BMJ Open

European Working Time Directive and doctors’ health: a systematic review of the available epidemiological evidence

Maria Cruz Rodriguez-Jareño,1,2 Evangelia Demou,3,4 Sergio Vargas-Prada,5 Kaveh A Sanati,4,6 Alenka Škerjanc,7 Pedro G Reis,8 Ritva Helimäki-Aro,9 Ewan B Macdonald,4 Consol Serra,1 5 10 11 on behalf of the UEMS Section of Occupational Medicine

ABSTRACT

Objective: To summarise the available scientific evidence on the health effects of exposure to working beyond the limit number of hours established by the European Working Time Directive (EWTD) on physicians.

Design: A systematic literature search was conducted in PubMed and EMBASE. Study selection, quality appraisal and data extraction were carried out by independent pairs of researchers using pre-established criteria.

Setting: Physicians of any medical, surgical or community specialty, working in any possible setting (hospitals, primary healthcare, etc), as well as trainees, residents, junior house officers or postgraduate interns, were included.

Participants: The total number of participants was 14 338.

Primary and secondary outcome measures: Health effects classified under the International Classification of Diseases (ICD-10).

Results: Over 3000 citations and 110 full articles were reviewed. From these, 11 studies of high or intermediate quality carried out in North America, Europe and Japan met the inclusion criteria. Six studies included medical residents, junior doctors or house officers and the five others included medical specialists or consultants, medical, dental, and general practitioners and hospital physicians. Evidence of an association was found between percutaneous injuries and road traffic accidents with extended long working hours (LWH)/days or very LWH/weeks. The evidence was assessed as low or insufficient for mood disorders and general health.

No studies on other health outcomes were identified.

Conclusions: LWH could increase the risk of percutaneous injuries and road traffic accidents, and possibly other incidents at work through the same pathway. While associations are clear, the existing evidence does not allow for an established causal or ‘dose–response’ relationship between LWH and incidents at work, or for a threshold number of extended hours above which there is a significantly higher risk and the hours physicians could work and remain safe and healthy. Policymakers should consider safety issues when working on relaxing EWTD for doctors.

INTRODUCTION

The European Working Time Directive (EWTD),1 applicable to all occupations across the European Union (EU), requires a maximum working week of 48 h and establishes rest periods.2 3 Since it came into force in healthcare in 2009, it has been associated with concerns about the provision of health services including continuity of care, lower staffing levels, introduction of shift working, a reduction in training time and the adequate supervision of junior doctors.4 5 The large intercountry variations in physicians’ working hours as well as the mandatory controls over working hours across occupations raise further questions as to the...
basis of selecting and setting these limits and restrictions. There is evidence of variable compliance to the EWTD within healthcare across Europe. The medical profession is also increasingly feminised with potential implications for women of childbearing age, which may coincide with the time in the profession where long working hours (LWH) are more likely to happen (ie, during residency). Constraint on public sector finances and increasing healthcare demands have stimulated the debate on physicians’ working hours, with some evidence showing that the quality of healthcare is positively correlated with the number of physicians available to deliver that care.

The Union of European Medical Specialists (UEMS) was created in 1958 to represent medical specialists in the EU, promote a high standard of training and practice at European level and facilitate the free movement of physicians across European countries. The Occupational Medicine Section was created in 1997 and is involved in issues related to specialist training, professional practice and research; elaborating and publishing reference documents, conducting and contributing to surveys and working in partnership with other key European groups.

The aim of this project was to systematically review whether LWH, defined as more than the 48 h/week limit imposed by the EWTD, are associated with health effects, classified under the International Classification of Diseases (ICD-10), in physicians, and to examine what these associations are. This review does not include the potential impact of LWH on patient care or physician training.

METHODS

The systematic review sought to respond to the following research question: Are LWH associated with health effects in physicians? For the purpose of this review, exposure to LWH was defined as working for more than 48 h/week. Health outcomes included any disease as defined in the ICD-10 and work-related injuries. If self-assessed tools on health status were used, only those validated were considered. Burnout, stress and outcomes such as individual symptoms, signs or biological markers (blood pressure, ECG, etc) were not considered. Physicians of any medical, surgical or community specialty working in any possible setting (hospitals, primary healthcare, etc) as well as trainees, residents, junior house officers or postgraduate interns, were included.

Study identification

Electronic searches were carried out using PubMed and EMBASE as search engines (December 2011). Our search strategy was similar in both databases and consisted of a combination of the following keywords and MeSH terms: night shift, morning shift, evening shift, afternoon shift, shift work, rotating shift, shift combination, shift duration or length, shift system, clockwise rotation, shift roster, extended shifts, night work, evening work, work schedule, work hours, starting time, early start, irregular working hours, direction of rotation, overwork, extended hours, shift rota, workload, work schedule tolerance, sleep deprivation, sleep disorders, chronobiology disorders, circadian rhythm, psychomotor performance, circadian disruption, vigilance, attention, wakefulness, drowsiness, fatigue, insomnia, hypersomnia, dyssomnia, eveningness, morningness, neurocognitive performance, concentration difficulties, arousal, health, morbidity, mortality, disease, illness, stress, strain, distress, accident, injury, death, suicide, education, medical, physician, medical staff, hospital, doctor, surgeon, house officers, medical school, surgery, surgical.

In addition, the reference lists of articles selected for inclusion were carefully reviewed to identify additional studies.

Study selection

Studies were included if they measured the association between the exposure to LWH and health effects in...
physicians, and used an observational epidemiological (ie, cohort, case–control, cross-sectional) or experimental design. Articles relating to on-call, night or shift work, but with unknown exposure or exposed to less than 48 working hours per week were excluded. They were also excluded if the working groups were other than physicians or addressed other exposures or outcomes (ie, patient safety, fatigue, sleep deprivation, social or family disruption). No limitation was set for languages.

A total of 2036 citations were obtained from PubMed and 980 non-duplicated citations using EMBASE, yielding a total of 3016 citations that were all screened by independent pairs of reviewers. All titles and, where necessary, abstracts were reviewed. Ninety-two potentially suitable publications were identified from the electronic searches, and 18 further studies were identified from the reference lists, yielding a total of 110 studies published in English, French, German, Italian, Slovenian and Spanish. For all of them, the full text was obtained and reviewed by independent pairs of reviewers. Disagreements within pairs were resolved by consensus or, where necessary, by a third reviewer who made the final decision. All authors participated as reviewers for screening the citations and full papers using well-defined and pre-established criteria. Ninety-nine articles were excluded at this stage. One further study was identified that was published after the search period. Finally, 12 papers were considered for quality assessment (figure 1).

The methodological quality of the 12 studies was assessed by independent pairs of reviewers. A standardised 16-point scale based on CONSORT and STROBE statements and adapted from a previous systematic review36 was used. It includes 16 items grouped into six areas: (1) objectives; (2) study design; (3) target population and sample; (4) variables; (5) data sources, collection and measurement and (6) statistical methods. Each item was rated as 1 (the requirement was met), 0.5 (the requirement was partially met) or 0 (the requirement was not or unclearly met). Disagreements within pairs of reviewers were resolved by consensus or, where necessary, by a third reviewer. For each of the 12 studies, a final score based on the sum of all items was assigned and the percentage was calculated based on a maximum score of 16. Study quality was rated as low, moderate or high if it scored less than 60%, between 60% and 79.9%, and 80% or more of the maximum score, respectively.

One article of low quality was excluded,37 and a total of 11 studies of intermediate or high methodological quality were included for the purpose of this review (figure 1).

Data extraction
Selected information was extracted from each paper, including publication year, country of origin, study design, setting, study population, sample size, response rate, measure of working hours, definition used for LWH, health outcomes and their measurement, main results on point risk estimates or frequencies of health outcomes, their corresponding 95% CI and whether the analyses were adjusted for potential confounders.

Evidence synthesis
To summarise the results on the relationship between LWH and health effects, levels of evidence synthesis was performed. This was based on the methodological quality, study design and consistency of the study outcomes. The following criteria were based on two previous relevant systematic reviews:38 39

Strong evidence: consistent results in more than two studies of high quality.

Moderate evidence: consistent results in one high-quality study and one intermediate, or between some studies of intermediate quality.

Insufficient evidence: identification of only one study or inconsistent results across studies.

Figure 1 Results of the search strategy, using search engines on PubMed and EMBASE (December 2011) and screening of references lists of identified full papers, study selection and quality assessment.
Evidence of no association: consistent results of a non-association in two or more studies.

A measure of the possible magnitude of the association was attempted using the following criteria based on the association point estimate (RR=relative risk, rate ratio or OR) when available; otherwise the comparison between two frequencies was used:

- High: RR >3.00;
- Intermediate: RR=1.50–3.00;
- Low: RR=1.01–1.49;
- No association: RR=1.00;
- Unclear.

RESULTS
We identified 11 studies related to LWH and health effects in physicians that were eligible for inclusion. The outcome of the methodological quality assessment is given in table 1.

Methodological quality was appraised as high for 3 of the 11 studies, with scores ranging from 84% to 94%. The other eight studies were considered as intermediate quality as their scores ranged from 63% to 78%. The majority of all included studies received positive scores on items describing the study objectives, design and the study population (items from areas 1, 2 and 3), as well as the description of exposure and outcome variables (item 4). However, although data sources and collection were well described, reliability and validity of exposure to LWH and health outcome measures had low scores (items 5b and 5c). The statistical methods were in general appropriate; however, the confounding variables were not measured in several studies and, as can be seen, the studies did not describe how they addressed missing data (items 6a and 6b).

Study characteristics
The characteristics of the included studies are described in table 2.

Five had been carried out in North America (four in the USA and one in Canada), five in European countries (Denmark, Germany, Norway and Germany, Sweden and the UK) and one study in Japan. Four used a prospective cohort design (follow-up ranging from 6 to 20 months), two were case-crossover, four cross-sectional and one was a before and after study. Overall, the total number of participants was 14,338 and included medical residents, junior doctors or house officers only working in hospitals, medical specialists or consultants, medical and dental practitioners, general practitioners, hospital physicians and one study included female physicians from a community service. Most studies (n=9) compared different working hour patterns within the same group of physicians, one compared physicians from two different countries and one included other healthcare workers as the comparison group.

IWH was defined as more than 48 h/week in most studies, and some did not provide a definition. Two studies considered LWH below 48 h/week. All the included studies investigated health effects associated with working more than 48 h/week, with number of hours ranging between <50 and 110 h/week. One paper studied the effects of an increasing number of hours from 30 to more than 50 h/week. Others studied more than 40, 47 and above 58 h/week (two studies), and six studies referred to more than 70 h/week, one of which compared 78–80 h/week with 90–110 weekly hours. Information on working hours was self-reported in nine studies, with reasonably good validity only in two studies and reliability in one. In the other two studies, it was obtained from established work schedules.

Identified health outcomes that fulfilled the inclusion criteria were percutaneous injuries, motor vehicle accidents, mood disorders and general health. Information was self-reported in all studies using well-known validated questionnaires, except three studies that used documented information on motor vehicle crashes, reported percutaneous injuries and antidepressant prescription data as a surrogate of depression.

Summary of findings
Table 3 shows the findings from each of the 11 studies.

Percutaneous injuries: One study of high quality and one of intermediate quality, with non-adjusted analyses, showed consistent results in medical trainees. Ayas et al found an increased risk of percutaneous injuries associated with working more than 20 h/day compared with non-extended hour periods, except in intensive care units. Fisman et al found an association between self-reported fatigue and the number of working hours at the time of injury and a threefold increased risk of sharp injuries in medical trainees compared with other healthcare workers (median of working hours/week before the injury: 70 vs 40; p<0.05).

Motor vehicle accidents: Two studies of high and intermediate quality showed that long working weeks were associated with an at least twofold increased risk. Barger et al found adjusted increased risks for car crashes (OR 2.3; 95% CI 1.6 to 3.3) and near miss accidents (OR 5.9; 95% CI 5.4 to 6.3) associated with working extended shifts. Kirkcaldy et al showed that non-adjusted incidence rates of traffic accidents on house visits was five times as much when physicians worked for a mean of 58 h/week compared with 38 h/week (p<0.05).

Mood disorders: Six studies of intermediate and high quality found contradictory results for mood disorders. Three of them, all of intermediate quality, provided evidence of an association between LWH and mental health problems. The prevalence of General Health Questionnaire (GHQ)-30 cases was higher for female physicians working above 50 h/week than for those working 30 or less hours/week (p<0.05), and an increased adjusted risk of impaired mental health was...
<table>
<thead>
<tr>
<th>Study ID</th>
<th>Ayas et al\textsuperscript{40}</th>
<th>Barger et al\textsuperscript{41}</th>
<th>Firth-Cozens\textsuperscript{\textdagger}</th>
<th>Fisman et al\textsuperscript{43}</th>
<th>Hayasaka et al\textsuperscript{44}</th>
<th>Kirkcaldy et al\textsuperscript{45}</th>
<th>Rosta and Aasland\textsuperscript{46}</th>
<th>Stamp et al\textsuperscript{47}</th>
<th>Sundquist and Johansson\textsuperscript{48}</th>
<th>Varma et al\textsuperscript{49}</th>
<th>Zahrai et al\textsuperscript{50}</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Objectives</td>
<td>Are the objectives or hypotheses of the research described in the paper?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>2 Study design</td>
<td>Is the study design presented?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>3a Target population</td>
<td>Do the authors describe the target population they wanted to research?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>3b Sample</td>
<td>Was a random sample of the target population taken? AND was the response rate 60% or more?</td>
<td>0.5</td>
<td>0.5</td>
<td>1</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>1</td>
<td>1</td>
<td>0.5</td>
</tr>
<tr>
<td>3c Sample</td>
<td>Is participant selection described?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0.5</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>3d Sample</td>
<td>Is participant recruitment described, or referred to?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>3e Sample</td>
<td>Are the inclusion and/or exclusion criteria stated?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0.5</td>
<td>0.5</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0.5</td>
</tr>
<tr>
<td>3f Sample</td>
<td>Is the study sample described? (minimum description: sample size, gender, age and occupation)</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>4 Variables</td>
<td>Are the measures of long working hours and the health outcome described?</td>
<td>1</td>
<td>1</td>
<td>0.5</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>5a Data sources, collection</td>
<td>Do authors describe the source of their data (eg, official registry, health survey) AND how were the data collected?</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>5b Measurement</td>
<td>Was reliability of the measure(s) of long working hours mentioned or referred to?</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>5c Measurement</td>
<td>Was the validity of the measure(s) of long working hours mentioned or referred to?</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>5d Measurement</td>
<td>Were health outcomes assessed by objective measures or validated self-reporting instruments?</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>6a Statistical methods</td>
<td>Were appropriate statistical methods used and described, including those for addressing confounders?</td>
<td>0.5</td>
<td>1</td>
<td>0.5</td>
<td>1</td>
<td>0.5</td>
<td>0.5</td>
<td>1</td>
<td>0.5</td>
<td>1</td>
<td>0.5</td>
</tr>
<tr>
<td>6b Statistical methods</td>
<td>Were the numbers/percentage of participants with missing data for long working hours and the health outcome indicated AND if more than 20% of data in the primary analyses were missing, were methods used to address missing data?</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0.5</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
</tbody>
</table>

| Total score* (%) | 12.0 (75) | 13.5 (84) | 11.0 (69) | 13.5 (84) | 11.0 (69) | 10.0 (63) | 10.5 (66) | 12.5 (78) | 12.0 (75) | 15.0 (94) | 11.0 (69) |
| Quality rate† (%) | Low=<60; intermediate (interm)=60–79; high=80–100. |

*Maximum score=16. †Quality rate=16.
Table 2  Characteristics of the included studies

<table>
<thead>
<tr>
<th>Study ID</th>
<th>Country of study population</th>
<th>Design*</th>
<th>Setting</th>
<th>Participants and sample size (response rate)</th>
<th>Working hours</th>
<th>Health outcomes</th>
<th>Quality score (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ayas et al</td>
<td>USA</td>
<td>Cohort, prospective (1 year)/case cross-over</td>
<td>Hospital</td>
<td>Interns in postgraduate residency programs Sample size: 2737 (80%)</td>
<td>Mean (SD) hours worked/month: 249.8 (75.3). Self-reported, monthly survey. Strong correlation (Pearson r=0.76; p 0.001) with hours worked 244 (69.3) from work diaries of randomly selected subset of 192 interns. Extended periods (20 or more consecutive hours) vs non-extended periods (12 h or less consecutive hours)</td>
<td>Self-reported percutaneous injuries</td>
<td>12.0 (75)</td>
</tr>
<tr>
<td>Barger et al</td>
<td>USA</td>
<td>Cohort, prospective (1 year)</td>
<td>Hospital</td>
<td>Residents first postgraduate year (interns) Sample size: 2737 (80%)</td>
<td>Mean (SD) hours worked/week 70.7 (26.0). Extended shift (≥24 h) vs non-extended shift (&lt;24 h). Self-reported, validated</td>
<td>Self-reported and documented motor vehicle crashes</td>
<td>13.5 (84)</td>
</tr>
<tr>
<td>Firth-Cozens</td>
<td>UK</td>
<td>Cohort, prospective (1 year)</td>
<td>Hospital</td>
<td>Junior house officers Sample size: 170 (72%)</td>
<td>Mean number of hours/week=90.6 h (include on call). Self-reported</td>
<td>GHQ-12 (case: score ≥2) and SCLDS, self-reported</td>
<td>11.0 (69)</td>
</tr>
<tr>
<td>Fisman et al</td>
<td>USA and Canada</td>
<td>Case-crossover</td>
<td>Hospital</td>
<td>Medical trainees vs other HCW. Sample size: 109 vs 241 (46%)</td>
<td>Median number of hours per week: medical trainees=70; other HCW=40 (p&lt;0.001). Self-reported, high reliability</td>
<td>Reported percutaneous injuries to employee healthcare service</td>
<td>13.5 (84)</td>
</tr>
<tr>
<td>Hayasaka et al</td>
<td>Japan</td>
<td>Cross-sectional</td>
<td>Hospital, clinics, other</td>
<td>Female physicians Sample size: 367 (63%)</td>
<td>Comparison of increasing number of hours/week from ≤30 to &gt;, self-reported</td>
<td>GHQ-30 (case: score ≥8), self-reported</td>
<td>11.0 (69)</td>
</tr>
<tr>
<td>Kirkcaldy et al</td>
<td>Germany</td>
<td>Cross-sectional</td>
<td>Not specified</td>
<td>Medical and dental practitioners. Sample size: 2500 (not specified)</td>
<td>Long hours: mean 58.36 h/week SD 9.16 Short hours: mean 38.17 h/week SD 7.72 Self-reported</td>
<td>Self-reported traffic accidents</td>
<td>10.0 (63)</td>
</tr>
<tr>
<td>Rosta and Aasland</td>
<td>Germany and Norway</td>
<td>Cross-sectional</td>
<td>Hospital</td>
<td>Hospital physicians Participants: 1917 (58%) in Germany and 1072 (65%) in Norway answered the questionnaire. From survey responders, 1260 (65.7%) Germans and 562 (52.4) Norwegians were included in the analysis after applying further inclusion criteria</td>
<td>German vs Norwegian physicians (%): Hours per day ≥9=58.8 vs 26.7 60 h on-call per month=63.4 vs 18.3 Self-reported</td>
<td>Validated questionnaire on self-rated health</td>
<td>10.5 (66)</td>
</tr>
</tbody>
</table>

Continued
### Table 2  Continued

<table>
<thead>
<tr>
<th>Study ID</th>
<th>Country of study population</th>
<th>Design*</th>
<th>Setting</th>
<th>Participants and sample size (response rate)</th>
<th>Working hours</th>
<th>Health outcomes</th>
<th>Quality score (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stamp et al 47</td>
<td>USA</td>
<td>Before-and-after</td>
<td>Hospital</td>
<td>Residents of general surgery Sample size: 28 (97%)</td>
<td>Changes of work patterns, after reduction from 90–110 to 78–80 work hours/week</td>
<td>SF-36, mental health; BDI, depression; self-reported</td>
<td>12.5 (78)</td>
</tr>
<tr>
<td>Sundquist and Johansson 48</td>
<td>Sweden</td>
<td>Cross-sectional</td>
<td>Primary care</td>
<td>General practitioners Sample size: 1004 (72%)</td>
<td>Overtime defined as working at least 47 h/week. Self-reported</td>
<td>Swedish SF-36, impaired mental health, self-reported</td>
<td>12.0 (75)</td>
</tr>
<tr>
<td>Varma et al 49</td>
<td>Denmark</td>
<td>Cohort, prospective (20 months)</td>
<td>Hospital</td>
<td>Senior medical consultants Sample size: 2790 (62%)</td>
<td>Long work hours (&gt;40 h/week). Self-reported</td>
<td>Depression: redemption of anti-depressive drug prescriptions</td>
<td>15.0 (94)</td>
</tr>
<tr>
<td>Zahrai et al 50</td>
<td>Canada</td>
<td>Cohort, prospective (6 months)</td>
<td>Hospital</td>
<td>Orthopaedic surgery residents Sample size: 16 (not specified)</td>
<td>Night float (n=9): 77.8% did &gt;80 h/week at baseline; 71.4% at follow-up Standard call (n=7): 57.1% did &gt;80 h/week at baseline; 80% at follow-up</td>
<td>SF-36, mental health score, self-reported</td>
<td>11.0 (69)</td>
</tr>
</tbody>
</table>

*Follow-up period in brackets for prospective cohort studies.
BDI, Beck Depression Inventory II; GHQ-12, General Health Questionnaire-12 items; HCW, Health Care Workers; SCLDS, Symptom Checklist Depression Scale.
<table>
<thead>
<tr>
<th>Health outcome</th>
<th>Main results</th>
<th>Adjustment by confounders</th>
<th>Study ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percutaneous injuries</td>
<td>OR (95% CI) for injuries during extended (&gt;20 h/day) vs non-extended periods (first 12 h at work): all percutaneous injuries: 1.61 (1.46 to 1.78); injuries reported to OH: 1.83 (1.48 to 2.28); injuries in the ICU: 1.87 (0.69 to 5.04); injuries in the operating room or labour and delivery: 1.77 (1.49 to 2.09); injuries in the ICU, non-ICU or ED: 2.17 (1.56 to 3.00)</td>
<td>Time of day and circadian influences</td>
<td>Ayas <em>et al</em>[^3]</td>
</tr>
<tr>
<td>Percutaneous injuries</td>
<td>Medical trainees vs other healthcare workers: total median working hours: 70 vs 40 h (p&lt;0.001); median previous working hours at the time of injury: 6.5 vs 5 h (p&lt;0.001). OR (95% CI) for self-reported fatigue: work &gt;40 h in the past week: 3.50 (2.06 to 5.92); work during more than 5d in past week: 4.20 (4.46 to 7.15); &gt;12 h at work before injury: 8.58 (3.7 to 19.86). Incidence rate ratio (95% CI) for association between fatigue and injuries: all workers: 1.40 (1.03 to 1.90); medical trainees: 2.94 (1.71 to 5.07); other HCWs: 0.97 (0.66 to 1.42; p=0.001 for heterogeneity)</td>
<td>Age and sex</td>
<td>Fisman <em>et al</em>[^3]</td>
</tr>
<tr>
<td>Motor vehicle accidents</td>
<td>ORs (95% CI), after extended shift (≥24 h) vs non-extended shift (&lt;24 h): Crashes=2.3 (1.6 to 3.3); near miss accidents=5.9 (5.4 to 6.3)</td>
<td>Age and sex</td>
<td>Barger <em>et al</em>[^1]</td>
</tr>
<tr>
<td>Motor vehicle accidents</td>
<td>Incidence rates of driving accidents on house visits: Long hours (mean 58.36 h/week)=0.10 (0.31) vs short hours (mean 38.17 h/week)=0.02 (0.18). p&lt;0.05 Not adjusted</td>
<td>Kirkcaldy <em>et al</em>[^45]</td>
<td></td>
</tr>
<tr>
<td>Mood disorders, depression</td>
<td>HR (95% CI) hours/week intervals and redemption of antidepressive drug prescription (reference group: 37–40 h/week): 25–36 h: 0.83 (0.24 to 2.82); 41–44 h: 0.95 (0.50 to 1.77); 45–49 h: 0.88 (0.43 to 1.78); 50–54 h: 0.83 (0.32 to 2.14); 55–59 h: 0.67 (0.15 to 2.94); &gt;60 h: 0.48 (0.06 to 3.68). Cox regression analysis of work hours as a continuous variable: HR=0.93 (0.76 to 1.13)</td>
<td>Age, sex, marital status, medical specialty, decision authority at work, social support at work, quantitative work demands and previous redemption of AD drug prescription</td>
<td>Varma <em>et al</em>[^9]</td>
</tr>
<tr>
<td>Mood disorders, GHQ-30</td>
<td>Prevalence of cases by working time (hours/week) ≤30 h: 35.7% (41/115); &gt;30–40 h: 39.0% (39/100); &gt;40–50 h: 37.7% (26/69); &gt;50 h: 56.8% (46/81); p=0.0179 Stepwise multiple logistic regression, &gt;50 vs ≤50 h/week: parameter 0.635 (p=0.0293)</td>
<td>Marital status, medical facility, position and night duty</td>
<td>Hayasaka <em>et al</em>[^44]</td>
</tr>
<tr>
<td>Mood disorders, GHQ-12 and SCLDS</td>
<td>No association was found between number of hours worked in a week and depression Not adjusted</td>
<td>Firth-Cozens[^42]</td>
<td></td>
</tr>
<tr>
<td>Mood disorders, SF-36</td>
<td>Scores when night float vs standard call at baseline and follow-up: Mental health mean score (SD): baseline=57.33 (22.63) vs 65.71 (7.61); follow-up=52.00 (15.49) vs 60.80 (11.45); p=0.72 Mental health component summary (SD), baseline=34.84 (14.06) vs 40.21 (7.61); follow-up=30.15 (10.71) vs 42.40 (6.23); p=0.39 Regression analysis: increased number of hours in hospital correlated with significantly lower SF-36 scores in almost all domains</td>
<td>Not adjusted</td>
<td>Zahrai <em>et al</em>[^60]</td>
</tr>
</tbody>
</table>

*Continued*
Table 3 Continued

<table>
<thead>
<tr>
<th>Health outcome</th>
<th>Main results</th>
<th>Adjustment by confounders</th>
<th>Study ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mood disorders, SF-36 and BDI</td>
<td>Improvement of levels before and after duty hours limitations from 90–110 to 78–80 h/week: Mental health SF-36: no statistically significant improvement Individual questions BDI before and after duty hours limitations, only energy level statistically significant</td>
<td>Not adjusted</td>
<td>Stamp et al(^{47})</td>
</tr>
<tr>
<td>Mood disorders, SF-36</td>
<td>OR (95% CI) for working 47 h/week or more: Impaired mental health, men: 1.59 (0.95 to 2.66); women: 1.86 (1.03 to 3.37)</td>
<td>Age and amount of time in practice</td>
<td>Sundquist and Johansson (^{48})</td>
</tr>
<tr>
<td>General health, SF-36</td>
<td>SF-36: Scores when night float vs standard call at baseline and follow-up: General health mean score (SD): baseline=62.11 (17.47) vs 77.57 (24.25); follow-up=56.43 (24.89) vs 84.20 (16.50); p=0.41 Physical health component summary (SD): baseline=46.16 (13.15) vs 52.01 (13.33); follow-up=39.32 (9.80) vs 56.15 (2.18); p=0.015 Vitality mean score (SD): baseline=51.67 (14.58) vs 51.43 (15.74); follow-up=48.57 (14.92) vs 51.00 (10.84); p=0.20 Regression analysis demonstrated that the increased number of hours spent in hospital correlated with significantly lower general health, physical function, mental health, role emotional, social function and mental component summary scale scores (all p&lt;0.05)</td>
<td>Not adjusted</td>
<td>Zahrai et al(^{50})</td>
</tr>
<tr>
<td>General health, SF-36</td>
<td>SF-36, BDI: Improvement of levels before and after duty hours limitations from 90–110 to 78–80 h/week: Physical health: no statistically significant differences</td>
<td>Not adjusted</td>
<td>Stamp et al(^{47})</td>
</tr>
<tr>
<td>General health, SF-36</td>
<td>OR (95% CI) adjusted for age and time in practice, for working 47 h/week or more: Impaired general health: men=1.66 (1.00 to 2.77); women=1.59 (1.00 to 3.17)</td>
<td>Age and amount of time in practice</td>
<td>Sundquist and Johansson (^{48})</td>
</tr>
<tr>
<td>General health</td>
<td>Self-rated health: OR (95% CI) of good self-rated health (logistic regression): Norwegian work time pattern+=1.35 (1.03 to 1.77); working in Norway=4.17 (3.02 to 5.73)</td>
<td>Age, sex and country of work</td>
<td>Rosta and Aasland (^{46})</td>
</tr>
</tbody>
</table>

\(^{*}\)Not working more than 9 h a day and having more than 60 h a month on-call. AD, anti-depressive (Mood disorders, depression); BDI, Beck Depression Inventory II; ED, Emergency Department; GHQ-12, General Health Questionnaire-12 items; ICU, intensive care unit; OH, Occupational Health (Percutaneous Injuries); SCLDS, Symptom Checklist Depression Scale.
found for male and female general practitioners who worked for 47 h/week or more.48 Zahrai et al49 found that increased number of hours spent by residents in hospital correlated significantly with lower mental health and Mental Component Summary Scale scores (SF-36). Three other studies, one of which is of high quality, did not find an association between depression or mood disorders and LWH. Varma et al50 did not find higher adjusted risks of depression associated with increased number of hours in senior medical consultants, even when they worked for more than 60 h/week. Depression or mood disorders measured with GHQ-12, Symptom Checklist Depression Scale (SCLDS) or Beck Depression Inventory (BDI), all validated tools for mental health, were not associated with increased number of working hours42 nor improved after decreasing the number of hours from 90–110 to 78–80 h/week in residents.47

**General health:** Four studies of intermediate quality analysed the association between LWH and general health. Zahrai et al49 found that increased number of hours spent by residents in hospital correlated significantly with poorer general health, physical function and vitality using SF-36. A comparative study of physicians in two different countries found that Norwegians showed higher non-adjusted prevalence of self-reported good health compared with physicians in Germany who worked longer hours.46 An increased adjusted risk of impaired general health was found for male and female general practitioners who worked for more than 47 h/week.48 However, following the implementation of reducing the number of hours from an average of 90–110 to 78–80 h/week did not lead to an overall improvement of residents’ self-reported physical health.47

**Discussion**

This review found moderate scientific evidence for a positive association of intermediate magnitude between exposure to LWH and percutaneous injuries and motor vehicle accidents in physicians. Evidence was assessed as low or insufficient for mood disorders and general health.

To the best of our knowledge, this is the first systematic review on the effects of LWH on physicians’ health and safety. The extensive searches were restricted to indexed journals. We systematically identified, selected and assessed the methodological quality of studies by means of independent pairs of reviewers. The quality assessment form and the levels of evidence used in this review were based on CONSORT and STROBE statements and on previous systematic reviews.36 38 39 To give some estimate of the potential magnitude of the effect, we added levels based on the point estimates of measures of association provided by the studies. However, such levels had not been previously established.

**Strengths and limitations of the studies**

Four studies had a longitudinal prospective design, however only two of them used a robust methodology.

**Synthesis of the evidence**

The levels of evidence synthesis obtained from the analysis of the 11 studies included in this review are shown in table 4.

<table>
<thead>
<tr>
<th>Health outcome</th>
<th>Degree of evidence*</th>
<th>Magnitude of the association†</th>
<th>Studies ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percutaneous injuries</td>
<td>++</td>
<td>++</td>
<td>Ayas et al,40 Fisman et al,43</td>
</tr>
<tr>
<td>Motor vehicle accidents</td>
<td>++</td>
<td>++</td>
<td>Barger et al,41 Kirkcaldy et al,45</td>
</tr>
<tr>
<td>Mood disorders</td>
<td>+</td>
<td>±</td>
<td>Varma et al,49 Hayasaka et al,44 Firth-Cozens,42</td>
</tr>
<tr>
<td>General health</td>
<td>+</td>
<td>±</td>
<td>Zahrai et al,50 Stamp et al,47 Sundquist and Johansson,46</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Zahrai et al,50 Stamp et al,47 Sundquist and Johansson,48,49 Rosta and Aasland,50</td>
</tr>
</tbody>
</table>

*Strong evidence (++): consistent results in more than two studies of high quality; moderate evidence (+): consistent results in two studies of high quality, or one high-quality study and one intermediate, or between some studies of intermediate quality; insufficient evidence (+): identification of only one study or inconsistent results across studies; evidence of no association (−): consistent results of a non-association in two or more studies.
†Magnitude of the association: it refers to the magnitude of the association point estimate (RR=relative risk, rate ratio or OR): high (+++) if RR >3; intermediate (++) if RR=1.5–3; low (+) if RR <1.5; no association (−); unclear (±).
Two other studies used a crossover design, which reduces the likelihood of individual variability and confounding, and were considered well designed. Four studies, all rated as intermediate methodological quality, were cross-sectional in design and therefore it is difficult to draw causal relationships from them. Although reverse causality cannot be ruled out from cross-sectional designs, it is unlikely that poorer health determines longer working hours than their healthier peers. One study used a quasi-experimental design but lacked a control group. Also, because of different designs used across the included studies, different estimates of the effects were used that may not be directly comparable.

Other limitations include the variability, validity and reliability of working hours, which in most studies was self-reported. Although only those papers specifying the number of working hours were accepted, shift and night work might have worked as confounders too, as it is difficult to acknowledge which proportion of the health effects observed was due exclusively to the LWH component and not due, for instance, to stress, fatigue or sleep deprivation. Moreover, the studies did not discriminate between time spent actively working and asleep while on call, though the EWTD considers all hours on call as working time. Likewise, it is possible that participants were aware that the study was conducted to assess associations between the number of hours worked and health outcomes. Therefore, there might be a volunteer bias in those doctors who work extended hours. To minimise this bias, a case-crossover analysis has been performed in three of the included studies.

Health outcomes were measured mainly based on self-report, with the exception of documented motor vehicle crashes, reported percutaneous injuries and register-notified antidepressive drug prescriptions. No studies with other objective measures of health (eg, mortality) or mental disease (eg, hospital data) were identified. However, self-reported or perceived health was assessed by validated and widely used instruments, especially when health status (general, mental or physical health) or ill-health symptoms were evaluated. Nonetheless, this raises the possibility that the observed associations might reflect differences in propensity to report health problems when they occur rather than true differences in the risk of worse health status. We cannot rule this out, but the higher propensity for reporting among those who work for longer hours and poorer work schedule patterns would have to relate specifically to the reporting of ill health rather than non-ill health. It seems unlikely that major differences in propensity to report would extend to a more concrete outcome.

The majority of the studies of this review took into account confounding variables in their analyses, such as sex and age. However, none of the studies analysed the potential effect of modifying factors, such as psychosocial aspects at work, including attitudes, motivation, job requirements, demands and content, organisational climate, social relationships at work, work satisfaction, supportive organisation or the relative number of physicians and other staff available for patient care. Neither were important aspects outside of work, such as life events or lifestyles, taken into account. The study of Japanese female physicians found significant differences by marital status with higher GHQ scores for those who were married. There are significant differences in the number of physicians per capita across different countries and this together with the structure and organisation of health services are potential confounders too that were not addressed in the selected studies across countries.

A relevant outcome from this review would be to establish a threshold number of extended hours above which there is a significantly higher risk. There is a lack of evidence for a dose–response relationship and nor does the evidence give any indication for a threshold number of hours that physicians could work and remain safe and healthy.

Finally, the heterogeneity of the included studies did not allow a more quantitative synthesis, and a level of evidence approach was used instead.

Comparison with other studies

Our findings of an association between LWH and injuries and accidents in physicians are consistent with research in other occupational groups. Dembe et al carried out a longitudinal survey in the USA including more than 10,000 workers from a variety of occupations and settings. A strong dose–response effect (adjusted for age, gender, occupation, industry and region) was found between LWH and injuries above 40 h/week and 8 h/day; jobs with overtime schedules were associated with a 61% higher injury hazard rate compared with jobs without overtime; working at least 12 h/day was associated with a 37% increased hazard rate, and working at least 60 h/week was associated with a 23% increased hazard rate. They concluded that LWH might indirectly precipitate workplace injuries through a causal process by inducing fatigue or stress in workers. In a separate study assessing injury risks to healthcare personnel, Dembe et al demonstrated that the risk of injury when working overtime or at least 60 h/week among physicians and nurses was statistically significant. In our review, we found that information about the length of shift work varied across studies, some analysing shifts of up to 24 h. Work that included such shifts was more strongly associated with accidents and injuries. We found insufficient evidence of an association between LWH and mood disorders, other diseases and general health in physicians, despite some scientific evidence suggesting that LWH increase morbidity and mortality in other occupations.

Possible explanations for these inconsistencies might be the scarcity of high-quality research, and the fact that some of the included papers did not fully address the effects of confounding factors, and the possibility that working as a physician might have a protective effect by
itself, for instance, through higher motivation and social recognition relative to other occupations.

Stamp et al.\textsuperscript{47} in the only before and after study, did not find significant differences for mood disorder or general health before and after implementing requirements to reduce working schedules to 80 h/week in residents. It is debatable, however, whether any conclusions can be drawn from the impact of reducing extremely high working hours (90–110 h/week) to very high working hours (78–80 h/week).

Although we searched for and identified literature describing associations between LWH and increased mortality in doctors,\textsuperscript{52,53} none of those studies specified the number of hours worked, and therefore were not included.

**Recommendations for future research**

Further research, including well-designed prospective and retrospective longitudinal studies, is needed in this field to establish on scientific grounds what number of hours are safe, to patients as well as to physicians. Also, studies on long-term health effects should be considered.

This review was aimed at studying exclusively possible detrimental health effects of LWH (more than 48 h/week) on physicians. Other systematic reviews should be conducted addressing issues such as the impact of working hours on the well-being and quality of life of doctors, the training quality of junior doctors, and the quality and continuity of care and their impact on patient safety. Also, from a staffing perspective, the economic and social burden of sick leave and physicians leaving the profession are important issues that were beyond the scope of this review and that should be considered in future research. Other established criteria,\textsuperscript{54} such as that working hours should be ‘family friendly’, promote gender equality, enhance productivity or facilitate worker choice and influence over working hours, were not addressed. Further research is needed in these other areas.

**Conclusions**

The findings of this review have confirmed that LWH are associated with an increased risk of needlestick injuries and road traffic accidents. The fact that these associations are labelled moderate in this review should not preclude safety procedures and measures being assessed and reassessed to ensure maximum physician and patient safety. The reduction of needlestick injuries requires improved use of safe sharp devices, compliance with safe working procedures, training and regular audits of working practices, particularly for junior doctors who are at greatest risk of needlestick injuries, possibly due to inadequate training.\textsuperscript{55,56} The risk of road traffic accidents could be reduced by discouraging driving after very long shifts (more than 16 h) and where there is sleep deprivation.\textsuperscript{57} Transport should be provided by the employer in these situations. This is a pragmatic recommendation which recognises that LWH have not been eliminated as a result of the EWTD.

This review uncovered the lack of literature on the effect of LWH on the general health of doctors, a topic that has important health ramifications not only for physicians but for patients as well. However, the paucity of available studies, which does not allow for a causal or dose–response relationship to be established, raises further questions about the evidence base of the current 48 h limit, especially as this is not enforced strictly and junior doctors, for example, in the UK, may opt-out.\textsuperscript{58} It may be relevant that the EWTD was not based entirely on science but also took into account, as does all EU legislation, the views of ‘social partners’.\textsuperscript{3}

It is likely that the mechanism for these increased health and safety risks is fatigue, and our finding of increased risk of accidents to staff may also be associated with increased risk of clinical errors. Patient safety was not the purpose of this review but LWH are well recognised to cause decrement in performance in healthcare and other professions, where the performance of staff can be safety critical, such as being airline pilots or professional drivers.\textsuperscript{59}

In addition, the EWTD has been associated with real concerns about the adequacy of the training of doctors given their reduced exposure to patient care, and the lack of experience of the patient journey, imposed by adherence to the EWTD.\textsuperscript{4} This is potentially exacerbated at the same time by the constraints on the overall duration of training imposed by other EU directives.\textsuperscript{60} Further reviews of the EWTD, with regard to this unique group of workers, need to take into account other social factors such as the impact on patient care. This is particularly the case when all healthcare systems are under strain because of burgeoning demands and limited resources.\textsuperscript{4}

The findings of this systematic review lead to the suggestion of the following guidelines in the event of non-compliance with the EWTD: (1) long shifts should be avoided to protect physicians’ health and patient safety and no shift should be longer than 16 h (and then exceptionally)\textsuperscript{60}; (2) physicians should be discouraged from driving after long shifts to reduce the risk of motor accidents\textsuperscript{41,42}; (3) rigorous attention should be paid to reduce the risk of sharp injuries\textsuperscript{50,48}; (4) organisational aspects such as workload and job control, as well as the pattern and distribution of working hours, breaks and recovery periods should be carefully taken into account, in order to avoid fatigue and sleep deprivation that could lead to mistakes and accidents\textsuperscript{61,62} and (5) physicians should work in supportive psychosocial environments, in teams and with adequate training and supervision as their well-being is important also for the health of their patients.\textsuperscript{61,62}

Further research is required: to establish any longer term effects, such as on mortality and mental health; to assess the impact of the nature of the work organisation taking into account the psychosocial aspects of the physicians’ working and non-working lives and to determine how many hours are safe for physicians to work.
Author affiliations

1 Catalan and Spanish Societies of Occupational Medicine, Barcelona, Spain
2 Department of Medical Sciences, School of Medicine, Universitat de Girona, Girona, Spain
3 MRC/CSO Social and Public Health Sciences Unit, University of Glasgow, Glasgow, UK
4 Healthy Working Lives Group, Institute of Health and Wellbeing, College of Medical, Veterinary and Life Sciences, University of Glasgow, Glasgow, UK
5 CISAL—Centre for Research in Occupational Health, Usetat Universitat Pompeu Fabra, Barcelona, Spain
6 Occupational Health Department, Epson & St Helier University Hospitals NHS Trust, Surrey, UK
7 Clinical Institute of Occupational, Traffic and Sports Medicine, University Medical Centre, Ljubljana, Slovenia
8 College of Occupational Medicine, Portuguese Medical Association, Lisbon, Portugal
9 Helsinki City Occupational Health Centre, Helsinki, Finland
10 Occupational Health Service, Parc de Salut MAR, Barcelona, Spain
11 CIBER of Epidemiology and Public Health, Spain
12 UEMS Section of Occupational Medicine

Acknowledgements

Leena Isolato, Trial Search Coordinator at the Cochrane Occupational Safety and Health Review Group, designed and conducted the final search of the evidence using MEDLINE and EMBASE databases.

Contributors

MCR-J, ED, SV-P, AS, KAS, EBM and CS conceived and submitted the design, carried out the acquisition of data, analysis and interpretation of data, drafted the article and revised it. PGR and R H A contributed to acquisition of data, analysis and interpretation of data, drafted the article and revised it. Authors gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Funding

The Section of Occupational Medicine of the UEMS provided limited support to the work of MCR-J, ED, SV-P, KAS, AS, PGR and RH-A. The Section of Occupational Medicine of the UEMS provided limited support to the work of MCR-J, ED, SV-P, KAS, AS, PGR and RH-A.

Contributors

MCR-J, ED, SV-P, AS, KAS, EBM and CS.

Role of funding source

None.

Contributing to the research in Occupational Medicine.

were also funded by the Section.

Competing interests

None.

Ethics approval

Not commissioned; externally peer reviewed.

Provenance and peer review

Not commissioned; externally peer reviewed.

Data sharing statement

No additional data are available.

Open Access

This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 3.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided they cite the original work and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/3.0/.

REFERENCES