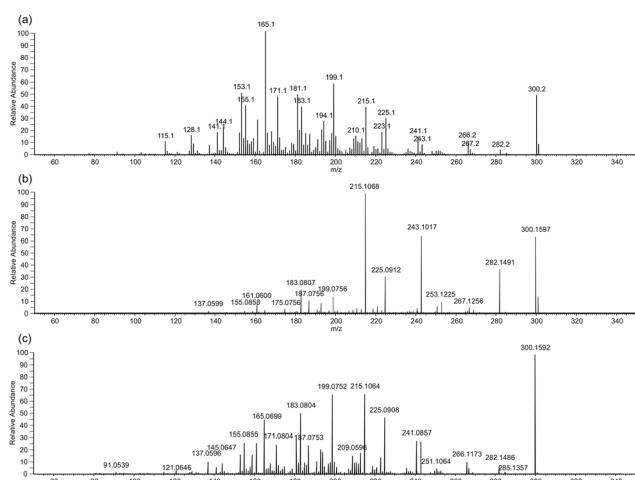


Fig. 3 MS/MS spectra of pure codeine, precursor ion $[M + H]^+$ is m/z 300. Spectrum (a) is the Nanospray-CID-MS/MS obtained on a Synapt instrument, spectrum (b) is the HESI-CID-MS/MS and spectrum (c) is the HESI-HCD-MS/MS both obtained on an Orbitrap instrument. All spectra are recorded at 30 eV collision energy.

a range of degraded opiates also cluster with masses different by 2 Da.

Fig. 3 clearly shows the need for different collision modes in MS/MS. Spectrum (a), is a classic Q-ToF type CID-MS/MS spectrum where the multiple fragmentation routes and sub fragmentations are fully accessible creating the classic opiate MS/MS spectrum with clusters of peaks with 2 m/z spacings. Spectrum (b), is the CID-MS/MS run on an Orbitrap instrument which demonstrates how ion-traps can access only a limited number of fragmentations – mainly it is the sub fragmentations that are missing from this spectrum. However, in spectrum (c) the HCD collision mode is used, and this demonstrates that the sub fragmentation routes can still be accessed by using the alternative collision mode. With opiates, the increased stability is bought about by the bridging *N*-methyl ethanamine. This moiety has been lost in the degraded opiates as demonstrated by the Rapport in 1948⁸ and more recently by Carlin *et al.* in 2017.¹⁴ The result of this is that this enables an increased amount of fragmentation to be observed in the CID-MS/MS mode when compared to the undegraded opiates.

Fig. 4 shows the UPLC-HESI-MS chromatogram of the paper extract. Chromatogram (a) is the base peak chromatogram (BPC) of the whole extract whereas chromatogram (b) is the extracted ion chromatogram (EIC) of the mass range of interest (m/z 220–300). The BPC shows the total chemical make-up of the extract in the full mass range from m/z 50–1000. More polar components will elute at the lower retention times with the hydrophobic components making up the bulk of chromatogram after 20 min. The EIC shows the same chromatogram as (a) but only showing masses in the range from m/z 220–300 which covers the mass range of interest for degraded opiates. This clearly demonstrates how the EIC is far more useful than

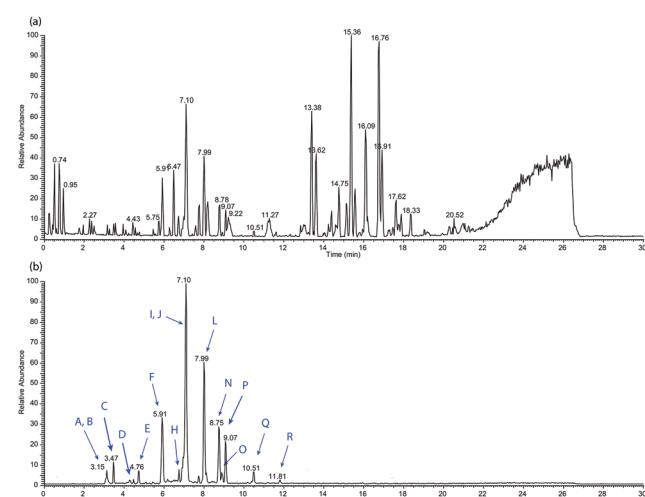


Fig. 4 UPLC-HESI-MS of the paper extract. Chromatogram (a) is the base peak chromatogram (BPC) and chromatogram (b) is the extracted ion chromatogram (EIC) for the region of interest (m/z range 220 to 300). Peaks labelled with letters are the identified laudanum degradation peaks (degradation of either morphine, codeine or thebaine).



Table 2 Table showing degraded opiate identity (see Fig. 4), retention time, relative peak area (peak I at 7.10 min is the base peak set to 100%), molecular formula, opiate origin, theoretical isotopic mass, measured accurate mass with mass measurement error (ppm). For the origin, Mor = Morphine, Cod = Codeine and The = Thebaine

ID	Retention time (mins)	Peak area (%)	Molecular formula	Origin	Theoretical mass $[M + H]^+$	Measured mass $[M + H]^+$ (error in ppm)
A	3.09	1.8	$C_{16}H_{20}O_3$	Mor	261.1485	261.1486 (0.38)
B	3.15	4.0	$C_{16}H_{18}O_3$	Mor	259.1329	259.1327 (0.77)
C	3.47	8.0	$C_{14}H_{14}O_3$	Mor	231.1016	231.1013 (1.30)
D	4.49	0.9	$C_{17}H_{22}O_3$	Cod	275.1642	275.1644 (0.73)
E	4.73	3.0	$C_{16}H_{16}O_3$	Mor	257.1172	257.1169 (1.17)
F ^a	5.91	36.7	$C_{16}H_{14}O_2$	Cod	239.1067	239.1072 (2.09)
G	6.27	0.5	$C_{13}H_{18}O_3$	Mor	245.1172	245.1171 (0.41)
H	7.04	41.4	$C_{17}H_{20}O_3$	Cod	273.1485	273.1487 (0.73)
I	7.10	100	$C_{17}H_{18}O_3$	Cod	271.1329	271.1326 (1.11)
J	7.10	1.6	$C_{18}H_{22}O_3$	The	287.1642	287.1639 (1.04)
K	7.21	1.4	$C_{18}H_{18}O_3$	The	283.1329	283.1325 (1.41)
L	7.99	56.4	$C_{14}H_{12}O_3$	Mor	229.0859	229.0860 (0.44)
M	8.11	1.2	$C_{18}H_{20}O_3$	The	285.1485	285.1482 (1.05)
N	8.75	6.8	$C_{14}H_{16}O_3$	Mor	233.1172	233.1173 (0.43)
O	8.91	4.5	$C_{16}H_{14}O_3$	Mor	255.1016	255.1018 (0.78)
P	9.07	17.6	$C_{17}H_{20}O_3$	The	273.1485	273.1488 (1.10)
Q	10.51	6.6	$C_{18}H_{26}O_3$	The	291.1955	291.1953 (0.69)
R	11.81	1.2	$C_{17}H_{16}O_3$	Cod	269.1172	269.1164 (2.97)

^a Structure F could be a degradation product from either morphine or codeine.¹⁴

the BPC as it only shows the components of the extract which might be degraded opiates. The BPC will show everything extracted from the paper which is ionisable under the HESI conditions used. The EIC is labelled with letters from A to R which represent the possible opiate degradation products – see Table 2 for further details.

To identify the degradation products, tandem mass spectrometry (MS/MS) is performed. In this study, all ions in the mass range of interest were fragmented in the ion trap of the Orbitrap instrument as they eluted from the chromatographic system. In the 1940s, two independent studies of the degradation products of opiates were published^{7,8} – this is the starting point for the identification of the opiate degradation products found in the brown stain. Fig. 5 shows the structures of the degradation products identified or postulated in the previous studies along with their identification in this study.

In the early studies, nine opiate degradation products were generated or postulated from chemical degradation methods used for structural elucidation. It is the authors view that environmental degradation should result in some, if not all, of the same degraded structures. In fact, there are peaks corresponding to the mass and formulae of all nine species observed in this analysis and it is fair to assume that these are the species observed. However, to confirm their identities, a thorough and systematic study of their MS/MS spectra was performed. Fig. 6 shows the LC-MS/MS spectrum and proposed fragmentation scheme for degradation product O (m/z 254) postulated as existing by Rapaport.⁸ The ESI $^+$ shows similar spectra and schemes for the other eight known and postulated degradation products (B, E, H, I, J, K, M, R in Table 2). The schemes make reasonable deductions about the neutral losses occurring leading to a consistent set of competing fragmentation routes.

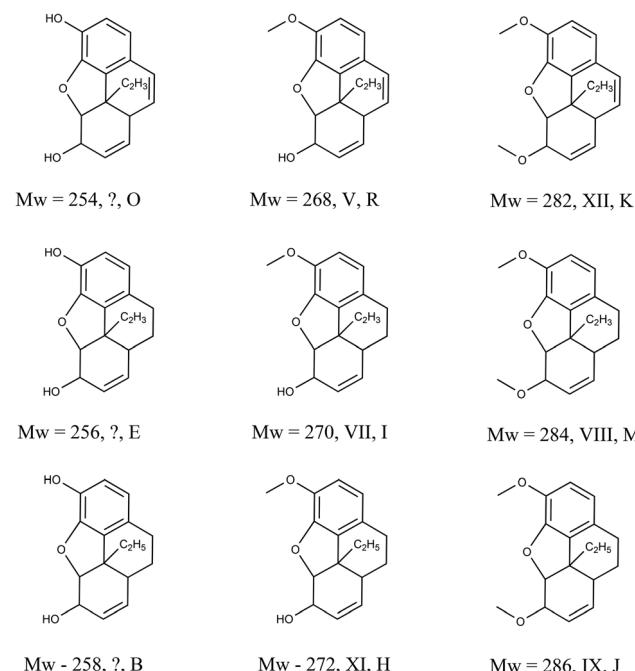


Fig. 5 Previously identified (and postulated) degradation products of morphine, codeine and thebaine and their identities. The Roman numeral represents the identity from the original work by Rapaport⁸ ('?' represents a postulated structure based on those elucidated for codeine and thebaine), the second letter corresponds to the identity used in this study (see Fig. 4 and Table 2).

For degradation product O, fragmentation can proceed *via* 3 competing initial losses – that of oxygen (mass 16), water (H_2O mass 18) or ethyne (C_2H_2 , mass 26) resulting in product ions m/z 239, 237 or 229 respectively. All these losses can occur



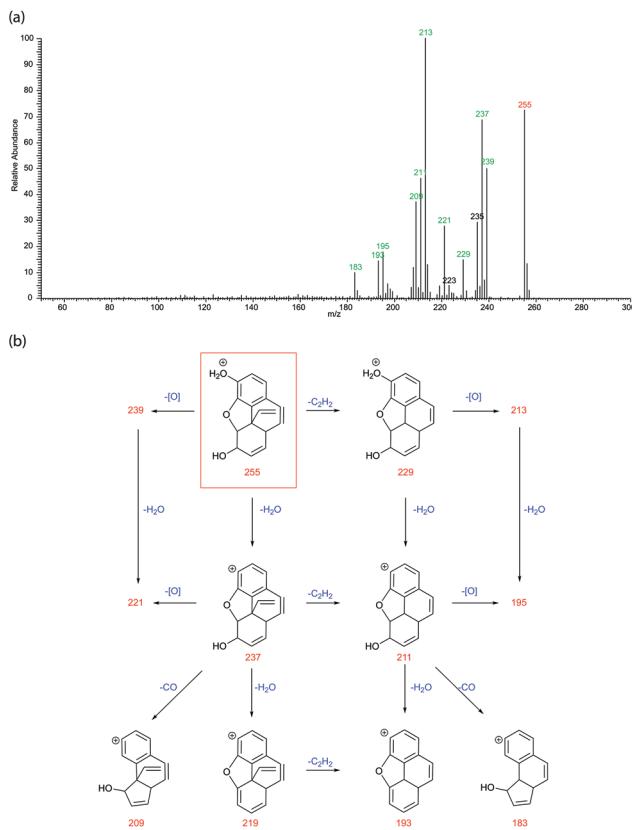


Fig. 6 Analysis of the previously identified degradation product O (m/z 255). (a) is the LC-MS/MS spectrum and (b) is the proposed fragmentation scheme – the precursor ion is boxed in red. Peaks labelled in green in the spectrum are those that are identified in the fragmentation scheme (b).

in any sequence and so all three of these initial product ions can then lose one of the other two neutrals to generate product ions m/z 221, 213 or 211. Product ions m/z 237 and 211 can also lose a molecule of carbon monoxide (CO, mass 28) or undergo another water loss to generate ions m/z 219, 209, 193 or 183. Product ion m/z 209 can then also lose ethyne to generate m/z 183 (not shown on the scheme for clarity). The loss of oxygen is quite unusual but is present in every sample in this study. This plethora of possible fragmentation routes and neutral losses leads to the characteristic MS/MS spectra with groups of peaks spaced by 2 m/z . All product ions have had their formulae confirmed by accurate-mass analysis to within 2 ppm maximum error (within 1 ppm for a vast majority of ions). Similar fragmentation schemes can be drawn up for all the previously identified degradation products and these can be found, along with their MS/MS in the ESI.†

As well as the nine previously identified degradation products, another ten were also observed. To propose structures for these new molecules, fragmentation schemes were drawn up based on those observed for the known products leading to plausible structures for the precursor ions. The fact that almost all product ions can be accounted for within these schemes leads us to have a high degree of confidence in their

structures, however these can only be proposals. A good example is for degradation product 'I' (m/z 271) from codeine. See Fig. 7 for the LC-MS/MS spectrum and proposed fragmentation scheme.

As degradation product I is from codeine, it has additional fragmentation routes due to the possible loss of formaldehyde (CH_2O , mass 30) from the methoxy group. Product I also has a saturated middle ring which enables the loss of 2 hydrogens to occur – this is a very characteristic loss for opiates^{10,11,13} and seen in this study for every instance where the middle ring is saturated. The precursor ion (m/z 271) can undergo losses of either 2 hydrogens, an oxygen, or a molecule of water to generate product ions m/z 269, 255 or 253 respectively. All these product ions can then also undergo one of the other losses to generate product ions m/z 253, 251 or 235. After the loss of water to generate m/z 253 or 2H and water to generate m/z 251, a loss of a molecule of carbon monoxide followed by formaldehyde occurs to generate product ions m/z 225, 223, 195 and 193. Product ions m/z 225 and 195 can also lose 2 hydrogens to generate m/z 223 and 193. An additional route is for the loss

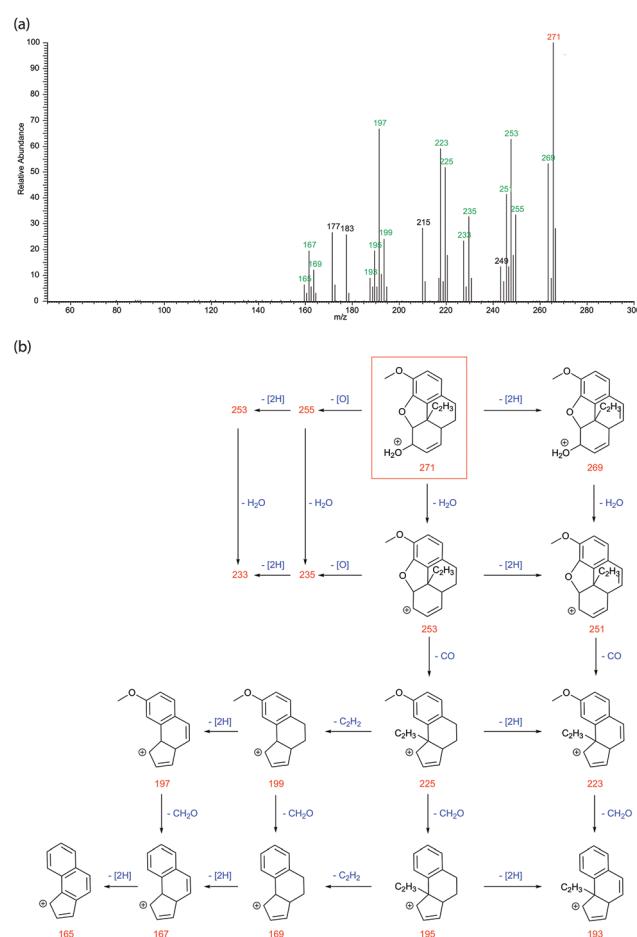


Fig. 7 Analysis of a previously unidentified degradation product I (m/z 271). (a) Is the LC-MS/MS spectrum and (b) is the proposed fragmentation scheme – the precursor ion is boxed in red. Peaks labelled in green in the spectrum (a) are those that are identified in the fragmentation scheme (b).

of ethyne (C_2H_2 , mass 26) from m/z 225 or 195 to generate m/z 199 or 169. These can then undergo additional losses of 2 hydrogens leading finally to the fully unsaturated product ion m/z 165.

Degradation products of thebaine (J, K, M, P and Q) all have 2 methoxy groups which makes one of these groups the likely site of protonation resulting in the competing losses of formaldehyde and methanol (CH_3OH , mass 32) in place of the losses of water. See ESI† for more details.

Degradation product 'F' is structurally similar to the named opioid morphenol and can result from the degradation of either morphine or codeine. The degradation route of codeine to morphenol is previously published¹⁵ and so F is tentatively also assigned as a degradation product of codeine. Fig. 8 is the LC-MS/MS spectrum and proposed fragmentation pathway for 'F'. The precursor ion (m/z 239) can undergo losses of either 2 hydrogens, oxygen, or a water molecule leading to product ions m/z 237, 223 and 221 respectively. In similar ways to the previous examples above, these three product ions can then undergo another neutral loss resulting in product ions m/z 219 or 205. Further losses of ethyne or ethene (C_2H_4 , mass 28) can then occur leading to product ions m/z 195, 193, 191 or 177.

The discovery of the morphenol analogue amongst the opiate degradation products lead on to the further examination

ation of the LC-MS/MS to see if any other opioids were also present. The EIC for m/z 209 – representing the $[M + H]^+$ of morphenol itself, generates a peak at 3.47 min co-eluting with degradation product 'C' but having a very different MS/MS spectrum (see ESI info Fig. S16†). Morphenol is not an opiate and is fully unsaturated and as a result has a relatively simple MS/MS spectrum consisting of only a water loss followed by aromatic like ring contractions (peaks spaced by CH_2). The existence of morphenol in the sample is not unexpected due to it being totally unsaturated preventing dehydration or any further environmental degradation. Morphenol was first proposed to be a degradation product of morphine in 1897¹⁶ and although is a very minor product here, it is clearly present amongst the other opiate degradation products.

Conclusions

The ultrahigh resolution UPLC-MS/MS analysis of the stain on Thomas Chatterton's memorandum book successfully identified 18 opiate and 1 opioid proposed degradation products from morphine, codeine or thebaine. The full unequivocal identification of the proposed degraded structures is not possible within the confines of this study. However, in a study published in the 1940s, Rapoport identified a series of chemical degradation products of morphine and codeine that had the *N*-methyl ethanamine moiety eliminated.⁸ A recent study in 2017 also showed similar results for the acid/base degradation of thebaine.¹⁴ Several of the structures proposed for the identity of the degradation products in this study match the masses and formulae of those observed previously. From there, a set of structures for all the other degradation products have been proposed based on their similar and highly characteristic MS/MS spectra.

The conclusion of this study is that the stain is most certainly the result of laudanum having been spilled. Laudanum was an expensive medicinal treatment in the Georgian era and any user would have been highly careful to not have accidental spilled it – especially on their memorandum book, which itself would have been a treasured possession. This would lead on to the speculation that Chatterton was self-medicating with Laudanum, took too much and spilled or knocked over the rest of the bottle as a result. Paregoric is another opiate medication which was widely taken at the time but was approximately 20 times more dilute than laudanum. It is quite possible that Chatterton mistook laudanum for paregoric – but this is purely speculation. The scientific evidence is overwhelming that the stain is the result of opiates, anything else is purely speculation, but the assertion that he committed suicide by arsenic poisoning should be in, the authors view, called into serious doubt.

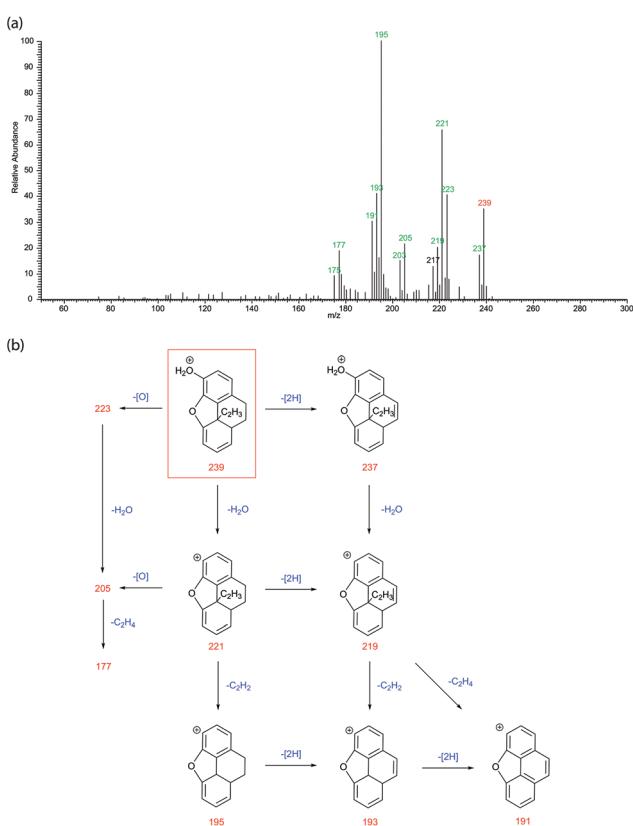


Fig. 8 Analysis of degradation product F (m/z 239). (a) Is the LC-MS/MS spectrum and (b) is the proposed fragmentation scheme – the precursor ion is boxed in red. Peaks labelled in green in the spectrum (a) are those that are identified in the fragmentation scheme (b)

Conflicts of interest

The authors declare no conflicts of interest.

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