

Managing severe cancer pain: the role of transdermal buprenorphine: a systematic review

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Abstract: Pain is a frequent and important symptom in cancer patients. Among the available strong opioids, transdermal buprenorphine has been licensed in Europe since 2002, and results from a few clinical studies suggest that it may be a good alternative to the other oral or transdermal opioids. To assess the best available evidence on its efficacy and safety, we carried out a systematic literature review with the aim of pooling relevant studies. We identified 19 eligible papers describing 12 clinical studies (6 randomized controlled trials and 6 observational prospective studies), including a total of about 5000 cancer patients. Given the poor quality of reports and the heterogeneity of methods and outcomes, pooling was not feasible as the type of data was not appropriate for combining the results statistically. A meta-analysis based on individual data is ongoing in the context of the Cochrane Collaboration. In conclusion, although the narrative appraisal of each study suggests a positive risk benefit profile, well designed and statistically powered controlled clinical trials are needed to confirm this preliminary evidence.

Keywords: systematic review, transdermal buprenorphine, cancer pain

Introduction

Advances in prevention, diagnosis, and therapy have extended the life expectancy of cancer patients, but improvement in the survival is still disappointing for most common tumors, mostly when the disease is diagnosed at advanced or metastatic stages. Despite the availability of new and innovative anti-cancer drugs, metastatic patients are indeed unlikely to have important benefit from these treatments either in terms of quantity and quality of life. For most of these patients, the later part of their lives is impaired by pain, fatigue, depression, and other symptoms related to the disease and treatments, which become prominent contributors to suffering. Pain, in particular, is one of the most important problems. Most patients with advanced or metastatic cancer experience pain during their life^{1,2} and despite effective treatments being available, undertreatment has been documented in nearly one of two patients with cancer pain, with a few differences according to geographical, economic, and cultural factors.³ The use of analgesic pharmacotherapy is the mainstay of all guidelines on pain management which are available since 1986.⁴⁻⁷ The World Health Organization (WHO) suggested that the choice of analgesics should be based on pain intensity, and not simply on its etiology; the preferred route of administration should be oral, for drugs ranging from paracetamol and NSAID to strong opioids, morphine being the first choice.⁴ Since then, a broad spectrum of analgesics has become available, such as morphine, methadone, hydromorphone, oxycodone, fentanyl, and buprenorphine, which have been shown

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to be effective in treating moderate to severe cancer-related pain. In addition to the traditional analgesics that are deliverable orally or parenterally, a few transdermal delivery systems (TDS) containing fentanyl or buprenorphine have been introduced onto the market, suggesting a potential for improvement in view of their advantages over the oral and parental routes, in terms of non-invasiveness and slow and continuous release of the compound. A recent review has pointed out the advantages and disadvantages of available TDS for cancer pain, focusing on the problem of transdermal dosing conversion.⁸ Another very recent review updated the pharmacological properties of transdermal buprenorphine with a focus on its ceiling effect,⁹ and also qualitatively summarized the clinical efficacy and tolerability of TDS containing buprenorphine, using results from clinical studies including cancer patients, as primary evidence. As this work combined the early and late randomized placebo and comparative trials with retrospective and prospective studies, it was difficult to obtain an overall quantitative estimate of the yield of TDS containing buprenorphine in terms of efficacy and safety.

Our paper is based on a systematic review of published literature with the aim of assessing the best available evidence for the effect of TDS containing buprenorphine in chronic cancer pain. We have searched, identified, appraised, selected, and integrated all relevant publications on this topic and attempted to combine statistically the valid studies. The results and comments are presented according to the study design (randomized clinical trials vs observational studies) and are based on 19 papers reporting data from 12 prospective studies involving more than 5000 patients.

Materials and Methods

Synthetic profile of buprenorphine

Excellent profiles of the pre-clinical and clinical pharmacology of buprenorphine have been reported elsewhere.⁹⁻¹¹

Briefly, buprenorphine is a semi-synthetic compound derived from tebaine, a natural opium alkaloid, which is structurally similar to morphine, even if several molecular differences confer a higher lipophilicity and a higher pharmacological potency.

Pharmacokinetically, buprenorphine shows very low bioavailability (about 15%) after oral administration,^{12,13} because of an intense metabolic degradation at the intestinal and hepatic level (first pass). After transdermal administration, the plasma concentration remains more or less steady as a consequence of continuous delivery of the drug. A very high percentage of available buprenorphine is strongly bound

to plasma proteins,¹⁴ and only the unbound fraction that consists of 3% to 5% of the total plasma buprenorphine may cross the blood–brain barrier. Transit across the blood–brain barrier is regulated by several factors, the most important being the free fraction of the drug, and lipophilicity. Among the opioids, buprenorphine has an intermediate to high lipophilic property. Once buprenorphine has crossed the blood–brain barrier, it must reach its biological target, the opioids receptors (ORs). Buprenorphine presents a high selectivity for the MOR (μ -ORs). In an *in vitro* study,¹⁵ the *K_i* values of buprenorphine were 0.08 for the MOR, 0.11 for the KOR, and 0.42 for the DOR. For pharmacological efficacy, buprenorphine presents 60% to 65% with respect to MOR (partial agonist).¹⁶⁻²² These results confirm that the analgesic activity of buprenorphine is mainly mediated by MOR; its partial agonist action reflects the potential existence of a ceiling effect that, in clinical practice, has been estimated to be about 15 to 25 mg daily,²³ a dosage that may not be compatible with the doses usually prescribed in clinical practice.

In terms of pharmacokinetics, two different metabolic pathways have been observed to work in parallel: the first consists of N-dealkylation by means of CYP3A4, and the second produces three glucuronized compounds. The first metabolic pathway generates nor-buprenorphine, the only molecule having some biological activity. Buprenorphine is eliminated through two routes: unchanged molecules are excreted mainly via the biliary system, whereas metabolites are eliminated via renal excretion, but metabolite accumulation is of minimal importance given that these agents are substantially inactive substances.²⁴ For these reasons buprenorphine may be considered as a safe opioid in cases of reduced renal function, which is a frequent clinical condition in cancer patients, especially in the far advanced phase.⁹

Transdermal buprenorphine in pain management

Buprenorphine was first synthesized in the late 1960, and introduced in clinical practice for parenteral and sublingual (SL) administration in 1978 and 1981, respectively. In the late 1990s, it was introduced as a transdermal formulation, contained in a matrix patch that can be applied to the skin for a duration of up to 4 days, which was indicated for moderate to severe cancer and chronic pain unrelated to cancer. The continuous release from the matrix across the skin and then into the systemic circulation is regulated mainly by the concentration gradient across the skin and the patch.²⁵ In the matrix patch, the drug is an integral part of the polymer structure,

making the delivery system more robust than the reservoir patch. This feature prevents “dose-dumping” and potential overdosing either intentionally or unintentionally, as damaging the patch does not interfere with the controlled release of medication. TDS containing buprenorphine is available with release rates of 35, 52.5, and 70 $\mu\text{g}/\text{hour}$ (Transtec[®]) that correspond to 0.8, 1.2, and 1.6 mg/day of buprenorphine or 60, 90, and 120 mg/day equivalent of oral morphine, respectively.^{25,26} Recently, low-dose patches (5 to 20 $\mu\text{g}/\text{hour}$ released for 7 days) have been marketed in a few countries.²⁶

Transtec[®] has been available since 2002 in 18 European countries including Russia and three South American countries.

Search strategy and study selection

MEDLINE (from 1966 to April 2009), EMBASE, and the Cochrane Library were searched, and only papers written in English were taken into account for data extraction; the main search terms were: ‘pain’, ‘neoplasms’, ‘buprenorphine’, ‘transdermal’, and ‘randomized controlled trial’ (RCT), as well as combinations of these terms. The complete search strategy can be obtained from the authors on request. References in the published articles, reviews, meta-analyses, and relevant organization websites were checked. No systematic attempt was made to identify unpublished studies.

Data extraction and synthesis

The eligibility assessment of the titles and abstracts was performed in a standardized manner by one reviewer (SD). Two reviewers (SD and GA) independently screened the full papers. Disagreements were resolved by discussion and consensus. Details of the study design, participants, inclusion criteria, clinical setting, study duration, definition of outcomes and endpoints, and results were recorded and summarized. Given the heterogeneity of the studies and the difficulty of extracting quantitative disaggregated information from each study, we undertook a narrative synthesis of each study. Eligible studies were then grouped based on the study design (randomized vs observational). Data from the studies reporting the use of standardized measures of pain intensity were grouped, described, and integrated. In this case, an attempt to summarize the results of the studies reporting the use of standardized measures of pain intensity has been attempted using appropriate statistical methods.²⁷

Results

The search of MEDLINE and EMBASE with the review of titles and abstracts, integrated by scanning the references and

consulting experts in the field, yielded 27 relevant articles. Four articles were excluded because they were published as abstracts and/or full text was not available, and the other four were excluded because they were written in languages other than English. Figure 1 presents a flowchart of manuscript selection.

The main characteristics of the remaining 19 papers^{28–46} are reported in Table 1. The 19 published articles described 16 different studies: Radbruch²⁹ presented the aggregated data from Bohme,²⁸ Sittl,³¹ and Sorge;³² Evans reported efficacy and safety data by pooling the 3 placebo-controlled RCTs;²⁵ and Radbruch³⁰ and Apolone³⁵ reported the preliminary information later published by Griessinger³³ and Apolone,⁴⁶ respectively. Nine were clinical trials^{28,31,32,37,39–42,44} and 7 were observational (non-intervention) studies.^{33,34,36,38,43,45,46}

Most of the papers reported data from a mixed population including both cancer and non-cancer patients (12/19).

Three experimental studies were subsequently excluded because they were early-phase feasibility trials: a rotation feasibility trial describing cross-over administration of two drugs,⁴¹ a pilot study describing the effect of pre-planned rotation administration of several drugs,³⁹ and a dose escalation administration trial.³⁷ The characteristics of the remaining 6 studies,^{28,31,32,40,42,44} involving 499 cancer patients (range 9 to 189) are reported in Table 2A.

One observational study was also excluded, because it was focused on assessing the effect of buprenorphine TDS across pre-planned age groups, to test its effectiveness in a special population:⁴³ the characteristics of the remaining 6 studies (4599 cancer patients, range 6 to 3690) are reported in Table 2B.

RCTs

Six RCTs were available for analysis and discussion. Details of the papers and the studies are summarized in Tables 2A and 3. Studies may be classified into 3 types: 3 preliminary pivotal studies comparing TDS buprenorphine vs placebo, 2 testing different schedules and a maintenance regimen, respectively, and 1 comparative trial vs morphine.

The first Phase III studies were 3 multicenter, randomized, double-blind clinical trials vs placebo (Bohme,²⁸ Sittl,³¹ Sorge)³² in patients with chronic cancer and non-cancer pain, initiated and sponsored by the industry marketing in Europe the drug under evaluation. In the first and the third study, only patients whose pain was satisfactorily relieved after a run-in phase with buprenorphine SL were randomized. In the study by Sorge,³² only 35 $\mu\text{g}/\text{hour}$ dose strength was allowed; the 3 dose strengths (35, 52.5, and 70 $\mu\text{g}/\text{hour}$)

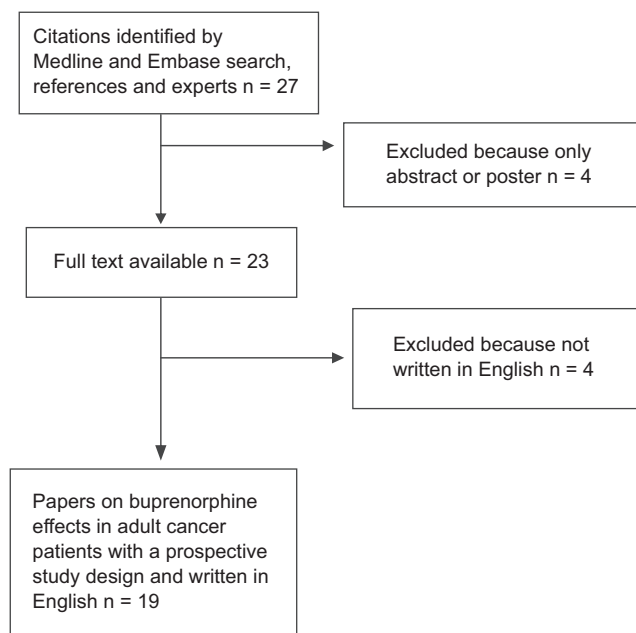


Figure 1 Flowchart of manuscript selection.

were used by Bohme²⁸ and Sittl.³¹ In the first 2 studies, the primary outcome was the responder's status defined as 'patients whose pain relief was at least satisfactory at all determination points (excluding the final examination) and who took a mean of 0.2 mg/day or less of SL buprenorphine on days 7–12' (Bohme)²⁸ and 'any patient who required no more than 1 SL tablet of buprenorphine as rescue medication per day from day 2 until the end of the study and who recorded at least satisfactory pain relief at each application of a new patch' (Sittl).³¹ The third study, on the contrary, used only the rescue therapy as the primary outcome, in terms of the number of buprenorphine SL tablets required each day. Bohme²⁸ reported a percentage of responder patients of 34% for 35 µg/hour, 37% with for 52.5 µg/hour, and 50% for 70 µg/hour, but these response rates failed to reach statistical significance. The proportion of patients reporting good to complete pain relief, none to mild pain on a verbal rating scale (VRS), and an uninterrupted sleep for more than 6 hours increased during the double-blind phase, but the authors do not report a test of statistical significance vs placebo. Overall, 23% of the patients reported adverse events, with no statistically significant differences between the treatment groups. In the study by Sittl,³¹ the percentage of responders was very similar to that in the Bohme²⁸ study (ie, 36.6%, 47.5%, and 33.3% for the 3 dosages); also, in this case, the difference vs placebo failed to reach statistical significance in the 70 µg/hour patch. Buprenorphine seemed to give better results than the placebo also in terms of rescue

therapy ($P = 0.002$), pain relief, pain intensity, and duration of sleep uninterrupted by pain (P value not reported). The percentage of patients reporting at least 1 adverse event was high for both buprenorphine (85.4%, 80.5%, 75.7% for the 3 patches) and placebo group (73.7%) with no statistical difference. In the study by Sorge,³² the mean daily requirement of buprenorphine SL tablets during the double-blind phase was lower than that required during the run-in for the buprenorphine TDS group (–55.4%, dose-dependent) than for placebo (–45.1%, $P = 0.01$), but was similar in terms of mean number of daily dose (in mg) consumed (0.5 for all patients, 0.4 vs 0.6 for cancer patients). Pain intensity, pain relief, and sleep quality evaluation suggested a better effect of buprenorphine, although a statistical significance was not reached or it was not evaluated or reported. Adverse events were reported in 54.4% of the buprenorphine TDS group and 42.6% of the placebo group with no significant difference. Radbruch²⁹ reported the aggregated results of these 3 RCTs and classified the patients according to the disease: among the cancer patients, 36% of those with 35 µg/hour patch and 42% and 40% of those with the 52.5 and 70 µg/hour patch were classified as responders vs 23% of the placebo.

Likar⁴⁰ compared the efficacy and tolerability of the 2 different delivery schedules in subjects who have already responded to buprenorphine: patients (only 9 with cancer) were randomized to a treatment sequence in which they changed the buprenorphine patch every 3 days during the first step of the study (3-day regimen) or every 4 days (4-day regimen). In the second phase, the patients crossed over to the alternative regimen. The 2 regimens failed to show any statistical difference in terms of patient's and physician's satisfaction, pain intensity assessed with a numerical rating scale (NRS) (3.73 for 3-day regimen and 3.88 for 4-day regimen), pain relief, and tolerability. The latter findings were presented only in an aggregated way, making it impossible to evaluate the effect of the 2 interventions in cancer patients.

To date, the study by Pace⁴² is the only comparative trial available. It compared the effects of buprenorphine of 35 µg/hour with the oral SR morphine of 60 mg/day in an open-label study. The patients treated with transdermal buprenorphine experienced significantly greater improvement in pain intensity (–2.5 for buprenorphine, –1.4 for morphine), quality of sleep, and quality of life. Adverse drug reactions (ADRs) were not reported and the side effects that were significantly more frequent for morphine than for buprenorphine, were vertigo, constipation, and nausea.

Finally, more recently, another RCT vs placebo was conducted to evaluate the maintenance of efficacy (Poulain)⁴⁴

Table 1 List and characteristics of the 19 papers selected

Reference	Year of publication	Country	Study design	Total number of patients (cancer:non-cancer)	Remarks
Böhme ²⁸	2003	Austria, Germany and Hungary	Parallel double-blind RCT vs placebo	151 (83:68)	
Radbruch ²⁹	2003	Germany	3 parallel double-blind RCTs vs placebo	445 (249:196)	Collects information from other studies ^{32,35,36}
Radbruch ³⁰	2003	Germany	Observational	3255 (846:2409)	Preliminary data from another study ³⁷
Sittl ³¹	2003	Austria, Germany and The Netherlands	Parallel double-blind RCT vs placebo	157 (121:36)	
Sorge ³²	2004	Germany and Poland	Parallel double-blind RCT vs placebo	137 (45:92)	
Griessinger ³³	2005	Germany	Observational	13179 (3690:9489)	Post-marketing surveillance
Muriel ³⁴	2005	Spain	Observational	1212 (207:1005)	Open-label
Apolone ³⁵	2006	Italy	Observational	not reported	Preliminary data from another study ⁵⁰
Likar ³⁶	2006	Austria, Germany, Hungary, Poland and The Netherlands	Observational	239 (134:105)	Long-term follow-up of patients recruited in previous RCTs ^{32,35,36}
Mercadante ³⁷	2006	Italy	Dose escalation feasibility trial	10	Describes the efficacy and tolerability of a dosage increase up to 140 µg/hour
Camba ³⁸	2007	Spain	Observational	762 (164:598)	
Freye ³⁹	2007	Germany	Rotation feasibility trial	42 (9:33)	Describes the switch from high-dose morphine to TDS buprenorphine
Likar ⁴⁰	2007	Austria	Crossover open-label RCT comparing 2 schemes	49 (9:40)	Compares the efficacy and tolerability of 3-day vs 4-day patch change schedule
Mercadante ⁴¹	2007	Italy	Crossover non-randomized clinical trial (vs fentanyl)	22	Describes the switch from one opioid to the other and then back to the previous one
Pace ⁴²	2007	Italy	Parallel open-label RCT vs morphine	52	
Likar ⁴³	2008	Austria	Observational	82 (6:76)	Compares efficacy and tolerability in different age groups
Poulain ⁴⁴	2008	Austria, Belgium, Croatia, France, Poland, The Netherlands	Parallel double-blind RCT vs placebo	189	
Wirz ⁴⁵	2008	Germany	Observational	174	Random selection of patients
Apolone ⁴⁶	2009	Italy	Observational	257	Outcome research study

Abbreviations: RCT, randomized controlled trial; TDS, transdermal delivery system.

in severe cancer pain. Opioid-tolerant patients were converted to buprenorphine TDS during a run-in phase and then randomized for the 2-week maintenance phase; of the 289 patients initially recruited, only 189 entered the second phase, because 100 discontinued the treatment for adverse

events or lack of efficacy. The primary efficacy outcome was the proportion of the responder patients (defined as patients completing at least 12 days of the double-blind period, with an average pain intensity score of <5 on the numerical rating scale during the last 6 days of treatment, and not using

Table 2A Characteristics of 6 randomized clinical trials investigating buprenorphine TDS effects in adult cancer patients

Reference	Number of patients with cancer	Inclusion criteria for randomization	Follow-up assessment	Primary outcome and endpoint	Secondary endpoints							
					PI	PR	AE	PS	SQ	RT	QoL	
Böhme ²⁸	83	Chronic pain at least satisfactorily relieved by sublingual buprenorphine	5 days run-in, 10 days double-blind	Responder status (composite endpoint)	x	x	x	-	x	-	x	-
Sittl ³¹	121	Chronic, severe pain related to cancer or other diseases	15 days	Responder status (composite endpoint)	x	x	x	-	x	-	x	-
Sorge ³²	45	Severe or very severe chronic pain related to cancer or other disorders justifying treatment with strong opioids and whose pain had been at least satisfactorily relieved during the run-in phase	6 days run-in, 3 days influx, 6 days double-blind	Rescue therapy (number of buprenorphine SL tablets)	x	x	x	-	x	-	x	-
Likar ⁴⁰	9	Had already responded to at least 4 weeks of TDS buprenorphine for chronic moderate or severe pain of malignant or nonmalignant origin and at steady state for at least 2 weeks before enrollment	24 days (12 for each schedule)	Patient satisfaction (composite endpoint)	x	x	x	-	-	-	x	x
Pace ⁴²	52	Having cancer chronic pain for a period of 1 to 3 years, diagnosis of abdominal neoplasm, a pain score equal to at least 40 mm on the VAS before the screening week and with a pain score average equal to at least 4 on the Likert scale and with at least 4 observations recorded in the daily diary during the screening week	8 weeks (1 week of screening before)	Pain intensity (NRS)	x	-	x	-	x	-	x	x
Poulain ⁴⁴	189	Documented malignant disease and insufficient pain relief from current opioid regimen who responded to buprenorphine TDS during the run-in	14 days run-in, 14 days maintenance	Responder status (PI lower than 5)	x	-	x	x	-	-	-	-

Abbreviations: AE, adverse events; PI, pain intensity; PR, pain relief; PS, patient satisfaction; QoL, quality of life; RT, rescue therapy; NRS, numerical rating scale; SL, sublingual; SQ, sleep quality; TDS, transdermal delivery system; VAS, visual analogue scale.

Table 2B Characteristics of 6 observational prospective studies investigating buprenorphine TDS effects in adult cancer patients

Name of the first author	Number of patients with cancer	Inclusion criteria	Follow-up assessment	Primary outcome and endpoint	Secondary endpoints							
					PI	PR	AE	PS	SQ	RT	QoL	
Griessinger ³³	3690	Unsatisfactory pain relief, moderate to severe pain or unacceptable side effects	10 weeks	Pain relief	-	-	-	-	-	-	-	-
Muriel ³⁴	207	Not responder to previous therapy and moderate-to-severe pain	3 months	Pain relief and tolerability (composite endpoint)	-	-	x	-	x	-	-	x
Likar ³⁶	134	Moderate-severe pain and participating to the three previous trials	6 months	Pain relief (VRS)	-	-	x	-	-	x	-	-
Camba ³⁸	164	Not responder to previous therapy and local or metastatic cancer	8 weeks	Pain intensity (VAS)	-	-	x	-	-	x	-	x
Wirz ⁴⁵	174	Already in therapy with that opioid for longer than 4 weeks	5 days	Safety	-	-	x	-	-	-	-	-
Apolone ⁴⁶	257	Persistent pain of any degree and life expectancy > 1 month	28 days	Pain intensity (worst and average pain on a NRS)	-	x	x	x	-	-	-	x

Abbreviations: AE, adverse events; PI, pain intensity; PR, pain relief; PS, patient satisfaction; QoL, quality of life; RT, rescue therapy; NRS, numerical rating scale; SQ, sleep quality; TDS, transdermal delivery system; VRS, verbal rating scale; VAS, visual analogue scale.

more than 2 tablets of rescue medication per day). More patients receiving transdermal buprenorphine of 70 µg/hour responded (74.5%) compared with those receiving placebo (50%) ($P = 0.0003$). Pain intensity was significantly lower in the TDS containing buprenorphine group (1.8) compared with placebo (2.7). In addition, the daily consumption of SL tablets remained stable during the double-blind phase for buprenorphine, but increased from 0.6 to 1.7 tablets for the placebo group. The incidence of adverse events was slightly higher for buprenorphine than for the placebo group, and the most commonly reported symptoms were nausea, vomiting, and constipation (P value not reported).

Observational prospective studies

The first observation prospective study was a post-marketing surveillance in Germany in 13,179 cancer and non-cancer patients with unsatisfactory pain relief or unacceptable side effects with previous therapy.³³ Patients with cancer-related pain were about 30% of the total sample. Seventy percent of them started with 35 µg/hour patch, 22% with 52.5 µg/hour, and 7% with 70 µg/hour; patch strength was changed at visit 1 (median 14 days) in 16% of the patients and at visit 2 (median 40 days) in 2% of the subjects. 'Good' or 'very good' pain relief was obtained in 84% of the cancer patients. Twenty-seven percent of the cancer patients experienced 1 adverse event (21.6% experienced serious adverse events).

Muriel³⁴ in Spain followed up for 3 months the cancer and non-cancer patients who were beginning buprenorphine for moderate to severe pain who had not responded to non-opioid analgesics. Patients began buprenorphine TDS at the lowest possible dosage, and the daily median amount at the baseline was 35 µg/hour (all the patients). At the end of the study, 22% of all the patients had been changed to a higher dosage. The study included 207 patients with cancer (17%). Sixty-five percent of these reported 'good' or 'very good' pain relief after 1 month, and 57% after 3 months. Other outcomes and adverse events were not reported for the cancer population alone: 63% of all the patients reported and improvement in the quality of sleep after 1 month, and 56% after 3 months, and the quality of life also improved, from a mean EQ-5D score of 40.6 at baseline to 56.8 at the end of the study ($P < 0.001$); 42% of the patients experienced at least 1 adverse event during the follow-up.

Some patients recruited in the 3 registered clinical trials mentioned earlier,^{28,31,32} were subsequently included in a follow-up study (Likar)³⁶ where they continued the treatment with 35 µg/hour patch and SL tablets (0.2 mg). There were 134 cancer patients who were followed up for a mean

of 119 days, but 107 discontinued after the first 2 months, mainly because of death and insufficient pain relief. Among the cancer patients, 86.6% reported an 'at least satisfactory' pain relief throughout the study period. Eighty-four cancer patients reported adverse events; 31 events were probably drug-related and 26 were ADRs (the more frequent were nausea and vomiting).

Camba³⁸ summarized 3 longitudinal multicentre studies conducted by the Sociedad Española del Dolor. The third study involved 164 cancer patients with moderate to severe pain, who did not respond to previous analgesics, and who were followed up for 8 weeks after the first administration of buprenorphine TDS. Fifteen percent of the patients started with 17.5 µg/hour and 82% started with 35 µg/hour of the drug. Mean pain intensity, measured with a visual analogue scale (VAS), at baseline was 7.4 and decreased to 3.2 at week 8 ($P < 0.001$). This study also included a quality of life questionnaire and parameters such as physical fitness, social and daily activities, and how they felt significantly improved during the course of the observation period. Fifty-one patients discontinued the study; the results of adverse events were not reported for the cancer study alone.

Wirz⁴⁵ conducted a controlled trial with the oral sustained-release hydromorphone, transdermal fentanyl, and buprenorphine on randomly selected 174 cancer patients, to assess the difference in terms of gastrointestinal symptoms in long-term treatment (all patients were pretreated with their current opioid therapy for more than 28 days). Patients were selected for participation by random selection from a sample of outpatients undergoing pain therapy with one of the study medications. Apart from a significantly higher incidence of stool-free periods of >72 hour for transdermal opioids, there were no differences in the effects of the 3 drugs in terms of nausea, emesis, and constipation.

Apolone⁴⁶ followed up 1801 advanced cancer patients with persistent pain of any degree, using several outcomes and endpoints, including pain intensity measured with a NRS, satisfaction, and quality of life: among the patients recruited, 257 were consuming buprenorphine at baseline, with a mean TDS dose of 43.2 µg/hour (median 35) and an average increase to 50.0 at day 28. All the outcome measures consistently improved over time, in terms of statistical significance and clinical relevance. For example, the worst pain differences were -1.4 points (95% confidence interval -1.1/-1.8, $P < 0.0001$). The effect-size estimates indicated that on average, the endpoints based on pain intensity were more responsive (range 0.2 to 0.6) than those measuring pain relief, satisfaction, or quality of life (range 0.2 to 0.3), and

the best measure was worst pain difference. About 34% of the patients had an improvement of at least 2 points in worst pain, nearly 48% improved less or were stable, 15% had a 20% improvement in pain relief, and 40% reported an increase in satisfaction. The most frequent side effects were constipation (56%) and sedation (51%), but the frequency of the symptoms rated by patients as 'a lot/very much' was always less than 25%.

Estimate of the overall treatment effect on pain

When studies were evaluated according to the type of outcomes and endpoints used to find a measure to pool studies and estimate the overall treatment effect, a substantial heterogeneity emerged that did not allow statistical pooling.

In the 6 RCTs, 3 studies^{28,31,44} shared the same primary outcome (the responder status), an endpoint that may be considered valid and relevant in this setting, but it was actually operationally defined in different ways (mentioned previously). The other primary outcomes were rescue therapy,³² patient satisfaction,⁴⁰ and pain intensity.⁴² When we expanded this assessment to the secondary outcomes, among the wide list of measures used (from indicators of pain intensity to safety and quality of life), the only one used in all 6 RCTs was pain intensity, which became our candidate for a formal attempt for a statistical combination.

In the 6 observational studies, the same evaluation again yielded a very heterogeneous result: in 3 cases the primary endpoint was pain relief,^{33,34,36} in 2 it was pain intensity,^{38,46} and in 1 it was safety.⁴⁵

As 8 studies, 6 clinical trials^{28,31,32,40,42,44} and 2 observational studies,^{38,46} have used patients' reported pain intensity, we focused on this subsample. Table 3 presents the details of the studies that were candidates for a pooling analysis.

A deeper analysis of the studies' characteristics and efficacy estimates reported in the papers showed that in this subsample also, the type of data was not appropriate for combining the results. For example, although all the studies used some standardized measures of pain intensity, the assessment tools were different: in 4 cases NRS was used,^{40,42,44,46} in 3 cases VRS was used,^{28,31,35} and in 1 case VAS was used.³⁸ In addition, for NRS different anchors were actually used to identify the least and worst pain intensity. In 4 cases^{28,31,32,40} it was not possible to distinguish the results between cancer and non-cancer patients. Finally, the period of time chosen by the authors to estimate the difference of pain intensity attributable to the intervention varied from 10^{28,32} to 56 days.³⁸

Table 3 Description of pain intensity outcomes and endpoints in 6 randomized clinical trials and 2 prospective observational studies investigating buprenorphine TDS effects in adult cancer patients

Reference	Primary outcome	Pain intensity measure	Details of the pain scale	Mean pain at baseline for cancer patients (sd)	Mean pain after treatment (sd)	Assessment time (days)
Böhme ²⁸	Responder status	5-point (VRS)	Very severe, severe, moderate, slight, absent	not reported	not reported	10
Sittl ³¹	Responder status	5-point (VRS)	Very severe, severe, moderate, slight, absent	not reported	not reported	15
Sorge ³²	Number of BP SL tablets required as rescue therapy	5-point (VRS)	Very severe, severe, moderate, slight, absent	not reported	not reported	10
Likar ⁴⁰	Patient satisfaction with treatment	11-point (NRS) MPQ	No pain- worst pain imaginable 15 items then categorized 0 (no pain) to 3 (worst pain imaginable)	not reported	not reported	12
Pace ⁴²	Pain severity	11-point (NRS)	0 = no pain; 10 = maximum possible pain	6.4 (0.2)	3.9 (0.3)	56
		SF-MPQ (VAS)	15 items for pain in the last week then categorized 0 (no pain) to 3 (acute pain)	67.7 (2.1)	36.9 (1.2)	
Poulain ⁴⁴	Responder status	11-point (NRS)	No pain to pain as bad you can imagine	1.5 (1.5)	1.5 (1.5)	14
Camba ³⁸	Pain intensity	VAS	0 cm = no pain to 10 cm = maximum pain	7.4 (1.2)	3.2 (not reported)	56
Apolone ⁴⁶	Worst and average pain intensity	11-point (NRS)	No pain to pain as bad you can imagine Worst, actual, least and average pain	6.4 (2.3) worst	4.8 (2.5) worst	28
				3.2 (2.6) actual	2.3 (2.2) actual	

Abbreviations: NRS, numerical rating scale; VRS, verbal rating scale; VAS, visual analog scale; SF-MPQ, Short-form McGill Pain Questionnaire.

Estimate of overall safety and tolerability

After the narrative description of safety and tolerability given for each paper and study, we tried to summarize and integrate these important characteristics; however, papers describing the 12 clinical studies reported the safety and tolerability of the drugs under evaluation using different and heterogeneous methods and terms. Most authors reported side effects in terms of adverse events, ADR, or simply using a list of most frequent symptoms reported by patients. Sometimes, adverse events and ADR were also classified as severe. Some authors reported side effects that were attributable to opioid administration only, while others classified them according to the system or organ involved, such as skin, central nervous system, or according to the type, such as nausea, vomiting, and constipation. Given the large variability of the methods used, we tried to classify them into the following groups: adverse events, ADR, gastrointestinal (including nausea, vomiting, constipation), central nervous system (including confusion, dizziness), and skin (including pruritus and local erythema). Table 4 shows the results of such reclassification and synthesis. Most of the times, it was not possible

to identify the exact quantity and type of the side effects, as the authors did not report the estimates according to the type of patients (cancer vs non-cancer). In addition, the unit of analysis was not homogeneous across the studies (as the authors reported either the number of events or the number of patients with at least 1 event), yielding a quite unreliable classification of the phenomenon.

Discussion

Buprenorphine TDS has been available in most European countries since 2002, to treat moderate to severe chronic pain. The TDS formulation that allows for a slow release by minimizing the typical opioid side effects makes it a good alternative to the other oral opioids in clinical practice. The results from 3 clinical pivotal trials^{28,31,32} including only 453 cancer patients, documented its analgesic efficacy, at least in terms of the responders' status, and the safety profile that was typically opioid in nature and not different from placebo in terms of incidence of reported events; local adverse events associated with TDS application were erythema and pruritus, while the most frequent systemic

adverse events were nausea and vomiting. Pharmacokinetics and pharmacodynamics data suggest that buprenorphine TDS may be safely used in aged patients and in renally impaired individuals, although the small sample size of the studies and the restriction in inclusion criteria did not allow any subgroup or interaction analysis. Post-marketing and outcome research of the prospective studies yielded satisfactory results in terms of effectiveness, despite the presence of a large variability in inclusion/exclusion criteria and outcomes assessment. Safety and tolerability results were often reported together for both cancer and non-cancer patients, making it difficult to extract and summarize the findings across the individual original studies. Despite these limitations, the type of adverse events was in line with that expected, given the opioid nature of the drug, and the reported incidence from observational studies, when evaluable, was low, mostly in terms of serious events.

The evidence for comparative efficacy and safety is indeed scanty, as most of the efficacy data are from placebo trials. Only one RCT⁴² was designed to compare it with morphine. This trial had an open-label design and a very small sample size. As pointed out by other researchers,⁹ well-designed and statistically powered controlled clinical trials focusing exclusively on treating cancer patients suffering from pain using buprenorphine TDS are still lacking. In addition, our analysis based on a systematic review that included published papers up to April 2009 confirmed that

the quality of the 19 reports describing 16 studies was poor and did not allow for safe data pooling. Studies, although stratified according to the study design, were very heterogeneous in terms of population included, interventions studied, comparators evaluated, and outcomes assessed. Even when the same outcome was formally assessed (eg, the responders' status or the patients' reported pain intensity), the differences in terms of operational definition of endpoints, timing of the assessment, or simply the type of measures used (NRS vs VRS vs VAS), made it impossible to identify a subsample for formal data synthesis and statistical pooling.

The results of this systematic review confirm the potential of this drug when delivered through a TDS, but it may be considered as a lost opportunity for a meta-analysis. The studies that we retrieved were very few, very heterogeneous, and reported results in a way that prevented an appropriate data extraction and data pooling. This fact might explain why TDS buprenorphine has not yet been recommended by national and international guidelines, and sometimes is not included in the list of reimbursable drugs at national or local levels.

In fact, the poor quality of pain studies is a well-known phenomenon throughout the field, across the type of interventions, countries, and clinical settings. A recent literature review carried out by the European Palliative Care Research Collaborative Group to assess the quality of pain assessment in palliative care documented that in the 230 papers retrieved and evaluated, the methods and tools

Table 4 Local and systemic safety and tolerability adverse events and adverse drug reactions from 12 selected studies

Name of the first author	Number of patients with cancer	Number of AEs	Number of ADRs	% of patients reporting side effects or AEs related to		
				GI	CNS	Skin
Böhme ²⁸	83	NA	NA	NA	NA	NA
Sittl ³¹	21	NA	6	NA	NA	NA
Sorge ³²	45	NA	NA	NA	NA	NA
Likar ⁴⁰	9	NA	0	NA	NA	NA
Pace ⁴²	52	15/26 TDS buprenorphine	0	19.2	38.4	NA
		37/26 oral morphine		73.0	69.2	
Poulain ⁴⁴	89	NA	2	18.0	4.8	NA
				15.8	0.0	
Griessinger ³³	3690	NA	18	NA	NA	NA
Muriel ³⁴	207	NA	NA	NA	NA	NA
Likar ³⁶	134	84	26	14.1	5.9	8.9
Camba ³⁸	164	NA	NA	NA	NA	NA
Wirz ⁴⁵	174	NA	NA	NA	NA	NA
Apolone ^{46,a}	257	NA	25%	33.5	25.4	2.5

^aIn this study side effects were reported as 'opioids related symptoms' rated by patients as 'a lot/very much' impact.

Abbreviations: AE, adverse event; ADR, adverse drug reaction; GI, gastrointestinal; CNS, central nervous system; NA, not available; TDS, transdermal delivery system.

used were very heterogeneous and did not meet the criteria identified by experts as minimal standards.^{47,48}

In conclusion, buprenorphine TDS has been observed to have good intrinsic (pharmacological) characteristics and promising clinical results, suggesting the need to appropriately review the evidence base for its utility. The results from this systematic literature review are not conclusive and suggest the need for 3 sequential initiatives: 1) a meta-analysis carried out on individual data to minimize the problems that we have found working on aggregated data; 2) an international initiative to find a consensus on definitions, methods, and measures of pain assessment in clinical research; and 3) a well-designed, large, multicenter, international comparative study to produce final evidence about the relative effectiveness and safety of the available third-level WHO opioids. These above-mentioned actions are now being planned.

For the Cochrane Collaboration, one group has published a protocol titled 'Buprenorphine for cancer pain'. The authors planned to carry out a systematic review based on an electronic search of 5 databases, including the contact of the original authors of the primary studies for clarification, or to obtain missing information or individual data.⁴⁹

A joint initiative of 3 non-profit organizations (Center for the Research and Evaluation of Pain, Norwegian University of Science and Technology, Italian National Cancer Institute) planned an expert panel meeting on the issues related to cancer-pain assessment in clinical research. The first meeting involving experts from the USA and Europe, including representatives of FDA and EMEA, will be held in Milan, Italy, in September 2009.

Finally, on the basis of the results of this review and according to the preliminary results of a prospective outcome research study carried out in Italy on 1800 cancer patients with pain,⁵⁰ a multicenter RCT to compare the efficacy of 4 strong opioids on cancer pain (buprenorphine, morphine, fentanyl, and oxycodone) will be launched at the end of this year, involving about 80 centers and 1000 patients.

Disclosures

Dr Apolone and Dr Corli have received consulting and lecture fees from Grunenthal-Italy.

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