Fact and fallacy in measurement of conduction velocity in motor nerves

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It is necessary to examine the possible fallacies before one can accept any statement as 'fact'. This paper will attempt to assess the importance of the more obvious fallacies inherent in the method of measurement of conduction velocity as determined by length and latency measurements and will not take account of possible biological variables such as age of the subject. Only the method of computation by length and latency differences between two stimulated points on a nerve will be considered, as the latency of muscle response from a single peripheral stimulus includes unknown variables, including slowing due to terminal branching, conduction by non-myelinated terminals (at a rate of the order of 0·3 m./sec. according to Katz and Miledi, 1963), junctional transmission, and propagation from motor end plates to the recording electrodes. Comparisons of latency may usefully be discussed, but not velocity (Simpson, 1956). These difficulties are eliminated if the evaluation is limited to a stretch of nerve which can be stimulated at two points, since the region of uncertainty is excluded from the calculations, but certain methodological difficulties remain.

MEASUREMENT OF LATENCY

The method of calculation requires accurate computation of the time of stimulation at some point on the nerve and of the response at some distant part,

\[ + \quad - \]

\[ \begin{array}{c}
1 \\
2 \\
3
\end{array} \]

\[ A \quad B \quad C \]

FIG. 1. A Triggering voltage too positive.
B Correct triggering voltage.
C Triggering voltage too negative. Note trigger artefact at 1 msec. which may be mistaken for the stimulus artefact.
1 Free-running time scale. Sweep triggered by stimulator.
2 Time scale and sweep both triggered by stimulator.
3 Time scale and sweep triggered by pre-pulse 3 msec. before stimulus.

Errors due to faulty triggering in oscilloscopes with variable trigger or synchronizing polarity and voltage control do not matter if the stimulus is delayed (3A, B, C). If the stimulus initiates the sweep (2, 3) errors are only obvious if the time scale is also triggered (2A, C). Only 2B is acceptable.

TEMPERATURE

The conduction velocity of human peripheral nerve varies widely with temperature (Helmholtz and Baxt, 1867, 1870). Henriksen (1956) calculated that the mean alteration in conduction velocity of the human ulnar nerve was 2·4 m./sec. per degree Centigrade change in temperature. In the conditions of clinical examination this could introduce an error of up to 6 m./sec. Fortunately the error is systematic and can be prevented by suitable precautions. There may, however, be a temperature gradient along the axis of the limb, especially with peripheral wasting. This may account for the common finding that conduction is slower in the distal segment of a limb and it is difficult to apply a correction for this factor.
either of a muscle fibre or of the compound action potential of the nerve. It is well known that the stimulus pulse is distorted by tissue capacitance but this should not introduce a significant error. Undetected difficulties may arise with inferior recording equipment when the oscilloscope sweep is triggered by the stimulator (Fig. 1). Even if there is no triggering delay, the start of the sweep may be blacked out, or the spot may be momentarily deflected in the wrong direction. These defects may obscure the first two milliseconds and will not be detected if the beam is blanked out between sweeps. Fortunately it is likely to be a systematic error which will not affect the difference between two latencies, but could prevent recognition of distal slowing as in the carpal tunnel syndrome. It is preferable that the stimulus should be delayed after the start of the sweep, and the artefact reduced to a small spike with clearly defined onset. If this is not possible the artefact will be more obvious if the time scale is not free-running but is also started by the triggering pulse.

The determination of the response is more difficult. Motor fibres can only be determined in a mixed nerve by using the muscle response as an index, but it is a complex action potential due to the different velocities of conduction of motor fibres of different calibres. The most readily identified motor units are those with the shortest latency and so the most reliable velocity measurements are for the fastest conducting fibres, but the problem that arises is to decide when the response starts. With surface recording the origin of the action potential of the muscle is curved, making it difficult to decide which point to measure. Recording with high gain minimizes the difficulty but does not remove it (Fig. 2). Recording with a needle electrode suitably positioned makes it easier to determine the onset of the recorded response and has the added advantage of localizing the recorded activity to the desired muscle, but it is theoretically possible that the shortest latency units might be too remote from the electrode to be recorded (Fig. 3). This type of measuring error is not systematic, leading to under- or overestimate of true latency, but has a tendency to cause apparent prolongation. The total error of time measurement may amount to 1 msec. but is usually less.

There are two other possible errors of timing. If the pickup electrode is not close to the end-plate zone of muscle there will be an added delay not exceeding 0·35 msec. due to conduction along muscle fibres (Henriksen, 1956). This is, however, a systematic variation affecting both readings equally and so not affecting the calculated velocity between two points of stimulation. The other difficulty is introduced by a small negative wave which is some-

![Figure 2. Stimulus artefact and initial deflection of muscle response at amplifier gain (a) suitable for displaying the whole of the response, (b) increased 18dB for inspection of initial deflection, (c) both traces superimposed. Surface electrodes.](http://jnnp.bmj.com/)

![Figure 3. Evoked response of muscle recorded by (a) coaxial needle and (b) surface electrode near the needle, and (c) on skin 2 cm. away. Each response is superimposed in (d). Time scale in milliseconds.](http://jnnp.bmj.com/)
times recorded immediately before the main muscle complex (Fig. 4). It may be a nerve action potential in the terminal fibres. This is uncertain, but its importance lies in the fact that it is not recorded from all parts of the muscle and may only be seen with high amplification. If the amplifier gain is changed because the muscle potential evoked from one point on the nerve is smaller, the early wave may be taken as the first response to one stimulus and not to the other. Provided it is included in both measurements no difficulty arises in calculating latency difference as this early wave appears to precede the main deflection by a constant amount. It will obviously affect the interpretation when only a single latency measurement is available as in the carpal tunnel syndrome. The nature of this wave requires further investigation.

A possible solution to the difficulty of precise measurement might be to use as a reference point a clearly defined peak of the compound muscle action potential. A moment's consideration will show that this reference point cannot be used. The muscle action potential evoked by indirect stimulation is 10 to 15 msec. in duration. The response is not synchronous in every unit because the contributing motor nerve fibres have different conduction velocities. As a result the compound muscle potential is more dispersed when its nerve is stimulated at a distance than when near the muscle (Bolzani, 1954). The difference is less than might be predicted, and Merton (1954) attributes this to artificial synchronization of propagation in different fibres of the muscles by the electrical stimulus. Even with direct recording of the nerve potential, the temporal dispersion is sufficient to make the latency differences significantly greater if measured to the first peak than when measured to the first trough (Dawson, 1956). The position of positive and negative peaks, and indeed the overall duration of the recorded nerve action potential depends on the distance between the recording electrodes. The same principle applies to surface recording of the compound muscle action potential but is minimized by placing the distal electrode over inactive tendon (belt-tendon derivation). More precise definition of the onset of the muscular response can be obtained by a suitably placed intramuscular needle electrode, with the additional advantage of greater certainty that the recorded response truly originates in the intended muscle. A possible fallacy is that there may be no motor unit innervated by one of the fastest nerve fibres within the pick-up range of the needle. However, the ability to recognize individual units makes it possible to study the conduction velocity of slower nerve fibres. This may be compromised by the possible muscle synchronization effect referred to above. It is never safe to consider any particular fibre as pathologically slow unless the muscle unit evoked by it fires later than the end of a normal compound action potential. There may simply be a fall-out of the fastest fibres, such as Hodes (1949) suggested occurred in poliomyelitis. This is particularly important with surface recording since it is a common experience that no 'slowing' can be observed until there is a marked fall in the amplitude of the compound muscle potential (Henriksen, 1956). Another difficulty is that nerve fibres may fire iteratively after a single shock stimulus, particularly in compressive neuropathies (Pinelli, 1954; Simpson, 1956). It is often impossible to be certain that the units recorded at the end of a compound muscle potential are not repetitions of units which have already fired with normal latency. This should be suspected if the late unit selected for measurement is separated from the previous unit by the same amount, irrespective of the site of stimulation, or if its latency from one stimulus site shows slight variability.

**THE STIMULUS**

If the only reliable units for study are those with the shortest latency, it is most important to ensure that the nerve stimulus is above threshold for all fibres
and in practice it should be supramaximal by at least 30%. If the stimulus intensity is gradually increased from below threshold it will be seen that the lowest threshold fibres, though usually rapid, are not always the fastest, since earlier units are recruited by increasing the stimulus (Fig. 5). At near-maximal stimulus intensity there may be seen a continuous decrease in latency and this becomes still shorter even after the muscle action potential has ceased to increase in size (Dawson, 1956). This is probably due to spread of the stimulus from the cathode.

The factors so far considered refer to the identification of a particular motor unit and the determination of its latency. It must be assumed for the purpose of discussion that an accurate time standard is available to be photographed simultaneously with the evoked potential, that the two spots synchronize in sweep speed when separated on the oscilloscope screen, and that accurate means of measuring the photograph are available. When all these factors of identification, recording, and measurement are taken into account, an error of 0-5 msec. in each latency measurement is well within the unavoidable error of the method, so that a latency-difference of 1-0 msec. might commonly occur. In the conditions of clinical recording it may well be greater.

**CONDUCTION DISTANCE**

The uncertainty of the length measurement is still more serious. Unless the stimulating cathode is placed exactly over the nerve (as identified by the point where a muscle response can be evoked with the least stimulus intensity) the measurement must be faulty. Even then there is the possibility that the true cathode is at some distance from the apparent one. Experiments by Henriksen (1956) suggest that this is not an important consideration and that the true cathode may be considered as lying beneath the centre point of a circular cathodal electrode, whatever its size. Nevertheless this need not be true for supramaximal stimulation where the ohmic spread of current may be sufficient to cause a virtual cathode some distance away, and the shortened latency described above suggests that this does occur. The uncertainty about cathodal position is trivial compared with the fact that it is not possible to make a direct measurement of the length of nerve stimulated. Uncertainty increases as the nerve assumes an irregular course, especially in the hand or foot. It is impossible to measure true velocity in this part of a limb nerve and only comparison of latencies is possible (Simpson, 1956). Fortunately the main nerve trunks in the limbs have comparatively straight courses. Length estimations made by surface measurement must only be approximate. Carpendale (1956) states that the surface measurement is surprisingly close to the true length of the nerve between cathodes provided that the measurement is made with the limb in the same position as it was when stimulated. This may be true (it requires confirmation), but a more serious difficulty is the observer error. Observers may differ by 1 cm. in measuring the distance between two points on a limb if instructed to lay the tape along the course of a nerve, especially one such as the ulnar which winds from dorsal to ventral surface of the limb.

**CALCULATED VELOCITY**

These factors have been investigated by most workers in the field including the writer. Henriksen (1956) and Carpendale (1956) concluded that none was significant except the temperature effect and even that would only modify the calculated velocity by about 2 m./sec., if reasonable precautions were taken. These conclusions assume that only one factor is operative at a time, but the picture is different if we consider that with reasonable technique there may be a simultaneous and unavoidable

<table>
<thead>
<tr>
<th>Latency-difference (msec.)</th>
<th>Length-difference (mm.)</th>
<th>Conduction velocity (m/sec.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>True</td>
<td>4-5</td>
<td>55-6</td>
</tr>
<tr>
<td>Measured</td>
<td>5-5</td>
<td>44-5</td>
</tr>
<tr>
<td>True</td>
<td>3-6</td>
<td>55-6</td>
</tr>
<tr>
<td>Measured</td>
<td>4-6</td>
<td>42-4</td>
</tr>
</tbody>
</table>
error of $+1.0$ msec. in latency-difference and 
$\pm 0.5$ cm. in length measurement.

It will be seen from the table that the unavoidable
error becomes proportionately greater as the length
of nerve is reduced. Fortunately errors acting in the
same direction will be comparatively rare; nevertheless
Henriksen (1956) found that measurements of
the conduction velocity of the ulnar nerve of normal
adults repeated on different days could vary by up
to 7.5 m./sec. (in 17 cases). In unfavourable condi-
tions the discrepancy could easily be as great as that
shown in the table. It may be concluded that no
‘borderline’ values should be accepted until con-
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