

1 **Progress in Low Field Benchtop NMR Spectroscopy in Chemical and Biochemical Analysis**

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13 **Abstract**

14 The employment of spectroscopically-resolved NMR techniques as analytical probes have previously
15 been both prohibitively expensive and logistically challenging in view of the large sizes of high-field
16 facilities. However, with recent advances in the miniaturisation of magnetic resonance technology,
17 low-field, cryogen-free “benchtop” NMR instruments are seeing wider use. Indeed, these miniaturised
18 spectrometers are utilised in areas ranging from food and agricultural analyses through to human
19 biofluid assays and disease monitoring. Therefore, it is both intrinsically timely and important to
20 highlight current applications of this analytical strategy, and also provide an outlook for the future,
21 where this approach may be applied to a wider range of analytical problems, both qualitatively and
22 quantitatively.

23 **Keywords**

24 Compact NMR, mobile NMR, benchtop NMR, biofluid analysis

1

2 **Abbreviations**

3	1D	One dimensional
4	2D	Two dimensional
5	bNMR	Benchtop nuclear magnetic resonance
6	COSY	Correlation spectroscopy
7	CPMG	Carr-Purcell-Meiboom-Gill
8	DEPT	Distortionless enhancement by polarization transfer
9	DOSY	Diffusion ordered spectroscopy
10	DNP	Dynamic nuclear polarisation
11	FID	Free induction decay
12	GC	Gas chromatography
13	HETCOR	Heteronuclear correlation spectroscopy
14	HMBC	Heteronuclear multiple bond correlation
15	HP	Hyperpolarisation
16	HPLC	High performance liquid chromatography
17	HSQC	Heteronuclear single quantum correlation
18	IR	Infra-red
19	LF	Low field
20	LLOD	Lower limit of detection
21	LLOQ	Lower limit of quantification

1	MAS	Magic angle spinning
2	MHz	Megahertz
3	MOF	Metal-organic framework
4	MRI	Magnetic resonance imaging
5	NMR	Nuclear magnetic resonance
6	NOE	Nuclear Overhauser effect
7	NP	Nanoparticle
8	NUS	Non-uniform sampling
9	NV	Nitrogen vacancy
10	PAT	Process analytical test
11	PCA	Principal component analysis
12	PLS	Partial least squares
13	QC	Quality control
14	SABRE	Signal amplification reversible exchange
15	SNR	Signal-to-noise ratio
16	T	Tesla
17	TOCSY	Total correlated spectroscopy
18	TSP	(Trimethylsilyl)[-2,2,3,3-d4] propionic acid /sodium salt
19	UV	Ultraviolet
20		
21		

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13 **1. Introduction**

14

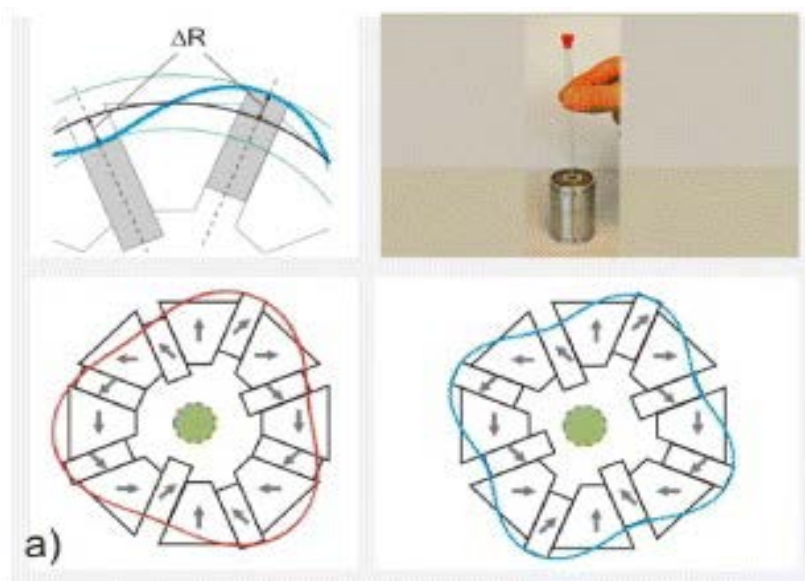
15 Nuclear magnetic resonance (NMR) spectroscopy has been widely applied in chemical and
 16 biochemical analysis for decades, with high-resolution instruments now possessing the necessary
 17 sensitivity to successfully probe large biological structures such as proteins [1]. Advances in the
 18 technology used to develop the very large, superconducting magnets employed for these high-
 19 resolution studies has led to successively increased spectral resolution, developments which have
 20 allowed us to consider even larger chemical systems. However, with these improvements in
 21 sensitivity and resolution come larger and more robust instruments requiring dedicated laboratories,
 22 with high purchase and operating costs, particularly those involving cryofluid replenishment for the
 23 latter; one major limitation of this approach is that such facilities are inherently inappropriate for field
 24 or benchtop studies [2,3], whilst being less sensitive to both charge effects in polyelectrolyte solutions
 25 and conducting samples. Indeed, although the advent of cryogenic probes has resulted in marked
 26 increases in sensitivity compared to earlier probeheads, this is only realised in practice if samples are
 27 in fact electrically insulating, such as in organic solvents. Cryogenic probes function by cooling the
 28 RF receiver coils to temperatures below 30 K. This lowers the resistance of the coils, thereby
 29 increasing signal amplitude and reducing thermal noise levels. The combination of these effects

1 results in an increase in the sensitivity, which is monitored by signal-to-noise ratio (SNR).
2 Conversely, conductive samples will therefore supplement the resistance to the RF receiver coil, and
3 hence decrease SNR values.

4 Commercial benchtop NMR spectrometers are most notably available from Thermofisher, Nanalysis,
5 Oxford Instruments, and Magritek, currently predominantly operating at 60 MHz (1.41 Tesla (T)) and
6 80 MHz (1.88 T) operating frequencies, with 190 MHz (4.45 T) Halbach-enhanced designs being
7 envisaged,[4] and 212 MHz (5 T) versions possible [5]. In very early experiments, NMR spectra were
8 recorded at only 40 or 60 MHz operating frequencies. However, current conventional high-field
9 spectrometer studies take place with magnets at 400 MHz or above, and therefore it is timely to
10 highlight the advantages of low-field NMR techniques [6].

11 Large NMR instruments based on permanent magnets and electromagnets were commonplace until
12 the 1970s, with the emergence of superconducting instruments; however, these were both expensive
13 and attained similar sizes to those of the superconducting instruments we see today. In order to
14 generate stable, sufficiently strong magnetic fields in instruments of manageable size for traditional
15 laboratories and benchtop work, small permanent magnets weighing approximately 500 grams, and
16 with homogeneous magnetic fields, were reported based on designs by Halbach [7] (Figure 1).
17 Considering that permanent magnets currently generate fields of up to 2 T, and that field strength
18 remains constant when magnet volume is scaled down, the observed sensitivity decrease would
19 approximate to a factor of 3 orders of magnitude when compared to the 7 T fields of instruments with
20 superconducting magnets [8]. Indeed, the construction and availability of these magnets, incorporated
21 within user-friendly, cryogen-free, low-field (LF) benchtop instruments, has led to an abundance of
22 recent applications of this technology [9].

23



1

2 **Figure 1.** Diagram from Ref. [10], originally published open access in the New Journal of Physics,
3 IOP. NMR magnets from permanent magnet materials. (a) Halbach magnet constructed from
4 trapezoidal magnet blocks with gaps between them in which magnetic plates are moved in and out in
5 order to shim the field, where ΔR is the shimming distance. Typical field strengths are 0.5 T for
6 imaging, and 1.0 T for spectroscopy.

7

8 Furthermore, with NMR spectroscopy representing one of the most widely used techniques in
9 analytical chemistry [11], the prospects are promising for small benchtop instruments to be applied in
10 areas ranging from medicine and clinical science through to food and environmental sciences. Herein,
11 we present a recent overview of the area of LF benchtop NMR spectroscopy (bNMR), and provide a
12 perspective on future applications. We also provide an update on the very recent novel employment of
13 LF bNMR technologies to the NMR-linked metabolomics area, and here we showcase the
14 applications of this technique to the diagnosis and metabolic monitoring of type 2 diabetes [12]. A
15 more extensive description of the fundamental principles of compact LF NMR can be found here [13–
16 15].

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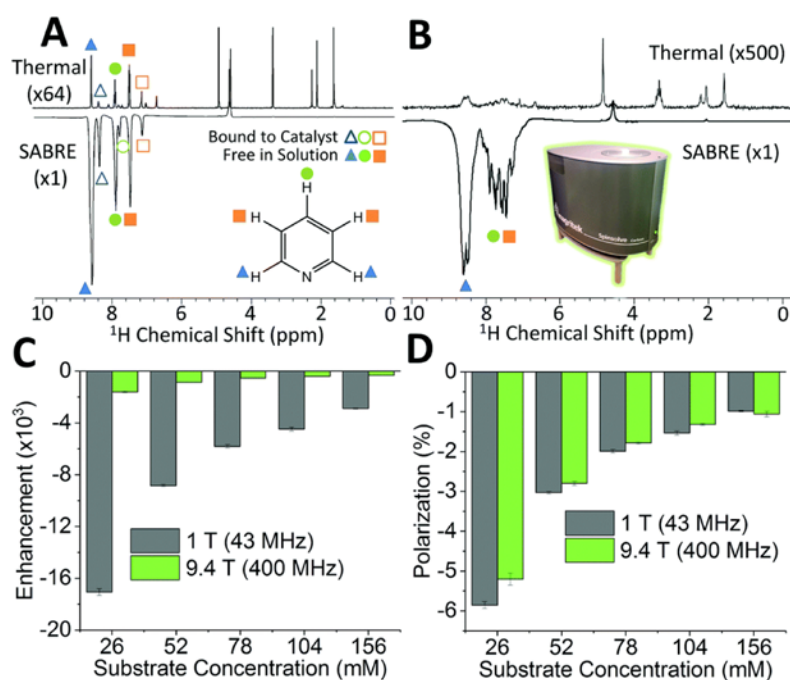
2. Experimental developments

Although NMR is ubiquitous in chemical analysis, it is less sensitive than alternative techniques [16]. As such, it is important to determine the SNR for analytes in NMR measurements. Indeed, the SNR scales to the square of the static magnetic field, B_0 , and this has led to significant investments in developing increasingly high-field magnets. However, with the relatively small magnetic fields produced in bNMR instruments, obtaining SNRs allowing for detection ($\text{SNR} \geq 3$) and quantification ($\text{SNR} \geq 10$) purposes can become challenging [17]. Hence, both sensitivity and selectivity barriers must be overcome for a number of bNMR applications.

The limit of detection (LOD) concentration values of LF instruments has long been described as an important barrier to overcome in order for the technique to have wider applications [18]. Indeed, a number of methods have been developed to enhance the spin effects observed in NMR for LF applications, amongst others [19]. Hyperpolarisation (HP) of nuclear spins greatly enhances signal intensities, with the potential to significantly expand the scope of both spectroscopic NMR and MRI systems, as well as sensing technologies [20]. The most well-known derivative of this technique, dynamic nuclear polarisation (DNP), engenders several hundred-fold increases in sensitivity for solid material analytes [21]. Moreover, in solution, DNP has been shown to increase sensitivity up to ten thousand times under standard conditions, opening up a variety of applications for lower frequency spectrometers [19].

DNP in compact, LF-NMR analysis has been applied to successfully study numerous processes such as lactate dehydrogenase activity in prostate carcinomas through hyperpolarisation of [$1\text{-}^{13}\text{C}$] pyruvate [22], as well as mouse brain metabolism [23], together with molecular dynamics in block copolymers [24]. In addition to DNP, signal amplification by reversible exchange (SABRE) is considered a rapid and efficient method of NMR hyperpolarisation that can enhance the sensitivity of proton and heteronuclear experiments [25]. SABRE does not require a hydrogenation reaction, and can occur in the liquid state under standard conditions with relatively cheap equipment (Figure 2); a transition

- 1 metal ion catalyst is employed to establish the contact between *para*-H₂, which is the polarisation
- 2 source, and nuclei of interest [26].



- 3
- 4 Figure 2. Reproduced from Ref [27] with permission from The Royal Society of Chemistry.
- 5 Comparison of thermally-polarized (top) and SABRE hyperpolarized (bottom) ¹H NMR spectra of 52
- 6 mM pyridine with 5.2 mM catalyst in methanol-d₄ for detection at (A) 9.4 T and (B) 1.0 T. (C)
- 7 SABRE enhancement factor, ϵ , and (D) polarization level, P , for the ortho resonance (blue triangle) as
- 8 a function of substrate concentration with NMR detection at 1.0 T (grey) and 9.4 T (green). Error bars
- 9 represent the standard deviation across 5 repeat measurements.

- 10
- 11 A heteronuclear alternative, described as SABRE-SHEATH, enables alignment transfer to
- 12 heteronuclei, and has been reported to successfully apply to the hyperpolarisation of ¹⁵N. It appears
- 13 that ¹⁵N-SABRE may be effectively applied to a wider range of substrates than its ¹H equivalent,
- 14 thereby generalising the technique. Additionally, T_1 values for ¹⁵N are prolonged up to 12 minutes,
- 15 allowing for increased levels of hyperpolarisation; this development is also appealing for lengthened
- 16 molecular imaging times, further opening up the field of hyperpolarised biomolecular imaging [28].

1 More recently, brute force HP has appeared as an alternative to DNP. It uses only two components in
2 order to hyperpolarise a molecule, specifically those involving a low temperature and high magnetic
3 field, and thereby not requiring free radical, microwave excitation or co-solvent processes necessary
4 for variants of DNP. Relying instead on the Boltzmann law, brute force hyperpolarisation also does
5 not require the metal ion catalysts inherent in parahydrogen hyperpolarisation techniques. Recently
6 reported applications of brute force hyperpolarisation NMR/MRI have reached thousand- fold gains in
7 sensitivity within the low-frequency bNMR range [29]. The simplicity of this low temperature, high
8 magnetic field approach to hyperpolarisation also opens up possibilities for the transport of HP nuclei.
9 Indeed, with slowly relaxing nuclei of low gyromagnetic ratios, it has been suggested that $T_1(^{13}\text{C})$
10 amounts to approximately 1 hour at 50 K and 24 hours at 10 K (or more than ten times the half-life of
11 ^{19}F used for positron emission tomography imaging). Hence, there is significant scope for this
12 technique to be developed further in order to drastically enhance the sensitivity of LF instruments,
13 which currently struggle somewhat with some analytes present at low concentrations. Indeed,
14 although HP techniques have substantial potential to increase the applicability of these instruments,
15 the current HP strategies described herein suffer from practical issues; dissolution DNP is frequently
16 used in developing contrast agents, but it suffers from a high cost, as well as complex hardware for
17 implementation. Whilst alternative parahydrogen techniques are effective, they are not as widely
18 applicable. Although the SABRE technique is a fast and reversible HP method, it is currently limited
19 to HP ^1H investigations, and therefore future work, most likely to be based on the aforementioned
20 SABRE-SHEATH methodology, will focus on HP transfer to ^{13}C and ^{15}N nuclei. It has also been
21 suggested that HP techniques will further encompass the applications of two-dimensional (2D) spectra
22 in the short-term [30].

23 In view of the low operating frequencies and low magnetic fields of bNMR instruments, resonances
24 often significantly overlap, which leads to less valuable and less easily interpretable spectra.
25 Moreover, such adverse overlap is amplified as the magnitudes of coupling constants approach the
26 range of chemical shift dispersion; this further complicates the characterisation of analytes. It has
27 therefore been valuable to apply 2D methods to LF spectra in order to enhance spectral resolution

1 through expanding the resonances over a 2D spectral map, whilst affecting sensitivity. Furthermore, a
2 number of pulse sequence variants have recently been developed in order to generate these enhanced
3 2D spectra [31]. However, these advantages offered by 2D LF NMR analyses are deleteriously
4 associated with long acquisition times, which somewhat reduce the viability of this technique for
5 high-throughput screening or reaction monitoring purposes. Until recently, hardware restricted the
6 application of ultra-fast 2D NMR analysis at LF. Fortunately, however, dedicated high-performance
7 gradient coils are becoming available for incorporation into LF bNMR instruments, with the first
8 application successfully achieved on a 43 MHz facility [32]. As such, robust methodologies in
9 ultrafast NMR have been developed at LF, and these rely on substitution of the t_1 period by a
10 spatially-dependent period of evolution. Within a single scan, the sample is therefore conceptually
11 separated into partitions subject to different periods of evolution, which depend on their relative
12 spectral frequencies.

13 Commonly-used resonance peak integration methods have been shown to suffer from certain
14 deficiencies, particularly when applied to LF bNMR data [33]. Specifically, these are limited in their
15 capability to resolve overlapping signals, and are inherently affected by noise, leading to the
16 development of several novel methods of corrections and FID data analysis [34]. A recently reported
17 Bayesian model involving lineshape and baseline corrections has been shown to account for chemical
18 shift, relaxation, lineshape imperfections and phasing, as well as baseline distortions. This model was
19 validated against simulation and experiment, using simple molecules for characterisation, and showed
20 an absolute accuracy of at least 0.01 mol/mol (in terms of mole fractions) for high SNR values of
21 more than 40 dB. In the context of this model, SNR is defined in Equation 1, where P_s and P_n are
22 represented as the energies of the resonance of interest, and the background noise, respectively. P_s is
23 obtained from the model-based method, whilst P_n can be derived from the Gaussian noise, or
24 alternatively the product of the total number of samples and the estimated noise variance, or from
25 direct measurement of signal-free regions in the spectrum.

$$26 \quad SNR = 10 \log \left(\frac{P_s}{P_n} \right) \quad (1)$$

1 For values of SNR less than 20 dB, accuracy was reported as 0.05-0.10 mol/mol. This has significant
2 implications for LF compact instruments which suffer from higher noise than their high-field
3 counterparts; however the simplicity of the test systems involved in the study have been highlighted,
4 and additional testing on more complex, inhomogeneous systems has been suggested [33].
5 Furthermore, additional work has focused on corrections to phase and set the baseline for signals
6 through complex optimisation strategies. This method applies these corrections simultaneously and in
7 an automated fashion, with no requirement for manual input, and has been reported to provide more
8 interpretable results than those arising from other techniques. Implementing a Pareto optimisation to
9 the concerted automated correction, as well as Whittaker smoothing to correct the baseline, this novel
10 approach was demonstrated on both high-field and bNMR spectroscopic data, suggesting
11 improvements in interpretation, particularly for spectra obtained from compact instruments [35].

12 The congestion in ^1H NMR spectra at LF arises mainly from overlapping multiplets, with a standard
13 $^3J_{\text{H-H}}$ coupling of 20 Hz for a *trans*-alkene occupying 0.50 ppm at a 40 MHz operating frequency.
14 COSY experiments can help to locate coupling pathways in congested parts of the NMR spectrum,
15 with selective-TOCSY approaches offering significant advantages by extracting a specific coupled
16 spin-system from a complex mixture. The pure-shift method can collapse a multiplet from a ^1H NMR
17 spectrum to yield a singlet, which offers significant advantages since overlap in spectra acquired is
18 vastly reduced. Recently, lanthanide shift reagents have been used to increase the frequency
19 dispersion in bNMR spectra through their abilities to form metal ion complexes, chelate or otherwise,
20 with target analytes. Whilst being frequently applied to high-field techniques, the implementation of
21 lanthanide shift reagents on a bNMR spectrometer through 1D ^1H and ^{19}F and 2D ^{19}F has only
22 recently been reported [36]. This presents an additional approach to the uncrowding of LF NMR
23 spectra, whilst effects on line-broadening must also be taken into account; high concentrations of
24 lanthanide complexes tend to induce increased bandwidth in spectra in view of decreases in T_2 values.
25 This effect is less pronounced for Eu(III) and Pr(III) complexes, a consequence of their inefficient
26 nuclear spin-lattice ratio properties.

1 Methods to improve acquisition efficiency have significant appeal to bNMR systems, and one robust
2 and widely applicable method is non-uniform sampling (NUS). This overcomes the inefficient
3 sampling method traditionally used for acquiring 2D NMR data, and allows equivalent quality data to
4 be recorded in just 25% of the time usually required to acquire a spectrum [37].

5 Materials such as foods, cosmetics, healthcare and oral healthcare products, polymers and further
6 chemical substances can be explored by relaxation measurements, for example determinations of the
7 solid fat contents of dairy products may be performed on MiniSpec benchtop NMR analysers (Bruker,
8 Billerica, Massachusetts, USA) via the differing relaxation behaviours of the solid and liquid
9 components of such non-solution-state matrices [38]. Such facilities have been available and
10 operational since the 1970's, and are readily applicable to the detection and quantification of ^1H , ^{19}F
11 and further NMR-active nuclei in a range of materials and other predominantly solid-phase products,
12 for example, determinations of the total fluorine contents of toothpastes [39]. However, since this
13 review is focussed on spectroscopic applications of LF bNMR instruments only, such relaxometry
14 analyses will not be discussed further here.

15 **3. Applications in materials chemistry**

16

17 NMR/MRI analysis has been widely applied to the study of materials and their interfaces [40]. More
18 recently, however, the availability and relatively low cost of LF benchtop instruments has led to a
19 plethora of applications in this field [13]. These include studies of rubber [3], nanoparticles [41], and
20 porous materials [42], with relaxation studies of solid-state materials and metal-organic frameworks
21 (MOFs) able to explore the pore-space of MOFs, and the determination of adsorption characteristics
22 [43].

23 LF NMR has been applied to the quantification of differences between varieties of styrene-butadiene
24 rubber. Individual components in the system, which differed between sites of origin, were detectable
25 with a benchtop instrument, on which 108 samples were analysed in ^1H and ^{13}C NMR experiments
26 performed on a 1.0 T instrument. Partial Least squares (PLS) regression allowed for a quality control
27 (QC) methodology to be developed for rubber varieties and their origins, and/or relative qualities.

1 Comparisons were made with a high-field 9.4 T instrument, which allowed for reliable assignment of
 2 signals in polymerised rubber systems, suggesting a defined role for LF NMR techniques combined
 3 with chemometric analysis in QC processes for raw rubber quality assessments [3].

4 Additionally, exploiting optical defects in crystals, such as nitrogen vacancy (NV) centres in diamond,
 5 can lead to signal enhancement compared to traditional methods of detection and excitation [44].

6 Indeed, these techniques can also be applied to transfer hyperpolarisation from the NV to nearby ^{13}C
 7 nuclei, leading to significant signal enhancements [45].

8

9 **Table 1.** Benefits, limitations and potential future applications of LF benchtop NMR techniques in the
 10 analysis of materials.

Criterion	Comments	Relevant References
Applicable to a range of materials and conditions	For geological methods, for example, NMR analysis is ideally performed on a portable instrument; however, a variety of facilities have been developed for this purpose. More traditional benchtop devices have been used to characterise materials ranging from magnetic nanoparticles to rubbers.	[3,10,41,46]
Simple sample preparation	Little or no sample preparation is required. No sample preparation at all is necessary for qualitative (identification) purposes performed by LF spectrometers not requiring added $^2\text{H}_2\text{O}$, nor for hand-held portable NMR devices. For the former class of facilities, in some cases only the direct transference of samples to NMR tubes is required.	[2,42]
Selectivity	Analytes such as complex polymers may require supplementary chemometric techniques to be applied in order for resonances to be robustly identified within multistep frameworks such as quality control. LF hand-held instruments rely instead on indirect secondary measurements.	[3]
Sensitivity	Superior to common alternatives for adsorption measurements in porous materials, such as gravimetric or volumetric analyses in which secondary effects such as pressure or mass changes are measured. Appropriate for the measurement of parameters such as Henry's constant, adsorption capacity, and adsorption separation factor in microporous materials. Measurement of transverse relaxivities (inverse T_1 and T_2) of nanoparticles are sensitive to their aggregation state, and appropriate for quantification purposes, respectively.	[41,43]
Versatility	A variety of instruments and techniques are applied in view of the diversity of media in the context of materials amenable to LF NMR characterisation. Hand-held sensors used in geological investigations are highly versatile, but with low sensitivity (when compared to the benchtop LF spectrometers or bespoke solid-state sensors).	[15,47]

Limitations	Solid-state NMR with benchtop instruments is limited to diffusion/relaxation studies through time-domain NMR. The chemical shift is not readily resolved in these inhomogeneous fields. However, it is possible to estimate indirect coupling, allowing for basic chemical analysis. Magnetic NPs currently also require characterisation with transmission electron microscopy or X-ray diffraction, which could support NMR analysis by providing additional information on relaxation behaviour, quantitative comparisons or relaxation theories.	[40,41]
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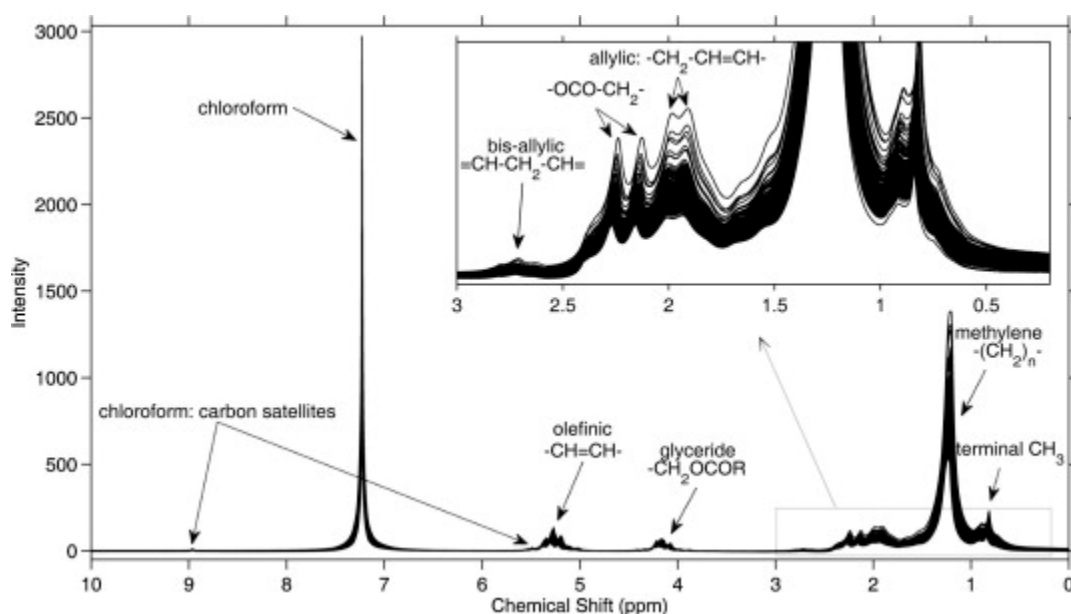
3 **4. Low-field NMR spectroscopy in food and agricultural chemistry**

4

5 The availability of compact LF NMR instruments has led to their rapid uptake in the food processing
6 and quality control areas, and the chemical analysis of agents in dietary products have been widely
7 reported [47,48]; recent media coverage of horsemeat product substitutions for beef highlighted
8 identifiable differences in their triacylglycerol contents in ^1H NMR spectra acquired on LF
9 instruments [49]. Moreover, the identification and purity of edible oils has generated much public
10 attention with adulterants being found in many commonly available products. Recent studies on olive
11 oil mixtures suggest an important role for compact, LF NMR instruments for quality control
12 evaluations. Indeed, since pressed hazelnut oil, a common olive oil adulterant, has oleoyl-, linoleoyl-,
13 and linolenoylglycerol contents which differ significantly from those of unadulterated olive oil, this
14 method serves as the basis of adulteration tests for commercial samplings of the latter. It has been
15 possible to use the lineshapes of selected methylene function resonances in order to identify
16 fingerprints for differing samples and mixtures (Figure 3) [50]. More recently, the authentication of
17 edible oils has formed part of a more methodological study to identify appropriate techniques for their
18 NMR characterisation at LF. Experiments were carried out on a 43 MHz instrument, and this study
19 reported that in terms of efficiency, a methodology based on 2D spectra obtained over less than 3
20 minutes was more efficient than 1D experiments performed on the same samples [51]. This was
21 rationalised in view of the significant resonance overlap in 1D spectra, and the facile discrimination
22 between six edible oils through principal component analysis (PCA) of 2D data. Furthermore, it has

1 been demonstrated that application of a partial least-squares (PLS) multivariate model permitted the
 2 detection of edible oil adulteration through the same method, with a similar procedure being applied
 3 in an authenticity and adulteration study of perilla oils distributed in Korea [52], and also for
 4 Patchouli essential oil [53].

5 Intriguingly, Grootveld *et al.* [54] recently employed LF NMR analysis using a 1.4 T LF benchtop
 6 NMR instrument in order to successfully detect 3 major classes of aldehydic lipid oxidation products
 7 (i.e. *trans*-2-alkenals, *trans,trans*-alka-2,4-dienals, and *n*-alkanals arising from the peroxidation of
 8 unsaturated fatty acid sources) in samples of repeatedly-used olive oils employed for deep-frying
 9 purposes in a restaurant. Consistent with this observation, frying time-dependent decreases in frying
 10 oil polyunsaturated fatty acid contents were also observable with this technique, specifically highly
 11 significant heat-induced decreases in the intensities of their *bis*-allylic-CH₂ function multiplet
 12 resonance ($\delta = 2.76$ ppm).



13

14 **Figure 3.** Extracted with permission from Ref [50]. Annotated 60 MHz ¹H NMR spectra for 20 extra
 15 virgin olive oils, 10 hazelnut oils and 144 olive oil–hazelnut oil mixtures. The inset shows an
 16 expansion of the chemical-shift region 0.20–3.00 ppm.

17

1 In addition to the study of edible oils, fuel and biodiesel analysis has formed part of the work
2 performed on compact NMR spectrometers in view of their relative portability and analysis cost-
3 effectiveness. Notably, there is a requirement for rapid and reliable quality control methods for diesel;
4 adulterants and contaminants substantially affect the properties of the fuel. A PLS regression model
5 has been developed through LF ^1H NMR experiments in order to determine a number of key
6 properties related to the fuel. Specifically, specific gravity, cetane number, flashpoint and distillation
7 temperatures could be compared through this PLS model, with errors less than or approaching those
8 of appropriate, more conventional analytical techniques. A univariate calibration curve was also
9 applied in order to derive the methyl and ethyl ester contents of biodiesel fuels from LF bNMR
10 experiments. It has been reported that compact NMR quality control and content experiments
11 generated comparable results to those of the current mid-IR reference technique; however, no prior
12 preparation of samples was necessary for the rapid scan times of 15 s, an advantage which expands
13 the possibilities for this technique as part of a wider QC process [55].

14 LF NMR analysis has also been applied to fields such as plant biotechnology. Indeed, secondary
15 biomolecules present in hops, which include bitter acids, volatile monoterpenes and sesquiterpenes,
16 are located in lupulin glands; extracellular trichomes are ideally suited to fast sample analyses with
17 little prior preparation. Screening of the composition of these glands would allow for the selection of
18 hops with particular aromatic properties, leading to the targeted breeding of particular strains. This
19 direct method of LF NMR analysis has been found to not only generate reliable quality indicators for
20 hops through combinations with chemometric techniques, but this approach also avoided the complex
21 extraction procedures usually employed for such analyses by other analytical techniques. This
22 highlights the simplicity of compact benchtop techniques, and also circumvents the use of
23 unsustainable and complex preparation practices [56].

24

25

- 1 **Table 2.** Benefits, limitations and potential future applications of LF bNMR analysis in food and
 2 agricultural chemistry. Limited references to relaxometry studies have been included for purposes of
 3 completion.

Criterion	Comments	Relevant References
Applicability to food standards and authenticity testing	Both relaxation parameters and amplitudes are powerful measures of the content and authenticity of intact foodstuffs. Standard protocols relying on fast, reliable measurements can be developed and transferred into a quality control pipeline.	[47]
Simple sample preparation	Little or no sample preparation is required. No sample preparation at all is necessary for qualitative (identification) purposes performed by LF spectrometers not requiring added $^2\text{H}_2\text{O}$ or alternative deuterated NMR solvents. For the latter class of facilities, in some cases only the direct transference of samples to NMR tubes is required. Simple preparation is essential, especially in the context of complex QC pipelines and frameworks, where rapid, reliable measurements are necessary.	[48,51]
Selectivity	For analyses of complex samples such as oils, with numerous resonances, it can be difficult to discern particular features which would distinguish between variants of the same types of sample (such as extra virgin olive oil <i>vs.</i> hazelnut oil comparisons). The purity of the sample will affect the level of selectivity, with pure samples of hazelnut oil and olive oil being distinguishable by the integrals of unsaturated fatty acid olefinic and olefinic-related resonances (of higher intensities in ^1H NMR spectra of hazelnut oils than in those of olive oils). This applies equally to analyses of alternative foodstuffs, with reasonable selectivity based on the complexity of the analytes investigated.	[50,51]
Sensitivity	Capable of detection of adulteration of olive oil with hazelnut oil at levels in excess of 11% (w/w). Improved sensitivity for the detection of oil adulteration at 60 MHz when compared to corresponding FTIR measurements.	[50]
Reproducibility of Analytical Data	95% confidence intervals for the analysis of edible oils are $\pm 13\%$ (w/w).	[50]
Reliability and Performance	Stability of measurements; repeat measurements across multiple weeks yield little variance. Application of chemometrics with supervised or unsupervised pattern recognition approaches is inherently applicable for quality control monitoring. Most facilities employ traditional 5-mm diameter NMR tubes. High sample throughput with rapid spectral acquisition. Simple operation which can be automated within a continuous flow set-up.	[47,48,50]
Versatility	Traditional LF NMR analysis is valuable for molecular determinations, whilst single-sided portable NMR sensors are more routinely used for whole foodstuff and packaged goods analysis.	[47]

Limitations	Solid-state NMR with benchtop instruments is limited to diffusion/relaxation studies through time-domain NMR. The chemical shift is not readily resolved in these inhomogeneous fields. However, it is possible to estimate indirect couplings, allowing for basic chemical analysis. The resolution of signals in complex samples such as edible oils may be limited, and hence leads to the implementation of rigorous chemometric techniques.	[13,48]
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2 5. Reaction monitoring and chemical transformations

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4 LF instruments have seen increased applications in process monitoring, in particular in
5 biotechnological engineering, where naturally heterogeneous disperse systems and their complex
6 optical behaviour engender numerous analytical challenges. *In-situ* reaction monitoring using NMR
7 flow cells has been available for high-field NMR systems for many decades, often as part of a
8 chromatographic or mass spectrometric set-up; however, this approach requires the employment of an
9 expert in several different areas of analytical science (NMR, chromatography, mass spectrometry
10 *etc.*), which is expensive and technically challenging. Benchtop NMR spectrometers therefore offer
11 low-cost and user-friendly options to monitor reactions in both teaching [57] and research laboratory
12 environments [58]. In 2015, Cronin and co-workers not only tracked reaction progress by monitoring
13 starting materials and products using a benchtop NMR system, but also incorporated a feedback loop
14 mechanism in order to modify the reaction composition, allowing reaction efficiency to be explored
15 and optimised in real-time [31]. This concept has been developed in order to incorporate a laboratory
16 plant design, such as in the synthesis of fluorinated compounds; the virtual 100% abundance, high
17 frequency and receptivity properties and half spin nucleus of ^{19}F render it a suitable nucleus for NMR
18 measurements in continuous flow systems, without the requirement for a solvent interference [59].
19 Additionally, Cortés-Borda and co-workers recently developed a modular, autonomous flow reactor,
20 synthesising the natural product carpanone as a model system [60]. As a self-optimising procedural
21 workflow, the equipment involved was designed to include either a high-performance liquid
22 chromatographic analysis system or LF NMR spectrometer. The developmental system required no

1 prior input regarding the reactions and systems in the experiment, leading to rapid optimisation of the
2 synthetic route to the carpanone product.

3 Recent work in process development has reported several appropriate set-ups for calibration
4 experiments, including connected 500 MHz and 43 MHz NMR facilities, together with Raman probes
5 and UV/Vis spectrophotometers in order to test the viability of compact instruments as substitutes for
6 high-field NMR spectrometers employed for the investigation of complex processes. Most notably,
7 data from variable temperature processes appear to show reasonable agreement between LF and HF
8 NMR instruments, although high temperature calibrations can cause issues with the internally-
9 regulated temperature of compact instruments [61]. Moreover, similar ideas can be applied to monitor
10 reaction progress [62], such as neutralisation of the mustard gas simulant, 2-chloroethyl ethyl sulphide
11 (CEES) [63]. This has been reported to take place using a portable continuous flow device, with fully
12 selective sulfoxidation by H₂O₂ being monitored with a compact LF NMR instrument, which allows
13 for real-time manipulation of reaction conditions. Alternatively, these compact NMR instruments
14 have been applied in stopped-flow analysis methods within microreactors, a strategy permitting the
15 optimisation of reaction times without the previously essential work-up procedures [64]. Additionally,
16 the incorporation of a mesofluidic reactor within a continuous flow synthesis as a means of optimising
17 acid-catalysed esterification, Knoevenagel condensation, Diels–Alder reactions, and alkylation
18 processes allowed for reaction monitoring without the conventional requirement of bespoke flow
19 cells. This was ascribable to the positioning of the LF NMR instrument after the back pressure
20 regulator, which also permitted the effect of temperature on reaction completion to be monitored [65].
21 Such applications miniaturise previously available, prohibitively large technologies, with the reported
22 system being the size of a suitcase and possibly also appealing to the defence sector [66].

23 Compact, LF spectrometers have also been used to compare different methods of reaction monitoring,
24 such as gas chromatography (GC) and high-field NMR analyses, as well as providing detection
25 methods for other techniques, such as size-exclusion chromatography [67]. Importantly, such compact
26 instruments can be applied directly within fume hoods, and have been used to monitor a variety of
27 reactions and processes; critical reviews dedicated to these strategies are available [68,69].

1 Recent work has employed a 1.0 T instrument in order to monitor the acid-catalysed acetalisation of
 2 *para*-nitrobenzaldehyde with ethylene glycol. Peak integration was used to monitor the concentrations
 3 of the reacting species, and reaction conditions were varied to determine kinetic parameters. Extracted
 4 samples at set intervals were also analysed on a high-field 9.4 T instrument for comparative purposes,
 5 each method appearing to provide similar kinetic data. It has therefore been suggested that compact
 6 NMR instruments can also reliably be used for these types of kinetic studies as viable, more cost-
 7 effective alternatives to those involving high-field spectrometers [70]. Fischer esterification, Suzuki
 8 cross-coupling, and oxime formation have also served as proof-of-concept reactions for monitoring
 9 with LF benchtop NMR analysis. These reactions were used as models for the consideration of such
 10 instruments as process analytical technologies (PAT), which are complementary to other methods
 11 such as IR and Raman spectroscopies. Reagents were required to be sufficiently concentrated for the
 12 dynamic range with solvent compositions minimised, whilst achieving a 10:1 SNR at 45 MHz
 13 operating frequency [71]. All reactions were performed at the magnet temperature of 42°C, limiting
 14 the technology in this study to the ambient temperature range, whilst variable temperature alternatives
 15 are now available [72].

16

17 **Table 3.** Benefits, limitations and potential future applications of LF benchtop NMR analysis in
 18 reaction monitoring and continuous flow systems.

Criterion	Comments	Relevant References
Incorporation within continuous flow systems	Although previously limited by the availability of variable temperature probes, a number of parameters can now be investigated by the incorporation of these instruments within a continuous flow framework, and the spatial position of the spectrometer within that protocol. In view of the variety of manufacturers and sample requirements, the fine-tuning of each system to a reaction monitoring process varies; however, LF NMR experiments can successfully monitor the evolution of both simple and complex chemical transformations.	[65,71,72]
Simple sample preparation	The requirement for deuterated solvents, for example, varies between spectrometers and their manufacturing sources. However, with the application of solvent suppression techniques, or the incorporation of $^2\text{H}_2\text{O}$ as a field frequency lock, this does not cause significant drawbacks in terms of reaction monitoring; ^{19}F and other nuclei have been used as probes in order to avoid interferences arising from the solvent.	[59,72]

	Little or no sample preparation is required. No sample preparation at all is necessary for qualitative (identification) purposes performed by LF spectrometers not requiring added $^2\text{H}_2\text{O}$. For the latter class of facilities, in some cases only the direct transference of samples to NMR tubes is required.	
Selectivity	The monitoring of complex molecular structures within continuous flow measurements becomes difficult with 1D techniques in view of the lower chemical shift dispersion and second-order coupling patterns, which give rise to an increased level of signal overlap.	[60,63,65,71]
Sensitivity	Concentrations of reaction materials must be sufficiently high in order to minimise dynamic range problems with solvents. However, SNRs of 10:1, which often represent the limit of quantification, are achievable with LF NMR analysis.	[14,71]
Reproducibility of analytical data	A high level of precision is obtainable both within- and between-assays. This is particularly relevant for the multiple simultaneous measurements performed within a continuous flow process.	[69]
Reliability and performance	Low cost, reliable determination of chemical structure and dynamic effects through the incorporation of variable temperature probes. Most facilities employ traditional 5-mm diameter NMR tubes. High sample throughput with rapid spectral acquisition. Simple operation which can be automated within a continuous flow set-up.	[31,65,71]
Versatility	Can be employed for reactants, intermediates and/or products which are not readily responsive to or accessible by alternative, non-NMR spectroscopic techniques such as UV/Visible spectrophotometric and IR probing systems. For example, those lacking chromophores and IR-active functions respectively..	[68,71]
Limitations	LF NMR techniques, particularly when using 1D ^1H experiments, may struggle to monitor complex molecular structures. Variable temperature processes necessitate specific equipment, whilst reports of bubbles within the flow cell, as well as precipitation, crystallisation, and the formation of phases during the reaction will affect the viability of the monitoring approach. Low SNRs are observed, with 10:1 remaining the recommended threshold. Automation is not yet fully implemented in many instruments, although some progress has been made in the development of bespoke algorithms and chemical robotics, enhancing its potential use as a robust PAT tool.	[71–73]

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6. Structural and forensic chemistry

Chemical analysis techniques such as bNMR spectroscopy are routinely applied in the detection and assessment of illicit drugs or agents [74], whilst also being used as experimental procedures in the teaching of analytical chemistry [75,76]. A comprehensive assessment of the forensic applications of LF compact, benchtop instruments recently described the structural characterisation and identification of strychnine adulteration in chemical identification tests [66]. Using a combination of 1D and 2D experiments, it was possible to identify strychnine and associated counter-ions through a consideration of chemical shift values as signatures in 1D ^1H , ^{13}C , and DEPT, as well as 2D COSY, HETCOR, HSQC and HMBC experiments at a magnetic field strength of only 1.0 T. In this extensive study, the molecular structure of strychnine was successfully identified through high-field 1D experiments, or alternatively via LF 1.0 T HMBC and HSQC approaches. This suggests that compact bNMR analysis strategies can be effectively used as a tool for structural elucidation, but also provides a viable and cost-effective solution to chemical forensic problems. Indeed, this 1.0 T facility was found to provide valuable information regarding the identity of several strychnine salts, employing their chemical shift signatures as structural descriptors based on the effects of heteroatoms on adjacent protons, as well as the identification of adulterants (Figure 4) [66].

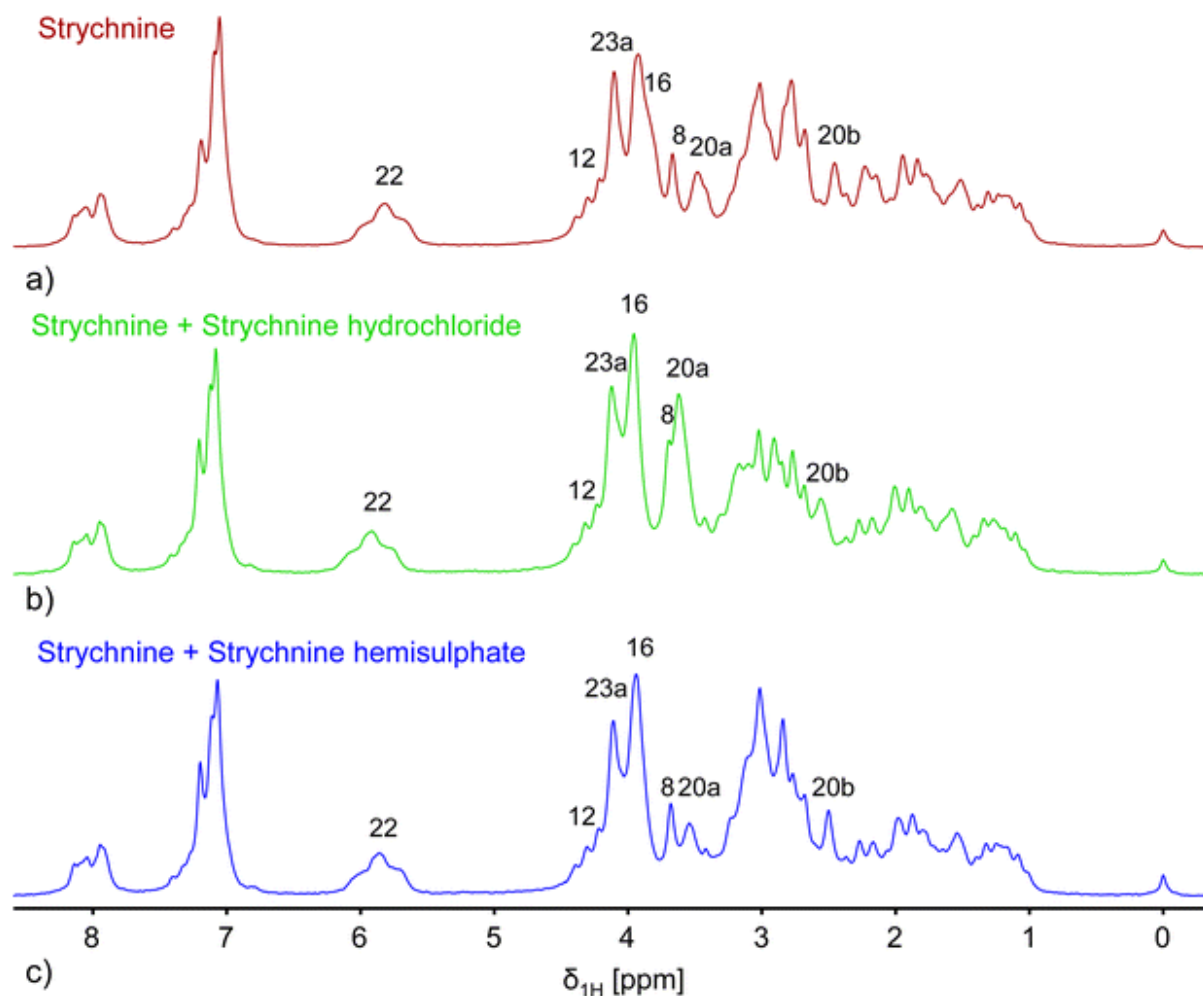


Figure 4. Reproduced from Ref [66] with permission from The Royal Society of Chemistry. 1D ^1H NMR spectra of strychnine and adulterated strychnine dissolved in CDCl_3 and recorded at 1.0 T. (a) Strychnine (135 mM), 4 scans. (b) Strychnine (135 mM) and strychnine hydrochloride (10 mM), 4 scans. (c) Strychnine and strychnine hemisulphate (10 mM), 4 scans. Assignments are provided in the ESI.

Benchtop NMR spectroscopy has been applied to the analysis of common illicit drugs such as methamphetamine. Organic and P-containing precursor molecules were identified in samples obtained from unregistered, so-called clandestine manufacturing laboratories. LF compact NMR analysis successfully characterised ephedrine, pseudoephedrine, and methamphetamine, whilst also detecting the P-containing precursors used during their synthetic routes [77].

Pharmaceutical and model systems were recently analysed using the first implementation of a DOSY pulse sequence on a LF NMR spectrometer. Indeed, Malet-Martino and co-workers [78] applied a

1 bipolar pulse pair-stimulated echo sequence with a longitudinal eddy current delay pulse sequence on
 2 esomeprazole, paracetamol, and hypromellose, as well as other model systems, and obtained a LF
 3 DOSY map. Although implementation of the technique was robustly established on a LF NMR
 4 spectrometer, the authors recognised the issues associated with the sensitivity of this system, leading
 5 to longer acquisition times or the need for higher concentrations in order to observe SNR values >5.
 6 This limits the range of application of this technique with current technology and methods, whilst the
 7 authors suggest this as a current approach for separating spectral profiles of different components in a
 8 mixture rather than obtaining accurate diffusion coefficients [78].

9 Therefore, in principal, the LF NMR analysis approach may soon be directly applied by forensic
 10 monitoring teams for the detection of illicit drugs and their synthetic precursors and by-products,
 11 together with traces of contaminating reaction solvents, ‘on-site’ at crime scenes, perhaps at police-
 12 raided clandestine laboratories or even roadsides. This will represent a major promising development
 13 in the forensic science area.

14
 15 **Table 4.** Benefits, limitations and potential future applications of LF benchtop NMR analysis in
 16 chemical forensics and structural chemistry.

Criterion	Comments	Relevant References
Mobile and field applications	Many forensic and point-of-interest assays will need to take place outside traditional laboratory spaces; therefore, there is a possibility of incorporating the technology of LF NMR spectrometers into portable boxes or mobile laboratories.	[66]
Simple sample preparation	The requirement for deuterated solvents, for example, varies between manufacturers; however, with the application of solvent suppression techniques, or the incorporation of $^2\text{H}_2\text{O}$ as a field frequency lock, this does not significantly limit the possibilities of this technology in forensic and chemical identification testing. Little or no sample preparation is required. No sample preparation at all is necessary for qualitative (identification) purposes in protocols performed by LF spectrometers not requiring added $^2\text{H}_2\text{O}$. For the latter class of facilities, in some cases only the direct transference of samples to NMR tubes is required.	[77,79]
Selectivity	Particularly for the assignment of areas of strong overlap in ^1H spectral profiles, ^{13}C experiments at 1.0 Tesla (chemical shift range > 200 ppm) can offer improved resolution of resonances. These experiments suffer from low sensitivity and the effects of the low abundance of ^{13}C (only 1.1%), yet at natural abundance, there is an absence of homonuclear couplings between ^{13}C nuclei.	[66,80]

	NOE signal enhancement and ^1H decoupling eliminates heteronuclear multiplet splitting leading to full resolution of all 22 ^{13}C resonances in the NMR spectrum of strychnine in 4,096 scans (the total acquisition taking 5.68 hours).	
Sensitivity	1D ^1H experiments offer much valuable qualitative, structural determination potential. However, 2D techniques provide additional information, in addition to improvements in sensitivity, in particular for structural elucidation purposes. HSQC experiments represent fast, spectroscopically-indirect measurements for ^1H nuclei. 2D HSQC NMR analysis of strychnine present at a level of 135 mmol per NMR tube was acquired in 128 scans for each time-step, and a total experimental time of 22.7 hours (repetition time of 10s). A HETCOR spectrum of the same molecule acquired within the same time period exhibited lower intensity and afforded lower SNR values than those of equivalent HSQC experiments.	[14,66]
Reproducibility of Analytical Data	A high level of precision is obtainable both within- and between-assays. Considering the high reproducibility/low repeat between-assay component of variance established in this study for biofluid measurements, reproducibility should be high for structural determination experiments.	[80]
Reliability and Performance	<p>Low cost, reliable determination of chemical structure and authenticity through 1D and 2D experiments.</p> <p>Adequate resolution, stability and sensitivity for both 1D and 2D experiments in order to permit the reliable determination of complex structures such as those of drugs and biomolecules. ^1H resonances are reliably assigned through 2D HETCOR, HSQC and COSY experiments, whilst a 2D <i>J</i>-resolved methodology will facilitate the identification of fine structure. ^1H and ^{13}C experiments are available, allowing for significant multivariate data to be obtained and correlated for robust structural determinations.</p> <p>Most facilities employ traditional 5-mm diameter NMR tubes.</p> <p>High sample throughput with rapid spectral acquisition.</p> <p>No major requirement for specialist NMR/technical staff for operation, i.e. there is no need for operating staff to understand and interpret spectral profile data, and this renders LF NMR facilities suitable for use by investigative crime scene field or forensic workers directly at their crime scene sites, i.e. no laboratory is required.</p> <p>Experimental time is longer than that of high-field techniques; however, the relative ease of use, absence of cryogenic facilities and compact nature render this technique is particularly suited to forensic and chemical forensic applications in the field.</p>	[81]
Versatility	The study of small, complex molecules with LF NMR spectroscopy, as well as the electrostatic interactions of dissolved ions, suggests a viable role in forensic analysis and mobile chemical assay systems.	[74]

Limitations	Long experimental times for robust structural analyses and 2D experiments, whilst 1D techniques suffer from some selectivity issues.	[66]
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2 **7. Potential biofluid screening and metabolomics applications of LF benchtop NMR analysis**

3

4 High- and very high-field NMR spectroscopic techniques continue to serve as extremely useful, and
5 now extremely-sensitive probing tools for the simultaneous multicomponent analysis of complex
6 biofluids collected from humans and other living systems. A very high level of valuable information
7 regarding the molecular nature and concentrations of a plenitude of endogenous biomolecules, along
8 with exogenous agents present in such fluids (e.g., human urine, blood plasma, knee-joint synovial
9 fluid, cerebrospinal fluid and salivary supernatants), can be acquired from such investigations. These
10 approaches offer much potential regarding the investigation of metabolic processes, and when linked
11 with multidimensional data analysis techniques in metabolomics strategies, serve as extremely
12 powerful means of probing, for example, the biomolecular basis of many human diseases. For
13 example, statistically significant differences between the concentrations of one or more biofluid
14 metabolites, or patterns of such metabolites, in human disease *versus* healthy control biofluid profile
15 comparisons, conceivably arise from disturbances or perturbations to key metabolic pathways
16 featured in disease aetiologies, developments and/or progressions. Such studies may also yield much
17 valuable diagnostic and/or prognostic screening information, including that involving the seeking,
18 identification and validation of dependable and potentially predictive biomarkers. NMR-based
19 metabolomics strategies may also be applied to the multicomponent analysis of many biomolecules in
20 tissue biopsies and cultured cells (either directly via the analysis of intact samples using ^1H high-
21 resolution magic angle spinning (MAS)-NMR analysis technologies, or indirectly as pre-selected
22 solution extract matrices).

23 Recently, Percival *et. al.*[12] demonstrated the very first application of LF benchtop NMR
24 technologies to the ^1H NMR-based metabolomics analysis of human urine, in this case for the
25 diagnosis and monitoring of type 2 diabetes patients. This study fully established that such an
26 analytical strategy was successful in detecting and quantifying key urinary biomarkers for this

1 condition, despite a series of resonance overlap problems arising from the low operating frequency
2 involved (resonances with similar linewidths appear increasingly broader at decreasing field strengths
3 within the overall *ca.* 10 ppm spectral width in ppm). Moreover, it also provided a bioanalytical
4 ‘blueprint’ protocol for the future performance of such LF metabolomics explorations.

5 In addition to expected upregulations in urinary glucose, and ketone bodies such as 3-D-
6 hydroxybutyrate, this investigation found that acetate, alanine, citrate, and creatine/creatinine also had
7 significantly higher concentrations in type 2 diabetic urine samples than those of healthy control
8 participants, together with downregulations in aromatic biomolecules, particularly hippurate and
9 indoxyl sulphate (which presumably arise from diabetic polyuria). These results were in accordance
10 with those obtained in a corresponding high-field (600 MHz) ^1H NMR study performed on human
11 urine [82]. Multivariate metabolomics analysis of these 60 MHz ^1H NMR profiles by PCA, partial
12 least squares and orthogonal partial least squares discriminatory analysis (PLS-DA and OPLS-DA
13 respectively), support vector machine (SVM)-based receiver operating characteristic (ROC) curve
14 analysis, and random forest (RF) models demonstrated a very high level of distinction between these
15 two sample classifications. There was very good agreement between estimations of urinary glucose
16 concentrations obtained at both 60 and 400 MHz operating frequencies, and those determined by both
17 approaches correlated very significantly with those estimated by a non-NMR spectrophotometric
18 method based on a glucose oxidase system.

19 Typical LF 60 MHz ^1H NMR profiles acquired on urine specimens collected from type 2 diabetic and
20 healthy control participants are shown in Figures 5(a) and (b) for comparative purposes. The clear
21 spectral differences observed between these profiles (particularly the observation of glucose signals in
22 the type 2 diabetic sample and their absence from the healthy control samples) are, at least in
23 principle, interpretable by non-specialist healthcare staff. It is also anticipated that the portable
24 benchtop NMR system employed in the laboratory for the diagnosis and prognostic monitoring of
25 diabetes by clinical and bioanalytical chemists, may be routinely managed and operated by such
26 untrained, non-NMR specialist staff when implemented in hospital or alternative point-of-care health
27 settings such as smaller general practice health centres, dental surgeries and community pharmacies.

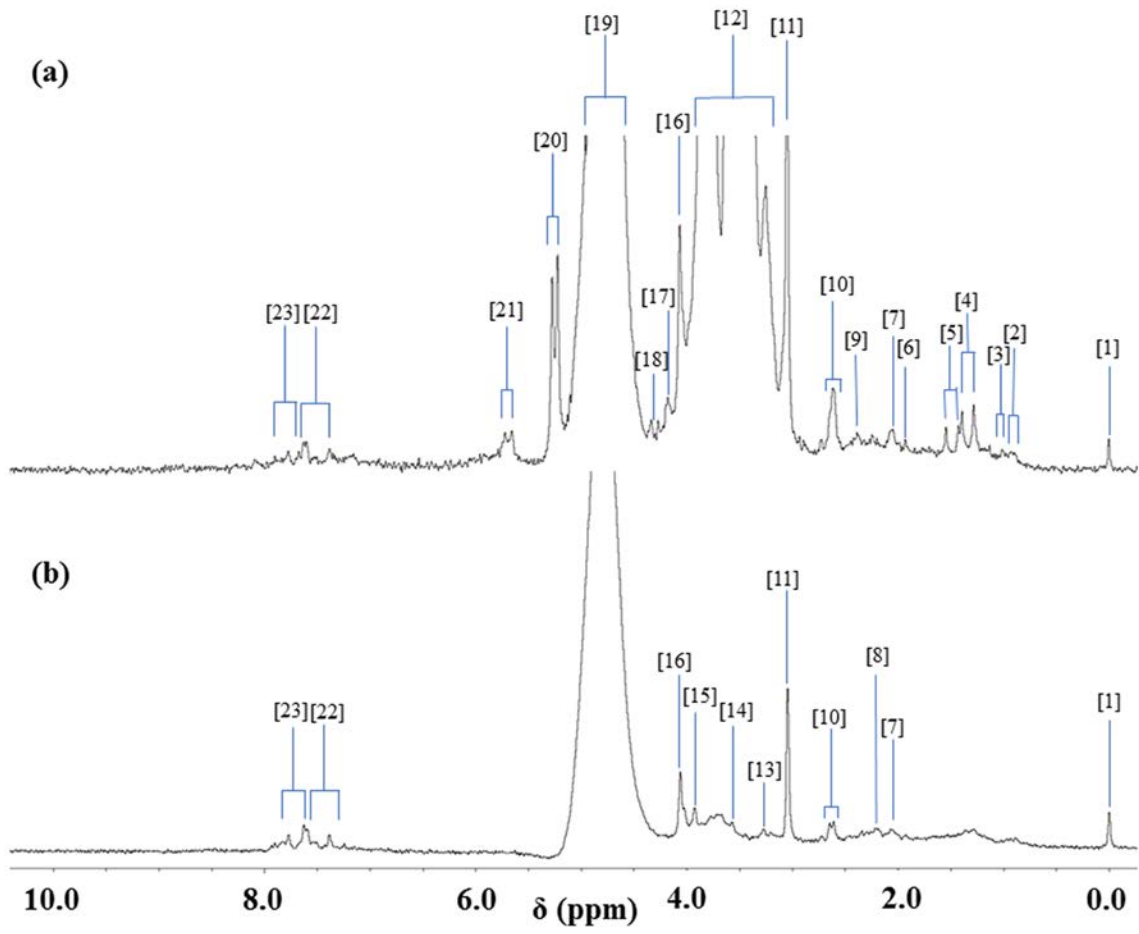
1 For one of the more prominent resonances visible in the benchtop 60 MHz ^1H NMR profiles acquired
2 on type 2 diabetic urine samples, *i.e.* that attributable to the $-\text{CH}_3$ function of lactate (d , $\delta = 1.33$ ppm),
3 the J value of this resonance is 7.00 Hz [83], and therefore it will envelop $2 \times 7.00/60 = 0.233$ ppm at
4 this operating frequency, but only $2 \times 7.00/400 = 0.035$ ppm at 400 MHz. Hence, at least some
5 caution is required regarding determinations of biomolecular analytes in biofluids via LF benchtop
6 NMR analysis, especially on consideration of inherent signal overlap complications. Notwithstanding,
7 these interferences are considered minimal for the determination of major urinary metabolites, *i.e.*
8 those with prominent resonances in spectra acquired which have only a limited level of overlap with
9 lower intensity signals, *e.g.* those of the citrate- CH_2 - and acetate- CH_3 functions at $\delta = 2.65$ (AB
10 coupling pattern) and 1.92 ppm (singlet) respectively, which include endogenous metabolites,
11 chemopathological diagnostic biomarkers, and xenobiotics present at high or very high concentrations
12 when expressed as a proportionate function of the total ^1H NMR spectral intensity, *i.e.* as in constant
13 sum normalisation (CSN) methodologies. For example, the mean \pm SEM urinary level of 3-D-
14 hydroxybutyrate in the type 2 diabetic urine samples investigated was 3.12 ± 0.99 mmol/L (upregulated
15 13-fold over that of healthy controls), and therefore it is readily monitored in LF 60 MHz NMR
16 spectra, provided that it is within the limit of quantification (LOQ) value estimated in samples to be
17 tested. Moreover, ^1H NMR-active molecules with resonances located within interferant-free regions
18 of the profiles obtained such as formate ($\delta = 8.46$ ppm) are also readily quantifiable. If focused on
19 biomolecule resonances with clearly-visible singlet, doublet or triplet resonances, the analytical
20 sensitivity of the LF benchtop spectrometer employed was excellent. Indeed, further experiments
21 established that the limit of quantification (LOQ) for the additional ketone body acetone (arising from
22 the decomposition of acetoacetate *in vivo*) was only 25 $\mu\text{mol/L}$ at an operating frequency of 60 MHz
23 [12].

24 Analytical ^1H NMR response problems encountered with the ‘intensity dampening’ of glucose’s α -
25 anomer C1- H resonance employed for quantification purposes ($\delta = 5.25$ ppm) by the $\text{H}_2\text{O}/\text{HOD}$
26 presaturation process employed in LF ^1H NMR determinations of total urinary glucose concentrations
27 (optimised at $\delta = 4.95$ ppm), were overcome by the rigorous employment of prepared calibration

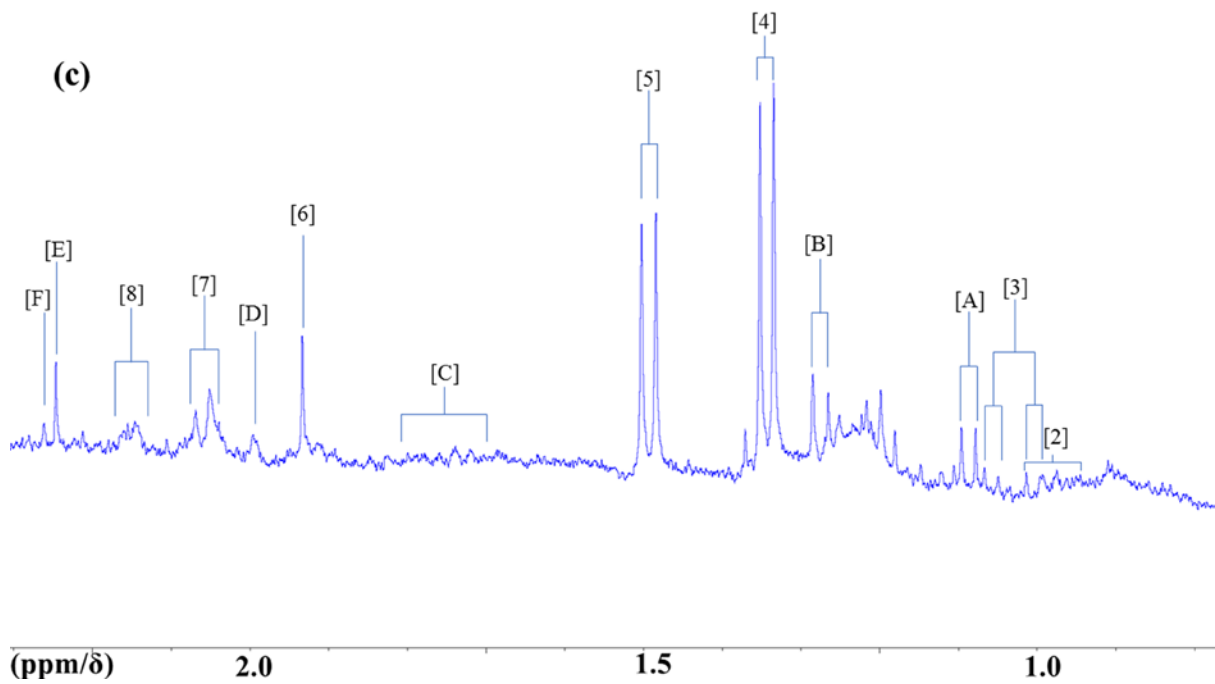
1 standards for all spectrometers utilised, both at 60 and 400 MHz operating frequencies. A further
2 limitation found was attributable to significant differences between intra-molecular relaxation times
3 for selected biomolecules with two or more magnetically-distinguishable ^1H nuclei, together with any
4 long-range coupling effects.

5 Prominent protein-based and low-molecular-mass metabolite resonances were also observable in
6 single-pulse ^1H NMR spectra of human blood plasma acquired on a 60 MHz LF benchtop facility, the
7 most intense being those assignable to lipoprotein-associated triacylglycerol fatty acid chains, and
8 ‘acute-phase’ glycoprotein carbohydrate side-chain N-acetylglucosamine and N-acetylneuraminate
9 residues (predominantly those of α_1 -acid glycoprotein for the latter), along with those of glucose’s
10 carbohydrate ring protons.

11 Overall, the authors recommended that in all future LF benchtop NMR-linked metabolomics
12 investigations, researchers should primarily focus on the most highly prominent signals which are
13 singlets, or those which have simple first-order coupling patterns (i.e. doublets, triplets, etc.) visible in
14 spectra acquired for quantification purposes. However, notable exceptions to this guideline include
15 targeted biomolecular analytes present at high or very high biofluid concentrations, for example
16 glucose and ketone bodies found in urine specimens collected from uncontrolled or poorly-controlled
17 diabetic patients. The technique is also likely to be applicable to the diagnosis of further metabolic
18 diseases, and further investigations to explore this are currently in progress.



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1 **Figure 5. (a) and (b)**, LF 60 MHz ^1H bNMR profiles of human urine samples collected from type 2
 2 diabetic and healthy control participants respectively. (c), Expanded 0.60-2.30 ppm region of the
 3 corresponding 400 MHz spectrum of that shown in (a). Typical spectra are shown. Abbreviations: [1],
 4 3-trimethylsilyl-(2,2,3,3- $^2\text{H}_4$)-1-propionate-Si(CH_3)₃ [TSP, internal chemical shift reference and
 5 quantitative ^1H NMR internal standard ($\delta = 0.00$ ppm)]; [2], isoleucine/leucine/valine- CH_3 ; [3],
 6 valine- CH_3 ; [4], lactate- CH_3 ; [5], alanine- CH_3 ; [6], acetate- CH_3 ; [7], glutamate- β - CH_2 /N-acetyl amino
 7 acids and N-acetylsugar-NHCO CH_3 ; [8], glutamine- β - CH_2 ; [9], glutamine- γ - CH_2 ; [10], citrate- CH_2 -
 8 CO_2^- ; [11], creatinine/creatinine-N- CH_3 ; [12], bulk glucose-C2H to C6H₂ carbohydrate ring protons;
 9 [13], trimethylamine-N-oxide-N(CH_3)₃; [14], glycine- CH_2 ; [15], creatine- CH_2 ; [16], creatinine- CH_2 ;
 10 [17], [18], unassigned; [19], residual $\text{H}_2\text{O}/\text{HOD}$; [20], α -glucose-C1-H; [21], urea-CO-NH₂; [22],
 11 indoxyl sulphate aromatic ring protons; [23], hippurate aromatic ring protons. [A] to [F] correspond to
 12 resonances which were detectable at an operating frequency of 400 MHz in this particular type 2
 13 diabetic urine sample, but not so at 60 MHz: [A], *iso*-butyrate- CH_3 groups; [B], 3-D-hydroxybutyrate-
 14 CH_3 ; [C], lysine- δ - CH_2 ; [D], proline- γ - and δ - CH_2 groups; [E], acetone- CH_3 ; [F], acetoacetate- CH_3 .

15

16 **Table 5.** Benefits, limitations and potential future applications of the LF benchtop NMR analysis of
 17 human or animal biofluids and liquid biopsies.

Criterion	Comments	Relevant References
Requirement for deuterated NMR solvents (particularly $^2\text{H}_2\text{O}$)	Many LF-NMR analysis systems do require deuterated NMR solvents [some exclusively so, and some only as a field frequency lock, <i>ca.</i> 10% (v/v) $^2\text{H}_2\text{O}$], but selected spectrometers operate effectively without such a requirement.	[12,84]
Sample preparation	Little or none required. No sample preparation at all is necessary for qualitative (identification) purposes performed by LF spectrometers not requiring added $^2\text{H}_2\text{O}$. For the latter class of facilities, in some cases only the direct transference of samples to NMR tubes is required.	[12,84]
Selectivity	Limited, but the most prominent resonances (i.e. those of relatively high concentration biomolecules), which are clearly visible and predominantly unaffected by minor potentially interfering signals, may be spectroscopically monitored. Those located in spectroscopically-clear regions are also accessible to quantification. Application of 2D COSY and TOCSY options available facilitates the rapid confirmation of identity assignments. However, these developments will markedly improve with the future availability of LF benchtop NMR spectrometers of higher operating frequencies.	[12,84]
Sensitivity	Surprisingly and disproportionately high: LLOD values of < 50 $\mu\text{mol./L}$ for selected metabolites with prominent - CH_3 and/or - CH_2 - function singlet resonances present in aqueous model solutions. SNR values > 10 achievable at concentrations of ≥ 25 $\mu\text{mol./L}$ for acetone.	[12,84]
Metabolite quantification	Particularly suited to the determination of acetate, alanine, branched-chain amino acids (BCAAs), lactate, N-acetyl storage	[12]

potential	<p>compounds, citrate, creatinine/creatinine, formate, 3-D-hydroxybutyrate, urea, indoxyl sulphate, hippurate and glucose in human urine.</p> <p>Excellent agreements were found between LF 60 MHz benchtop spectrometer determinations of glucose and those performed at high-field (400 MHz). There was also a very highly significant correlation between these 60 MHz values and those generated by a non-NMR method.</p> <p>Very reliable linear internal standard-normalised intensity <i>vs.</i> analyte concentration calibration standard plots readily facilitate the determination of a range of biofluid metabolite concentrations.</p> <p>Problems are encountered with the determination of metabolites involving the electronic integration of resonances located in close proximity to the H₂O/HOD presaturation frequency. However, such problems may be overcome via the integration of alternative signals of biomolecules of interest (if available and accessible), and/or the employment of suitable pre-validated calibration standards for the particular LF NMR spectrometer employed. Careful pre-selection and optimisation of the presaturation frequency and its power setting aids such analyses.</p> <p>At least some of these metabolites, together with propionate, and further ones of relatively high concentration and with ¹H NMR resonances with simple (first-order) coupling patterns such as glycine, methanol and succinate, are also quantifiable in human saliva (data not shown).</p>	
Reproducibility of analytical data	<p>A high level of precision is obtainable both within- and between-assays. Typical 'within-assay' mean±SD values obtained for n = 11 repeat citrate determinations made on a single 20.00 mmol/L calibration standard sample was 19.86±0.15 mmol/L (coefficient of variation only 0.75%).</p> <p>Typical 'between-assay' coefficients of variation for urinary citrate (AB coupling pattern, δ = 2.65 ppm), trimethylamine-N-oxide (<i>s</i>, δ = 3.23 ppm), glycine (<i>s</i>, δ = 3.60 ppm), creatine (-CH₂- function <i>s</i>, δ = 4.04 ppm) and indoxyl sulphate (2 x sets of aromatic proton <i>dds</i>, δ = 7.24 ppm) concentration determinations were 5.6, 4.1, 7.3, 4.7-5.6 and 2.9% respectively (n = 2 or 3 repeat measurements made).</p>	[12]
Reliability and performance	<p>Offers a very convenient, high-performance biomedical/clinical chemistry NMR analysis system at a cost which is much lower than that which involves more traditional high-field NMR spectrometers.</p> <p>Provides a rapid and reliable strategy for the multicomponent analysis of human biofluids or liquid aspirates/biopsies at their site of collection (e.g. at health centres, dental surgeries and/or pharmacies), or even outdoors in the field (e.g. at crime scene investigations). These facilities are quickly installed at such sites (the installation of HF NMR facilities with superconducting magnets is simply not feasible under these circumstances for</p>	[12,84]

	<p>logistical reasons). It has a small space ‘footprint’, and the methods developed involve little or no sample preparation.</p> <p>Most facilities employ traditional 5-mm diameter NMR tubes.</p> <p>High sample throughput (up to <i>ca.</i> 60 samples per 24 hr. period), with rapid spectral acquisition.</p> <p>No major requirement for specialist NMR/technical staff for management and operation, i.e. there is no need for operating staff to understand and interpret spectral profile data, and this renders LF NMR facilities suitable for use on-site by clinical nursing staff, or investigative field or forensic workers at their points of contact.</p> <p>Highly suited to the pre-screening of samples prior to their analysis at higher operating frequencies, i.e. at ≥ 500 MHz. For example, LF NMR instruments may be employed to simply detect glucose and/or ketone bodies in the urine of suspected diabetic or pre-diabetic patients so that a full urinary ^1H NMR screen can then be performed at HF.</p>	
Versatility	<p>Applicable to most, if not all, biofluids and liquid tissue biopsy extracts, but particularly valuable for those containing little or no protein and other macromolecules, e.g. human urine and salivary supernatants. Both single-pulse and Carr-Purcell-Meiboom-Gill (CPMG) pulse sequence-filtered spectra may be acquired on human blood plasma samples.</p>	[12,85]
Limitations	<p>With the exception of biomolecules present at very high biofluid concentrations (such as α- and β-glucose anomers in uncontrolled or poorly-controlled diabetic urine specimens), currently not readily suitable for the determination of metabolites containing ^1H nuclei with complex coupling patterns (which give rise to higher-order multiplets), nor to those with low intensity signals which are at least partially obscured by more prominent ones such as lactate- and acetate-CH_3 function resonances. However, this problem may also be experienced in high-field NMR analysis, especially at operating frequencies ≤ 500 MHz.</p> <p>Resonance ‘intensity-dampening’ problems experienced with the integration of those with chemical shift values similar to that of the $\text{H}_2\text{O}/\text{HOD}$ presaturation frequency are readily rectifiable (please refer to metabolite quantification potential column above). Similar, albeit less marked, complications were also encountered at an operating frequency of 400 MHz.</p> <p>An additional complication arises from significant differences between intra-molecular relaxation times for analytes with two or more magnetically-distinct ^1H nuclei; however, limitation of intensity integrations for quantification purposes to the most intense resonances with simple first-order coupling patterns is recommended in such cases.</p>	[12,84]

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Conclusions

The developing field of compact benchtop NMR instruments for chemical and biochemical analysis has recently rapidly expanded, both in terms of applications and advanced methodologies. Applications now range from biomedicine, through to advanced energy research and materials science, highlighting the versatility of the technique, together with the great strides that early innovators and current specialists are making in transitioning this technique into an over-arching, viable LF alternative to the expensive, larger instruments available in an increasing number of research areas and industry. Indeed, most recently, Cronin and co-workers have pioneered a novel machine-learning framework for probing chemical reactivity, in an ‘organic synthesis robot’, incorporating benchtop NMR within the analysis pathway [73]. Novel algorithms and analytical routines are being developed to treat the common SNR problems associated with LF instrumentations, paving the way to new gains in accuracy. More recently, LF NMR instruments have been employed in the analysis of biofluids, including applications of metabolomics techniques for the identification of diabetes in patient urine [12]. In this study, urine was collected from a cohort of type 2 diabetic patients and healthy individuals as controls. The study successfully detected markers of diabetic control such as glucose and acetone at concentrations of 2.8 mmol./L and 25µmol./L, respectively. Whilst demonstrating the application of common metabolomics strategies on data collected from LF NMR instruments, associated biofluid preparation and assay protocols were also presented, and optimised for use with bNMR spectrometers. Further work on the implementation of these compact instruments in the analysis of biofluids should involve large cohort studies to provide more statistically robust evidence for the use of these facilities in metabolomics investigations. Essentially coupled with the future application to low-field NMR measurements, is the concept of the ‘singlet’ spin state relaxation (T_s) in NMR as a tool to monitor chemical and biochemical processes.[86] Indeed, it is regularly suggested that T_1 is the longest relaxation process and will return the measurable magnetisation to equilibrium. More recently, however, it has been considered that T_s

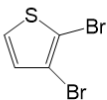

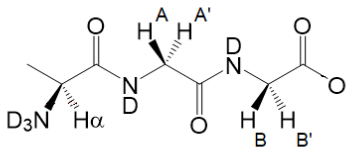
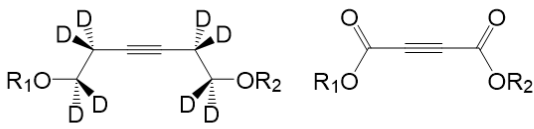
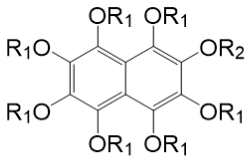
1 can in fact be longer than T_1 . Simple molecules with carefully selected spin-symmetry and often with
2 isotopic enhancement can display T_s values of the order of many minutes to approximately 1 hour in
3 the liquid state, with T_1 values that are considerably shorter.

4 The theory of relaxation processes was described in 1965 by Redfield,[87] and superoperators
5 described in 1982 by Jeener,[88] with the steady-state pulse sequences in 1992,[89] and a master
6 equation of relaxation networks in 1994 by Levitt.[90] Subsequently, application of long-lived singlet
7 states have been reviewed,[91,92] and highlight T_s values of up to 1 hour (Figure 6). This opens up
8 the possibility of monitoring reaction processes and metabolism over a much longer time period than
9 what was possible before, using NMR spectroscopy, as well as *in-vivo* MRI applications using
10 hyperpolarised material, and could indeed lead to further developments in the applications of LF
11 NMR analysis.

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Theory of NMR relaxation 1955 - 2017

1955	The Theory of Relaxation Process		Redfield
1982	SuperOperators in Magnetic Resonance		Jeener
1992	Steady State Magnetic Resonance Pulse Experiments		Levitt
1994	The Homogeneous Master Equation and Manipulation of Relaxation Networks		Levitt
2004		$T_1 = 16.8\text{s}$	$T_s = 104\text{s}$
2004		$T_1 = 7.8\text{s}$	$T_s = 141\text{s}$
2008	N_2O	$T_1 = 2\text{-}3\text{min}$	$T_s = 26\text{min}$
2011		$T_1 = 1\text{s}$	$T_s = 11\text{s}$
2012		$T_1 = 27\text{s}$	$T_s = 577\text{s}$
2015		$T_1 = 1\text{min}$	$T_s = 1\text{hour}$

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2 **Figure 6.** Progress in NMR relaxation theory from 1955 to 2017 describing singlet NMR3 methodology in spin- $\frac{1}{2}$ systems based on Ref [92].

4 We may also conclude that cryogen-free benchtop NMR systems do not require a dedicated,

5 customised laboratory space, nor also specialist staff, for their effective operation; no specialist

6 laboratory requirements, maintenance and management are necessary; multinuclear measurements,

7 e.g. ^1H , ^{13}C and ^{19}F , are possible with a single probe; as with higher field spectrometers; only 0.50 ml

8 of sample is required, and it employs common 5-mm diameter NMR tubes; rapid spectral acquisition

9 and therefore high sample throughputs are achievable; suitability for the on-line monitoring of

1 chemical reactions; avoidance of time-consuming and expensive acquisition of spectra on high
2 operating frequency spectrometers (i.e. those ≥ 400 MHz) via the rapid prior screening of samples in
3 analytical strategies targeted on the removal of those not suitable for this purpose; for some available
4 benchtop facilities, there is no requirement for expensive deuterated NMR solvents; and finally, the
5 technique is therefore highly promising for the 'point-of-care' diagnostic and prognostic screening of
6 biofluids for selected human or animal diseases, in any small laboratory space, directly at non-
7 laboratory health centre clinics (i.e. medical or dental surgeries, or pharmacies), or alternative point-
8 of-contact patient sites. Therefore, the applications of these techniques continue to grow, and will
9 likely develop further into personalised medicine and clinical chemistry-based disease monitoring
10 applications. Analytical parameters relating to articles considered in this study are summarised in
11 Table 6.

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1 **Table 6.** Summary of analytical parameters from studies employing low-field NMR spectroscopy,
2 referenced in this review, ordered by the section of this manuscript in which the reference appears,
3 and study name. Sensitivity expressed as SNR or LOD/Q, number of scans, acquisition time and
4 repetition time are featured where these appear in the text or supplementary information of the
5 relevant manuscripts. Abbreviations: h (hours), m (minutes), s (seconds).

Section	Study	Analyte/Reaction	Sensitivity (including LOD, LOQ, SNR) (lowest recorded)	Number of scans or total acquisition time	Repetition time	Reference
Experimental developments	SABRE hyperpolarization enables high-sensitivity ^1H and ^{13}C bNMR	Pyridine 4-methylpyridine	SABRE methodology leads to up to 17,000-fold ^1H and 75,500-fold ^{13}C NMR signal enhancements	^1H : 256 ^{13}C : 4096	Not reported	[27]
	A self optimizing synthetic organic reactor system using real-time in-line NMR	Diels-Alder cycloaddition Cyclopentadiene Acrolein Benzylamine	Not reported	^{13}C : 64 ^1H COSY: 9 m HSQC: 38 m	<10 s	[31]
	Compact NMR with Shift Reagents	Adamantan-2-ol Europium(III)-tris(1,1,1,2,2,3,4-heptafluoro-7,7-dimethyl-4,6-octanedionate)	Not reported	^1H : 16 ^{19}F : 32	10 s 5 s	[36]
Applications in materials chemistry	Compact low-field NMR spectroscopy and chemometrics: A tool box for quality control of raw rubber	Styrene 1,2-butadiene <i>cis</i> -1,4-butadiene <i>trans</i> -1,4-butadiene	Not reported	^1H : 4 ^{13}C : 4096 (5.7 h) ^{13}C DEPT: 4096 (17 h)	15 s 5 s	[3]
Applications in food and agricultural chemistry	Authentication of beef versus horse meat using 60 MHz ^1H NMR spectroscopy	Beef meat Horse meat glyceryl tristearate (C18:0) glyceryl trioleate (C18:1) glyceryl trilinolenate (C18:3) stock mixture: 15% w/w C18:0 and 85% w/w C18:1	Not reported	Total acquisition time was maintained at ~10 m	Training set: 30 s Test samples: 2-30 s	[49]

60 MHz ¹ H NMR spectroscopy for the analysis of edible oils	Extra virgin olive oil Hazelnut oil	Capable of detecting adulteration of olive oil with hazelnut oil at levels >11% w/w	¹ H: 16	Not reported	[50]
High-throughput authentication of edible oils with benchtop Ultrafast 2D NMR	Olive oil Hazelnut oil Sesame oil Rapeseed oil Corn oil Sunflower oil	1D: Not reported 2D: >20:1	1D: 16 (2.4 m), dwell-time: 200 μs 2D COSY: 72 (2.4 m)	1D: 15 s 2D COSY: 2 s	[51]
A 43 MHz LF Benchtop Nuclear Magnetic Resonance Method to Discriminate Perilla Oil Authenticity	Perilla Oil Linoleic acid Unsaturated fatty acids Fatty acids Glycerol	Not reported	¹ H: 32 (4-5 m)	7 s	[52]
Is LF NMR a Complementary Tool to GC-MS in Quality Control of Essential Oils? A Case Study: Patchouli Essential Oil	Patchouli oil (13 varieties) Amyris oil Benzyl alcohol Benzyl benzoate Cedar wood oil Copaiva balsam Diethyl phthalate Dioctyl phthalate Gurjun balsam Hercolyn Isobornyl acetate Methyl benzoate Paraffin Paraffin viscid Pepper oil Propyleneglycol Ricinus oil Vetiver oil 1R -(-) myretenol	LOD = Patchouli oil 13% w/w	¹ H: 32	4 s	[53]

	Mobile compact ^1H NMR spectrometer promises fast quality control of diesel fuel	Commercial diesel Mineral diesel Diesel/biodiesel mixtures: <i>Soybean methyl biodiesel</i> <i>Soybean ethyl biodiesel</i>	Not reported	^1H : 1 (15 s)	Not reported	[55]
	Fast Sampling, Analyses and Chemometrics for Plant Breeding: Bitter Acids, Xanthohumol and Terpenes in Lupulin Glands of Hops	Lupulin gland extracts Cohumulone Colupulone mixture of humulone and adhumulone mixture of lupulone and adlupulone xanthohumol Myrcene Humulene Caryophyllene β -Farnesene aromadendrene α - and β -selinene germacrene B	Cohumulone signal not resolved from that of adhumulone at 43 MHz	^1H : 60	20 s	[56]
Reaction monitoring and chemical transformations	Continuous processing and efficient in situ reaction monitoring of a hypervalent iodine (III) mediated cyclopropanation using bNMR spectroscopy	Cyclopropanation: Hypervalent Iodine (III)-mediated synthesis of 1,1-dicyanocyclopropane. 4- <i>tert</i> -butylstyrene malonitrile	Not submitted	1D ^1H : Reaction monitoring: 5-180 minutes	10 s	[58]

Continuous-flow synthesis of fluorine-containing fine chemicals with integrated bNMR analysis	Krapcho decarboxylation; Ruppert-Prakash perfluoroalkylation of benzaldehyde; Dye-sensitised visible light C-H arylation. Benzaldehyde in THF; TMS-CF ₃ ; TMS-C ₂ F ₅ ; 3-(Trifluoromethyl)phenyl diazonium tetrafluoroborate; Eosin Y; 4-(trifluoromethyl) benzyl alcohol		¹⁹ F: 4,8,16,64	2 s	[59]
		Lowest recorded SNR = 4.9:1 at a concentration of 0.15 mol dm ⁻³ and flow rate of 4 mL min ⁻¹ . SNR > 20 is achieved for higher numbers of scans (>4) and concentrations, and lower flow rates.			
Process spectroscopy in microemulsions-setup and multi-spectral approach for reaction monitoring of a homogeneous hydroformylation process	Hydroformylation reaction: 1-dodecene; To tridecanal; <i>Catalysed by:</i> Rh coordination complex with SulfoXantPhos ligand	Not reported	Reaction monitoring: 1 spectrum per minute	N/A	[61]
Online LF NMR spectroscopy for process control of an industrial lithiation reaction—automated data analysis	Aniline 1-fluoro-2 nitrobenzene 2-nitrodiphenylamine Lithium bis(trimethylsilyl) amide	Not reported	1 scan	15 s	[62]

A bNMR spectrometer as a tool for monitoring mesoscale continuous-flow organic synthesis: equipment interface and assessment in four organic transformations	Acid-catalysed esterification Acetic acid Sulfuric acid Base-catalysed Knoevenagel condensation Benzaldehyde/Ethyl acetoacetate Diels-Alder reaction Piperdine Maleic anhydride Isoprene Alkylation reaction Salicylaldehyde/DBU 1-bromobutane	Not reported	10 scans	9 s	[65]
Medium Resolution ¹ H NMR at 62 MHz as a New Chemically Sensitive Online Detector for Size-Exclusion Chromatography (SEC-NMR)	Polystyrene Polymethylmethacrylate	From 27:1 (-CH ₂ -) to 110:1 (OCH ₃) for polymethylmethacrylate	¹ H: 128	0.5 s	[67]
Desktop NMR spectroscopy for real-time monitoring of an acetalization reaction in comparison with gas chromatography and NMR at 9.4 T	Acetalization Ethylene glycol <i>p</i> -nitrobenzaldehyde <i>p</i> -toluenesulphonic acid 2-(4-Nitrophenyl)-1, 3-dioxolane	Not reported	¹ H: 1	15 s	[70]

	Monitoring chemical reactions by low-field benchtop NMR at 45 MHz: pros and cons	<p>Fischer esterification Methanol Acetic acid Sulfuric acid</p> <p>Suzuki coupling Phenyl boronic acid 5-bromo-2-methoxypuridine Palladium(II) chloride Tripotassium phosphate Dimethylformamide</p> <p>Oxime reaction Sodium ((R)-2,2-dimethyl-1,3-dioxolan-4-yl)(hydroxy)methanesulfonate 2-methyltetrahydrofuran</p>	<p>Methanol: 253.9:1 to 168.9:1 Methyl acetate 38:1 to 156.9:1</p> <p>Methoxy signal starting material: 17.4:1 to 0:1 Methoxy signal product: 0:1 to 17.3:1</p> <p>Aldehyde doublet signal: 1:0 to 10.4:1</p>	<p>¹H: 8</p> <p>¹H: 50</p> <p>¹H: 50</p>	<p>8 s</p>	<p>[71]</p>
Structural and forensic chemistry	Diffusion-ordered spectroscopy on a benchtop spectrometer for drug analysis	Paracetamol Hypromellose Esomeprazole	5:1 (lowest recorded)	¹ H: 80 scans per experimental increment (total experimental duration: 2.84h)	4 s	[78]
	Gradient-based solvent suppression methods on a benchtop spectrometer	Lactate Alanine	Not reported	¹ H: 4	11.5 s at 0.8ml/m 6.5s s at 2.0ml/m	[79]
	Screening of “spice” herbal mixtures: From high-field to low-field proton NMR	Synthetic cannabinoids	3:1	¹ H: 128 (7.6 m)	2 s	[81]

Desktop NMR for structure elucidation and identification of strychnine adulteration	Strychnine (S)	Not reported		¹ H: S 4 ¹ H SHY: 64 ¹ H SHE: 4	¹ H: 15 s	[66]
	Strychnine hydrochloride (SHY)			¹³ C: 4096 2D COSY: 4 2D HETCORE: 128 2D HSQC: 64 2D HMBC: 64 2D J-r: 4	¹³ C: 5 s 2D COSY: 2.2 s 2D HETCORE: 10 s 2D HSQC: 10 s 2D HMBC: 10 s 2D J-r: 2 s	
	Strychnine hemisulphate (SHE)					
Potential biofluid screening and metabolomics applications	LF bNMR Spectroscopy as a Potential Tool for Point-of-Care Diagnostics of Metabolic Conditions: Validation and Protocols	Glucose Acetone	11:1 >10:1	¹ H: 64	10 s, 15s	[12]

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